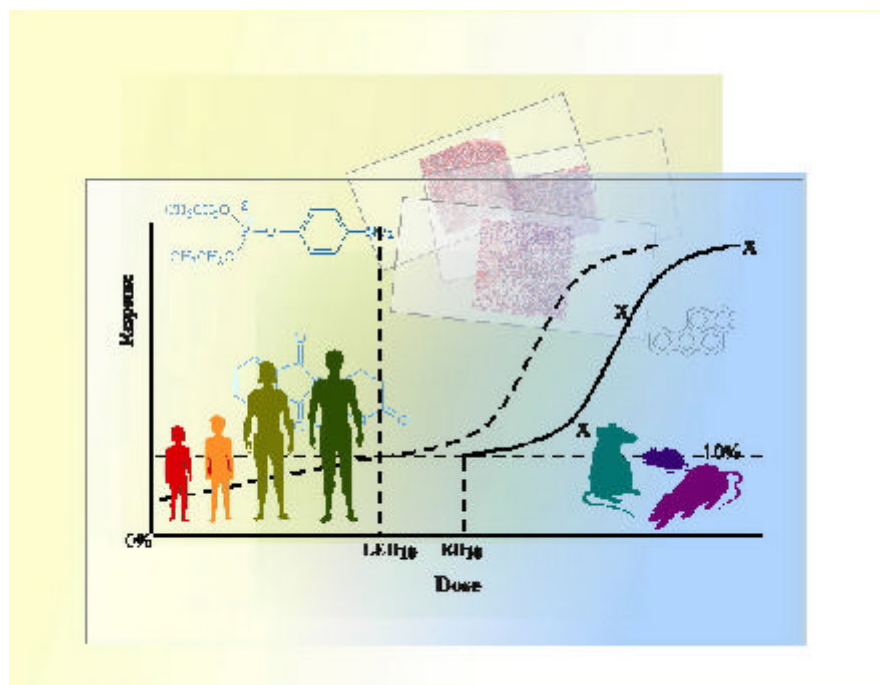


US EPA ARCHIVE DOCUMENT

# HUMAN HEALTH RISK ASSESSMENT

## CHLORPYRIFOS



U.S. Environmental Protection Agency  
Office of Pesticide Programs  
Health Effects Division (7509C)

Deborah C. Smegal, M.P.H., Risk Assessor  
June 8, 2000

# HUMAN HEALTH RISK ASSESSMENT

## CHLORPYRIFOS

### Phase 4

#### Risk Assessment Team:

**Lead Risk Assessor:** Deborah C. Smegal, M.P.H., Toxicologist

**Dietary Risk:** David Soderberg, Chemist

**Residue Chemistry:** Steven Knizner, Senior Scientist/Chemist

**Product Chemistry:** Steven Knizner, Senior Scientist/Chemist

**Agricultural,  
Occupational and  
Residential Exposure:** Timothy Leighton, Environmental Health Scientist

Deborah C. Smegal, M.P.H., Toxicologist

**Toxicology:** Deborah C. Smegal, M.P.H., Toxicologist

**Incident Review:** Jerome Blondell, Health Statistician

Virginia Dobozy, Veterinary Medical Officer

#### Management:

**Senior Scientist:** Steven Knizner

**Branch Chief:** Jess Rowland

**Division Director:** \_\_\_\_\_  
Margaret J. Stasikowski, June 8, 2000

## Background

Attached is HED's risk assessment of the organophosphate pesticide, chlorpyrifos, for purposes of issuing a Reregistration Eligibility Decision (RED) Document for this active ingredient. Cumulative risk assessment considering risks from other pesticides or chemical compounds having a common mechanism of toxicity is not addressed in this document. This risk assessment updates the October 18, 1999 version and addresses the Public Comments in accordance with Phase 3 of the Tolerance Reassessment Advisory Committee (TRAC) Organophosphate (OP) Pilot Process. EPA and the registrants have agreed to certain modifications to the use of chlorpyrifos to mitigate dietary, worker and residential risks. This risk assessment incorporates elements of the risk mitigation agreement in a number of its analyses in order to characterize post-mitigation risks. The disciplinary science chapters and other supporting documents for the chlorpyrifos RED are also included as attachments as follows:

- ' Report of the Hazard Identification Assessment Review Committee. D. Smegal (4/6/2000, HED Doc No. 014088)
- ' Report of the FQPA Safety Factor Committee. Brenda Tarplee (4/4/00; HED Doc No. 014077)
- ' Revised Product and Residue Chemistry Chapter. Steven Knizner (June 2000)
- ' Toxicology Chapter. Deborah Smegal (4/18/00; D263892)
- ' Occupational/Residential Handler and Post-Application Residential/Non-Occupational Risk Assessment. D. Smegal/T. Leighton (June 2000; D266562)
- ' Agricultural and Occupational Exposure Assessment: Tim Leighton (June 2000; D263893)
- ' Acute Dietary Risk Assessment for Chlorpyrifos. (D. Soderberg June 2000, D263890)
- ' Chronic Dietary Exposure Assessment for Chlorpyrifos. D. Soderberg (June 2000, D263889)
- ' Chlorpyrifos Incident Review Update: Jerome Blondell (4/20/00). Update of Incident Data on Chlorpyrifos for Domestic Animals. Virginia Dobozy (04/26/99; D255514)
- ' Analysis of Chlorpyrifos IDS Data for Domestic Animals. Virginia Dobozy (1/23/95)
- ' Drinking Water Assessment from the Environmental Fate and Effects Division (EFED). Michael Barrett (11/13/98)

- ' EFED Concerns over well contamination associated with termiticide use and EFED Recommended Concentrations for HED Drinking Water Assessment of Chlorpyrifos. Henry Nelson (10/6/99)
- ' Chlorpyrifos Preliminary Risk Assessment for Trichlorpyridinol (TCP) Metabolite. S. Knizner. D265035.

HED's Hazard Identification Assessment Review Committee (HIARC) reviewed the toxicological database for chlorpyrifos and selected toxicological endpoints for acute oral, chronic oral and for short-, intermediate and long-term dermal and inhalation exposure risk assessment in February 1999, and January 2000 (memorandum dated April 6, 2000). HED's FQPA Safety Factor Committee reviewed the hazard and exposure data for chlorpyrifos on January 24, 2000, and deferred to the Office of Pesticide Programs Division Directors and senior scientists (DD-SS). The DD-SS recommended that the 10X FQPA Safety Factor (as required by Food Quality Act of August 3, 1996) be retained in assessing the risk posed by this chemical (memorandum dated April 4, 2000).

In June 1997, the registrants of chlorpyrifos voluntarily agreed to measures designed to reduce household exposure to chlorpyrifos, as part of a risk reduction plan. This voluntary plan included deletion of indoor broadcast use, use as an additive to paint, direct application to pets (sprays, shampoos and dips), and indoor total-release foggers. The technical chlorpyrifos products have been amended to reflect the negotiated plan. The technical label limits end use product labeling to only those sites which are specified on its label. In addition, the registrants have implemented measures to improve education, training, and labels, and report and analyze incidents. In addition, as part of this agreement, the registrants agreed to work with EPA to develop broad, market-wide policies for all indoor insecticides for a number of areas.

EPA and the registrants have agreed to certain modifications to the use of chlorpyrifos to mitigate dietary, worker and residential risks. This risk assessment incorporates elements of this agreement in a number of its analyses in order to characterize post-mitigation risks. The agreement includes:

- ' Agricultural Uses
  - Restrict use on apples to pre-bloom (dormant) application only
  - Cancel use on tomatoes
  - Implement revised restricted-entry intervals for all agricultural crops.

## Non-Agricultural Uses

- Cancel all indoor residential uses (except fully contained ant baits in child resistance packaging).
- Cancel all outdoor residential uses (except limited public health uses).
- Cancel all indoor and outdoor non-residential uses (e.g. FHE) except:
  - Use on golf courses
  - Limited public health uses
  - Limited use in industrial settings (e.g. manufacturing plants, ship holds)
- Cancel whole house “post-construction” termiticide use.
- Phase out limited post-construction spot and local termiticide treatments
- Phase out pre-construction termiticide treatments
- Reduce the maximum application rate for phased-out termiticide treatments to a 0.5% concentration.
- Reduce the maximum application rate for use on golf courses to 1 lb. active ingredient per acre.

In addition to these agreed upon actions the Agency will also propose to revoke the tolerance on tomatoes and reduce the tolerances on apples and grapes to 0.01 ppm. These changes were also included in the analysis of post-mitigation dietary exposure.

# CHLORPYRIFOS REVISED RISK ASSESSMENT

## TABLE OF CONTENTS

|            |  |           |
|------------|--|-----------|
| <b>1.0</b> | <b>Executive Summary</b>                                   | <b>1</b>  |
| <b>2.0</b> | <b>Physical/Chemical Properties Characterization</b>       | <b>12</b> |
| <b>3.0</b> | <b>Hazard Characterization</b>                             | <b>13</b> |
| 3.1        | Hazard Profile   | 13        |
| 3.1.1      | TCP  | 13        |
| 3.1.2      | Neurotoxicity  | 14        |
| 3.1.3      | Subchronic Toxicity  | 14        |
| 3.1.4      | Carcinogenicity/Genotoxicity                               | 15        |
| 3.1.5      | Chronic Toxicity   | 15        |
| 3.1.6      | Developmental Toxicity                                     | 15        |
| 3.1.7      | Reproductive Toxicity                                      | 16        |
| 3.1.8      | Human Studies  | 16        |
| 3.1.9      | Metabolism/Pharmacokinetic Studies.                        | 18        |
| 3.1.10     | Sensitivity/Susceptibility of the Young                    | 19        |
| 3.1.11     | Paraoxonase  | 19        |
| 3.2        | Acute Toxicity   | 20        |
| 3.3        | FQPA Considerations  | 21        |
| 3.4        | Endpoint Selection   | 22        |
| 3.5        | Endocrine Disrupter Effects                                | 26        |
| <b>4.0</b> | <b>Exposure Assessment</b>                                 | <b>27</b> |
| 4.1        | Summary of Registered Uses                                 | 27        |
| 4.2        | Dietary Exposure   | 28        |
| 4.2.1      | Residue Chemistry Data Requirements                        | 28        |
| 4.3        | Dietary Exposure (Food Source)                             | 29        |
| 4.3.1      | Acute Dietary Exposure Assessment                          | 31        |
| 4.3.2      | Chronic Dietary Exposure Assessment                        | 35        |
| 4.3.3      | Drinking Water Exposure                                    | 41        |
| 4.3.3.1    | Groundwater Exposure Levels                                | 42        |
| 4.3.3.2    | Surface Water Exposure Levels                              | 43        |
| 4.3.3.3    | Drinking Water Exposure Concentrations                     | 44        |
| 4.3.3.4    | DWLOCs for Acute (Drinking Water) Exposure                 | 46        |
| 4.3.3.5    | DWLOCs for Chronic Drinking Water Exposure                 | 47        |
| 4.4        | Non-Dietary Exposure                                       | 48        |
| 4.4.1      | Occupational Handler Exposure Scenarios                    | 50        |
| 4.4.1.1    | Occupational Handler Exposure Data Sources and Assumptions | 51        |
| 4.4.1.2    | Occupational Handler Risk Characterization                 | 53        |

|                    |  |            |
|--------------------|--|------------|
| 4.4.2              | Occupational Postapplication Exposure Scenarios .....                        | 57         |
| 4.4.2.1            | Occupational Postapplication Exposure Data and Assumptions .....             | 57         |
| 4.4.2.2            | Occupational Postapplication Risk Characterization .....                     | 58         |
| 4.4.3              | Residential Handler Exposure .....   | 59         |
| 4.4.3.1            | Residential Handler Exposure Scenarios .....                                 | 60         |
| 4.4.3.2            | Residential Handler Exposure Data Sources and Assumptions .....              | 61         |
| 4.4.3.3            | Residential Handler Risk Characterization .....                              | 61         |
| 4.4.4              | Residential/Recreational Postapplication Exposures and Risks ..              | 63         |
| 4.4.4.1            | Postapplication Exposure Scenarios .....                                     | 64         |
| 4.4.4.2            | Data Sources and Assumptions for Postapplication Exposure Calculations ..... | 65         |
| 4.4.4.3            | Residential/Recreational Postapplication Risk Characterization .....         | 65         |
| 4.4.4.4            | Incident Reports .....   | 90         |
| 4.4.5              | Pet Incident Reports .....   | 93         |
| 4.5                | Chlorpyrifos Exposure Estimates in the U.S. Population .....                 | 94         |
| <b>5.0</b>         | <b>Aggregate Risk Assessments and Risk Characterization .....</b>            | <b>100</b> |
| 5.1                | Acute Aggregate Risk .....   | 100        |
| 5.2                | Short-Term Aggregate Risk .....  | 102        |
| 5.3                | Intermediate-Term Aggregate Risk .....                                       | 106        |
| 5.4                | Chronic Aggregate Risk .....   | 106        |
| <b>6.0</b>         | <b>Cumulative Exposure and Risks .....</b>                                   | <b>109</b> |
| <b>7.0</b>         | <b>Confirmatory Data .....</b>   | <b>110</b> |
| 7.1                | Toxicology Data for OPPTS Guidelines .....                                   | 110        |
| 7.2                | Product and Residue Chemistry Data for OPPTS Guidelines .....                | 110        |
| 7.2.1              | Product Chemistry .....  | 110        |
| 7.2.2              | Residue Chemistry .....  | 111        |
| 7.3                | Occupational Exposure Data for OPPTS Guidelines .....                        | 113        |
| <b>8.0</b>         | <b>References .....</b>  | <b>115</b> |
| <b>APPENDIX A:</b> | <b>Sensitivity/Susceptibility of the Young .....</b>                         | <b>120</b> |



# CHLORPYRIFOS

## 1.0 Executive Summary

### Background

The Health Effects Division (HED) has conducted a Human Health Risk Assessment for the active ingredient chlorpyrifos for the purposes of making a reregistration eligibility decision (RED). The toxicological database is complete and adequate to support reregistration in accordance with the Subdivision F Guidelines for a food use chemical. Residue chemistry requirements are substantially complete pending receipt of limited confirmatory data.

Chlorpyrifos, [O,O-diethyl O-(3,5,6-trichloro-2-pyridinyl)-phosphorothioate], is a broad-spectrum, chlorinated organophosphate insecticide that was first registered in 1965 to control foliage- and soil-borne insect pests on a variety of food and feed crops. Chlorpyrifos' most common trade names are Dursban, Empire 20, Equity, and Whitmire PT 270. Lorsban is a trade name for agricultural-use products. It is one of the most widely used organophosphate insecticides in the U.S., and is one of the major insecticides used in residential settings. Approximately 21 to 24 million pounds are used annually in the U.S, of which approximately 11 million pounds are applied in non-agricultural settings. There are approximately 800 registered products containing chlorpyrifos on the market. Registered uses include: variety of food crops (i.e., there are approximately 112 tolerances for food/feed commodities); turf and ornamental plants; greenhouses; sodfarms; indoor pest control products (e.g., crack and crevice); structural pest control (e.g., termites); and pet collars. It is registered for use in residential and commercial buildings, schools, daycare centers, hotels, restaurants and other food-handling establishments, hospitals, stores, warehouses, food manufacturing plants, vehicles, and livestock premises. In addition, it is used as a mosquitocide, and as impregnated in ear tags for cattle. In 1998, Dow AgroSciences (DAS) estimated that 70% of the urban chlorpyrifos use involved termite control. Chlorpyrifos products are widely used by homeowners and professionals.

The following are formulation types for chlorpyrifos: wettable powder, emulsifiable concentrate, dust, granular, bait, flowable concentrate, impregnated material, pelleted/tableted, pressurized liquid, and microencapsulated. Dry flowable and wettable powder formulations in open bags are no longer supported by the primary registrant, Dow AgroSciences (DAS). Therefore, these formulations are not assessed in this risk assessment and are not eligible for re-registration.

## Hazard

Chlorpyrifos is moderately toxic following acute oral, dermal and inhalation exposures (toxicity category II). Chlorpyrifos affects the nervous system by reversibly inhibiting the activity of cholinesterase (ChE), an enzyme necessary for the proper functioning of the nervous system. Inhibition of ChE is the most sensitive effect in all animal species evaluated and in humans, regardless of route or duration of exposure. In animals, significant inhibition of plasma and red blood cell (RBC) ChE occur at doses below those that cause brain ChE inhibition. Data from two human studies suggest that humans are similarly and possibly more sensitive than animals following acute and short-term oral exposure and acute dermal exposure based on plasma ChE inhibition and/or possible clinical signs. Females are slightly more sensitive than males based on ChE inhibition and acute toxicity (comparison of LD<sub>50</sub>'s). Studies in the scientific literature suggest that neonates are more sensitive to oral chlorpyrifos exposure than adults for ChE inhibition and behavioral effects. The increased sensitivity of the young may be attributed to a reduced capacity to detoxify chlorpyrifos.

Developmental and reproductive effects have been observed in rats, rabbits and/or mice, but only at doses that induced maternal or parental toxicity. In rats, chlorpyrifos causes delayed alterations in brain development in offspring of exposed mothers. Several studies in the peer reviewed literature and results of the guideline developmental neurotoxicity study are supportive of the possibility that chlorpyrifos exposure may affect brain development (e.g., altered synaptic development, alterations in DNA, RNA, and protein synthesis, inhibition of mitosis and mitotic figures, and disruption of the structural architecture of the brain). There are suggestive data that these effects may arise independent of cholinesterase inhibition.

Chlorpyrifos did not induce treatment-related tumors or provide evidence of carcinogenicity in two chronic rat or two chronic mouse studies. Chlorpyrifos was not mutagenic in bacteria, or mammalian cells, but did cause slight genetic alterations in yeast and DNA damage to bacteria.

For the purposes of this assessment, HED has concluded that the primary metabolite of chlorpyrifos, 3,5,6-trichloro-2-pyridinol (3,5,6-TCP), is not of toxicologic concern because 3,5,6-TCP does not induce cholinesterase inhibition (58 FR 19354, April 14, 1993). However, because of potential exposure to TCP in food and residential settings, and evidence of increased susceptibility of rabbit fetuses relative to dams based on the DAS-submitted rabbit developmental study, HED conducted a screening-level risk assessment for TCP. This assessment is attached in memorandum from S. Knizner to D. Smegal, D265035 June 5, 2000.

The toxicity endpoints used in this document to assess hazards include acute dietary and chronic dietary reference doses (RfDs), and short-, intermediate- and long-term dermal and inhalation doses. In light of the developing Agency policy on use of toxicology studies employing human subjects, HED selected doses and endpoints for risk assessment based solely on animal studies. Therefore, this document contains risk

assessments based on animal toxicity studies.

The acute dietary RfD of 0.005 mg/kg/day is based on a no-observed adverse effect level (NOAEL) of 0.5 mg/kg/day from an acute oral rat blood time-course study that observed 28-40% plasma cholinesterase (ChE) inhibition 3-6 hours after dosing male rats with a single dose of 1 mg/kg/day (the lowest-observable adverse effect level, LOAEL). This NOAEL is supported by statistically significant 30% RBC ChE inhibition 4 hours after a single 1.5 mg/kg/day exposure by a study in the scientific literature (Zheng et al. 2000). The chronic RfD of 0.0003 mg/kg/day is based on an oral NOAEL of 0.03 mg/kg/day for significant plasma and red blood cell (RBC) ChE inhibition at 0.22 to 0.3 mg/kg/day (LOAEL) based on a weight of the evidence consideration of 5 toxicity studies in dogs and rats. An uncertainty factor of 100 (10X for interspecies extrapolation and 10X for intraspecies variability) was applied to the NOAELs to obtain the RfDs.

A route-specific short-term dermal NOAEL of 5 mg/kg/day from a 21-day dermal rat study has been selected based on plasma and RBC ChE inhibition of 45% and 16%, respectively at 10 mg/kg/day (LOAEL). A dermal absorption adjustment is not necessary because a dermal study was selected. The intermediate- and long-term dermal NOAELs and long-term inhalation NOAEL are 0.03 mg/kg/day based on statistically significant plasma and RBC ChE inhibition that occurred at 0.22 to 0.3 mg/kg/day based on a weight of the evidence of 5 toxicity studies in dogs and rats. Because an oral NOAEL was selected, a 3 percent dermal absorption factor was used. Dermal absorption was estimated to be 3 percent based on the ratio of the oral LOAEL of 0.3 mg/kg/day from the rat developmental neurotoxicity (DNT) study to the dermal LOAEL of 10 mg/kg/day from the 21-day rat dermal study. This absorption factor is comparable to the dermal absorption estimated from human data of 1-3%.

The short- and intermediate-term inhalation NOAEL is 0.1 mg/kg/day from two separate 90-day rat inhalation studies that did not observe effects at the highest vapor concentration tested. HED selected a LOAEL of 0.3 mg/kg/day for 43% plasma and 41% RBC ChE inhibition from the oral developmental neurotoxicity study in rats to complete the dose-response assessment. A 100% default inhalation absorption factor (i.e., inhalation and oral absorption are equivalent) was used.

### FQPA Safety Factor

The Food Quality Protection Act (FQPA) Safety Factor Committee re-evaluated the previous FQPA safety factor recommendation based on new hazard information, and deferred to the OPP Division Directors and several Agency senior scientists (DD-SS group) for the recommendation. The Division Directors and senior scientists (DD-SS group), recommended that the FQPA safety factor should be **retained at 10X** for the protection of infants and children from exposure to chlorpyrifos. The FQPA safety factor is applicable to **females 13-50, and infants and children** population subgroups for acute and chronic dietary risk assessments and residential and other non-occupational risk assessments of all durations. The safety factor was retained because new data in the

literature (Zheng et al. 2000) demonstrated increased neonatal sensitivity following a low-level single oral exposure, and a registrant submitted developmental neurotoxicity (DNT) study showed a clear qualitative difference in response (i.e., susceptibility) between adult rats and their offspring. Cholinesterase inhibition was observed in dams versus structural effects in the developing brain of the offspring.

In addition, the new data in the literature also gave rise to uncertainties such as the suggestion that the inhibition of cholinesterase may not be essential for adverse effects on brain development; and the lack of an offspring NOAEL in the DNT based upon structural alterations in brain development as the toxicity endpoint of concern (i.e., effects were seen at the lowest dose evaluated).

## Dietary Exposure and Risk

HED conducted the most highly refined acute probabilistic and chronic deterministic dietary (food) exposure analyses possible using the Dietary Exposure Evaluation Model (DEEM). Both the acute and chronic dietary analyses incorporate monitoring data obtained from U.S. Department of Agriculture's (USDA's) Pesticide Data Program (PDP), the Food and Drug Administration's (FDA's) Surveillance Monitoring Program, in addition to monitoring data from Dow AgroSciences' (DAS') 1993 National Food Survey (NFS) (a market basket survey), and field trial data for a limited number of crops. Percent crop treated data and processing and cooking factors were also used to refine the exposure estimates. The Agency's acute and chronic analyses incorporated PDP and FDA monitoring data to the greatest extent possible, and NFS data for seven of the nine commodities included in the survey (milk, apple juice, applesauce, orange juice, ground beef, pork sausage and peanut butter). The NFS data for fresh apples were also included in a sensitivity analysis. The NFS tomato data were not included because only 54 samples were collected from Florida, while more extensive and recent data for fresh tomatoes are available from PDP (881 samples, collected in 1996 and 1997). PDP monitoring data also reflect the use of chlorpyrifos on imported fresh tomatoes (a significant source of fresh tomatoes). Therefore the PDP fresh tomato residue data were used exclusively in all analyses.

Three data sets are available for estimating residues on fresh apples: PDP data for analysis of individual single apples; PDP "decomposed" apple data; and NFS "decomposed" apple data. Use of each of these three data sets for fresh apples leads to a different exposure estimate. The dietary exposure analysis has been performed using all commodities having chlorpyrifos uses and each of the apple data sets separately: PDP data for single apples; PDP "decomposed" apple data; and NFS "decomposed" apple data.

In both acute and chronic risk assessments, exposure was compared to a population adjusted dose, (PAD), which is the reference dose (RfD) reflecting retention of the FQPA 10x factor for females and children. HED considers dietary residue contributions greater than 100% of the PAD to be of concern. The acute and chronic PADs are 0.0005 and 0.00003 mg/kg/day, respectively for children and females 13-50 years. The acute and chronic PADs are 0.005 and 0.0003 mg/kg/day, respectively for all

other population groups. The Agency's highly refined **acute dietary exposure** estimates at the 99.9<sup>th</sup> percentile were greater than 100% of the aPAD for all child subpopulations based on the 1999 PDP single apple data, the decomposited 1994-1997 PDP apple data, and/or the decomposited 1993-1994 NFS apple data. Children 1-6 years old were the most highly exposed population subgroup, regardless of which data set is used for fresh apples. Apples contribute most to the child risk estimates. For **children 1-6 years old**, risk estimates ranged from **170% to 355% of the aPAD** depending on which fresh apple data set was used. Use of PDP's 1999 single apple data resulted in the highest exposure estimates. Use of the decomposited NFS fresh apple data resulted in the lowest exposure estimates. Because the PDP single apple data are the most recent and do not require decompositing, these data are expected to provide the most reliable exposure and risk estimates. However, no matter which of the three data sets is used for fresh apples, the critical exposure commodity (CEC) analysis indicated that residues on fresh apples were the major contributor to dietary exposure estimates for children 1-6 years old at the 99.9th percentile exposure. Residues on whole tomatoes and grapes were the next major contributors to exposure.

Various risk mitigation measures were examined to reduce acute dietary exposure and risk estimates. Risk estimates could be reduced to less than 100% of the aPAD for children 1-6 years old only with mitigated exposures from consumption of fresh apples, grapes and tomatoes. Acute dietary risk estimates for children 1-6 years old were reduced to 82% of the aPAD based on the following mitigation measures: reduction of the apple tolerance to 0.01 ppm based on dormant application only; reduction of the grape tolerance to 0.01 ppm based on the domestic use pattern; and deletion of the use and removal of the tolerance on tomatoes. Ingestion of residues detected on a number of commodities (spinach, squash and carrots) that lack chlorpyrifos tolerances does not impact the acute dietary risk estimates. Because chlorpyrifos is not registered for use on these crops, these residues represent chlorpyrifos misuse or possibly spray drift.

The Agency's average **chronic dietary exposure** estimates for the U.S. population and all subgroups, with or without consideration of food handling establishment use, **are below HED's level of concern**. Without consideration of the food handling establishment (FHE) use, the average exposure estimates comprised 3% of the cPAD for the general population and 61% of the cPAD for the most highly exposed subgroup, children 1-6 years old. The Agency average exposure estimates including the food handling establishment use comprised 4% of the cPAD for the general population and **81% of the cPAD** for the most highly exposed subgroup, children 1-6 years old. The risk mitigation measures designed to reduce acute dietary risk also reduce chronic dietary risk. Children 1-6 years old remain the most highly exposed subpopulation, with risk estimates of 51% and 31% of the cPAD, including the FHE use or using zero residues for the FHE use, respectively. Ingestion of residues on a number of commodities (spinach, squash and carrots) that lack chlorpyrifos tolerances does not impact the chronic dietary risk estimates.

## Drinking Water Exposure and Risk

The available environmental fate data suggest that chlorpyrifos has a low potential to leach to groundwater in measurable quantities from typical agricultural uses, however, there have been instances of well contamination following termiticide use. The available data indicate that the primary metabolite of chlorpyrifos, 3,5,6-TCP is more mobile, and significantly more persistent in many soils, especially under anaerobic conditions. The Agency has provided a screening-level drinking water assessment based on simulation models and an analysis of available monitoring data to estimate the potential concentrations of chlorpyrifos in ground and surface water.

The Agency conducted an analysis of over 3000 filtered groundwater monitoring well data available in U.S. Geological Survey's National Water Quality Assessment (NAWQA) Program databases, and in the Agency's Pesticides in Ground Water Data Base (PGWDB). Chlorpyrifos was infrequently detected in groundwater (< 1% of the 3000 wells), with the majority of concentrations reported to be <0.01 ppb, and a maximum detected concentration of 0.65 ppb in the PGWDB. Groundwater concentrations following termiticide use are potentially much higher, with a maximum reported concentration of 2090 ppb because of well contamination. The Agency also performed screening-level model estimates of chlorpyrifos concentrations in groundwater using SCI-GROW. Inputs to the models included high exposure agricultural scenarios for major crops (alfalfa, corn, citrus, and tobacco) at the maximum application rates. The estimated concentrations of chlorpyrifos in groundwater using the SCI-GROW screening model range from 0.007 to 0.103 ppb.

The Agency also evaluated more than 3000 samples from 20 NAWQA study units for surface water. In surface water, chlorpyrifos was detected at frequencies up to 15% of 1530 agricultural streams, 26% of 604 urban stream samples in 1997 and in 65% of 57 urban stream samples from Georgia, Alabama and Florida in 1994. The maximum reported dissolved chlorpyrifos concentration in surface water is 0.4 ppb, with the 95<sup>th</sup> percentile at 0.026 ppb, and the majority of concentrations < 0.1 ppb. However, the Agency notes that the monitoring data are not available for the most vulnerable watersheds or groundwater where chlorpyrifos use is pervasive. The Agency also performed screening-level model estimates of chlorpyrifos concentrations in surface water using Tier I GENECC or Tier II PRZM/EXAMS. Estimated maximum 90 day average and peak concentrations of chlorpyrifos in surface water using the PRZM/EXAMS screening model are 6.7 Fg/L and 40.6 ppb, respectively.

Based on the monitoring data and model estimates the Agency used a range of upper-bound estimated environmental concentrations (EECs) in water for the water assessment. For the acute and chronic groundwater assessment an EEC range of 0.007 to 0.103 ppb was used based on screening-level model estimates. For the acute surface water assessment a range of 0.026 to 0.4 ppb was used, based on the 95<sup>th</sup> percentile and maximum reported concentrations from monitoring data. For the chronic surface water assessment, the 95<sup>th</sup> percentile concentration from monitoring data of 0.026 ppb was used. For termiticide use, the Agency had upper-bound groundwater concentrations of 30 to 2090 ppb for the acute exposures, based on well remediation efforts and monitoring data, respectively, and 8.3 to 578 ppb (acute values adjusted for partial environmental

degradation) for chronic exposures. The SCIGROW model and the monitoring data do not reflect actual drinking water concentrations after dilution (from source to tap) or drinking water treatment.

HED calculated drinking water levels of comparison (DWLOCs) assuming mitigation measures for diet and residential uses. Except for possible contamination resulting from termiticide use, the acute and chronic DWLOCs are greater than the EECs and thus do not exceed HED's level of concern.

Exposures to chlorpyrifos from groundwater because of well contamination as a result of the termiticide use for either acute or chronic durations may result in exposures that are potentially of concern. However, implementation of PR-96-7 has reduced the reported incidents of groundwater contamination resulting from termiticide treatment.

### Occupational and Residential Exposure and Risk

Occupational and residential exposures to chlorpyrifos can occur during handling, mixing, loading and application activities. Occupational postapplication exposure can occur for agricultural workers re-entering treated fields such as during scouting, irrigation and harvesting activities.

Residential postapplication exposure can occur following treatment of lawns, or residences for cockroaches, carpenter ants, termites, and other insects. In addition, there is a potential for inadvertent oral exposure to children from eating chlorpyrifos-treated turf and soil or licking fingers following contact with treated areas. Postapplication exposure to children can occur in locations other than the home, including schools, daycare centers, playgrounds, and parks.

There is insufficient use information and exposure data to assess exposure resulting from use in vehicles (i.e., planes, trains, automobiles, buses, boats) and other current label uses such as treatment of indoor exposed wood surfaces, supermarkets, theaters, furniture, and draperies, etc. HED has concern for these uses based on the residential scenarios assessed within this document, which show that nearly all current uses evaluated result in exposures that exceed HED's level of concern. HED has requested additional exposure data for all registered uses not evaluated in this assessment. Although there is concern for these uses, the Agency believes that exposure to these uses will not be higher than the scenarios evaluated in the risk assessment.

HED has conducted dermal and inhalation exposure assessments for: occupational and residential handlers; occupational postapplication; and residential postapplication dermal and inhalation exposure to adults and children as well as inadvertent oral exposure to children. The exposure duration for short-term assessments is defined as 1 to 30 days. Intermediate-term durations are 1 month to six months, and long-term exposures are durations greater than six months. The duration of exposure is expected to be: short-term for agricultural handlers; intermediate and long-term for the occupational handler in residential settings (i.e., lawn care operator and pest control operator); intermediate-term

for occupational postapplication; and short-term for the residential handler. The postapplication residential exposures evaluated in this assessment are considered short-term, except for exposures from termiticide treatment which is considered a long-term exposure.

For the dermal and inhalation risk assessment, risk estimates are expressed in terms of the Margin of Exposure (MOE), which is the ratio of the NOAEL selected for the risk assessment to the exposure level. For occupationally exposed workers, MOEs >100 (i.e., 10x for interspecies extrapolation and 10x for intraspecies variability) do not exceed HED's level of concern. For residential populations, MOEs >1000, which includes the 10x FQPA safety factor for females 13-50 and children, do not exceed HED's level of concern. The target MOE of 1000 is applicable for residential handlers.

The **majority of occupational risk estimates do not exceed HED's level of concern** with appropriate personal protective equipment (PPE) or engineering controls. The results of the short-term handler assessments indicate that only 1 of the 16 potential exposure scenarios did not provide at least one application rate with a total MOE(s) greater than or equal to 100 at either the maximum PPE (i.e., coveralls over long pants, long sleeved shirts, and chemical resistant gloves while using open systems) or using engineering controls (i.e., closed systems). In the majority of cases, dermal exposure contributes more significantly to the total MOE than inhalation exposure.

In total, exposure and risk estimates were calculated for 56 scenarios. Based on the maximum level of protection (i.e., various levels of PPE or engineering controls) 2 MOEs are estimated to be less than 10; 6 MOEs are between 10 and 50; 9 MOEs are between 50 and 100, and 39 MOEs are greater than 100. Fourteen of the scenarios were evaluated based on data obtained from five chemical-specific studies submitted by DAS. The agricultural handler assessments are believed to be reasonable high end exposure representations of chlorpyrifos uses.

There is insufficient information (e.g., dermal and inhalation exposure data) to assess 3 scenarios: seed treatment uses, dip applications (e.g., preplant peach root stock, and nursery stock), and dry bulk fertilizer applications to citrus orchard floors. Given the results from the other scenarios assessed, these scenarios may also need to be mitigated. HED has requested data for these scenarios.

The results of the **Pest Control Operator (PCO)/Lawn Care Operator (LCO) handler assessment in residential/recreational settings** for short-, intermediate and/or long-term exposure scenarios indicate that **most** of the MOEs are less than 100, and therefore **exceed HED's level of concern**. The only scenarios that result in MOEs above 100, and do not exceed HED's level of concern are: (1) lawn care professionals that wear PPE and mix and load liquid lawn products (but do not apply) (total MOEs 100-820), (2) workers who mix/load or apply chlorpyrifos for aerial mosquito applications of less than 30 days with the use of engineering controls (closed systems)(total MOEs 160-240); (3) workers who mix/load or apply chlorpyrifos for ground-based fogger mosquito



applications up to several months with the use of PPE or engineering controls (total MOEs 100-560), and (4) most golf course workers who use the typical rate of 1 lb ai/acre or mixer/loaders of wettable powder that handle product to treat 4 lb ai/acre for less than 30 days (total MOE 100-400).

A number of risks were estimated based on chemical-specific biomonitoring studies submitted by DAS (i.e., indoor crack and crevice treatment, broadcast turf application, and pre- and post-construction termiticide treatment) in which the LCOs/PCOs wore label-specified PPE or PPE in addition to that specified on labels. Several of these studies did not apply the product at the maximum label rate, or only evaluated exposures for a few hours (i.e. 1-3 hours) of the work day, and consequently could underestimate exposures and risks to LCOs/PCOs. Overall, the exposures and risk estimates for LCOs/PCOs based on the chemical-specific biomonitoring studies are considered to be central tendency estimates because they evaluated less than a full day's exposure at the maximum label rate. In the absence of chemical-specific data, LCO/PCO exposures were estimated using data from Pesticide Handlers Exposure Database (PHED) or the Draft Residential SOPs.

The results of the **short- and intermediate-term postapplication** assessments for **workers at agricultural use sites** indicate that restricted entry intervals (REIs) need to be established. REIs represent the duration in days which must elapse before the Agency would not have a concern (MOE \$100) for a worker wearing a long-sleeved shirt and long pants to enter the treated area and perform specific tasks. The **REIs range from 24 hours** for the low, medium, and high crop grouping matrix **to 10 days** for harvesting cauliflower. In short, REIs are 24 hours for all crops except the following: cauliflower (10 days), all nut trees (2 days), all fruit trees (4 days), and citrus (5 days). The occupational postapplication assessment is believed to be reasonable high end representations of chlorpyrifos uses. Four registrant-submitted dislodgeable foliar residue (DFR) studies are included in this assessment. Specifically, data are available for sugar beets, cotton, sweet corn, almonds, pecans, apples, citrus, cauliflower, and tomatoes. The short-term MOEs for postapplication exposure for mow/maintenance workers at golf courses are above 100 (110-210) and therefore, do not exceed HED's level of concern, even at the maximum label rate of 4 lb ai/acre.

**All nine short-term residential handler** exposure scenarios evaluated have total dermal and inhalation MOEs (based on typical, and maximum usage rates) that **exceed HED's level of concern** defined by a target MOE of 1000. MOEs for the residential handler ranged from 3 to 900 for dermal risk, from 120 to 57,000 for inhalation risk, and from 3 to 880 for total dermal and inhalation risk. The following scenarios were evaluated: (1) indoor crack and crevice treatment, (2) lawn treatment with liquid products, (3,4,5) lawn treatment with granular formulations via push-type spreader, belly grinder and hand application, (6) application of ready to use products, (7) dust product applications, (8) paintbrush application, and (9) treatment of ornamentals. In some instances, when the product is not applied at the maximum label rate, the MOEs are above 1000 (i.e., 2 oz crack and crevice spot treatment with a MOE of 1600). Only one of the residential handler

scenarios was evaluated using chemical-specific data submitted by DAS, the remaining scenarios were evaluated using the Residential SOPs or PHED.

The results of the **residential postapplication** exposure scenarios indicate that **seven of the nine** scenarios evaluated have MOEs that are less than 1000, and therefore **exceed HED's level of concern**. These scenarios include exposures following indoor crack and crevice treatment, pet collars, termiticide treatments, liquid and granular lawn treatments and yard and ornamental sprays. In addition, for post application exposure to children following perimeter applications to homes, it was estimated that more than seven hand-to-mouth events or more than 8 minutes of play on treated turf the day of treatment could result in potential exposures that could exceed the Agency's level of concern (i.e., MOE < 1000). An additional scenario could not be quantitatively evaluated (post application exposure to insecticidal dust product use) due to an absence of chemical-specific data and recommended procedures in the residential SOPs. MOEs that exceed HED's level of concern ranged from 6 to 980 for total dermal, inhalation and inadvertent oral (in the case of children) risk. The only residential/recreational scenarios that resulted in a MOE above 1000 are the aerial and ground-based fogger adult mosquitocide application (MOEs 15,000 to 42,000) and adolescent and adult golfers for the typical application rate of 1 lb ai/acre (MOEs 1500 - 2400). Several of the residential postapplication risks were estimated based on chemical-specific studies submitted by DAS (i.e., crack and crevice treatment of the kitchen and bathroom, broadcast treatment of turf with chlorpyrifos spray or granules, and termiticide treatment). The exposure and risk estimates based on the chemical-specific studies are considered to be reasonable central-tendency estimates (i.e., arithmetic mean or median exposure was used to calculate risk). Because these studies were conducted in adults, standard EPA assumptions were used to estimate child exposures.

### Poisoning Incidents

Because of its widespread use in residences, chlorpyrifos is often involved in unintentional exposures. About 6% of all pesticide-related calls (estimated at 7,000 annually) received by the poison control centers are related to chlorpyrifos. The overwhelming majority of cases experience only minor symptoms, but about 200 cases per year are serious enough to require special medical attention. Although only a small proportion of cases involve products used by pest control operators, these exposures often involve exposures to concentrated chemical, which can lead to more serious health effects.

### Aggregate Exposure and Risk

As mandated by the FQPA amendments to the Federal Food, Drug and Cosmetic Act (FFDCA), the Agency must consider total aggregate exposure from food, drinking water, and residential sources of exposure to chlorpyrifos. Based on the mitigation plan, this aggregate assessment considers exposure to chlorpyrifos from food, drinking water and residential uses. In addition, the Agency has concerns about possible residential exposures from chlorpyrifos spray drift. The Agency is currently developing methods to

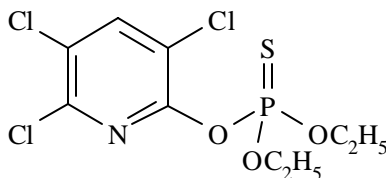
assess residential exposures from spray drift, and these will be assessed in the future when new methods are available. The **acute aggregate risk estimates do not exceed HED's level of concern** because combined exposure to chlorpyrifos through food and drinking water sources are <100% aPAD. The **short-term aggregate risk estimates do not exceed HED's level of concern** based on concurrent exposure to chlorpyrifos from golfing, mosquito abatement activities, in addition to food and drinking water. The **chronic food and drinking water aggregate risk estimates do not exceed HED's level of concern**.

Although not all of the risk estimates for termiticide use achieve a margin of exposure of 1000, the Agency believes that individuals are unlikely to experience adverse health effects from the termiticide use of chlorpyrifos. This conclusion is based on: the public health protective assumptions; the 1000 fold safety factor; and the additional 3 to 10 fold cushion between the NOAEL and the LOAEL. Mitigation measures will further reduce exposures and risk associated with the termiticide use. For example, the removal of whole house barrier treatment addressed the exposures of most concern. It is expected that the limited spot and localized treatment, and pre-construction treatments would represent less exposure and risk. In conclusion, based on the mitigation plan, and best professional and scientific judgement, the Agency concludes that the chronic aggregate risk including termiticide use, does not raise a concern.

Because of its extensive use, the majority of the U.S. population is exposed to chlorpyrifos or its environmental breakdown product, 3,5,6-trichloro-2-pyridinol (3,5,6-TCP). Epidemiology data have reported measurable concentrations of 3,5,6-TCP, which is also the primary metabolite of chlorpyrifos, chlorpyrifos-methyl and trichlorpyr in the urine of individuals. These data represent potential aggregate exposure to chlorpyrifos and/or 3,5,6-TCP from all exposure routes. 3,5,6-TCP was detected in the urine of 82% of 993 adults from the National Health and Nutrition Examination Survey III conducted between 1988 and 1993 (NHANES III). Preliminary results from the recent Minnesota Children's Exposure Study found that 92% of the 89 children evaluated had measurable urinary concentrations of 3,5,6-TCP. A 1998 biomonitoring study of 416 children in North and South Carolina found 3,5,6-TCP in urine of 100% of the children evaluated. TCP was found at higher average levels than all previous epidemiological studies of the general population. HED believes that chlorpyrifos contributes significantly more to urinary TCP than chlorpyrifos-methyl and trichlorpyr based on relative usage of 21-24 million pounds chlorpyrifos versus 92,000 pounds chlorpyrifos-methyl, and 700,000 pounds for trichlorpyr. Because chlorpyrifos, chlorpyrifos-methyl and trichlorpyr degrade to 3,5,6-TCP in the environment, exposure to TCP per se also contributes to the urinary 3,5,6-TCP residues to an unknown degree. As noted previously, HED conducted a screening-level risk assessment for TCP. This assessment is attached in memorandum from S. Knizner to D. Smegal, D265035 June 5, 2000.

## 2.0 Physical/Chemical Properties Characterization

Technical chlorpyrifos is a white crystalline solid with a melting point of 41.5-42.5° C. Chlorpyrifos is stable in neutral and acidic aqueous solutions; however, stability decreases with increasing pH. Chlorpyrifos is practically insoluble in water, but is soluble in most organic solvents (i.e., acetone, xylene and methylene chloride). Chlorpyrifos is not particularly volatile based on its low vapor pressure of  $1.87 \times 10^{-5}$  mmHg at 20°C (Merck Index, 11<sup>th</sup> Edition). Its maximum attainable vapor concentration is 25 ppb at 25° C.



|                    |   |
|--------------------|---|
| Empirical Formula: | C <sub>9</sub> H <sub>11</sub> Cl <sub>3</sub> NO <sub>3</sub> PS |
| Molecular Weight:  | 350.6   |
| CAS Registry No.:  | 2921-88-2   |
| Chemical No.:      | 059101  |

The persistence of chlorpyrifos in soil varies depending on soil type, and environmental conditions. The typical aerobic soil metabolism half life ( $T_{1/2}$ ) ranges from 11 to 180 days, with a mean of 28.7 days. Much longer soil half lives of 175 to 1576 days have been reported for termiticide application rates (Memorandum from M. Barrett to S. Knizner, Drinking Water Assessment of Chlorpyrifos, November 13, 1998, and memorandum from H. Nelson to D. Smegal/M. Hartman, October 6, 1999). The soil/water partition coefficient ( $K_{oc}$ ) value ranges from 360 to 31000, indicating that it is not very mobile in soils.

Technical Grade Active Ingredient (TGAI) data requirements concerning the DAS 99% T (EPA Reg. No. 62719-44) and the 97% T (EPA Reg. No. 62719-15) are satisfied. Guideline 830.6314 (oxidation/reduction) data requirements remain outstanding for the DAS 99% T. There are 45 chlorpyrifos Manufacturing-Use Products (MPs). Data remain outstanding for many MPs. Product chemistry data requirements will be complete, provided that the registrants submit the data required as identified in the Revised Product and Residue Chemistry Chapter (Memorandum from S. Knizer to M. Hartman, October 1, 1999, D259613) for the chlorpyrifos MPs. In addition, the registrants must either certify that the suppliers of starting materials and the manufacturing processes for the chlorpyrifos technicals and manufacturing-use products have not changed since the last comprehensive product chemistry review or submit complete updated product chemistry data packages.

### 3.0 Hazard Characterization

#### 3.1 Hazard Profile

The toxicological database is complete and adequate to support reregistration. in accordance with the Subdivision F Guidelines for a food use chemical.

Chlorpyrifos is moderately toxic following acute oral, dermal and inhalation exposures and is classified in toxicity category II for all exposure routes. Chlorpyrifos affects the nervous system by reversibly inhibiting the activity of cholinesterase (ChE), an enzyme necessary for the proper functioning of the nervous system. Inhibition of ChE is the most sensitive effect in all animal species evaluated and in humans, regardless of exposure duration. In animals, significant inhibition of plasma and red blood cell (RBC) ChE occur at doses below those that cause brain ChE inhibition. In animals, significant plasma and RBC ChE have been observed at oral doses as low as 0.025 to 0.3 mg/kg/day following exposure for two weeks to two years, while significant brain ChE inhibition has been observed at oral doses as low as 1 mg/kg/day following exposure for two weeks in pregnant rats (Hoberman 1998a,b). Female rats and especially pregnant rats appear to be more sensitive than adult male rats to cholinesterase inhibition (Moser et al. 1998, Hoberman 1998a,b, Mattsson et al. 1998). Data from two human studies suggest that humans (adult males) are similarly sensitive and possibly more sensitive than rats and dogs following acute and short-term oral exposure and acute dermal exposure based on plasma ChE inhibition and/or possible clinical signs. It is likely that the human sensitivity for ChE inhibition relative to rats (but not dogs) is due to species differences in the constituents of plasma ChE between rats and humans. For example, in rats, plasma ChE consists of approximately a 60:40 ratio of acetyl cholinesterase (AChE) and butyryl cholinesterase (BuChE), while in most humans and dogs, plasma ChE is predominately as BuChE, which is more sensitive to inhibition than AChE.

##### 3.1.1 TCP

HED has concluded that the primary metabolite of chlorpyrifos, 3,5,6-trichloro-2-pyridinol (3,5,6-TCP), does not induce cholinesterase inhibition, and therefore is less toxic than chlorpyrifos (58 FR 19354, April 14, 1993). However, because of the potential exposure to TCP in food and residential settings, and evidence of increased susceptibility of rabbit fetuses relative to dams, HED conducted a screening-level risk assessment for TCP. This assessment is attached in a memorandum from S. Knizner to D. Smegal, D265035 June 5, 2000.

### 3.1.2 Neurotoxicity

Adult male rats acutely exposed to chlorpyrifos exhibited peak plasma ChE inhibition of 28-40% 3-6 hours after exposure at 1 mg/kg (Mendrala and Brzak 1998), while significant 30% RBC ChE inhibition was noted 4 hours following a single oral dose of 1.5 mg/kg (Zheng et al. 2000). Plasma, RBC and heart ChE inhibition of 45%, 17% and 19%, respectively were observed in female rats 24 hours following a single dose of 5 mg/kg (Dittenber 1997). The acute oral NOAEL for plasma ChE inhibition in male rats is 0.5 mg/kg/day. Clinical signs of neurotoxicity, in the absence of neuropathology, were observed in rats exposed to a single oral dose of 50 mg/kg as evidence by decreased motor activity, and increased incidence of clinical signs consistent with organophosphate intoxication. Chlorpyrifos was negative in the delayed neurotoxicity study in hens at single doses of 50, 100 or 110 mg/kg. Acute oral exposure to hens at 60 to 150 mg/kg caused 59-87% inhibition of neurotoxic esterase (NTE) 4-6 days after exposure (Capodicasa et al. 1991). In addition, delayed neuropathy was noted at 60-90 mg/kg which corresponded to 4-6 times the LD<sub>50</sub> and required aggressive antidotal treatment. In rats, chlorpyrifos failed to inhibit NTE at single doses up to 100 mg/kg. There is evidence that NTE inhibition is related to organophosphate-induced delayed neuropathy (OPIDN).

Following longer-term exposures, there was no evidence of neurotoxicity or neuropathology in rats exposed at doses up to 15 mg/kg/day for 13 weeks. However, in the developmental neurotoxicity study, pregnant dams exposed to chlorpyrifos for approximately 2 weeks exhibited 43% and 41% inhibition of plasma and RBC ChE activity at 0.3 mg/kg/day, significant 18% brain ChE inhibition at 1 mg/kg/day, and clinical signs of neurotoxicity, including fasciculations (muscle twitching), hyperpnea (increased respiration), and hyperactivity in addition to decreased body weight gain at 5 mg/kg/day (Hoberman 1998a,b). Cholinesterase inhibition (68% plasma, 56% RBC and 8% brain) was also noted in rats exposed to 1 mg/kg/day chlorpyrifos for 4 weeks in the cognitive study, while clinical signs of toxicity were not observed until higher doses of 3 mg/kg/day for miosis (pupil contraction) and 10 mg/kg/day for salivation and tremors (Maurissen et al. 1996).

### 3.1.3 Subchronic Toxicity

Several subchronic studies are available for chlorpyrifos including two oral rat studies, one oral dog study, a 21 day dermal toxicity study in rats, and two inhalation studies in rats. The most sensitive effect following subchronic oral exposure is inhibition of plasma ChE in rats and dogs at 0.025 to 0.03 mg/kg/day, and RBC ChE inhibition in dogs and rats at 0.22 to 0.3 mg/kg/day. Rats exposed to higher doses exhibited hematological

effects at doses of 10 mg/kg/day and increased brain and heart weight, adrenal gland effects and decreased body weight gain at 15 mg/kg/day. No adverse effects were noted in rats exposed via inhalation to the highest attainable vapor concentration of 20.6 ppb (287 Fg/m<sup>3</sup>) (0.1 mg/kg/day). No adverse effects were observed in the 21-day dermal study in rats at doses as high as 5 mg/kg/day. However, in a 4-day dermal probe study, rats dermally exposed to doses of 0, 1, 10, 100, or 500 mg/kg/day exhibited reductions in plasma and RBC ChE activities at doses of 10 to 500 mg/kg/day. The 21-day dermal NOAEL is 5 mg/kg/day based on a 45% and 16% inhibition of plasma and red blood cell cholinesterase, respectively in rats dermally exposed to 10 mg/kg/day for 4 days.

### 3.1.4 Carcinogenicity/Genotoxicity

Chlorpyrifos was evaluated for carcinogenic potential in both rats (2 studies), and mice (2 studies). There was no evidence of carcinogenicity. Chlorpyrifos is not mutagenic in bacteria, or mammalian cells, but did cause slight genetic alterations in yeast and DNA damage to bacteria. In addition, chlorpyrifos did not induce chromosome aberrations *in vitro*, was not clastogenic in the mouse micronucleus test *in vivo*, and failed to induce unscheduled DNA synthesis in isolated rat hepatocytes.

### 3.1.5 Chronic Toxicity

Chlorpyrifos was evaluated for chronic toxicity in rats, mice and dogs. In all animal species, the most sensitive effect is inhibition of plasma, RBC and brain ChE that occurred at levels in the range of 0.03 to 3 mg/kg/day. Following chronic exposure dogs appear to be the most sensitive species for cholinesterase inhibition and systemic effects, as noted by increased liver weights in dogs exposed to 3 mg/kg/day that could be an adaptive response. Rats exposed to 7-10 mg/kg/day had decreased body weight and decreased body weight gain, ocular effects, adrenal gland effects and altered clinical chemistry and hematological parameters. Mice appear to be the least sensitive to chronic oral doses of chlorpyrifos, as exposure to 45-48 mg/kg/day resulted in decreased body weight and an increased incidence of non-neoplastic lesions (i.e., keratitis, hepatocyte fatty vacuolation).

### 3.1.6 Developmental Toxicity

Chlorpyrifos was evaluated for developmental toxicity in rats, mice and rabbits. In one rat study, developmental effects (increased post-implantation loss) were noted at 15 mg/kg/day (highest dose tested, HDT), that were also associated with maternal toxicity, while another rat study failed to observe developmental effects at 15 mg/kg/day. Developmental effects were also noted at higher doses in mice at 25 mg/kg/day (minor skeletal

variations, delayed ossification and reduced fetal weight and length) and rabbits at 140 mg/kg/day (decreased fetal weights and crown rump lengths, and unossified xiphisternum and/or 5<sup>th</sup> sternebra). However, in both mice and rabbits, the developmental effects occurred at maternally toxic doses as indicated by reduced weight gain, and food consumption in both species, and increased mortality in mouse dams.

In the rat developmental neurotoxicity study, chlorpyrifos was associated with delayed alterations in brain development in offspring of exposed mothers. Specifically, pups of the 1 mg/kg/day group exhibited significant dose- and treatment-related decreases in measurements of the parietal cortex in female offspring at postnatal day 66. The only maternal effect at this dose was plasma and RBC ChE inhibition. At higher doses, pups of the 5 mg/kg/day group exhibited decreased body weight/body weight gain and food consumption in both sexes, reductions in pup viability, delays in development, decreased brain weight and morphometric alterations in the brain. However, these effects were observed in the presence of maternal toxicity as evidenced by fasciculations, hyperpnea and hyperactivity, in addition to reduced body weight gain.

Several studies in the peer reviewed literature and results of the guideline developmental neurotoxicity study are supportive of the possibility that chlorpyrifos exposure may affect brain development (e.g., altered synaptic development, alterations in DNA, RNA, and protein synthesis, inhibition of mitosis and mitotic figures, and disruption of the structural architecture of the brain) (Whitney et al. 1995, Campbell et al. 1997, Song et al. 1997, Johnson et al. 1998, Das and Barone 1999, Dam 1999, Roy et al. 1998, Hoberman 1998a,b). There are suggestive data that these effects may arise independent of cholinesterase inhibition.

### **3.1.7 Reproductive Toxicity**

Chlorpyrifos induced reproductive toxicity in one generation of rats, but only at dose levels that induced parental toxicity. Reproductive effects included reduced pup weights and increased pup mortality that corresponded to slightly but significantly reduced body weight gain in F0 dams during lactation days 1-21, in addition to parental toxicity as evidenced by inhibition of plasma, RBC and brain cholinesterase activities as well as histological lesions of the adrenal gland (vacuolation of cells of the zona fasciculata).

### **3.1.8 Human Studies**

HED has reviewed two human studies conducted with chlorpyrifos submitted by the registrant (MRID 95175, Accession No. 249203). A third



human study (Kisicki et al. 1999) that evaluated a single dose exposure was submitted on April 27, 1999 but is an incomplete submission because two Appendices with critical data were omitted. In the first study (MRID No. 95175; Coulston et al., 1972), male volunteers from Clinton Correctional Facility (4/dose group) were given daily oral (tablet) doses of 0, 0.014, 0.03, or 0.1 mg/kg chlorpyrifos technical for 7 weeks, 9 days, 21 days and 28 days, respectively. Significant 36-82% plasma ChE inhibition relative to baseline was observed after 9 days of treatment with 0.1 mg/kg/day chlorpyrifos. In addition, one of the four men in the 0.1 mg/kg/day developed blurred vision, runny nose and a feeling of faintness on day 9. Exposure was discontinued on day 9 in this dose group however, due to plasma cholinesterase inhibition that exceeded the study investigator's guideline of 20%-30%. No significant plasma ChE inhibition was observed in the men exposed to 0.03 mg/kg/day for 21 days or at any other dose that could be attributed to treatment. No effects on RBC ChE were found at any dose that could be attributed to treatment. A gradual recovery was observed in plasma ChE values equaling baseline values by day 25 of the recovery period. The registrant and study director contend that the clinical signs were attributed to a cold, and not chlorpyrifos exposure. HED believes that blurred vision is a typical cholinergic sign of ChE inhibition, and can not be attributed to a common cold (February 2, 1998 HIARC Report, HED Doc No. 012471). In addition, there is no reason to believe that other clinical signs would not have appeared if the dosing had continued for 21 or 28 days as it did for the other groups. While the study director claims that exposure to the high dose group was discontinued on day 9 because plasma ChE inhibition was 20-30%, rather than because of concern for the clinical signs, this reason is inconsistent with the study findings of 46% mean plasma ChE inhibition following day 6 of treatment in the 0.1 mg/kg/day group, and 41% plasma ChE inhibition in one individual on day 3. HED notes that the relatively long recovery period of 25 days is unusual for plasma ChE, and is more characteristic of recovery for RBC acetyl ChE inhibition based on the 2 year dog data (McCollister et al. 1971, Kociba et al. 1985).

An acute oral and dermal pharmacokinetic study (Nolan et al. 1982, Accession No. 249203) dosed six men once with 0.5 mg/kg orally and four weeks later dosed five of these same men with 5 mg/kg dermally, and one man with 0.5 mg/kg dermally. No clinical signs or symptoms were observed in any of the subjects, but unlike the previous study, the primary focus of this study was pharmacokinetics. Men orally exposed to 0.5 mg/kg chlorpyrifos exhibited peak plasma ChE inhibition of 64-85%, 12 to 24 hours post-exposure. Peak RBC ChE inhibition of 11-52% occurred on post-exposure day 4. Men dermally exposed to 5 mg/kg chlorpyrifos exhibited peak plasma ChE inhibition of 27-45% on day 3, and mean RBC ChE inhibition of 8.6% on day 4. The return of plasma ChE activity to pre-dose levels required about 30 days. The registrant stated that the inhibition noted on days 3 and 4 is an analytical artifact based on chlorpyrifos

pharmacokinetics. If this is the case, it raises concerns about the quality and reliability of the study data. Again, HED notes that the relatively long recovery period of 30 days is unusual for plasma ChE, and is more characteristic of recovery for RBC acetyl ChE inhibition based on the 2 year dog data (McCollister et al. 1971, Kociba et al. 1985). On the basis of urinary excretion of the 3,5,6-trichloro-2-pyridinol (3,5,6-TCP) metabolite, the minimum oral absorption of chlorpyrifos was estimated at 70% and the minimal dermal absorption at 1-3%. Because the proportion of the administered dose metabolized to this pyridinol is unknown, these estimates are considered minimum values (i.e., absorption could be higher). The mean pharmacokinetic half-life for 3,5,6-TCP in the urine was approximately 27 hours following both oral and dermal exposure.

As noted previously, data from the two human studies suggest that humans are as sensitive and possibly more sensitive than animals based on plasma ChE inhibition and possible clinical signs. For example, in animals (rats), the acute oral (single dose) NOAEL is 0.5 mg/kg/day, while humans exposed to a single oral 0.5 mg/kg/day dose exhibited 64-85% plasma ChE inhibition. Based on an overall assessment of the plasma and RBC ChE inhibition data, the HIARC identified an animal NOAEL and LOAEL of 0.03 mg/kg/day and 0.22-0.3 mg/kg/day, respectively for longer term exposures (several months), while humans exposed to 0.1 mg/kg/day for only 9 days exhibited 36-82% plasma ChE inhibition and possible clinical signs (blurred vision). The short-term dermal NOAEL in rats is 5 mg/kg/day based on plasma and RBC ChE inhibition observed at 10 mg/kg/day, while humans exposed dermally for one day to 5 mg/kg/day exhibited 27-45% plasma ChE inhibition. For all endpoints based on rat data, it is likely that this sensitivity can be attributed to species differences in plasma ChE between the rat and humans. For example, in rats, plasma ChE consists of approximately a 60:40 ratio of acetyl cholinesterase (AChE) and butyryl cholinesterase (BuChE), while in most humans and dogs, plasma ChE is predominately as BuChE, which is more sensitive to inhibition than AChE.

### **3.1.9 Metabolism/Pharmacokinetic Studies.**

In the rat, chlorpyrifos is excreted primarily in the urine (84%) with lesser amounts excreted in the feces (5%) within 72 hours. The metabolism of chlorpyrifos was extensive, and no unchanged parent compound was found in the urine. The major urinary metabolites were 3,5,6-TCP, as well as glucuronide and sulfate conjugates of TCP.

As noted previously, in humans (adult males) approximately 70% of chlorpyrifos is excreted in the urine as TCP within 5 days following acute oral exposure, and the minimum dermal absorption is 1 to 3% (Nolan et al. 1982, Accession No. 249203). The mean pharmacokinetic half-life for 3,5,6-TCP in the urine was approximately 27 hours following both oral and dermal

exposure.

### 3.1.10 Sensitivity/Susceptibility of the Young

A number of studies published in the scientific literature have also been considered by the Agency and are discussed in the Hazard Identification and Assessment Review Committee (HIARC) April 6, 2000 report (HED No. 014088), February 2, 1998 report (HED No. 012471) and December 7, 1998 report (HED No. 013004). Summaries of several of these studies are presented in the attached Toxicology Chapter memorandum from D. Smegal to M. Hartman, April 18, 2000, D263892, and in the report "Chlorpyrifos Children's Hazard: Sensitivity and Susceptibility" March 28, 2000, HED No. 014074 (which is an appendix to the April 6, 2000 HIARC report). The HIARC concluded that there is sufficient evidence in the scientific literature to suggest that exposure to chlorpyrifos results in increased sensitivity and susceptibility to neonates as compared to adult rats. The Weight of Evidence Characterization and Conclusions of the "Chlorpyrifos Children's Hazard: Sensitivity and Susceptibility" document (March 28, 2000, HED No. 014074) are presented in Appendix A.

### 3.1.11 Paraoxonase

Chlorpyrifos, and some other organophosphate (OP) compounds, are detoxified via a two-step pathway involving bioactivation of the parent compound to an oxon by the cytochrome P450 systems, and then hydrolysis of the resulting oxon compounds by esterases such as liver or serum paraoxonase (PON1) (located in the plasma) (Davies et al. 1996, Furlong et al. 1998, Shih et al. 1998). In the human population, serum PON1 activity is genetically determined (polymorphic) and individuals express widely different levels of this enzyme (Davies et al. 1996). Therefore, it is possible that some individuals may be more sensitive to chlorpyrifos toxicity based on genetic factors that regulate serum PON1 activity resulting in a reduced capacity to detoxify chlorpyrifos-oxon. Paraoxonase data were collected for individuals in a recent single dose human study (Kisicki et al. 1999). HED will evaluate these data once they are submitted to the Agency.

In animals, there is evidence that serum paraoxonase is protective against poisoning by OPs. Animals with low PON1 levels were more sensitive to specific OP compounds than animals with high enzyme levels. For example, birds, which have very low to undetectable PON1 activity are more sensitive than various mammals to the acute toxicity of oxons for other OPs (paraoxon, diazinon oxon and pirimiphos oxon). Further rabbits, which have a sevenfold higher serum PON1 activity than rats, are more resistant to the acute toxicity of chlorpyrifos (approximately 9 and 25 fold for acute oral and dermal toxicity, respectively). Rabbit paraoxonase hydrolyzes chlorpyrifos-oxon with a much higher turnover number than does rat paraoxonase (Costa et al. 1999, Li et al. 1993).

### 3.2 Acute Toxicity

Chlorpyrifos is moderately toxic following acute oral, dermal and inhalation exposures, and is classified in toxicity category II for all three routes of exposure for rats. The oral LD<sub>50</sub> values for technical chlorpyrifos are higher in rats (223 mg/kg) than mice (62.5 mg/kg, toxicity category II) or chicks (32 mg/kg, toxicity category 1). Female rats are more sensitive (i.e., lower LD<sub>50</sub>) than male rats for both technical chlorpyrifos and formulated products. Guinea pigs and rabbits are less sensitive to acute toxicity than rats as noted by the oral LD<sub>50</sub> values of 504 mg/kg and 1000-2000 mg/kg, respectively (both category III), and the rabbit dermal LD<sub>50</sub> value of >5000 mg/kg (category IV). Chlorpyrifos was not acutely neurotoxic when given to hens at a single oral dose of 50 mg/kg (the LD<sub>50</sub>), 100 or 110 mg/kg. In rats, the LC<sub>50</sub> was greater than 0.2 mg/L (or 200 mg/m<sup>3</sup>), which is normally assigned toxicity category II. This study is classified as Supplementary because only nominal concentrations were measured. Acute toxicity values and categories for the technical grade of chlorpyrifos are summarized in the following table.

| Table 1. Acute Toxicity Results for Technical Chlorpyrifos |                                   |  |          |
|--|-----------------------------------|--|----------|
| STUDY  | MRID Number                       | RESULTS  | CATEGORY |
| Acute Oral LD <sub>50</sub> - rat                          | 44209101                          | 223 mg/kg M&F  | II       |
| Acute Dermal LD <sub>50</sub> - rat                        | Accession No. 112115              | 202 mg/kg  | II       |
| Acute Dermal LD <sub>50</sub> - rabbit                     | 44209102                          | >5000 mg/kg  | IV       |
| Acute Inhalation LC <sub>50</sub> ; rat Supplementary      | 00146507 and Accession No. 257590 | LC <sub>50</sub> > 0.2 mg/L (200 mg/m <sup>3</sup> ) (nominal concentration) | II       |
| Eye Irritation - rabbit                                    | 44209103                          | slight irritation resolved within 24 hours                                   | IV       |
| Dermal Irritation - rabbit                                 | 44209104                          | mild irritant; (irritation resolved within 7 days)                           | IV       |
| Dermal Sensitization - guinea pig                          | 44209105                          | non-sensitizing  | NA       |
| Acute Delayed Neurotoxicity in hens                        | 00097144<br>00405106              | not neurotoxic at 50, 100 or 110 mg/kg                                       | NA       |

NA = not applicable

### 3.3 FQPA Considerations

In March 1999, the FQPA Safety Factor Committee (SFC) recommended that an FQPA safety factor was needed due to concern for increased sensitivity seen at high doses in a literature study comparing adults and neonates, and for the qualitative increased susceptibility occurring at the high dose in the developmental neurotoxicity study. Nonetheless, the FQPA safety factor was reduced to 3X because of lack of data addressing whether or not these differences would also occur at lower doses. A re-evaluation of this recommendation was conducted by the FQPA SFC on January 24, 2000. The new evaluation was undertaken in order to consider the possible impact of new hazard information received in the last year (Slotkin 1999, Zheng et al. 2000). At the January 24<sup>th</sup> meeting, however, the Committee members were unable to reach consensus on the safety factor recommendation. Subsequently, arguments for retention of the safety factor at 10X or reduction of the safety factor to 3X were presented, with supporting information for review, to the OPP Division Directors and several Agency senior scientists at a February 7, 2000 meeting. The Division Directors and senior scientists (DD-SS group), recommended that the FQPA safety factor should be **retained at 10X** for the protection of infants and children to exposure resulting from chlorpyrifos. The details of this decision are presented in the attached memo from B. Tarplee 4/4/00 HED Doc No. 014077. The DD-SS group recommended that a 10X safety factor be retained for chlorpyrifos due to:

In February 2000, new data (Zheng et al. 2000, Hoberman 1998a,b) demonstrated that the increased sensitivity and susceptibility was not only a high dose phenomenon since:

- < increased sensitivity following a single oral exposure to neonates was seen at substantially lower doses (Zheng et al. 2000, in press); and
- < a clear qualitative difference in response (i.e., susceptibility) between adult rats and their offspring was demonstrated in the developmental neurotoxicity (DNT) study (cholinesterase inhibition in dams versus structural effects on developing brain of the offspring) (Hoberman 1998a,b).

New data in the literature also gave rise to uncertainties such as:

- < the suggestion that the inhibition of cholinesterase may not be essential for adverse effects on brain development; and
- < the lack of an offspring NOAEL in the DNT based upon structural alterations in brain development as the toxicity endpoint of concern.

Therefore, the DD-SS group concluded that their evaluation of the available hazard and exposure databases for chlorpyrifos, including the information received and

reviewed in the past year, results in an overall **higher** degree of concern regarding the potential consequences of chlorpyrifos exposure to infants and children than was determined during the FQPA safety factor evaluation in March 1999. Consequently, they recommended that the FQPA safety factor should be Retained at 10X for the protection of infants and children to exposure resulting from the use of chlorpyrifos.

The FQPA SFC determined that the FQPA safety factor would be applicable to **Females 13-50** and **Infants and Children** population subgroups for **all exposure durations**:

Acute Dietary Assessment - The FQPA safety factor is applicable for Females 13-50 and Infants and Children population subgroups due to the concern that adverse effects could result from a single exposure to chlorpyrifos (as demonstrated in several open literature studies including Zheng et al.).

Chronic Dietary Assessment - The FQPA safety factor is applicable for Females 13-50 and Infants and Children population subgroups due to the concern that potential adverse effects could result from repeated exposure to chlorpyrifos (as demonstrated, for example, in the developmental neurotoxicity study in rats).

Residential and other Non-occupational Exposure Assessment - The FQPA safety factor is applicable for Females 13-50 and the Infants and Children population subgroups for all exposure durations due to the adverse effects resulting from single or repeated exposure(s) to this organophosphate insecticide in or around residential (non-occupational) settings.

### 3.4 Endpoint Selection

It is current Agency policy that a regulatory decision can not be made based on a human study until a formal decision has been made concerning the ethical aspects of such use. The ethics decision regarding the use of toxicology studies employing human subjects has not yet been made. Therefore, the Agency selected doses and endpoints to calculate dietary and non-dietary risk in the current assessment based solely on animal studies.

There are three human studies available for chlorpyrifos, however one of these studies is an incomplete submission (Kisicki et al. 1999). The HED HIARC met on January 5, 1999 to evaluate the scientific quality of the two human studies which were the basis of the previous RfDs and dermal and inhalation risk assessment endpoints. This re-evaluation was initiated because of a joint Science Advisory Panel/Science Advisory Board (SAP/SAB) meeting held in December 1998 that discussed issues surrounding the scientific and ethical concerns for human toxicity testing. The HIARC committee concluded that both human studies (Coulston et al. 1972 MRID No. 00095175, Nolan et al. 1982, MRID No. 00249203)

provided useful scientific information that can be used as supportive data along with the results of animal studies. However, these studies alone are not sufficient for endpoint selection or use in risk assessment primarily because of the small sample size (n=4-6/dose group), evaluation of only adult males (when females tend to be more sensitive), insufficient information on study protocol, and lack of control for confounding factors. In addition, the Nolan et al. (1982) pharmacokinetic study only tested one dose level. Furthermore, the registrant contends that the plasma and RBC ChE activity data results on day 3 and 4 of the Nolan et al. (1982) study are analytical artifacts, which raises concerns about the quality and reliability of the study data.

The HIARC met on February 2, 1999 and re-assessed the toxicology database to select toxicology endpoints based on animal studies for dietary and non-dietary exposure risk assessments. On January 20, 2000, and March 28, 2000 the Committee re-convened to address issues raised during the Phase 3 public comment period. The Committees decisions are presented in the attached HIARC memorandum dated April 6, 2000 (D. Smegal to S. Knizner, HED Doc No. 014088). The doses and toxicological endpoints selected for various exposure scenarios based on animal toxicity studies with chlorpyrifos are summarized in Table 2.



**Table 2  
Summary of Doses and Endpoints Selected for Chlorpyrifos Risk Assessment**

| EXPOSURE SCENARIO   | DOSE (mg/kg/day)  | ENDPOINT   | STUDY  | Target MOE for Workers | Target MOE for Non-Occupational                          |
|---------------------|---|--|--|------------------------|--|
| Acute Dietary       | NOAEL=0.5<br><br>UF = 100<br>FQPA = 10<br>(infants,children and females 13-50)  | Significant (28-40%) plasma cholinesterase inhibition at peak time of inhibition (3-6 hours post exposure) at 1 mg/kg (Mendrala and Brzak 1998).<br><br>Significant 30% RBC ChE inhibition 4 hours post exposure to 1.5 mg/kg/day (Zheng et al. 2000). | Acute Blood Time Course Study in male rats (Mendrala and Brzak 1998) with support from Zheng et al. (2000)   | NR                     | NR   |
|                     | <b>Acute RfD =0.005 mg/kg/day</b><br><b>Acute PAD (children and females 13-50) = 0.0005 or 5x10<sup>-4</sup> mg/kg/day</b><br><b>Acute PAD (general population) = 0.005 or 5x10<sup>-3</sup> mg/kg/day</b>          |  |  |                        |  |
| Chronic Dietary     | NOAEL= 0.03<br>UF= 100<br>FQPA = 10<br>(infants,children and females 13-50)   | Significant plasma and RBC cholinesterase inhibition at 0.22 to 0.3 mg/kg/day  | Weight of Evidence from 5 studies:<br>2 year dog<br>90 day dog<br>2 year rat<br>90 day rat<br>developmental neurotoxicity (DNT) study (at 2 weeks) | NR                     | NR   |
|                     | <b>Chronic RfD =0.0003 mg/kg/day</b><br><b>Chronic PAD (children and females 13-50) = 0.00003 or 3x10<sup>-5</sup> mg/kg/day</b><br><b>Chronic PAD (general population) = 0.0003 or 3x10<sup>-4</sup> mg/kg/day</b> |  |  |                        |  |
| Short-Term (Dermal) | Dermal NOAEL =5<br><br>Absorbed Dermal NOAEL = 0.15 (for biomonitoring) (a)   | Plasma and RBC cholinesterase inhibition of 45 and 16%, respectively at 10 mg/kg/day after 4 days. (Dermal absorption factor not necessary for administered dermal NOAEL)  | 21-day dermal rat study  | 100                    | 1000 (infants,children and females 13-50)<br>100 (males) |

**Table 2  
Summary of Doses and Endpoints Selected for Chlorpyrifos Risk Assessment**

| EXPOSURE SCENARIO                          | DOSE (mg/kg/day)   | ENDPOINT  | STUDY  | Target MOE for Workers | Target MOE for Non-Occupational                           |
|--|--|---|--|------------------------|---|
| Intermediate- and Long-Term (Dermal)       | Oral<br>NOAEL =0.03 (3% dermal absorption)                                       | Significant plasma and RBC cholinesterase inhibition at 0.22 to 0.3 mg/kg/day   | Weight of Evidence from 5 studies:<br>2 year dog<br>90 day dog<br>2 year rat<br>90 day rat<br>DNT study (at 2 weeks) | 100                    | 1000 (infants, children and females 13-50)<br>100 (males) |
| Short-, and Intermediate-Term (Inhalation) | Inhalation<br>NOAEL= 0.1   | Lack of effects in 2 rat inhalation studies at the highest dose tested; 43% plasma and 41% RBC cholinesterase inhibition following oral doses of 0.3 mg/kg/day for 2 weeks in the DNT study | Two 90 day rat inhalation studies (NOAEL) and DNT (LOAEL)  | 100                    | 1000 (infants, children and females 13-50)<br>100 (males) |
| Long-Term (Inhalation)                     | Oral<br>NOAEL= 0.03<br>(assume inhalation absorption is 100% of oral absorption) | Significant plasma and RBC cholinesterase inhibition at 0.22 to 0.3 mg/kg/day   | Weight of Evidence from 5 studies:<br>2 year dog<br>90 day dog<br>2 year rat<br>90 day rat<br>DNT (at 2 weeks)       | 100                    | 1000 (infants, children and females 13-50)<br>100 (males) |

RBC = red blood cell

NR = not relevant

UF = Uncertainty Factor

MOE = Margin of Exposure

PAD = Population Adjusted Dose (includes UF and FQPA safety factor)

(a) Use absorbed dermal NOAEL of 0.15 mg/kg/day (5 mg/kg/day \* 0.03 dermal absorption factor) for comparison with absorbed biomonitoring exposure.

### 3.5 Endocrine Disrupter Effects

The Food Quality Protection Act (FQPA; 1996) requires that EPA develop a screening program to determine whether certain substances (including all pesticides and inerts) “may have an effect in humans that is similar to an effect produced by a naturally occurring estrogen, or such other endocrine effect....” EPA has been working with interested stakeholders, including other government agencies, public interest groups, industry and research scientists to develop a screening and testing program as well as a priority setting scheme to implement this program. The Agency’s proposed Endocrine Disrupter Screening Program was published in the Federal Register of December 28, 1998 (63 FR71541). The Program uses a tiered approach and anticipates issuing a Priority List of chemicals and mixtures for Tier 1 screening in the year 2000. As the Agency proceeds with implementation of this program, further testing of chlorpyrifos and its end-use products for endocrine effects may be required.

## 4.0 Exposure Assessment

### 4.1 Summary of Registered Uses

Chlorpyrifos is a broad-spectrum, organophosphate insecticide that was first registered in 1965 to control foliage- and soil-borne insect pests on a variety of food and feed crops. It is one of the most widely used organophosphate insecticides in the U.S. and is one of the major insecticides used in residential settings. There are approximately 822 registered products containing chlorpyrifos on the market (REFs 9/14/99). Registered uses include: a wide variety of food crops (i.e., there are approximately 112 tolerances for food and/or feed commodities such as citrus, vegetable crops, tree fruits, etc); turf and ornamental plants; greenhouses; sodfarms; indoor pest control products (e.g., crack and crevice); structural pest control (e.g., termites); and in pet collars. Indoor uses include residential and commercial buildings, schools, daycare centers, hotels, restaurants and other food handling establishments, hospitals, stores, warehouses, food manufacturing plants, vehicles, livestock premises, and mushroom houses. In addition, it is used as an adult mosquitocide and is registered for ear tag treatment of cattle (beef and lactating and non-lactating dairy). Chlorpyrifos products are widely used by both homeowners and LCOs/PCOs.

BEAD estimates that the annual total domestic usage of chlorpyrifos is approximately 21 to 24 million pounds ai for 8 million acres treated in the U.S. Approximately 11 million pounds are applied annually in non-agricultural settings (i.e., residences, schools, golf courses, parks). Chlorpyrifos has the largest agricultural market in terms of total pounds ai allocated to corn (26%). The largest non-agricultural markets in terms of total pounds ai applied are PCOs, termite control (24%), and turf (12%). Crops with a high average percentage of their total

U.S. planted acres treated include brussel sprouts (73%), cranberries (46%), apples (44%), broccoli (41%) and cauliflower (31%).

Comprehensive lists of chlorpyrifos end-use products (EPs) and of use patterns with food/feed uses which are subject to re-registration appear are summarized in the Revised Product and Residue Chapter (Memorandum from S. Knizner to M. Hartman, June 2000).

The formulations registered for use on food and feed crops include the granular (G), wettable powder (WP), impregnated material (Impr), dry flowable (DF), and emulsifiable concentrate (EC). Dry flowable and wettable powder in open bags are not assessed and no longer are eligible for re-registration. These formulations may be applied as foliar, bark, seed, and soil-incorporated band or broadcast treatments using ground, sprinkler irrigation, or aerial equipment. The different crop growth stages or timings as to when chlorpyrifos formulations may be applied are dormant, delayed dormant, preplant, at-planting, transplanting, postplant, post-transplant, preemergence, and postemergence. The impregnated material formulation is registered for ear tag use on cattle. The chlorpyrifos formulations registered for food-handling establishments include the microencapsulated (Mcap), emulsifiable concentrate, and liquid ready-to-use (RTU) and soluble concentrate (SC/L) [Source: REFS 9/99].

## 4.2 Dietary Exposure

OPP has determined that TCP is not of toxicological concern and can be excluded from the tolerance expression because it does not inhibit cholinesterase (PP3F2884 and 3F2947 and FAP3H5396 and 3H5411/R1191, Final Rule, D.Barolo, 4/1/93). The conclusions specified in the "Tolerance Reassessment Summary" section of the Revised Product and Residue Chemistry Chapter (Memorandum from S. Knizner to M. Hartman, June 2000) reflect this decision and recommendation to consider only chlorpyrifos *per se* as the residue of concern. HED conducted a screening-level TCP assessment (memorandum from S. Knizner to D. Smegal, June 5, 2000, D265035).

### 4.2.1 Residue Chemistry Data Requirements

Plant and Animal Metabolism. The qualitative nature of the residue in plants and animals is adequately understood based on acceptable metabolism studies with a cereal grain (corn), a root and tuber vegetable (sugar beets), and acceptable poultry and ruminant metabolism studies. The residue of concern in plants and animals is chlorpyrifos *per se*. There are presently no direct application uses of chlorpyrifos on meat- and milk-producing animals, except for ear tag treatment of cattle (beef and lactating and non-lactating dairy).

Residue Analytical Methods - Plants and Animals. The requirements for residue analytical methods are fulfilled for purposes of re-registration. In consideration of HED's decision to regulate only the parent chlorpyrifos, acceptable methods are available for enforcement and data collection purposes. The behavior of chlorpyrifos using FDA's multi residue protocols has also been investigated and reported.

Storage Stability. The requirements for storage stability data are fulfilled for purposes of reregistration. Acceptable storage stability studies have been conducted on representative oil seeds, non-oily grains, root crops, fruits and fruiting vegetables, and low moisture content forage and hay. Additional studies have also been conducted to investigate the frozen stability of chlorpyrifos in selected processed food/feed commodities and in animal tissues and milk.

Magnitude of the Residue. The reregistration requirements for magnitude of the residue in plants (crop field trials and processed food/feed commodities) are fulfilled for the majority of crops. There are minor data gaps for asparagus, corn, cotton, crops grown solely for seed (clover and grasses), mint, peppers, sorghum, tomatoes, tree nut group and wheat. The reregistration requirements for magnitude of the residue in food-handling establishments are fulfilled. Sufficient data exist to determine that when registered formulations are used according to label directions, no detectable residues (<0.01-<0.025 ppm) are likely to occur in food items. Bait and insecticidal strip uses would not result in residues greater than those resulting from spray applications. Therefore, the outstanding data are considered confirmatory.

The reregistration requirements for magnitude of the residue in animals are fulfilled. There are presently no registered direct application uses of chlorpyrifos on livestock animals except for ear tag treatment of cattle (beef and lactating and non-lactating dairy). An acceptable residue transfer study of chlorpyrifos to milk and cream from dairy cows wearing chlorpyrifos-impregnated tags has been submitted; data from this study indicate that residues in whole milk and fat resulting from ear tag use should not be a significant fraction of the residues resulting from intake of animal feeds containing chlorpyrifos. Cattle and poultry feeding studies have been evaluated and found adequate to satisfy feeding study requirements.

Confined/Field Rotational Crops. Provided that the Registrant modifies all labels for its chlorpyrifos containing products to limit application to 5 lb ai/A/season on those crops where rotation to another crop could occur (as was stated in their letter to the Agency dated 8/12/94), HED will not require field rotational crop studies. Furthermore, a 30 day plant back interval for rotational crops would then be appropriate.

### 4.3 Dietary Exposure (Food Source)

As noted previously, chlorpyrifos is registered for use on a wide variety of food crops, and has approximately 112 tolerances for food and/or feed commodities (which translates to approximately 700 food forms in the dietary analysis). Food uses evaluated in this analysis were those reflected by the established tolerances in/on raw agricultural, animal, and processed food/feed commodities for chlorpyrifos as listed in 40 CFR §180.342. Food handling establishment (FHE) tolerances were also included as cited in 40 CFR §185.1000 for the chronic dietary analysis (i.e., as a result of the registered use in FHE, all foods have an established tolerance of 0.1 ppm, unless they are covered by higher tolerances). The tolerances published for chlorpyrifos under 40 CFR §180.342, 185.1000 and 186.1000 have been reassessed (HED Revised Product and Residue Chemistry Chapter, memorandum from S. Knizner to M. Hartman, June 2000). The established tolerances in/on raw agricultural, animal, and processed food/feed commodities are expressed either in terms of the combined residues of chlorpyrifos and its metabolite 3,5,6-trichloro-2-pyridinol (TCP) or as chlorpyrifos *per se*. HED has determined that TCP is not of toxicological concern and concluded that TCP can be excluded from the tolerance expression. Reassessed tolerances are in terms of chlorpyrifos *per se*. Thus, for purposes of this analysis, only residues of chlorpyrifos *per se* were considered, when data were available. Whenever possible, data for anticipated residues (ARs) reflect levels of chlorpyrifos *per se*. HED has conducted a screening-level risk assessment for TCP, which is in the attached memorandum from S. Knizner to D. Smegal, D265035 June 5, 2000.

Highly refined acute and chronic dietary exposure assessments were conducted using the Dietary Exposure and Evaluation Model (DEEM™) system. DEEM can be used to estimate exposure to residues in foods comprising the diets of the U.S. population, including population subgroups. The software contains food consumption data from the USDA Continuing Survey of Food Intake by Individuals (CFSII) from 1989-1992. For chronic dietary risk assessments, the 3-day average of the consumption data for each sub-population is combined with average residues in commodities to determine the average exposure in mg/kg/day. For acute dietary risk assessment, the entire distribution of single day food consumption events is combined with a distribution of residues (probabilistic analysis, referred to as "Monte Carlo") to obtain a distribution of exposures in mg/kg/day.

For chlorpyrifos, inputs to the DEEM analysis include DAS' National Food Survey (NFS, 1993 - 1994), U.S. Department of Agriculture's Pesticide Data Program (PDP) monitoring data (1994-1999), the Food and Drug Administration (FDA) Surveillance Monitoring Program data (1992-1998), and to a much lesser extent, field trial residue data. Percent crop treated data were supplied by the Biological and Economic Analysis Division (Quantitative Usage Analysis for Chlorpyrifos dated 3/30/00). Where percent crop treated estimates indicated no

chlorpyrifos use, a default minimum assumption of 1% crop treated was applied. In general, when residues on commodities were nondetectable, one-half the limit of detection (LOD) was assumed. All available processing and cooking factors were incorporated into the dietary exposure analysis.

At their own initiative, DAS conducted a market basket survey (NFS), with samples collected from the Fall of 1993 to the Fall of 1994, to better determine the dietary exposure of consumers to chlorpyrifos. The results of this survey have been reviewed by HED (L. Cheng, 5/19/98, D217707). Samples of fresh apple, applesauce, apple juice, orange juice, peanut butter, whole milk, ground beef and pork sausage were collected from grocery stores located in the 48 contiguous states; for fresh tomatoes, sampling was conducted in Florida only over a period of 9 months, because the domestic use of chlorpyrifos was restricted to Florida at the time of sampling. Approximately 200 samples were collected for each commodity, except for tomatoes, where 55 samples were collected. The nine food items were selected because of their significant contributions to dietary exposure in general (and in infants and children), and the potential for high residues based on modes of application and the percentage of crop treated. The apple and tomato samples were composite samples consisting of six apples and four tomatoes, respectively.

The Reference Dose (RfD) is derived from an exposure level at which there are no statistically or biologically significant increases in the frequency or severity of adverse effects between the exposed population and its appropriate control, along with the application of uncertainty factors. The percent of the RfD is calculated as the ratio of the exposure value to the RfD ( $\text{exposure/RfD} \times 100 = \% \text{ RfD}$ ). The population adjusted dose (PAD) is the adjusted RfD reflecting the application of the FQPA safety factor. The FQPA safety factor for females and children is 10X, for all other populations subgroups it is 1X. For females and children, the population adjusted doses for acute and chronic dietary risk assessment are 0.0005 mg/kg/day and 0.00003 mg/kg/day, respectively. For all other population subgroups, the population adjusted doses for acute and chronic dietary risk assessment are 0.005 mg/kg/day and 0.0003 mg/kg/day, respectively. Exposures less than 100% of the PAD do not exceed HED's level of concern.

#### **4.3.1 Acute Dietary Exposure Assessment**

The HED probabilistic acute dietary exposure estimates used PDP, and FDA monitoring data to the greatest extent possible, in conjunction with the DAS's NFS data for all commodities included in the survey except apples and tomatoes. NFS data were used for milk, apple juice, applesauce, orange juice, ground beef, pork sausage, and peanut butter. A summary of the acute dietary analysis can be found in the attached memorandum from D. Soderberg to M. Hartman, June, 2000, D263890.

Three data sets are available for estimating residues on fresh apples: PDP data for analysis of individual single apples; PDP "decomposed"

apple data; and NFS “decomposed” apple data. Use of each of these three data sets for fresh apples leads to a different exposure estimate. The dietary exposure analysis has been performed using all commodities having chlorpyrifos uses and each of the apple data sets separately: PDP data for single apples; PDP “decomposed” apple data; and NFS “decomposed” apple data.

In 1999 PDP collected data on residues of chlorpyrifos on individual single apples. A total of 377 single apple samples were analyzed. Of these, 75 (20%) had measurable chlorpyrifos residues, ranging from 0.005 to 0.54 ppm. In an acute exposure analysis, results of analyses on single items of produce for a non-blended food are generally preferable to analyses of composite samples because they can be used without decomposing.

During 1994 - 1997, PDP also collected a total of 1908 composite apple samples, of which 425 samples (22%) had measurable chlorpyrifos residues, ranging from the ½ LOD for each laboratory (average 0.0026 ppm) to 0.4 ppm. Because fresh apples are considered to be a non-blended commodity, these results were decomposed using the Allender method (Allender, H. “Use of the Pesticide Data Program (PDP) in Acute Dietary Assessment”, August 1998) to estimate single serving acute exposure.

DAS also submitted a market basket survey for fresh apples. All composite samples were collected from Fall 1993 - Fall 1994. There were 200 composite samples in this survey. A total of 68 samples (34%) had measurable chlorpyrifos residues, ranging from the LOD of 0.001 to 0.052 ppm.

Other programs have also analyzed fresh apples for chlorpyrifos. The FDA Surveillance Monitoring Program analyzed 1152 fresh apples (composites) between 1993 - 1998. FDA found 151 (13%) samples with measurable residues, ranging from 0.0005 ppm to 0.31 ppm.

FDA Total Diet Study (TDS) data are also available for chlorpyrifos, and in the case of apples these data also support use of the PDP data for risk assessment purposes. Measurable residues of chlorpyrifos (> 0.001 ppm) were found in apples for 14 of the 18 TDS surveys conducted from 1991 to 1997. Residues ranged from less than 0.001 ppm to 0.103 ppm, with a mean value of 0.012 ppm. Samples analyzed in the TDS are purchased at grocery stores and prepared according to standard consumer practices prior to analysis (in the case of apples this means washing). Samples are broadly composited in that composites are formed from samples purchased in three different cities from a given geographic region.

In summation, the maximum residue level found on composite apples in the NFS data is less than the maximum found in all other monitoring



programs, including the TDS, which most closely approximates NFS sampling.

NFS data on fresh tomatoes were submitted. However, only 54 samples were collected and all samples were from FL. More extensive and recent data for fresh tomatoes are available from PDP (881 samples, collected in 1996 and 1997). As was the case for apples, the highest reported detectable residue in the PDP data (0.31 ppm) was greater than that reported in the NFS data (0.0565 ppm). PDP monitoring data also reflect the use of chlorpyrifos on imported fresh tomatoes (a significant source of fresh tomatoes). Therefore the PDP fresh tomato residue data were used exclusively in all analyses. For commercially processed tomato commodities, PDP data were used but data obtained from FL grown tomatoes and fresh imported tomatoes were excluded, as these tomatoes are not used for processing. Appropriate processing residue reduction factors were incorporated for tomato juice, puree, catsup, and paste.

Exposure (consumption x residues) was compared to the acute population adjusted doses (aPAD) of 0.0005 mg/kg/day for children and females and 0.005 mg/kg/day for all other populations. The acute dietary risk analysis estimates the distribution of single day exposures for the overall U.S. population and certain subgroups. The analysis evaluates exposure to the chemical for each food commodity.

Table 3 summarizes the acute probabilistic dietary risk estimates for the U.S. Population and most highly exposed sub-populations. At the 99.9th percentile exposure, risk estimates based on the PDP single apple data, the decomposited PDP apple data, and/or the decomposited NFS apple data, were greater than 100% of the aPAD for the following population subgroups: all infants less than one-year old; children 1-6 years old; and children 7-12 years old. Children 1-6 years old were the most highly exposed population subgroup, regardless of which data set is used for fresh apples. For children 1-6 years old, risk estimates ranged from 170% to 355% of the aPAD depending on which fresh apple data set was used. Use of PDP's 1999 single apple data resulted in the highest exposure estimates. Use of the decomposited NFS fresh apple data resulted in the lowest exposure estimates.

Because the PDP single apple data are the most recent and do not require decompositing, these data are expected to provide the most reliable exposure and risk estimates. However, no matter which of the three data sets is used for fresh apples, the critical exposure commodity (CEC) analysis indicated that residues on fresh apples were the major contributor to dietary exposure estimates for children 1-6 years old at the 99.9th percentile exposure. Residues on whole tomatoes and grapes were the next major contributors to exposure.

Various risk reduction measures were examined to reduce acute dietary exposure and risk estimates. As was previously noted, fresh apples, fresh grapes and fresh tomatoes were the major contributors to acute dietary exposure for children 1-6 years old, the highest exposed subpopulation. Risk estimates could be reduced to less than 100% of the aPAD for children 1-6 years old only with mitigated exposure for all three of these commodities.

To mitigate exposure from fresh apples, the effect of deleting the late season foliar applications was examined. Currently, chlorpyrifos can be applied to apple trees when they are dormant or later in the season as a foliar treatment (up to 8 applications, with 21 days between the final two applications, and a 28 day PHI). In contrast to apples, chlorpyrifos can only be applied to pear trees as a dormant/delayed dormant application. PDP monitoring data are available for analysis of single pears. In the dietary exposure assessment, these data were translated to apples to determine the effect of deleting the apple foliar applications. Using this comparison,

residues on apples as a result of the dormant spray application are expected to be non-detectable (i.e., not expected to exceed 0.01 ppm). As part of risk mitigation, the tolerance for apples will be reassessed at 0.01 ppm, reflecting retention of only the pre-bloom application.

An examination of the PDP monitoring data for fresh grapes indicated that imported samples contained higher residues than domestic grapes. The current domestic use pattern limits application to a directed spray soil treatment to the base of dormant vines. Residues as a result of this application scenario are expected to be non-detectable (i.e., not exceed 0.01 ppm). The higher residues found on imported samples are most likely arising from later season foliar applications. As part of risk mitigation, the tolerance grapes will be reassessed at 0.01 ppm, reflecting the current domestic use pattern.

For tomatoes, PDP monitoring data again indicated that samples containing high residues were from imported fresh tomatoes. Chlorpyrifos is currently registered for use only in Florida (the state with the largest domestic production of fresh tomatoes) and Georgia. Information obtained from grower groups in FL indicates that chlorpyrifos is not used. Therefore, to mitigate dietary exposure the chlorpyrifos use on tomatoes will be deleted (i.e., tolerances revoked).

Based on these mitigation measures, risk estimates for all population subgroups are less than 100% of the aPAD as shown on Table 3. Children 1-6 years old remain the most highly exposed sub-population at 82% of the aPAD.

**Table 3**  
**Summary of Chlorpyrifos Acute Dietary Probabilistic Exposure and Risk Analysis (99.9th percentile)**

| Population Subgroup        | PDP single apple monitoring data from 1999 |            | “decomposed” PDP monitoring results for apples collected from 1994-1997 |            | “decomposed” NFS monitoring results for apples collected from 1993-1994 |            | Assuming Risk Mitigation (apples, tomatoes and grapes) |            |
|----------------------------|--|------------|---|------------|---|------------|--|------------|
|                            | Exposure (mg/kg/day)                       | % aPAD (a) | Exposure (mg/kg/day)  | % aPAD (a) | Exposure (mg/kg/day)  | % aPAD (a) | Exposure (mg/kg/day)                                   | % aPAD (a) |
| US Population              | 0.000790                                   | 16         | 0.000602  | 12         | 0.000453  | 9.1        | 0.000240   | 4.8        |
| All Infants (< 1 year old) | 0.000648                                   | 130        | 0.000548  | 110        | 0.000517  | 100        | 0.000258   | 52         |
| Children 1-6 years old     | 0.001779                                   | 355        | 0.001247  | 250        | 0.000855  | 170        | 0.000410   | 82         |
| Children 7-12 years old    | 0.001288                                   | 258        | 0.000939  | 190        | 0.000607  | 120        | 0.000319   | 64         |
| Females 13-50 years old    | 0.000635                                   | 127        | 0.000484  | 97         | 0.000375  | 75         | 0.000201   | 40         |
| Males 20+ years old        | 0.000580                                   | 12         | 0.000456  | 9.1        | 0.000359  | 7.2        | 0.000205   | 4.1        |

(a) The acute population adjusted dose (aPAD) is 0.0005 mg/kg/day for females and children and 0.005 mg/kg/day for all other sub-populations. Values rounded to two significant figures.

The uncertainties in the acute dietary exposure estimates are discussed below following the chronic dietary exposure assessment discussion.

#### 4.3.2 Chronic Dietary Exposure Assessment

A refined chronic exposure analysis was performed using the DEEM™ exposure modeling software. The input values included the PDP, FDA and DAS' NFS data, in addition to average residues from field trials and percent of the crop treated information from BEAD. All NFS data available were used except for fresh apples and tomatoes, for which PDP monitoring data were used. An additional analysis was conducted using NFS data for apples. Exposure (consumption) was compared to the chronic population adjusted dose (cPAD) of 0.00003 mg/kg/day for females and 0.0003 mg/kg/day for all other subpopulations. A summary of the residue information included in this analysis can be found in the attached memorandum from D. Soderberg to M. Hartman, June, D263889.

As shown in Table 4, for both risk estimates based on PDP or NFS data for fresh apples, the average chronic dietary residue contributions with or without the food handling establishment use are less than 100% of the cPAD and thus do not exceed HED's level of concern. Based on PDP monitoring data for fresh apples, without consideration of the food handling establishment use, the average exposure estimates comprised 3% and 61% of the cPAD for the general population and the most highly exposed subgroup, children 1-6 years old, respectively. The average exposure estimates including the food handling establishment use comprised 4% and 81% of the cPAD for the general population and for the most highly exposed subgroup, children 1-6 years old, respectively.

For the dietary exposure analysis using NFS fresh apple data, dietary risk estimates ranged from 3% to 57% for the general population and children 1-6 years of age, respectively without the food handling establishment tolerance. With food handling establishment tolerances, the dietary risk estimates ranged from 3% to 63% for the general population and children 1-6 years of age, respectively.

The effect of the risk mitigation measures discussed above, on the chronic dietary risk estimates was examined. Based on the mitigation measures (i.e., reduction of apple tolerance to 0.01 ppm based on pre-bloom application, reduction of grape tolerance to 0.01 based on domestic use pattern, and deletion of the use on tomatoes), chronic dietary risk estimates were also reduced, as shown on Table 4. Children 1-6 years old remain the most highly exposed subpopulation, with risk estimates of 51% and 36% of the cPAD, including the FHE use or using zero residues for the FHE use, respectively.

**Table 4  
Summary of Chlorpyrifos Chronic Dietary  
Exposure Analysis(a)**

| Population Subgroup   | Estimate w/PDP Apple Data                |        |  |        | Estimate w/NFS Apple Data                |        |  |        | Assuming Risk Mitigation (apples, tomatoes and grapes) |        |  |        |
|-----------------------|--|--------|--|--------|--|--------|--|--------|--|--------|--|--------|
|                       | Excludes Food Handling Establishment Use |        | Includes Food Handling Establishment Use |        | Excludes Food Handling Establishment Use |        | Includes Food Handling Establishment Use |        | Excludes Food Handling Establishment Use               |        | Includes Food Handling Establishment Use |        |
|                       | Average exposure (Fg/kg BW/day)          | % cPAD | Average Exposure (Fg/kg BW/day)          | % cPAD | Average exposure (Fg/kg BW/day)          | % cPAD | Average Exposure (Fg/kg BW/day)          | % cPAD | Average exposure (Fg/kg BW/day)                        | % cPAD | Average Exposure (mg/kg BW/day)          | % cPAD |
| US Population         | 0.008                                    | 3      | 0.012                                    | 4      | 0.008                                    | 3      | 0.008                                    | 3      | 0.004  | 1.4    | 0.008                                    | 2.5    |
| All infants (< 1 yr)  | 0.007                                    | 23     | 0.014                                    | 45     | 0.007                                    | 24     | 0.008                                    | 28     | 0.003  | 11     | 0.01                                     | 33     |
| Children (1-6 years)  | 0.018                                    | 61     | 0.024                                    | 81     | 0.017                                    | 57     | 0.019                                    | 63     | 0.009  | 31     | 0.015                                    | 51     |
| Children (7-12 years) | 0.013                                    | 45     | 0.018                                    | 59     | 0.012                                    | 41     | 0.014                                    | 46     | 0.006  | 21     | 0.011                                    | 36     |
| Females 13-50 years   | 0.006                                    | 21     | 0.009                                    | 30     | 0.006                                    | 20     | 0.006                                    | 22     | 0.003  | 11     | 0.006                                    | 20     |

(a) Values based on DEEM output, and are based on non-rounded exposure results.

## Uncertainties of Dietary Exposure Estimates

The Agency believes the risk assessment presented is the most refined to date for acute and chronic dietary exposure to chlorpyrifos. However, there are some uncertainties associated with these exposure estimates as follows:

- (a) Residues were detected in PDP over several years for a number of commodities that lack chlorpyrifos tolerances (i.e., chlorpyrifos is not registered for use on these commodities). These include spinach, squash, and carrots as shown below in Table 5:

| <b>Table 5<br/>Commodities with Detected Residues in PDP and Frequently Fed to Children<br/>that Lack Established Chlorpyrifos Tolerances</b> |               |  |  |   |   |
|---|---------------|--|--|---|---|
| <b>Commodity</b>  | <b>Year</b>   | <b># Samples<br/>with<br/>Detections</b> | <b>% Samples<br/>with<br/>detections</b> | <b>Minimum<br/>Residue<br/>Detected<br/>(ppm)</b> | <b>Maximum<br/>Residue<br/>Detected<br/>(ppm)</b> |
| Carrots   | 1994          | 2  | 0.3                                      | 0.005   | 0.005   |
|   | 1995          | 6  | 0.9                                      | 0.005   | 0.019   |
|   | 1996          | 7  | 1.4                                      | 0.005   | 0.074   |
| Spinach   | 1995          | 46                                       | 7.5                                      | 0.005   | 0.11  |
|   | 1996          | 26                                       | 5.0                                      | 0.003   | 0.030   |
|   | 1997          | 11                                       | 2.1                                      | 0.005   | 0.026   |
|   | 1998 (canned) | 4  | 0.6                                      | 0.007   | 0.014   |
| Squash  | 1997          | 4  | 1.8                                      | 0.005   | 0.005   |
|   | 1998          | 6  | 1.1                                      | 0.005   | 0.022   |

Residues were also detected in celery (4 samples in 1994, 0.005 - 0.045 ppm), potatoes (1 sample in 1994, 0.024 ppm), and lettuce (1 sample in 1994 at 0.01 ppm).

The FDA Total Diet Study also contains data indicating that chlorpyrifos residues in/on spinach may occur. Measurable chlorpyrifos residues have been found on cooked spinach in 10 of 18 market basket surveys (56%) conducted from 1991 to 1997.

These residue results were not included in the Agency's dietary exposure assessment as they represent misuse of chlorpyrifos. However, because these violations have occurred over the years,

excluding them might have under-represented potential dietary exposure, especially for infants and children. Therefore, an additional set of dietary exposure assessments have been performed including results for squash, spinach and carrots - three commodities frequently fed to infants and children. Celery, lettuce and potatoes were not included. These additional assessments were not significantly different from the mitigated acute or chronic dietary assessments.

- (b) The consumption database used in the dietary exposure analysis (CSFII, 1989-1992) has a limited number of individuals in the age group infants less than one year old (approximately 100). The USDA is currently conducting the Supplemental Children's Survey (approximately 5000 children).
- (c) The dietary exposure analyses relied primarily on monitoring data obtained either "at the farmgate" in the case of FDA or in regional distribution warehouses for PDP data. The NFS results are for samples obtained at supermarkets, but only represent one year of data. Residues potentially present on items purchased at roadside produce stands or farmer's markets are not represented in this analyses.
- (d) The acute dietary analysis does not include FHE use, in accordance with current policy.
- (e) Potential exposure to chlorpyrifos residues from consumption of fish was not addressed. No tolerances for fish are currently established. In 1992 the Agency's Office of Water (OW) published a report (EPA 1992) that summarized chlorpyrifos residues found in freshwater fish in lakes and rivers at that time. The primary focus of the study was monitoring for dioxin/furan in fish. However, chlorpyrifos residues were detected in 26% of the 388 sites tested, with median, mean, and maximum concentrations of non-detect, 4.09, and 344 ppb respectively. This study indicated that consumption of freshwater fish (i.e., sport fisherman and their families, or others) could contribute to dietary exposure to chlorpyrifos. FDA also has monitored farm-raised fish for chlorpyrifos. Of all fish and crustacean samples tested between 1992 to 1998, FDA found residues of chlorpyrifos in one trout (1994) and twelve catfish (four catfish in each year 1992 - 1994). FDA has found no detectable residues of chlorpyrifos in any farm-raised fish from 1995 to 1998. This is discussed in more detail below.



## Chlorpyrifos Screening-Level Exposures and Risks from Freshwater Fish Consumption

In 1992, the EPA Office of Water (OW) published a report that summarized the chlorpyrifos residues in freshwater fish, and evaluated the health risks to individuals that consume freshwater fish as part of a National Screening Assessment (EPA 1992). The results of the EPA OW Assessment were not included in HED's dietary analysis because of the screening-level nature of this investigation (i.e., limited fish samples collected in areas of chlorpyrifos use, and a greater focus on bottom feeding fish such as carp and white sucker that do not contribute significantly to the diet). Nevertheless, this study indicates that consumption of freshwater fish could also contribute to the dietary exposures and risks of chlorpyrifos for sports fisherman and their families. The results of this assessment are presented below.

In the OW study, game and bottom feeding fish were collected from 388 sites, of which 314 were near point and non point sources of pollution, 39 locations were from the U.S. Geological Survey (USGS) National Stream Quality Accounting Network (NASQAN), and 35 locations represented background levels. The selection of sites was biased toward sites where dioxin/furan concentrations in fish are expected (i.e., near pulp and paper mills and industrial sources), because the original intent of study was to investigate these compounds. Consequently, few of the sites (n=15) investigated were near agricultural areas, where chlorpyrifos use is pervasive.

Chlorpyrifos was detected in fish from 26 percent of the 388 sites, with median, mean and maximum concentrations of non detect, 4.09 and 344 Fg/kg (ppb), respectively. (The second highest concentration was 64.5 Fg/kg). Over 70 percent of the fish concentrations at all sites were below detection. The highest concentrations were observed primarily in bottom feeding fish such as carp near agricultural facilities. The mean concentration from agricultural areas was 24.46 Fg/kg. In general, chlorpyrifos concentrations were detected in whole-body samples of bottom feeders and in fillet samples of game fish at roughly the same average concentration.

Health risks were calculated using fillet samples of game fish collected from 106 sites. Risk estimates were calculated using standard EPA risk assessment procedures, an average fish consumption rate of 6.5 g/day for the U.S. population, daily fish consumption over a lifetime of 70 years, and the chlorpyrifos RfD on EPA's Integrated Risk Information System (IRIS) of  $3 \times 10^{-3}$  mg/kg/day (which is an order of magnitude higher than the RfD developed by HED). The resulting hazard indices associated with ingestion of the maximum and mean chlorpyrifos fillet concentrations were

$2.4 \times 10^{-3}$  and  $6.4 \times 10^{-5}$ , respectively for the U.S. population. These risk estimates are both  $< 1\%$  of the EPA RfD on IRIS, and would represent 24% and  $< 1\%$  of the HED chronic PAD, respectively for chronic consumption of the maximum and mean fillet concentrations. However, it is unlikely that an individual would chronically consume the maximum detected residue of 344 Fg/kg, therefore, it may be more appropriate to compare this dose estimate to the acute PAD than the chronic PAD. In this case, consumption of fish containing 344 Fg/kg reflects only 1.4% of the aPAD.

The potential chlorpyrifos exposures could be higher for Native Americans or other subsistence populations that typically consume more freshwater fish than the general U.S. population. USEPA (1997) reports average and 95<sup>th</sup> percentile fish consumption rates of 70 g/day and 170 g/day, respectively for Native American Subsistence Populations. Consequently, potential exposures and risks could be 11 to 26 times higher than those reported for the general population of sport fisherman and their families. Risk estimates could potentially exceed HED's level of concern if chlorpyrifos fish fillet residues of 344 Fg/kg were ingested daily for 70 years at rates of 70 to 170 g/day. However, subsistence populations are not expected to have exposures or risks that exceed HED's level of concern following chronic ingestion of fish fillets with mean chlorpyrifos concentrations of 4.08 Fg/kg (up to 26% of the aPAD).

### 4.3.3 Drinking Water Exposure

The Environmental Fate and Effects Division (EFED) conducted a drinking water assessment for chlorpyrifos based on an analysis of existing ground and surface water monitoring data in conjunction with conservative Tier 1 and Tier 2 modeling (using GENECC 1.2, PRZM 2.3-EXAMS, and SCI-GROW) (Attached memo from H. Nelson to D. Smegal/M. Hartman, October 6, 1999 and M. Barrett to S. Knizner, November 13, 1998). The drinking water exposure estimates are discussed in greater detail below by water source.

The available environmental fate data suggest that chlorpyrifos has a low potential to leach to groundwater from most typical agricultural uses in measurable quantities, except following termiticide use. Chlorpyrifos is persistent in concentrated applications used in termiticide treatments. The available data indicate that the primary metabolite of chlorpyrifos, 3,5,6-TCP is more mobile, and significantly more persistent in many soils, especially under anaerobic conditions.

Currently, HED uses Drinking Water Levels of Comparison (DWLOCs) as a surrogate to capture risk associated with exposure to pesticides in drinking water. A DWLOC is the concentration of a pesticide in drinking water that would be acceptable as a theoretical upper limit in light

of the total aggregate exposure to that pesticide from food, water, and residential uses. HED uses DWLOCs in the risk assessment process as a surrogate measure of potential exposure associated with pesticide exposure through drinking water. In the absence of reliable monitoring data for a pesticide, the DWLOC is used as a point of comparison against the conservative estimated environmental concentrations (EECs) provided by computer modeling (SCI-GROW, GENECC, PRZM/EXAMS). A DWLOC may vary with drinking water consumption patterns and body weights for specific subpopulations.

HED back-calculates DWLOCs by a two-step process: exposure [food + (if applicable) residential exposure] is subtracted from the PAD to obtain the maximum exposure allowed in drinking water; DWLOCs are then calculated using that value and HED default body weight and drinking water consumption figures. In assessing human health risk, DWLOCs are compared to EECs. When EECs are **greater** than DWLOCs, HED considers the aggregate risk [from food + water + (if applicable) residential exposures] to exceed HED's level of concern.

#### 4.3.3.1 Groundwater Exposure Levels

EFED conducted an analysis of over 3000 filtered groundwater monitoring well data available in U.S. Geological Survey's National Water Quality Assessment (NAWQA) Program databases, and in EFED's Pesticides in Ground Water Data Base (PGWDB). Chlorpyrifos was infrequently detected in groundwater (< 1% of the 3000 wells). The majority of concentrations were reported to be <0.01 Fg/L, with only occasional contamination at a maximum level of 0.026 Fg/L. Although the available monitoring data represent a large part of the U.S., it is not clear that they represent the most vulnerable groundwater where chlorpyrifos is used most intensively. The Pesticides in Ground Water Database (PGWDB) reports a maximum detected concentration of 0.65 Fg/L.

EFED also performed screening-level model estimates of chlorpyrifos concentrations in groundwater using SCI-GROW for four crops (corn, cotton, alfalfa and citrus). The estimated chlorpyrifos concentrations in groundwater using the SCI-GROW screening model range from 0.007 Fg/L (typical application to alfalfa) to 0.103 Fg/L (maximum multiple applications to sweet corn). Therefore, based on an analysis of both monitoring and modeling data, EFED concludes the large majority of the country (>99%) will not have potable groundwater that contains chlorpyrifos at levels greater than 0.1 Fg/L. EFED recommends a range of 0.007 to 0.103 Fg/L as conservative EECs to be used to evaluate both acute and chronic exposures. The

NAWQA monitoring data support that the SCI-GROW modeling estimates are conservative.

Chlorpyrifos use as a termiticide is significant, with a recent estimate of seven million pounds ai applied annually constituting about 30% of the total annual use. Chlorpyrifos groundwater exposure from termiticidal use is highly localized and usually only in wells located within 100 feet of the treatment area. For this use, the maximum detected dissolved concentration is 2090 Fg/L with unknown chronic exposure levels that are presumably significantly lower, but that can persist at detectable levels for at least 6 months. EFED recommends an upper bound range of 30 to 2090 Fg/L to evaluate acute groundwater exposures following termiticide use. The 30 Fg/L represents the concentration that DAS recommends before resuming the use of a contaminated well (i.e., current USEPA Health Advisory for a child), while the 2090 Fg/L concentration represents the maximum detected value. EFED recommends a range of 8.3 to 578 Fg/L to be used to evaluate upper bound chronic groundwater exposures for termiticide use. These values are the acute groundwater termiticide concentrations with adjustments for partial environmental degradation (abiotic hydrolysis at pH 7). DAS states that this exposure only occurs in homes where the well casing has a crack in it, and the well is near or in the foundation. HED has determined that the Label Improvement Process for Termiticides (PR notices 96-7 for termiticides) have reduced the potential for this exposure. For example, reported incidents associated with termiticide use were 28.2 per 100,000 homes in 1997 (pre PR-96-7), and were 8.3 per 100,000 homes in 1998 (post PR-96-7).

#### **4.3.3.2 Surface Water Exposure Levels**

EFED conducted an analysis of over 3000 samples from 20 NAWQA study units for flowing surface water collected from rivers and streams over the last several years. Chlorpyrifos was detected at frequencies up to 15% of 1530 agricultural streams, 26% of 604 urban stream samples in 1997 and in 65% of 57 urban stream samples from Georgia, Alabama and Florida in 1994. The maximum reported dissolved chlorpyrifos concentration in surface water was 0.4 Fg/L, with the majority of detected concentrations < 0.1 Fg/L. EFED notes that although the available monitoring data represent a large part of the U.S., the monitoring data may not represent the most vulnerable watersheds where chlorpyrifos use is pervasive. EFED notes that a limited number of watersheds in the U.S. may have chlorpyrifos concentrations higher than 0.4 Fg/L due to higher usage rates or greater pesticide runoff. In particular, acute exposure levels could be higher for streams draining watersheds with more intense chlorpyrifos use or for lakes and reservoirs for which there are little data.

EFED also performed screening-level model estimates of chlorpyrifos concentrations in surface water such as lakes and reservoirs using Tier I GENECC or Tier II PRZM/EXAMS. Inputs to the models included high exposure agricultural scenarios for major crops (alfalfa, corn, citrus, and tobacco) at the maximum application rates. Estimated maximum 90 day average and peak concentrations of chlorpyrifos in surface water using the PRZM/EXAMS screening model were 6.7 Fg/L and 40.6 Fg/L, respectively. These estimated concentrations should be highly conservative for most surface waters and all drinking water because they are based on a pond draining an adjacent 100% treated field model (it is highly unlikely that 100% of a watershed constituting a major drinking water source would be treated with chlorpyrifos in a given year).

Based on an analysis of the NAWQA monitoring and EFED modeling data, an upper-bound EEC range of 0.026 to 0.4 Fg/L was selected to assess acute risks associated with non-termiticide uses of surface water. The 0.026 Fg/L concentration represents the 95<sup>th</sup> percentile dissolved concentration, while the 0.4 Fg/L concentration is the maximum detected dissolved chlorpyrifos concentration from streams and rivers reported in the first phase of the NAWQA study. The 95<sup>th</sup> percentile concentration of 0.026 Fg/L was used to assess chronic surface water exposures. The Agency concluded that the 0.4 Fg/L estimate (a high acute exposure level for streams) is more reasonable than the conservative PRZM/EXAMS maximum peak EEC of 40.6 Fg/L for lakes and reservoirs. This is because multi-month or annual mean concentrations in a reservoir are expected to be less than the maximum reported concentrations in the flowing water feeding the reservoir. The monitoring data also demonstrate that chronic concentrations of chlorpyrifos are unlikely to exceed 0.1 Fg/L. These estimates only apply to drinking water because residues of lipophilic pesticides, such as chlorpyrifos, bound to sediment and suspended solids could contribute to exposure following consumption of unfiltered water.

#### **4.3.3.3 Drinking Water Exposure Concentrations**

The estimated environmental concentrations (EECs) are shown on Table 6. As noted previously, the groundwater EECs are based on conservative modeling, with support from monitoring data, while the surface water EECs are based on upper-bound levels from monitoring data.

| Table 6<br>ESTIMATED ENVIRONMENTAL CONCENTRATION (EECs)           |                      |            |
|---|----------------------|------------|
| Drinking Water Source   | Concentration (Fg/L) |            |
|   | Acute                | Chronic    |
| Groundwater, except for well contamination<br>SCI-GROW (Fg/L) (a) | 0.007 to 0.103       |            |
| Groundwater as a result of well contamination (Fg/L)              | 30 to 2090           | 8.3 to 578 |
| Surface Water Monitoring Data (Fg/L)                              | 0.026 to 0.4 (b)     | 0.026 (c)  |

- (a) SCI-GROW (Screening Concentration in Ground Water) is an empirical model for predicting pesticide levels in ground water. The value from SCI-GROW is considered an upper bound concentration estimate.
- (b) Based on the 95<sup>th</sup> percentile and maximum detected surface water concentrations.
- (c) Based on the 95<sup>th</sup> percentile surface water concentration from monitoring data

In comparison, the one-day, 10-day, and longer-term USEPA health advisories for a 10-kg child are 30 Fg/L. The lifetime health advisory for a 70-kg adult has been established at 20 Fg/L; the adult longer-term health advisory is 100 Fg/L.

EFED notes that there are significant uncertainties associated with the EECs which are as follows:

- (1) The estimates are intended to be as realistic as possible but apply only to the most vulnerable populations because existing monitoring data imply that the majority of the U.S. population will not be exposed at these levels (for surface water note that the 95<sup>th</sup> percentile estimate is 15 times less than the maximum detected value in monitoring data);
- (2) All of these estimates are for unfinished water, and could be lower in finished drinking water that has received treatment; and
- (3) The exposure estimates are highly conservative (i.e., exceed actual exposure by several-fold) for the majority of the U.S. population, based on the existing monitoring database, which covers a large part of the U.S. However, chlorpyrifos residues in surface waters could be higher in some areas where chlorpyrifos usage is more pervasive in the watershed.

#### 4.3.3.4 DWLOCs for Acute (Drinking Water) Exposure

Acute DWLOCs were not calculated for chlorpyrifos initially because the acute dietary risks alone exceed HED's level of concern based on currently registered uses. Therefore, in effect, the DWLOCs would be zero. However, acute DWLOCs were calculated based on risk mitigation measures that reduce the acute dietary risk estimates to below 100% of the aPAD.

The acute DWLOC values are presented in Table 7. For each population subgroup listed, the acute PAD and the acute dietary (food) exposure (from Table 3) for that subgroup were used to calculate the acute DWLOC for the subgroup, using the formulas in footnotes of Table 7. The EECs are less than the DWLOCs for all populations (highest EEC of 0.4 Fg/L is less than the lowest DWLOC of 0.9 Fg/L), indicating that acute food and drinking water exposures (except possible well contamination) do not exceed HED's level of concern. It should be noted that neither the SCI-GROW model nor the monitoring data reflect concentrations after dilution (from source to treatment to tap) or drinking water treatment.

| <b>Table 7<br/>DWLOCs for Chlorpyrifos Acute Dietary Exposure<br/>Considering Mitigation Measures</b> |                       |                                      |                                     |  |  |                            |
|---|-----------------------|--------------------------------------|-------------------------------------|--|--|----------------------------|
| Population Subgroup (a)   | Acute PAD (Fg/kg/day) | Food Exposure 99.9th (Fg/kg/day) (b) | Max. Water Exposure (Fg/kg/day) (c) | Surface Water (Monitoring Data) (Fg/L) | Ground Water SCI-GROW, (excluding well contamination) (Fg/L) | Acute DWLOC (Fg/L) (d,e,f) |
| U.S. Population   | 5                     | 0.24                                 | 4.76                                | 0.026 to 0.4                           | 0.007 to 0.103   | 166                        |
| All Infants (< 1 Year)  | 0.5                   | 0.258                                | 0.242                               |  |  | 2.4                        |
| Children (1-6 years)  | 0.5                   | 0.410                                | 0.09                                |  |  | 0.9                        |
| Females (13-50 years)   | 0.5                   | 0.201                                | 0.299                               |  |  | 9                          |

- (a) In addition to the U.S. population (all seasons), the most highly exposed subgroup within each of the infants, children, female groups is listed.
- (b) 99.9th percentile exposure. Values are from Table 3 (and rounded).
- (c) Maximum Water Exposure (Fg/kg/day) = Acute PAD (Fg/kg/day) - [Acute Food Exposure (Fg/kg/day)].
- (d) DWLOC (Fg/L) = Maximum water exposure (Fg/kg/day) x body wt (kg) ÷ water consumed daily (L/day)]
- (e) HED default body weights are: general U.S. population, 70 kg; adult females, 60 kg; and infants/children, 10 kg.
- (f) HED default daily drinking water rates are 2 L/day for adults and 1 L/day for children.



Acute exposure to chlorpyrifos in groundwater as a result of well contamination from termiticide use could potentially result in exposures of concern. However, as noted previously, the groundwater exposures from well contamination resulting from termiticide use are highly localized. The implementation of PR 96-7 for termiticides has reduced reported incidents of groundwater contamination resulting from termiticide treatments. For example, reported incidents associated with termiticide use were 28.2 per 100,000 homes in 1997 (pre PR-96-7), and were 8.3 per 100,000 homes in 1998 (post PR-96-7).

#### **4.3.3.5 DWLOCs for Chronic Drinking Water Exposure**

The chronic DWLOC is effectively zero because the long-term residential postapplication risks alone exceed HED's level of concern. However, DWLOCs were calculated based on food (including food handling establishment uses) and water exposure alone. The chronic DWLOC values are presented in Table 8. For each population subgroup listed, the chronic PAD and the chronic dietary (food) exposure (from Table 4) for that subgroup were used to calculate the chronic DWLOC for the subgroup, using the formulas in footnotes of Table 8. As shown, the EEC for surface water (which represents the 95<sup>th</sup> percentile concentration from monitoring data) is less than the DWLOCs, and therefore does not exceed HED's level of concern. It should be noted that neither the SCIGROW model nor the monitoring data reflect actual drinking water concentrations after dilution (from source to tap) or drinking water treatment.

**Table 8  
DWLOCs for Chlorpyrifos Chronic Dietary Exposure  
Includes Mitigation**

| Population Subgroup (a) | Chronic PAD (Fg/kg/day) | Chronic Food Exposure with FHE (Fg/kg/day) (b) | Max. Water Exposure (Fg/kg/day) (c) | Surface Water Monitoring Data (Fg/L) | Ground Water SCI-GROW (excluding well contamination) (Fg/L) | Chronic DWLOC (Fg/L) (d,e,f) |
|-------------------------|-------------------------|--|-------------------------------------|--------------------------------------|---|------------------------------|
| U.S. Population         | 0.3                     | 0.008  | 0.292                               | 0.026                                | 0.007 to 0.103  | 10                           |
| All Infants (< 1 Year)  | 0.03                    | 0.01   | 0.02                                |                                      |   | 0.2                          |
| Children (1-6 years)    | 0.03                    | 0.015  | 0.015                               |                                      |   | 0.15                         |
| Females (13-50 years)   | 0.03                    | 0.006  | 0.024                               |                                      |   | 0.72                         |

- (a) In addition to the U.S. population (all seasons), the most highly exposed subgroup within each of the infants, children, female groups is listed.
- (b) Values are from Table 4 (and rounded).
- (c) Maximum Water Exposure (Fg/kg/day) = Chronic PAD (Fg/kg/day) - [Chronic Food Exposure + Chronic Residential Exposure (Fg/kg/day) (if applicable)]. Chronic residential uses were not considered based on mitigation options.
- (d) DWLOC (Fg/L) = Maximum water exposure (Fg/kg/day) x body wt (kg) ÷ water consumed daily(L/day)]
- (e) HED default body weights are: general U.S. population, 70 kg; adult females, 60 kg; and infants/children, 10 kg.
- (f) HED default daily drinking water rates are 2 L/day for adults and 1 L/day for children.

Long-term exposure to chlorpyrifos as a result of well contamination from termiticide use could potentially result in exposures of concern. However, as noted previously, the groundwater risk estimates from well contamination resulting from termiticide use are highly localized. The implementation of PR 96-7 for termiticides has reduced the reported incidents of groundwater contamination resulting from termiticide treatments.

#### 4.4 Non-Dietary Exposure

Chlorpyrifos is an organophosphate insecticide used extensively in residential settings by both residents and PCOs, and for agricultural use (e.g., citrus, vegetable crops, tree fruits, etc.), greenhouse uses, outdoor ornamental uses, and sodfarm uses. It is one of the top five insecticides used in residential settings. There are approximately 800 registered products containing chlorpyrifos on the market (REFs 9/14/99). Registered uses include a wide variety of food, turf and ornamental plants, as well as indoor products, structural pest control, and in pet collars. It is used in residential and commercial buildings, schools, daycare

centers, hotels, restaurants, hospitals, stores, warehouses, food manufacturing plants and vehicles. In addition, it is used as an adult mosquitocide. In 1998, the DAS estimated that 70% of the urban chlorpyrifos use involved termite control. Approximately 11 million pounds a.i. are applied annually in non-agricultural settings (i.e., residences, schools, golf courses, parks).

Chlorpyrifos, is formulated as a wettable powder packaged in water soluble packets (containing 50% a.i.), emulsifiable concentrates (41.5-47%), dust (containing 0.1-7% a.i.), granular (containing 0.075%-15% a.i.), bait (containing 0.5% a.i.), flowables (containing 30% a.i.), impregnated material (containing 0.5-10% a.i.), pelleted/tableted (containing 0.5-1.0% a.i.), pressurized liquids (0.9-3.8% a.i.), microencapsulated (0.5-20% a.i.) and soluble concentrate/liquids (0.5 to 62.5% ai). Dry flowables and wettable powder in open bags are not supported by the registrant, and therefore, the assessment of these formulation types/packaging is not included in this document. According to DAS, formulations with concentrations greater than one pound a.i. per gallon (approximately 13% a.i.) are sold to licenced pest control or turf and ornamental professionals only. Lower concentrations are available to homeowners from other suppliers for over-the-counter purchase. Except aerosols, granules and dusts, all formulations for application are diluted in water to a concentration of 1 percent a.i. or less (Dow AgroSciences 1998). However, HED is aware of at least one company that sells concentrated chlorpyrifos products (i.e., >13% up to 44.8% ai) to the public on the Internet ([www.ADDR.com/~pestdepo/gizhome.htm](http://www.ADDR.com/~pestdepo/gizhome.htm)) as of March 1, 2000.

Occupational and residential exposures to chlorpyrifos can occur during handling, mixing, loading and applying activities. Occupational postapplication exposure can occur for agricultural workers during scouting, irrigation and harvesting activities. Residential postapplication exposure can occur following treatment of lawns, or residences for cockroaches, carpenter ants, termites, and other insects. In addition, there is a potential for inadvertent oral exposure to children from eating chlorpyrifos-treated turf and soil or hand to mouth activities following contact with treated surfaces or turf. Postapplication exposure to children can occur in locations other than the home, including schools, daycare centers, playgrounds, and parks. There is insufficient use information and exposure data to assess exposure resulting from use in vehicles (i.e., planes, trains, automobiles, buses, boats) and other current label uses such as treatment of indoor exposed wood surfaces, supermarkets, theaters, furniture, and draperies. However, HED has concern for these uses based on the scenarios assessed within this document, and has requested exposure data for all uses of registered products not currently assessed in this document. Although there is concern for these uses, the Agency believes that exposure from these uses will not be higher than the scenarios evaluated in this assessment.

Based on toxicological criteria and potential for exposure, HED has conducted dermal and inhalation exposure assessments for the occupational and residential handlers, occupational postapplication, in addition to residential postapplication dermal, inhalation to adults and children and inadvertent oral exposure to children.

Details of the agricultural and ornamental exposure scenarios are presented in the attached memorandum from T. Leighton to D. Smegal/M. Hartman, D263893, June 2000. Details of the occupational/residential handler assessment for residential settings and the postapplication residential risk assessment are presented in the attached memorandum from D. Smegal/T. Leighton to M. Hartman, D266562, June 2000.

#### **4.4.1 Occupational Handler Exposure Scenarios**

HED has identified 26 major exposure scenarios (resulting in 56 assessments) for which there is potential occupational handler exposure during mixing, loading, and applying products containing chlorpyrifos to agricultural crops and ornamentals (16 scenarios) and to non-agricultural use sites (10 scenarios) such as residential or recreational settings. These occupational scenarios reflect a broad range of application equipment, application methods and use sites. For agricultural uses, application techniques include tractor-drawn equipment, open and closed mixing/loading, and hand held equipment. The application rates used in the assessment are intended to reflect the upper range of rates on the labels. Maximum rates are always included in the assessment to provide a hazard evaluation for those individuals that may use the label as approved by the Agency. In some instances, the rates also include values Dow AgroSciences (DAS) specifically requested to be included as "typical" (e.g., a variety of sod farm rates, corn, citrus, greenhouse, and various nursery rates).

DAS has recently submitted a market survey (Mar-Quest) and the Agency is currently reviewing the results before including additional characterization of chlorpyrifos typical use conditions. HED also included the typical, or median use rates of 1 and 2 lb ai/acre for treatment of surface and subsurface-feeding insects on turf, respectively based on lawn care data submitted by the Registrant and TruGreen/ChemLawn (Jefferson Davis Associates, 1999, TruGreen/ChemLawn 1999). Examples of the application rates used in this assessment include, but are not limited to the following: liquid turf treatment from 1 to 4 lb ai/acre, granular turf treatment at 2 lb ai/acre, vegetable crops range from 1 to 2 lb ai/acre; maximum citrus rate is 6 lb ai/acre; the maximum rates for tree nuts and fruits is 2 lb ai/acre; outdoor ornamental rates for wettable powders are up to 4 lb ai/acre and up to 0.16 lb ai/gallon for liquid formulations; and up to 8 lb ai/acre for fire ant control in sodfarm turf just prior to harvest. The predominant maximum application

rates are defined as those rates which are most frequently cited in the labels and are also believed to be representative of the maximum allowable rates that would not underestimate exposure. Even though an attempt was made to include rates requested by DAS, some of the rates assessed do not necessarily reflect all of the typical rates used on those crops such as the tobacco rate (i.e., only maximum rate of 5 lb ai/A assessed).

The scenarios were classified as short-term (1 to 30 days), intermediate-term (1 to 6 months) and in some cases long-term (greater than 6 months) based primarily on frequency of exposure. The occupational handler scenarios for agricultural use are expected to be of a short-term duration only. It is believed that if there are any agricultural applicators applying chlorpyrifos daily for over a month, those individuals will represent a very small segment of the population. Moreover, those individuals would not be applying the amount of chemical estimated to be handled at the maximum rates in the short-term assessment. On the other hand, several of the LCO/PCO handler scenarios in residential settings (i.e., treatment of homes for insect infestations) were considered to be long-term duration. For the agricultural handlers, the estimated exposures considered personal protective equipment (PPE, which includes a double layer of clothing and gloves and/or a dust/mist respirator), and engineering controls (closed mixing/loading systems for liquids and granulars and enclosed cabs/trucks). Baseline attire (long pants, long sleeved shirt, no gloves) is not presented in this assessment to conserve resources and because of the need for additional PPE and/or engineering controls for all scenarios, and the labels currently require PPE. For LCO/PCO exposure scenarios in residential settings, in most cases only exposures associated with the label-recommended clothing were considered (i.e., scenarios with additional PPE or engineering controls could not be evaluated) based on chemical-specific studies submitted by DAS (many of which include biological monitoring).

#### **4.4.1.1 Occupational Handler Exposure Data Sources and Assumptions**

Multiple chemical-specific handler exposure studies were conducted by the registrant and submitted to the Agency. The handler data collected included biological monitoring of urinary 3,5,6-TCP, the primary metabolite of chlorpyrifos, and passive dosimetry data. These chemical-specific exposure data are used by the Agency to assess the potential handler exposures to chlorpyrifos. However, of the five agricultural monitoring studies submitted by DAS, only two of the studies measured at least 15 replicates (minimum as per the Pesticide Assessment Guideline criteria) of a specific activity (one measuring 15 replicates of both mixer/loader and airblast applicators, the other study measuring 16 replicates of a

combined mixer/loader/applicator for a granular formulation). As for the other three studies, one study measured 13 replicates of an applicator applying chlorpyrifos with various types of high pressure handwands in a greenhouse, 1 replicate of a low pressure handwand, and 2 replicates of a backpack sprayer; the second study measured 9 replicates of an open cab groundboom applicator, 6 replicates of an open mixing/loading EC formulation, and 3 replicates of an open bag WP formulation (open bag WP formulation no longer supported by DAS); and the final study measured 14 replicates of an open mixing/loading of liquids for aerial applicators. Therefore, three of the five DAS studies contain an insufficient number of replicates (as specified by Subdivision U Guidelines) to support the exposure scenarios. Moreover, the total of five agricultural studies submitted by DAS in support of the chlorpyrifos reregistration do not encompass all of the uses of the chemical on the labels nor do they all provide sufficient mitigation (e.g., PPE or engineering controls) to meet an occupational target MOE of 100.

In the absence of applicable chemical-specific data, agricultural handler and LCO/PCO potential exposures resulting from handling and applying chlorpyrifos were estimated using data from the Pesticide Handlers Exposure Database (PHED) Version 1.1 or the Draft Residential SOPs. PHED was designed by a Task Force of representatives from the U.S. EPA, Health Canada, the California Department of Pesticide Regulation, and member companies of the American Crop Protection Association. PHED is a software system consisting of two parts -- a database of measured exposure values for workers involved in the handling of pesticides under actual field conditions and a set of computer algorithms used to subset and statistically summarize the selected data. Currently, the database contains values for over 1,700 monitored individuals (i.e., replicates). PHED's policy is to supplement chemical-specific data with available surrogate data in PHED to increase the sample size (U.S. EPA and HC 1995a - PHED V1.1 Evaluation Guidance). This policy is in effect because individual chemical-specific studies, even when fulfilling the Guideline minimum number of replicates, do not necessarily encompass the variety of equipment in use throughout the country and the large variability of exposures among handlers. While data from PHED provides the best available information on handler exposures, it should be noted that some aspects of the included studies (e.g., duration, acres treated, pounds of active ingredient handled) may not accurately represent labeled uses in all cases.

The PHED data used for the mixer/loader for lawn treatment, and granular bait application (hand, belly grinder and push-type spreader) scenarios in residential settings are representative of the chlorpyrifos uses as the surrogate data were monitored for the same uses.

Potential exposures and internal doses were calculated using unit exposures (i.e., normalized to amount of active ingredient handled -- mg/lb ai handled) from both passive dosimetry and biological monitoring data extrapolated to be representative of the maximum rates on the label (in some instances to typical rates). The normalized exposure data are extrapolated by multiplying by the amount of chlorpyrifos handled per day (i.e., lb ai/day). The amount of chlorpyrifos assumed handled per day was derived from the various application rates and the number of acres (or gallons of spray solution) that could be applied in a single day. Dermal and inhalation margins of exposure (MOEs) are presented separately along with a combined total MOE.

#### **4.4.1.2 Occupational Handler Risk Characterization**

A summary of the short- and intermediate-term risks estimates for PPE and engineering controls is presented in Table 9 for agricultural uses. Table 9 also provides a summary of the range of application rates assessed for chlorpyrifos. Table 10 presents a summary of the short-, intermediate, and long-term risk estimates for LCOs/PCOs at non-agricultural use sites, such as residential and recreational settings.

MOEs for occupational handlers were derived by dividing the appropriate NOAEL, shown on Table 2, by the daily dermal or inhalation exposure estimate. As noted previously, the short-term dermal NOAEL of 5 mg/kg/day is from a dermal rat study, and therefore, no dermal absorption adjustment is necessary. However, both the intermediate- and long-term dermal NOAELs of 0.03 mg/kg/day are based on the weight of evidence from 5 oral toxicity studies in dogs and rats for plasma and red blood cell cholinesterase inhibition, and consequently, dermal exposures were adjusted to absorbed dermal doses using an 3% dermal absorption factor. Inhalation exposure estimates were compared directly to the short- and intermediate-term inhalation NOAEL of 0.1 mg/kg/day, and to the long-term NOAEL of 0.03 mg/kg/day based on the weight of evidence from 5 oral studies in dogs and rats, assuming inhalation absorption is 100% of oral absorption. In evaluating biomonitoring data, which represents total chlorpyrifos exposure via dermal, inhalation and oral exposure, an adjusted absorbed dermal NOAEL of 0.15 mg/kg/day was used (i.e., 5 mg/kg/day \*0.03) to estimate MOEs because most

of the total exposure is from the dermal route. Details of this assumption are presented in the HIARC report (D. Smegal April 6, 2000, HED doc no. 014088). For occupationally exposed workers, MOEs >100 (i.e., 10x for interspecies extrapolation and 10x for intraspecies variability) do not exceed HED's level of concern. MOEs below this level would represent a risk concern. A total dermal and inhalation MOE was also calculated because there is a common dermal and inhalation toxicity endpoint (i.e., cholinesterase inhibition).

### **Agricultural and/or Ornamental/Greenhouse Uses**

The results of the short-term handler assessments as shown on Table 9 indicate that only 1 of the 16 potential exposure scenarios did not provide at least one application rate with a total MOE(s) greater than or equal to 100 at either the maximum PPE (i.e., coveralls over long pants, long sleeved shirts, and chemical resistant gloves while using open systems) or using engineering controls (i.e., closed systems). There are no data, chemical-specific or surrogate, to assess 3 of the 16 scenarios. For specific details and calculations of inhalation, dermal, and total exposures and MOEs see the attached memorandum from T. Leighton to D. Smegal/M. Hartman, D263893, June 2000. In the majority of cases, it is dermal exposure rather than the inhalation exposure driving the total MOEs.

Within the other 12 scenarios, not all of the application rates/crops have MOEs greater than or equal to 100. More specifically, the total dermal and inhalation MOEs for the 12 scenarios evaluated range from 6 to 10,000. In total, 56 iterations of potential exposures and total MOEs were calculated for the various application rates. Based on the maximum level of protection (i.e., various levels of PPE or engineering controls) 2 MOEs are estimated to be less than 10; 6 MOEs are between 10 and 50; 9 MOEs between 50 and 100 and 39 of the MOEs are greater than 100. There are insufficient information (e.g., dermal and inhalation exposure data) to assess the seed treatment uses, dip applications (e.g., preplant peach root and nursery stock), and dry bulk fertilizer applications to citrus orchard floors. These scenarios are of concern given the results from the other scenarios assessed, and HED has requested data for these uses. Fourteen of the scenarios were based on data obtained from five chemical-specific studies submitted by DAS. Of the 14 MOEs calculated using the biological monitoring results, only two reach the target MOE of 100 using PPE. The test subjects' absorbed dose levels indicate the need for additional risk mitigation measures such as closed systems for loading liquids and enclosed cabs for groundboom and airblast applicators. The results and discussion for each of the 16 exposure



scenarios are presented in greater detail in attached memorandum from T. Leighton to D. Smegal/M. Hartman, D263893, June 2000.

The agricultural handler assessments are believed to be reasonable high end representations of chlorpyrifos uses. There are, however, many uncertainties in these assessments. The uncertainties include but are not limited to the following:

- C extrapolating exposure data by the amount of a.i. handled or applied; and
- C not all of the exposure data are of high confidence because of the lack of replicates and/or inadequate QA/QC in the studies.

These uncertainties are inherent in most pesticide exposure assessments. The conservative nature of the assessments, however, are believed to be protective of the handlers.

**Occupational/Non-Agricultural Uses (e.g., Residential/Recreational Settings)**

The following scenarios (by number presented on Table 10) result in total MOEs that exceed HED's level of concern (i.e., MOE less than 100 for LCOs/PCOs):

- (1) Indoor Crack and Crevice Treatment by a PCO;
- (2) Broadcast Turf Treatment by a LCO (intermediate and long-term applicator/ mixer/loader);
- (3) Golf Course Treatments by workers (maximum label rate of 4 lb ai/acre for: mixer/loaders of liquids, and mixer/loaders and applicators for greens and tees) and typical and maximum label rates of 1 and 4 lb ai/acre for groundboom applicators);
- (5) Application of Insecticidal Dust Products by a worker;
- (6) Application of Granular Formulations by a LCO by hand;
- (7) Application of Granular Formulations by a LCO with a belly grinder;

- (8) Application of Granular Formulations by a LCO with push-type spreader;
- (9) Termiticide Treatments for Pre-Construction by a PCO;
- (10) Termiticide Treatments for Post-Construction by a PCO; and
- (13) Mosquitocide mixer/loader or applicator for aerial applications of more than 30 days, even with engineering controls

The following scenario results in a total MOE greater than or equal to 100 that does not exceed HED's level of concern for occupational pesticide handlers in residential settings:

- (2) Mixer/loader of lawn care products wearing PPE (total MOEs 100-820);
- (3) Golf Course Treatments by workers (typical label rate of 1 lb ai/acre for: mixer/loaders of liquid and wettable powders, and mixer/loaders and applicators for greens and tees; maximum label rate of 4 lb ai/acre for mixer/loaders of wettable powders) (total MOEs 100-400),
- (13) Workers who mix/load or apply chlorpyrifos for aerial mosquitocide applications of less than 30 days with the use of engineering controls (closed systems)(total MOEs 160-240); and
- (13) Workers who mix/load or apply chlorpyrifos for ground-based fogger mosquitocide applications up to several months with the use of PPE and/or engineering controls (total MOEs 100-560).

The results of the LCO/PCO handler assessment in residential/recreational settings for short-, intermediate and/or long-term exposure scenarios indicate that most of the MOEs are less than 100, and therefore exceed HED's level of concern. Exposure for four of the scenarios were estimated based on chemical-specific biomonitoring studies submitted by DAS (i.e., indoor crack and crevice treatment, broadcast turf application, and pre- and post-construction termiticide treatment) in which the LCOs/PCOs wore label-specified PPE, or PPE in addition to that specified on the labels. Several of these studies did not represent the maximum label application rates, or only evaluated exposures for a few hours (i.e. 1-3 hours) of the work day, and consequently could underestimate exposures and risks to LCOs/PCOs. Overall, the exposures and

risks for LCOs/PCOs based on the chemical-specific biomonitoring studies are considered to be central tendency estimates because they evaluated less than a full day's exposure at the maximum label rate or they exclude accidental exposure (e.g., exposures resulting from equipment malfunction).

All risk assessments involve the use of assumptions, judgement and available reliable data to varying degrees. Often, the available data are not the ideal data for evaluating potential exposure scenarios. This results in uncertainty in the numerical estimates of risk. Consideration of the uncertainty inherent in the risk assessment process permits better evaluation of the risk assessment and understanding of the human health impacts. Risks estimates may be overestimated or underestimated to varying degrees. Table 10 characterizes the exposure and risk estimates as low-end, central-tendency and high-end based on the assumptions used in the assessment, and identifies the most significant uncertainties.

#### **4.4.2 Occupational Postapplication Exposure Scenarios**

EPA has determined that there is potential exposure to persons entering treated sites (e.g., scouts and harvesters) after application is complete. Postapplication exposure data were required during the chlorpyrifos Data Call In (DCI) of the reregistration process, since, at that time, one or more toxicological criteria had been triggered for chlorpyrifos.

##### **4.4.2.1 Occupational Postapplication Exposure Data and Assumptions**

Multiple chemical-specific postapplication exposure studies were also conducted by the registrant and submitted to the Agency. These studies included biological monitoring and passive dosimetry data, along with dislodgeable foliar residue (DFR) data. Data were submitted by DAS for sugar beets, cotton, sweet corn, almonds, pecans, apples, citrus, cauliflower, and tomatoes. The residue decline for these crops indicate that chlorpyrifos quickly dissipates in the first few days after application and then the decline is more subtle. For instance, in most of the crops monitored, the half life of chlorpyrifos for the first part of the curve [i.e., 0 to 7 days after treatment (DAT)] is less than 1 day. However, the second part of the decline curve exhibits a half life of more than 10 days using data from sampling intervals of 7 up to 43 days after treatment (DAT). Based on the initial rapid dissipation of chlorpyrifos as shown in the DFR studies, most of the crops were analyzed using the first part of the decline curve for the short-term endpoint (i.e., up to 1 month) to

establish the restricted-entry interval (REI). The second part of the decline curve was used to assess the intermediate-term duration to assure that workers exposed in treated fields for 1 to 6 months are adequately protected. If the intermediate-term MOEs at the initially assessed short-term REI were less than 100, then the intermediate-term MOEs were used to determine the appropriate length of the REI.

Specific transfer coefficients were also monitored and submitted for citrus harvesting, citrus tree pruning, cauliflower scouting, and tomato scouting. Additional transfer coefficients for other crops/activities are currently being researched by the Agricultural Reentry Task Force (ARTF). In the mean time, HED's standard values for transfer coefficients are used to estimate potential reentry exposure because the ARTF data are not available. Once available, the ARTF data may impact the REIs for tree nuts, tree fruits, and cauliflower. In addition, chemical-specific DFR data are not available for all crops that are potentially treated with chlorpyrifos. Therefore, the assessment of postapplication exposures in this document is based on a grouping of activities associated with various representative crops. The potential for dermal contact during postapplication activities (e.g., harvesting) is assessed using a matrix of potential dermal contact rates by activity and associated crops with groupings of "low", "medium", and "high". In addition to this matrix, citrus, cauliflower, tree nuts and tree fruits are assessed separately. Table 11 summarizes the crops characterized as "low", "medium", and "high".

Maintenance workers and mowers for golf courses were also considered in this assessment and were considered to contact treated turf the day of treatment for short-term durations (i.e., less than 30 days). Although the golf course workers may be working up to 12 months a year, chlorpyrifos levels on the turf will not be available for an appreciable length of time (e.g., residues declining, irrigation, mowing of the turf).

#### **4.4.2.2 Occupational Postapplication Risk Characterization**

The results of the short- and intermediate-term postapplication assessments indicate that REIs need to be established. The REIs are presented on Tables 12 and 13. The REIs range from 24 hours for the crop grouping matrix to 10 days for harvesting cauliflower. In short, REIs are 24 hours for all crops except the following: cauliflower (10 days), all nut trees (2 days), all fruit trees (4 days) and citrus (5 days). The timing of the applications are noteworthy because most

of the applications to trees are to the bark during the dormant to early season. There is insufficient information (e.g., timing of applications - - dormant/bark versus foliar treatments) and exposure data to assess postapplication activities for ornamental and soil incorporated uses. The data needed to assess these areas include ornamental dislodgeable foliar residues in greenhouses and biological monitoring data for reentry into areas with soil directed applications. Details of this assessment are presented in memorandum from T. Leighton to D. Smegal/M. Hartman, June 2000, D263893.

Postapplication risks to golf course workers during mow/maintenance activities are presented on Table 14. The short-term MOEs are above 100 (MOE 110 to 210) and therefore, do not exceed HED's level of concern, even at the maximum label rate of 4 lb ai/acre. These risks are conservative because they assume contact with golf course turf the day of treatment.

The occupational postapplication assessments are believed to be reasonable high end representations of chlorpyrifos uses. There are, however, many uncertainties in these assessments. The uncertainties include but are not limited to the following:

- C extrapolating exposure and DFR data by the amount of active ingredient handled or applied;
- C not all of the exposure data are of high confidence because of the lack of replicates and/or inadequate QA/QC in the studies;
- C translating crop-specific DFR data to assess other crops; and
- C application timing in comparison to actual potential postapplication exposure scenarios.

These uncertainties are inherent in most pesticide exposure assessments. The conservative nature of the assessments, however, are believed to be protective of the worker.

#### 4.4.3 Residential Handler Exposure

Potential chlorpyrifos residential handler exposures can result from treatment of turf and ornamental plants, as well as indoor use (i.e., for cockroaches, carpenter ants, etc), and structural pest control (i.e., termites). Residential handler exposures to chlorpyrifos can occur via dermal and inhalation routes during handling, mixing, loading and applying activities. All residential handler exposure durations are classified as short-term (1-30 days). As noted previously, in 1997 DAS agreed to work with EPA in

limiting household consumer use to only products packaged as ready-to-use in order to minimize exposure to concentrates that require mixing.

#### **4.4.3.1 Residential Handler Exposure Scenarios**

EPA has determined that there is potential exposure to residents during application of chlorpyrifos products. Based on residential use patterns, nine major residential/non-occupational exposure scenarios (by number presented on Table 10) were identified and evaluated for chlorpyrifos:

- (1) indoor crack and crevice treatment using an aerosol can;
- (2) broadcast turf mixing/loading/application using either a hose end sprayer or a low pressure hand wand;
- (4) application of a 0.5% ready-to-use formulated product in a screw top bottle;
- (5) application of an insecticidal dust product using a shaker can or bulbous duster;
- (6) application of granular formulation by hand;
- (7) application of granular formulation with a belly grinder;
- (8) application of granular formulation with a push-type spreader;
- (11) paintbrush application to wood for an insect infestation; and
- (12) treatment of ornamentals (mixing/loading/application) using a low pressure hand wand.

#### 4.4.3.2 Residential Handler Exposure Data Sources and Assumptions

For most cases, residential handler exposure assessments were completed by HED assuming an exposure scenario for residents wearing the following attire: short-sleeved shirt, short pants, shoes and socks, and no gloves or respirator. The only exception is the application of a ready-to-use formulated product, which was evaluated based on a chemical-specific biomonitoring study in which the volunteers wore long pants. Daily unit exposure values were obtained from the Draft Standard Operating Procedures (SOPs) for Residential Exposure Assessments (December 1997) or PHED. Eight of the nine scenarios were evaluated based on data obtained from PHED.

For broadcast turf application, the area treated per day was assumed to be 0.5 acre for hose end sprayer and 1000 ft<sup>2</sup> for spot treatment using a low pressure hand wand or hand application of a granular formulation. Recent lawn size survey data suggest that up to 0.5 acre lawn size represents 73% of 2300 respondents, while nearly 16% of the respondents had lawn sizes that ranged from 0.57 to 1 acre (Outdoor Residential Use and Usage Survey and National Gardening Association Survey 1999). For application of the granular formulation with a belly grinder or push-type spreader, it was assumed that an average of 0.97 lbs active ingredient was handled (i.e., 0.5 acre at 2 lb ai/acre), based on a chemical-specific study of a granular formulated product and the average of 55 replicates from the studies cited in PHED for this use pattern. For a number of scenarios, multiple evaluations were conducted using application rates less than the maximum label rate, or application using different equipment or methods (i.e., ornamental treatment via low pressure hand wand and hose-end sprayer, and granular application via hand, belly grinder and push-type spreader) to assist in risk mitigation and management decisions.

#### 4.4.3.3 Residential Handler Risk Characterization

A summary of the short-term risk estimates, method of evaluation and risk characterization/uncertainties for residential handlers is presented on Table 10. MOEs for residential handlers were derived by dividing the appropriate short-term NOAEL, shown on Table 2, by the daily short-term dermal or inhalation exposure estimate. As noted previously, the short-term dermal NOAEL of 5 mg/kg/day is from a dermal rat study, and therefore, no dermal absorption adjustment is necessary. For inhalation, the short-term

NOAEL is 0.1 mg/kg/day based on two inhalation studies conducted in rats. Evaluation of adult biomonitoring data was conducted two ways, first the total chlorpyrifos dose was compared to an adjusted dermal NOAEL of 0.15 mg/kg/day (i.e., 5 mg/kg/day \* 0.03 dermal absorption), because based on available data the majority of exposure is via the dermal route. In addition, HED segregated the total biomonitoring dose into dermal, inhalation, and oral, for comparison with the route-specific toxicity endpoints.

For residential applicators, MOEs > 1000 (i.e., 10x for interspecies extrapolation, 10x for intraspecies variability and 10x for the FQPA factor) do not exceed HED's level of concern. MOEs below this level would represent a risk concern. A total dermal and inhalation MOE was also calculated because there is a common dermal and inhalation toxicity endpoint (i.e., cholinesterase inhibition).

The results of the residential handler assessment for short-term exposure scenarios indicate that all nine scenarios evaluated have total dermal and inhalation MOEs that exceed HED's level of concern defined by a target MOE of 1000. The residential handler MOEs ranged from 3 to 900 for dermal risk, from 120 to 57,000 for inhalation risk, and from 3 to 880 for total dermal and inhalation risk for the maximum, typical and even minimum label-recommended application rates. Dermal exposure contributes most to total exposure. For a number of scenarios, multiple evaluations were conducted using application rates less than the maximum label rate, or application using different equipment or methods (i.e., ornamental treatment via low pressure hand wand and hose-end sprayer, and granular application via hand, belly grinder and push-type spreader, spot treatment for crack and crevice). These additional analyses were conducted to provide information for risk mitigation and management decisions. The following scenarios (by scenario number shown in Table 10) result in total MOEs that exceed HED's level of concern (i.e., MOE < 1000) for the typical and/or maximum application rate:

- (1) indoor crack and crevice treatment using an aerosol can;
- (2) broadcast turf mixing/loading and application using either a hose end sprayer or a low pressure hand wand (spot treatment);
- (4) Application of a 0.5% ready to use formulated product in a screw top bottle;
- (5) application of an insecticidal dust product using a shaker can



or bulbous duster;

- (6) application of granular formulation by hand;
- (7) application of granular formulation with a belly grinder;
- (8) application of granular formulation with a push-type spreader;
- (11) paintbrush application to wood for an insect infestation; and
- (12) mixing/loading and treatment of ornamentals using a low pressure hand wand.

As noted previously, all risk assessments involve the use of assumptions, judgement and available reliable data to varying degrees. Often, the available data are not the ideal data for evaluating potential exposure scenarios. This results in uncertainty in the numerical estimates of risk. Consideration of the uncertainty inherent in the risk assessment process permits better evaluation of the risk assessment and understanding of the possible human health impacts. Risks estimates may be overestimated or underestimated to varying degrees. Table 10 characterizes the exposure and risk estimates as low-end, central-tendency and high-end based on the assumptions used in the assessment, and identifies the most significant uncertainties.

#### **4.4.4 Residential/Recreational Postapplication Exposures and Risks**

EPA has determined that there are potential postapplication exposures to residents/individuals entering treated areas both indoors following residential/commercial/institutional treatment (i.e., homes, schools, day care centers, etc) for cockroaches, termites or other insects and outdoors following turf treatment (i.e., homes, schools, parks, playgrounds, ball fields, etc) or mosquitocide use. In addition, there is a potential for inadvertent oral exposure to children from eating chlorpyrifos-treated soil, grass and/or granules, or placing their fingers in their mouths. For residential postapplication activities, the exposure duration is expected to be short-, intermediate- and long-term (1 days to several years) depending on the scenario. Adolescent and adult golfers were considered to contact treated turf the day of treatment for short-term durations (i.e., less than 30 days). Details of this assessment are presented in a memorandum from D. Smegal/T. Leighton to M. Hartman, June 2000, D266562.

#### 4.4.4.1 Postapplication Exposure Scenarios

HED identified a total of eleven scenarios likely to result in postapplication exposures to residents/recreational users, and quantitatively evaluated the following ten scenarios:

- (1) Indoor Crack and Crevice Treatment of kitchen and bathroom (inhalation exposure in treated room);
- (2) Indoor Crack and Crevice Treatment of other rooms (dermal and oral exposure from deposition in untreated room based on registrant data);
- (3) Pet Collar Products;
- (4) Termiticide Treatments for Basement, Plenum, Slab and Crawlspace Construction Homes;
- (6) Broadcast Lawn Treatment Using a Liquid Spray;
- (7) Broadcast Lawn Treatment Using a Granular Formulation;
- (8) Golf Course Exposure (adolescent and adult golfer);
- (9) Aerial and ground-based fogger adult mosquitocide application;
- (10) Yard and Ornamental Spray Products, and
- (11) Perimeter treatment of residence.

An additional scenario, insecticidal dust product use (scenario 5) was considered, but could not be quantitatively evaluated due to an absence of chemical-specific information and residential SOPs. HED requests exposure data for this, as well as all other scenarios not evaluated.

HED is in the process of revising the Residential Exposure Assessment SOPs. This process may identify specific areas of further concern with respect to chlorpyrifos and exposure to the general population. For example, some of the secondary exposure pathways that EPA is currently examining include exposures resulting from residue tracked into homes from outdoor use, indoor dust, and spray drift. In a recent study, polycyclic aromatic hydrocarbons (PAHs) that are abundant in house dust were shown to increase the toxicity of chlorpyrifos *in vitro*, particularly at low levels (i.e., 2-50 FM

PAHs with 1-180 nM chlorpyrifos-oxon, a metabolite of chlorpyrifos that inhibits acetyl cholinesterase) (Jett et al. 1999). Currently, there are no SOPs available to evaluate these potential exposure pathways. These scenarios however, may be evaluated in the future pending revisions to the residential SOPs.

#### **4.4.4.2 Data Sources and Assumptions for Postapplication Exposure Calculations**

HED evaluated four of the eleven residential postapplication exposures scenarios based on chemical-specific studies submitted by DAS (i.e., crack and crevice treatment of the kitchen and bathroom (1), broadcast treatment of turf with chlorpyrifos spray (6) and granules (7), and termiticide treatment (4)). Three of these studies (crack and crevice, and two lawn studies) included biomonitoring of the urinary metabolite 3,5,6-TCP, in addition to environmental measurements to quantify chlorpyrifos exposures. In the absence of chemical-specific data, the other exposures (scenarios 2, 3, 8, 9 and 11) were evaluated using the equations and assumptions presented in the Draft SOPs for Residential Exposure Assessments guidance document or revised assumptions from the SOPs to be released in 2000 (i.e., indoor crack and crevice treatment of other rooms, mosquitoicide uses, golfer exposures, pet collar uses and perimeter treatments), which are generally considered to result in high-end exposure estimates, except for the crack and crevice treatment. Scientific literature studies, the AgDrift Model and assumptions from the updated and Draft Residential SOPs were used to evaluate adult mosquitoicide uses.

#### **4.4.4.3 Residential/Recreational Postapplication Risk Characterization**

A summary of the postapplication risk estimates, method of evaluation, and risk characterization/ uncertainties is presented in Table 15. MOEs for residential/recreational postapplication exposures were derived by dividing the appropriate NOAEL, shown on Table 2, by the daily dermal, inhalation or oral exposure estimate. As noted previously, biomonitoring data was evaluated two ways, first the total chlorpyrifos dose was compared to an adjusted dermal NOAEL of 0.15 mg/kg/day (i.e., 5 mg/kg/day \* 0.03 dermal absorption), because the majority of exposure is via the dermal route. In addition, because there is no scientifically valid method to extrapolate from adult biomonitoring data to child exposure, HED segregated the total biomonitoring dose into dermal, inhalation, and oral exposure estimates, for comparison with the route-specific

toxicity endpoints. This extrapolation was conducted only for the post application exposures from lawn treatment. For residents, the acceptable MOE is 1000 (i.e., 10x for interspecies extrapolation, 10x for intraspecies variability and 10x for the FQPA factor). MOEs below this level would represent a risk estimate of concern for the Agency. A total dermal and inhalation MOE was also calculated because there is a common dermal and inhalation toxicity endpoint (i.e., cholinesterase inhibition). For child exposures, oral exposure also contributed to the total MOE. The following scenarios result in MOEs less than 1000, or potential exposures that exceed HED's level of concern:

- (1,2) Indoor Crack and Crevice Treatment of kitchen and bathroom (inhalation exposure in treated room, dermal and oral exposure in untreated room);
- (3) Pet Collar Products;
- (4) Termiticide Treatments for Crawlspace, Basement, Plenum and Slab Construction Homes;
- (6) Broadcast Turf Treatment Using a Liquid Spray;
- (7) Broadcast Turf Treatment Using Granular Formulation;
- (8) Golf Course Exposure (adolescent and adult golfer) following treatment at the maximum rate of 4 lb ai/acre, and
- (11) Perimeter Treatments of Residences.

In addition, by analogy, HED evaluated yard and ornamental spray products (Scenario 10) and concluded that these products result in comparable doses and short-term MOEs with the lawn care products based on label uses and application rates. Therefore, use of many of these products is likely to result in MOEs that exceed HEDs level of concern.

The following scenarios result in MOEs greater than 1000 that do not exceed HED's level of concern for post-application residential/recreational exposures:

- (8) Golf Course Use (adolescent and adult golfer) following treatment at the typical rate of 1 lb ai/acre; and
- (9) Aerial and ground-based fogger adult mosquitocide application.

In conclusion, seven of the nine scenarios evaluated quantitatively have MOEs that are less than 1000, and therefore exceed HED's level of concern. In addition, for post application exposure to children following perimeter applications to homes, it was estimated that more than seven hand-to-mouth events or more than 8 minutes of play on treated turf the day of treatment could result in potential exposures that could exceed the Agency's level of concern (i.e., MOE < 1000). Total MOEs for the residential postapplication exposures that exceed HED's level of concern ranged from 6 to 980. The only postapplication scenario that resulted in a MOE consistently above 1000 was from the aerial and ground-based fogger adult mosquitocide applications (MOEs are 17,000 and 29,000 for children and adults, respectively). In addition, MOEs for adolescent and adult golfers are above 1000 following treatment of golf courses at the typical, or median rate of 1 lb ai/acre (MOEs 1500-2400). A summary of the termiticide postapplication exposure and risk estimates is presented in greater detail below.

As noted previously, all risk assessments involve the use of assumptions, judgement and available reliable data to varying degrees. Often, the available data are not the ideal data for evaluating potential exposure scenarios. This results in uncertainty in the numerical estimates of risk. Consideration of the uncertainty inherent in the risk assessment process permits better evaluation of the risk assessment and understanding of the possible human health impacts. Risks estimates may be overestimated or underestimated to varying degrees. Table 15 characterizes the exposure and risk estimates as low-end, central-tendency and high-end based on the assumptions used in the assessment, and identifies the most significant uncertainties. As noted on Table 15, the exposure and risk estimates based on the chemical-specific studies are generally considered to be reasonable central-tendency estimates (i.e., arithmetic mean, or median exposure was used to calculate risk). Because three of the chemical-specific studies were conducted in adults, conservative assumptions were used to estimate child exposures. However, because adult activity patterns differ from children, i.e., hand-to-mouth activity, some of the registrant-submitted chemical-specific studies could under-estimate a child's exposure (e.g., lawn studies are not designed to reflect any potential for incidental ingestion of residues from treated turf, soil and/or granules).

An additional scenario, postapplication exposures associated with insecticidal dust product use (scenario 5) could not be quantitatively evaluated due to an absence of chemical-specific data or recommended procedures in the Residential SOPs. Nevertheless,

HED has concerns about the use of these products based on the low MOEs calculated for residents or workers that could apply dust products. HED recommends that the registrant provide additional information on the potential post-application residential exposures associated with dust products.

HED identified a number of data gaps for assessing post application exposure, and these data gaps are discussed in Section 6.0.

HED has concerns for the potential for children's exposure in the home as a result of residential and/or agricultural uses of chlorpyrifos. Environmental concentrations of chlorpyrifos in homes may result from residential uses, spray drift, track-in, or from redistribution of residues brought home on the clothing of farm workers or pesticide applicators. Potential routes of exposure for children may include incidental ingestion and dermal contact with residues on carpets/hard surfaces, in addition to inhalation of vapor and airborne particulates. There are several literature studies that quantify the levels of chlorpyrifos in household dust, indoor and outdoor air, dermal wipe (hands) and soil samples. These residues may persist and the resulting exposures are of a potential chronic nature. Currently, there are no SOPs available to evaluate potential exposures from spray drift and track-in. The Agency is currently in the process of revising its guidance for completing these types of assessments. Modifications to this assessment shall be incorporated as updated guidance becomes available. This will include expanding the scope of the residential exposure assessments by developing guidance for characterizing exposures from other sources already not addressed such as from spray drift; residential residue track-in; and exposures to farm worker children.

### **Termiticide Risk Characterization and Uncertainty Analysis**

Because of chlorpyrifos' extensive use as a termiticide, HED has provided a detailed summary of the risks and uncertainties associated with termiticide treatments. The Agency conducted an assessment of termiticide postapplication risks based on a chemical-specific exposure study submitted by DAS. This study collected air measurements from the basement, kitchen and bedroom of 31 homes for up to 1 year following a termiticide treatment. Four types of housing structures were evaluated: basement, plenum, slab and crawlspace. Chlorpyrifos was applied according to the label-recommended rate of approximately 1% active ingredient.

The Agency calculated incremental time-weighted average (TWA) air concentrations for the entire house, assuming an individual could be in any room. Based on this assessment, risks from inhalation exposure was the primary concern. Based on the mitigation plan, the TWA concentrations were normalized to a reduced application rate of 0.5% ai. As part of risk characterization, the Agency evaluated risks for both intermediate and chronic exposures because of uncertainties in the toxicity endpoints for both durations. Details of this analysis are presented in the Occupational/Residential Handler and Post-Application Residential/Non-Occupational Risk Assessment (memo from D. Smegal/T. Leighton, June 2000, D266562). The MOEs are presented on Table 15.

Similar to the dietary assessment, children 1-6 years of age have higher potential exposures than adults, primarily because of to a higher breathing rate per body weight, and data that indicate young children spend more time at home than adults. For children, all the 90-day median MOEs are greater than 1000 (median MOEs range from 1,900 to 3,800), and therefore do not exceed HED's level of concern. However, some of the 1-year median MOEs are below 1000, and therefore exceed HED's level of concern (median MOEs range from 530 to 1,100). As shown on Table 15, the lowest 90-day and 1-year MOEs for an individual house are 440 and 270, respectively.

The median MOEs for adults were greater than 1000 for all housing types for both the 90-day and 1-year analysis, and therefore, do not exceed the Agency's level of concern (MOEs range from 1,800 to 13,000).

There are however, a number of uncertainties in the risk assessment that arise from the following sources: choice of toxicological data used to establish the inhalation toxicity endpoint, chlorpyrifos air concentrations, and exposure assumptions. The most significant uncertainties will be discussed below.

Toxicity Endpoints: There are uncertainties associated with both the intermediate and long-term inhalation NOAELs used to calculate the MOEs. The intermediate-term NOAEL of 0.1 mg/kg/day is based on two 90-day inhalation studies, in which the rats were exposed 6 hours/day, 5 days/week (nose-only) to the highest attainable vapor concentration of chlorpyrifos (287 Fg/m<sup>3</sup>). HED could not identify an inhalation LOAEL because no adverse effects were noted at the highest dose tested. Therefore, HED selected an oral LOAEL of 0.3 mg/kg/day to use in the dose-response



assessment. The 3 fold difference between the NOAEL and LOAEL, adds an extra buffer of safety to the intermediate-term inhalation endpoint for a total MOE of at least 3000. Although the inhalation route of exposure is ideal for this assessment, the exposure regimen does not fully mimic the potentially continuous inhalation exposure for children associated with a termiticide treatment (i.e., up to 20 hours/day).

The long-term NOAEL of 0.03 mg/kg/day is based on oral animal studies that observed cholinesterase inhibition at 0.2 to 0.3 mg/kg/day (the LOAEL). HED notes that the large difference between the NOAEL and LOAEL (i.e., factor of 6.7 to 10), adds an extra buffer of safety to the long-term inhalation endpoint. Therefore, relative to the LOAEL, the MOE is actually at least 6,000 to 10,000 for a target MOE of 1000. In addition, there are significant uncertainties associated with route-to-route extrapolation due to differences in pharmacokinetics. Following oral exposure, chlorpyrifos is absorbed in the gastrointestinal tract and is transported to the liver, where it can undergo biotransformation to a potent cholinesterase inhibitor (chlorpyrifos-oxon), and be further detoxified. However, following inhalation exposure, chlorpyrifos is absorbed directly into the systemic circulation and initially bypasses the liver. These pharmacokinetic differences may play an important role in the route-specific toxicity of chlorpyrifos. In the absence of inhalation pharmacokinetic data, it is difficult to predict whether use of an oral NOAEL would over- or under-estimate inhalation risks.

Air Concentrations: There are also a number of uncertainties associated with the chlorpyrifos air concentrations used to assess termiticide risks, which affect both the 90 day and 1 year MOEs calculations. Measured chlorpyrifos air concentrations may be overestimated because of use of other chlorpyrifos-containing products. For example, more than half (55% or 17/31) of the homes in the DAS study had detectable chlorpyrifos air concentrations prior to termiticide treatment, indicating that residents may have used other chlorpyrifos products in the home, or had a previous chlorpyrifos termiticide treatment. Several studies in the scientific literature reported chlorpyrifos air concentrations up to 8 years following termiticide treatments (Wright et al. 1988, 1994). However, these studies did not control for use of other chlorpyrifos products (i.e., lawn treatment, flea control, or other indoor uses, etc) (personal communication by D. Smegal with G. Dupree 5/17/2000), and therefore, may also overestimate potential exposures and risks.

In addition, spills inside the home can contribute to higher airborne concentrations of chlorpyrifos. In the DAS study, one of the

homes had elevated basement air concentrations because of a spill. The elevated basement measurements were excluded from the analysis (i.e., only kitchen and bedroom air data were used). This is considered reasonable because spills are likely to be an infrequent occurrence, and because pest control operators (PCOs) are trained to promptly clean spills that occur during application. However, possible applicator error, unreported, undetected or unremediated spills can contribute to air concentration measurements.

The available data suggest that temperature influences indoor chlorpyrifos concentrations resulting from termiticide treatments (i.e., warmer temperatures are associated with higher concentrations). In the DAS study, 26 of 31 homes were from the South or warm climates. Therefore, it is possible that the air concentrations used in this assessment represent high-end estimates, that could overestimate exposures for treated houses in more temperate climates.

There are uncertainties associated with the incremental TWAs air concentration calculations. Based on the mitigation plan, HED calculated the incremental TWAs by adjusting the air measurements associated with a 0.7-1% ai product application to 0.5% assuming that there is a linear relationship between percent ai and resulting air concentrations. This assumption is considered reasonable, although it could under- or over-estimate the air concentrations associated with 0.5% a.i. product application. In addition, the 1-year incremental TWA concentration may be overestimated for two basement homes, because one year air concentration measurements were not available. HED assumed the 90 day air concentration remained constant from 90 to 365 days. This assumption only impacts two basement homes (B1 and B2), both of which had 1 year MOEs less than 1000, but 90 day MOEs greater than 1000.

Exposure Assumptions. The assumptions used to estimate exposures are based on USEPA recommended values (Exposure Factors Handbook), and are designed to be conservative for the majority of the population. These estimates could be conservative for children that do not spend their entire day at home (i.e., those that attend day-care, pre-school, and/or school). This assessment assumed that children aged 1-6 years are exposed to chlorpyrifos air concentrations in a treated home for 20 hours/day, 7 days/week, for up to 1 year.

Summary: In summary, HED believes that individuals are unlikely to experience adverse health effects from termiticide use of chlorpyrifos, even though a few of the child MOEs are below 1000.

Based on the uncertainties described above, the 90 day risk estimates may be underestimated, while the 1 year risk estimates may be overestimated. Overall, HED believes that the risk estimates are bounded by the ranges presented in Table 15. As shown on Table 15, the lowest 90-day and 1-year MOEs for an individual house are 440 and 270, respectively and the highest estimates are 13,000 and 9,500, respectively. Although some MOEs are less than 1000, there is an additional 3 to 10 fold buffer because of the difference between the NOAEL and the LOAELs. In addition, a number of conservative assumptions were incorporated into these MOEs, such as assuming that all children spend 20 hours/day, 7 days/week for up to 1 year in a treated home.

Mitigation measures will further reduce exposures and risk. For example, the removal of whole house barrier treatment addressed the exposures of most concern. It is expected that the limited spot and localized treatment, and pre-construction treatments would represent less exposure and risk. Based on the mitigation plan, and best professional and scientific judgement, HED concludes that the termiticide risk does not raise a concern and that individuals are unlikely to experience adverse health effects from termiticide treatments conducted according to the label. This conclusion is based on the conservative assumptions, the risk mitigation measures, coupled with the uncertainties of the toxicity endpoints and the air measurements.

**Table 9  
Exposure Variables and MOEs for Agricultural Uses  
(Including Non Worker Protection Standard Ornamental Uses) of Chlorpyrifos**

| Exposure Scenario (Scenario#)                                  | Are Biological Monitoring Data Available? (a) | Application Rates (lb ai/acre) (b)        | Daily Acres Treated (c) | Short-Term PPE MOEs                                       |            |       | Short-Term Eng. Control MOEs |            |          |
|--|---|---|-------------------------|---|------------|-------|------------------------------|------------|----------|
|  |   |   |                         | Dermal  | Inhalation | Total | Dermal                       | Inhalation | Total    |
| <b>Mixer/Loader Exposure</b>                                   |   |   |                         |   |            |       |                              |            |          |
| Mixing/Loading Liquids for Aerial/Chemigation Application (1a) | Yes<br>MRID No. 44739302                      | 1.5 cranberries, corn                     | 350                     | 39  | 56         | 23    | 78                           | 160        | 52       |
|  |   | 3.5 citrus (d)                            | 100                     | 59  | 83         | 34    | 120                          | 240        | 78       |
| Mixing/Loading Liquids for Groundboom Application (1b)         | Yes<br>MRID No. 42974501                      | 1.5 predominant max                       | 80                      | 170   | 240        | 100   | Target MOE reached at PPE    |            |          |
|  |   | 5.0 tobacco max                           | 80                      | 51  | 73         | 30    | 100                          | 210        | 69       |
|  |   | 2 Sodfarm (includes tobacco/potatoes)     | 80                      | 130   | 180        | 75    | 250                          | 530        | 170      |
|  |   | 4 Sodfarm                                 | 80                      | 64  | 91         | 38    | 130                          | 260        | 86       |
|  |   | 8.0 sodfarm fire ants                     | 10                      | 260   | 360        | 150   | Target MOE reached at PPE    |            |          |
| Mixing/Loading Liquids for Airblast Application (1c)           | Yes<br>MRID No. 43138102                      | 2.0 predominant max such as Fruits & Nuts | 40                      | 260   | 360        | 150   | Target MOE reached at PPE    |            |          |
|  |   | 6.0 citrus                                | 20                      | 170   | 240        | 100   | Target MOE reached at PPE    |            |          |
| Mixing WP for Aerial/Chemigation Application (2a)              | No  | 2.0 predominant max (orchards)            | 350                     | DAS is not supporting the open bag formulation for the WP |            |       | 51                           | 42         | 23       |
|  |   | 3.5 citrus (d)                            | 100                     |   |            |       | 100                          | 83         | 46       |
| Mixing WP for Groundboom Application (2b)                      | Yes<br>MRID No. 42974501                      | 1.0 predominant max (brassica)            | 80                      |   |            |       | 450                          | 360        | 200      |
|  |   | 4.0 soil treatment ornamentals outdoors   | 10                      |   |            |       | 890                          | 730        | 400      |
|  |   | 1.3 & 3.0 Sodfarm                         | 80                      |   |            |       | 340 / 150                    | 280 / 120  | 150 / 67 |

**Table 9  
Exposure Variables and MOEs for Agricultural Uses  
(Including Non Worker Protection Standard Ornamental Uses) of Chlorpyrifos**

| Exposure Scenario (Scenario#)                 | Are Biological Monitoring Data Available? (a) | Application Rates (lb ai/acre) (b)   | Daily Acres Treated (c) | Short-Term PPE MOEs   |            |       | Short-Term Eng. Control MOEs |            |       |
|---|---|--------------------------------------|-------------------------|---|------------|-------|------------------------------|------------|-------|
|   |   |                                      |                         | Dermal  | Inhalation | Total | Dermal                       | Inhalation | Total |
|   |   | 8.0 sodfarm fire ants (harvest only) | 10                      |   |            |       | 4500                         | 3600       | 200   |
| Mixing WP for Airblast Application (2c)       | No  | 2.0 predominant max                  | 40                      |   |            |       | 450                          | 360        | 200   |
|   |   | 6.0 citrus                           | 20                      |   |            |       | 300                          | 240        | 130   |
| Loading Granulars for Aerial Application (3a) | No  | 1.95 maximum aerial rate             | 350                     | 150   | 30         | 25    | 3000                         | 300        | 270   |
| Loading Granulars for Ground Application (3b) | Yes<br>MRID No. 44483501 (3b and 8)           | 1.0 typical corn                     | 80                      | 1300  | 260        | 210   | Target MOE reached at PPE    |            |       |
|   |   | 2.0 max corn                         | 80                      | 640   | 130        | 110   | Target MOE reached at PPE    |            |       |
|   |   | 3.0 maximum ground rate (tobacco)    | 80                      | 430   | 86         | 71    | 8600                         | 860        | 780   |
| <b>Applicator Exposure</b>                    |   |                                      |                         |   |            |       |                              |            |       |
| Aerial (Spray) -- Enclosed Cockpit (4a)       | No  | 2.0 orchards                         | 350                     | No Open cockpit data available  |            |       | 100                          | 150        | 60    |
|   |   | 3.5 citrus (d)                       | 100                     |   |            |       | 200                          | 290        | 120   |
| Aerial (Granulars) -- Enclosed Cockpit (4b)   | No  | 1.95                                 | 350                     | No Open cockpit data available  |            |       | 320                          | 8          | 8     |
| Groundboom Tractor (5)                        | Yes<br>MRID No. 42974501                      | 1.5 predominant max                  | 80                      | The biological monitoring results (Table A4) indicate that open cabs provide insufficient protection . Therefore, only the enclosed cab MOEs are presented. |            |       | 580                          | 1400       | 410   |
|   |   | 5.0 tobacco max                      | 80                      |   |            |       | 180                          | 410        | 120   |
|   |   | 4 Sodfarms                           | 80                      |   |            |       | 220                          | 510        | 150   |
|   |   | 8.0 sodfarm fire ants                | 10                      |   |            |       | 880                          | 2000       | 610   |

**Table 9  
Exposure Variables and MOEs for Agricultural Uses  
(Including Non Worker Protection Standard Ornamental Uses) of Chlorpyrifos**

| Exposure Scenario (Scenario#)           | Are Biological Monitoring Data Available? (a) | Application Rates (lb ai/acre) (b)   | Daily Acres Treated (c) | Short-Term PPE MOEs   |                 |                | Short-Term Eng. Control MOEs   |            |       |
|---|---|--|-------------------------|---|-----------------|----------------|--|------------|-------|
|   |   |  |                         | Dermal  | Inhalation      | Total          | Dermal   | Inhalation | Total |
| Airblast Applicator (6)                 | Yes<br>MRID No. 43138102                      | 2.0 predominant max  | 40                      | The biological monitoring results indicate that open cabs are insufficient. |                 |                | 230  | 190        | 110   |
|   |   | 6.0 citrus   | 20                      |   |                 |                | 150  | 130        | 70    |
| Tractor-Drawn Granular Spreader (7)     | Yes<br>MRID No. 44483501 (3b and 8)           | 1.0 typical corn   | 80                      | 1000  | 360             | 270            | Target MOE reached at PPE  |            |       |
|   |   | 2.0 max corn   | 80                      | 520   | 180             | 140            | Target MOE reached at PPE  |            |       |
|   |   | 3.0 maximum ground rate (tobacco)  | 80                      | 350   | 120             | 90             | 690  | 130        | 110   |
| Seed Treatment (8)                      | No  | No Data  | No Data                 | No Data   |                 |                | No Data  |            |       |
| Dip Application (Preplant Peaches) (9)  | No  | No Data  | No Data                 | No Data   |                 |                | No Data  |            |       |
| <b>Flagger Exposure</b>                 |   |  |                         |   |                 |                |  |            |       |
| Spray Applications (10)                 | No  | 2.0 predominant max  | 350                     | 50  | 140             | 37             | 2300   | 1400       | 880   |
|   |   | 3.5 citrus (d)   | 100                     | 100   | 290             | 74             | 4500   | 2900       | 1800  |
| Granular Applications (11)              | No  | 1.95   | 350                     | 320   | 340             | 170            | Target MOE reached at PPE  |            |       |
| <b>Mixer/Loader/Applicator Exposure</b> |   |  |                         |   |                 |                |  |            |       |
| Backpack Sprayer (12)                   | Yes<br>MRID No. 43027901                      | 0.0417 lb ai/gal predominant max / 0.08 lb ai/gal bark beetle treatment / 0.03 lb ai/gal stump treatment | 40 gal/day              | 130 / 68 / 180  | 700 / 360 / 970 | 110 / 58 / 150 | Target MOE reached at PPE, except for the higher concentration for the beetle bark treatment |            |       |
|   |   | 3.5 citrus bark  | 1 A/day                 | 63  | 330             | 53             | Not feasible   |            |       |
|   |   | 0.039 lb ai/gal /750 ft2   | 1000 ft2                | 4200  | 22000           | 3500           | Target MOE reached at PPE  |            |       |

**Table 9  
Exposure Variables and MOEs for Agricultural Uses  
(Including Non Worker Protection Standard Ornamental Uses) of Chlorpyrifos**

| Exposure Scenario (Scenario#)                                       | Are Biological Monitoring Data Available? (a) | Application Rates (lb ai/acre) (b)   | Daily Acres Treated (c) | Short-Term PPE MOEs |                 |                 | Short-Term Eng. Control MOEs |            |       |
|---|---|--|-------------------------|---------------------|-----------------|-----------------|------------------------------|------------|-------|
|   |   |  |                         | Dermal              | Inhalation      | Total           | Dermal                       | Inhalation | Total |
| Low Pressure Handwand (13)  | Yes<br>MRID No.<br>43027901                   | 0.0417 lb ai/gal predominant max / 0.08 lb ai/gal bark beetle treatment / 0.03 lb ai/gal stump treatment | 40 gal/day              | 570 / 300 / 790     | 700 / 360 / 970 | 310 / 160 / 440 | Target MOE reached at PPE    |            |       |
|   |   | 3.5 citrus bark  | 1 A/day                 | 270                 | 330             | 150             | Target MOE reached at PPE    |            |       |
|   |   | 0.039 lb ai/gal / 750 ft2 animal prem.   | 1000 ft2                | 18000               | 22000           | 10,000          | Target MOE reached at PPE    |            |       |
| High Pressure Handwand (greenhouse uses) (14)                       | Yes<br>MRID No.<br>43027901                   | Min. 0.0033 lb ai/gal  | 1000 gal/day            | 66                  | 88              | 38              | Not feasible                 |            |       |
|   |   | Max. 0.0066 lb ai/gal  |                         | 33                  | 44              | 19              | Not feasible                 |            |       |
| Hydraulic Hand-held Sprayer for Bark / Pine Seedling Treatment (15) | No  | 3.5 citrus bark  | 10                      | 16                  | 100             | 14              | Not feasible                 |            |       |
|   |   | 0.08 lb ai/gal bark beetle treatment / 0.16 lb ai / gal pine seedling treatment /                        | 1,000                   | 14 / 7              | 88 / 44         | 12 / 6          | Not Feasible                 |            |       |
|   |   | 0.039 lb ai/gal / 750 ft2 animal prem  | 10000 ft2               | 2,200               | 13,000          | 1,900           | Target MOE reached at PPE    |            |       |
| Dry Bulk Fertilizer Impregnation                                    | No  | 1.0 lb ai / 200 lb fertilizer / acre   | No Data                 | No Data             |                 |                 | No Data                      |            |       |

- (a) Biological monitoring data are available from several chemical-specific studies. Although biological monitoring scenarios are available for some of the scenarios as indicated in this table, passive dosimetry data are presented for comparison because insufficient replicates and/or additional risk mitigation measures were necessary.
- (b) Application rates are the maximum labeled rates found on EPA Reg. Nos. 62719-38, -221, -245, -34; -79, -72, -166, -220, 34704-66 (Clean Crop Chlorpyrifos 4E -- sodfarm fire ant rate), 499-367 (499-367 is the only greenhouse label identified), and 10350-22 for animal premise treatments. **“Predominant max”** in this table refers to the most **frequently identified maximum** application rate found on the labels for the specific formulation and equipment type. Typical rates are also included to characterize the chlorpyrifos uses. Not all application rates are included for all crops, instead, a cross-section of rates are used to represent the uses of chlorpyrifos.
- (c) Daily acres treated are based on HED’s estimates of acreage (or gallonage) that would be reasonably expected to be treated in a single day for each

exposure scenario of concern. The sodfarm fire ant rate is restricted on the label for harvest only, therefore, this rate is limited to the amount of sod that may be harvested in a reasonable time frame. Therefore, using the limited data available, approximately 10 acres treated per day are assumed to be the upper range.

- (d) The application rates on the Lorsban 4E (EPA Reg. No. 62719-220) and 50W (EPA Reg. No. 62719-39 discontinued as of 1995 and sold as -221) labels indicate that for citrus at the 6.0 lb ai/A rate it is necessary to use 100 to 2,400 gallons per acre dilute spray. Therefore, this rate is not expected to be feasible for an aerial applicator. The label language should be clarified so that the 6.0 lb ai/A rate is for ground only. Additionally, citrus orchards are believed to be relatively small plots and 100 acres per day is assumed in the assessment for aerial applications.



**Table 10. Estimates of Risks to Commercial Applicators and Residents Applying Chlorpyrifos in the Residential/Recreational Environment**

| Application Scenario  | Clothing   | Method of Evaluation   | MOE   |   |   | Risk Characterization/<br>Uncertainties   |
|---|--|--|---|---|---|---|
|   |  |  | Dermal  | Inhalation  | Total   |   |
| <b>(1) Indoor Crack &amp; Crevice Treatment</b>   |  |  |   |   |   |   |
| Long term PCO Applicator (0.29% Dursban Pro; EPA Reg. 62719-166)  | double layer clothes, chemically-resistant boots and gloves, eye protection  | Biomonitoring study MRID No. 44444801 (minimum, mean and maximum amount handled)   | 17 (max)<br>59 (mean)<br>5900 (min)                 | 58 (max)<br>200 (mean)<br>20,000 (min)              | 13 (max)<br>45 (mean)<br>4500 (min)                 | Central-tendency risk estimates for applicators; MOEs less than 100 for workers that could handle \$0.02 lb ai/day (the mean amount handled in the study). Only two of 15 replicates reflect the maximum label concentration of 0.5% ai. (avg of 0.29% ai was handled in study). Underestimates exposure to workers that mix/load and apply chlorpyrifos because study only evaluated applicators.                        |
| Short-term Residential Applicator (EPA Reg 026693-00003 for 1% ai; 239-2619 for 0.5% ai)                            | SS, SP, no gloves  | Residential SOPs (PHED V1.1)   | 159 (1%)<br>318 (0.5%)<br><br>2540 (spot treatment) | 292 (1%)<br>584 (0.5%)<br><br>4700 (spot treatment) | 100 (1%)<br>200 (0.5%)<br><br>1600 (spot treatment) | High-end risk estimates for 1% ai; central tendency for 0.5% ai; assumes application of one 16 oz. aerosol can for both; low-end to central tendency risk for spot treatment which assumes 2 oz application of 0.5% ai. product   |
| <b>(2) Broadcast Turf Application (Intermediate and Long-Term for PCOs; Short-Term for Residential Applicators)</b> |  |  |   |   |   |   |
| Applicator (1 or 4 lb ai/Acre of Dursban Pro, EPA Reg. 62719-166)   | single layer clothes, chemically-resistant knee high boots and gloves, hat (knee high boots not required by label) | Biomonitoring Study MRID No. 44729401 (25% of label maximum rate or adjustment for label-recommended max application rate) | Biomonitoring: 75 (IT&LT) (1 lb ai/acre)            |   |   | Central-tendency risk estimates for 1 lb ai/acre; product applied at 25% of label maximum. High-end risk estimates for 4 lb ai/acre (label maximum for subsurface soil treatment). Study evaluated an average 1.5 hour spray time over a 6 hour work day which may underestimate worker exposure based on TruGreen/ChemLawn data for 193 workers that show an average spray time of 2.75 hours over a 8.75 hour work day. |
|   |  |  | Label Max: 20 (IT&LT) (4 lb ai/acre)                |   |   |   |
| Mixer/Loader (liquid) (Dursban Pro, EPA Reg. 62719-166)   | single layer clothes, gloves   | PHED V1.1 (biomonitoring study rate and 25% of maximum label rate)   | 260-1032  | 500-1980 (IT)<br>150 -600 (LT)                      | 170-680 (IT)<br>100-380 (LT)                        | Central-tendency to High-end risk estimates; maximum ai handled in study with maximum (4 lb ai/acre) and 25% of maximum label rate (1 lb ai/acre), respectively   |
|   | double layer clothes, gloves   |  | 350 -1400   |   | 200-820 (IT)<br>100 -420 (LT)                       |   |

**Table 10. Estimates of Risks to Commercial Applicators and Residents Applying Chlorpyrifos in the Residential/Recreational Environment**

| Application Scenario   | Clothing          | Method of Evaluation                                      | MOE     |            |         | Risk Characterization/<br>Uncertainties   |
|--|-------------------|---|---------|------------|---------|---|
|  |                   |   | Dermal  | Inhalation | Total   |   |
| Residential Mixer/Loader/ Applicator Broadcast with Hose End Sprayer (Dursban 1-12 Insecticide EPA Reg 62719-56)           | SS, SP, no gloves | Residential SOPs (PHED V1.1) (min and max dilution rates) | 6-23    | 368-1470   | 6-23    | Central-tendency to High-end risk estimates; Low confidence in exposure estimates from PHED V1.1; assumes resident handles 22 gallons of minimally and maximally diluted product  |
| Residential Mixer/Loader/ Applicator Spot treatment with Low Pressure Handwand (Dursban 1-12 Insecticide EPA Reg 62719-56) | SS, SP, no gloves | Residential SOPs  | 37-150  | 2490-9960  | 37-150  | Central-tendency to High-end risk estimates; Low confidence in dermal exposure estimates, and medium confidence in inhalation exposure estimates; assumes resident handles 1 gallon of minimally and maximally diluted product to treat 1000 ft <sup>2</sup> .  |
| <b>(3) Golf Course Use (Dursban Turf Insecticide; EPA Reg. 62719-35) (Short-term)</b>                                      |                   |   |         |            |         |   |
| Mixer/Loader (Liquid)  | LS, LP, gloves    | PHED V1.1   | 95-380  | 36-150     | 26-100  | High-end for 4 lb ai/acre and central tendency for 1 lb ai/acre; assumes handling product to treat 40 acres at 1-4 lb ai/acre. Using PHED only 4 lb ai/acre results in MOEs < 100 for liquid mixer/loader (MOE=26). For groundboom applicator, MOE < 100 based on biomonitoring at both 1 and 4 lb ai/acre. HED has more confidence in the biomonitoring results than PHED. |
| Mixer/Loader (Wettable Powder in water soluble bags)   | LS, LP, gloves    | PHED V1.1   | 220-820 | 180-730    | 100-400 |   |
| Groundboom Applicator  | LS, LP, no gloves | PHED V1.1   | 160-630 | 59-240     | 43-170  |   |
|  |                   | Biomonitoring (MRID 42974501)                             | 15-63   |            | 15-63   |   |
| Mix/Load/Apply via Handgun (greens/tees) (Liquid)  | LS, LP, gloves    | PHED V1.1   | 49-190  | 130-540    | 36-140  | High-end for 4 lb ai/acre and central tendency for 1 lb ai/acre; assumes handling product to treat 5 acres at 1-4 lb ai/acre. Only 4 lb ai/acre results in MOEs < 100   |

**Table 10. Estimates of Risks to Commercial Applicators and Residents Applying Chlorpyrifos in the Residential/Recreational Environment**

| Application Scenario   | Clothing          | Method of Evaluation                          | MOE                       |            |                           | Risk Characterization/<br>Uncertainties   |
|--|-------------------|---|---------------------------|------------|---------------------------|---|
|  |                   |   | Dermal                    | Inhalation | Total                     |   |
| <b>(4) Ready-to-Use 0.5% a.i. Formulated Product (Ortho Ant Stop)</b>                                    |                   |   |                           |            |                           |   |
| Short-term Residential Applicator  | SS, LP, no gloves | Outdoor Biomonitoring Study MRID No. 44739301 | 625 (biomonitoring)       |            | 625                       | Central-tendency to high-end risk estimate; assumes resident applies five 24 oz bottles of product/day, however, resident wore long pants and current HED policy is to evaluate exposures for short pants. Risks calculated two ways, one using total exposure based on biomonitoring, and second by comparing estimated route-specific exposure to appropriate toxicity endpoints. |
|  |                   |   | 714                       | 3,400      | 590                       |   |
| <b>(5) Insecticidal Dust Product (Shaker Can or Bulbous Duster)</b>                                      |                   |   |                           |            |                           |   |
| <b>Residential Applicator (1% ai chlorpyrifos; 2.83 g ai) (EPA Reg. 62719-66, 62719-54, and 192-171)</b> |                   |   |                           |            |                           |   |
| Short-term   | SS, LP, no gloves | Scientific Literature Study                   | 250                       | NE         | 250                       | Central-tendency to High-end risk estimates; assumes an individual applies a 10 oz can of 1% ai chlorpyrifos dust; neglects inhalation exposure due to an absence of data.  |
| <b>Worker (7% ai chlorpyrifos; 7.91 or 198 g ai) (EPA Reg. 13283-17, Rainbow Kofire Ant Killer)</b>      |                   |   |                           |            |                           |   |
| Short-term   | LS, LP, gloves    | Scientific Literature Study                   | 98 (7.9 g)<br>3.9 (198 g) | NE         | 98 (7.9 g)<br>3.9 (198 g) | Central-tendency short term risk assessments for 7.9 and 198 g ai; High-end intermediate-term risk estimates for 7.9 and 198 g ai (based on size of dust container); Neglects inhalation exposure due to an absence of data.  |
| Intermediate term  |                   |   | 20 (7.9 g)<br>0.8 (198 g) | NE         | 20 (7.9 g)<br>0.8 (198 g) |   |

**Table 10. Estimates of Risks to Commercial Applicators and Residents Applying Chlorpyrifos in the Residential/Recreational Environment**

| Application Scenario   | Clothing                      | Method of Evaluation | MOE       |            |           | Risk Characterization/<br>Uncertainties   |   |
|--|-------------------------------|----------------------|-----------|------------|-----------|---|---|
|  |                               |                      | Dermal    | Inhalation | Total     |   |   |
| <b>(6) Granular Formulation (Hand Application) (EPA Reg. 672719-14, 62719-210) (2 lb ai/acre)</b>  |                               |                      |           |            |           |   |   |
| LCO (intermediate-term)  | LS, LP, gloves                | PHED V1.1            | 21        | 324        | 20        | <p>High-end risk estimates; medium confidence in PHED unit exposure estimates which are based on a single study in which a test subject wearing chemical-resistant gloves spread the granular formulation around the outside of the residence and over 90 percent of the samples contained no detectable material. Therefore, residents also evaluated wearing long pants, long sleeved shirt and gloves. Assumes treatment of 1000 ft<sup>2</sup>. Could underestimate exposure because PHED data excludes head and neck area.</p> |   |
|  | Double layer clothing, gloves |                      | 38        | 324        | 34        |   |   |
| Residential Applicator (short-term)  | SS, SP, no gloves             | Residential SOPs     | 18        | 327        | 17        |   |   |
|  | LS, LP, gloves                |                      | 106       | 330        | 80        |   |   |
| <b>(7) Granular Formulation (Belly Grinder) (EPA Reg. 672719-14, 62719-210) (2 lb ai/acre)</b>     |                               |                      |           |            |           |   |   |
| LCO (intermediate-term)  | LS, LP, gloves                | PHED V1.1            | 8         | 120        | 7         |   | <p>Central-tendency risk estimates for worker; High-end risk estimates for residents, except for spot treatment. Low and high confidence in the dermal and inhalation exposure estimates, respectively. Assumes treatment of 0.5 acre at typical rate of 2 lb ai/acre for subsurface feeding insects. Could underestimate exposure because PHED data excludes head and neck area. Workers could treat more than 0.5 acre/day.</p> |
|  | Double layer clothing, gloves |                      | 12.5      | 120        | 11        |   |   |
| Residential Applicator (short-term)  | SS, SP, no gloves             | Residential SOPs     | 3         | 120        | 3         |   |   |
|  |                               |                      | 69 (spot) | 36 (spot)  | 24 (spot) |   |   |
| <b>(8) Granular Formulation (Push-type Spreader) (EPA Reg. 672719-14, 62719-210)(2 lb ai/acre)</b> |                               |                      |           |            |           |   |   |
| LCO (intermediate-term)  | LS, LP, gloves                | PHED V1.1            | 57        | 1150       | 54        | <p>Central-tendency risk estimates for worker; High-end risk estimates for residents. Low and high confidence in the dermal and inhalation exposure estimates, respectively. Assumes treatment of 0.5 acre at typical rate 2 lb ai/acre for subsurface feeding insects. Could underestimate exposure because PHED data excludes head and neck area. Workers could treat more than 0.5 acre/day.</p>   |   |
|  | Double layer clothing         |                      | 100       | 1150       | 92        |   |   |

**Table 10. Estimates of Risks to Commercial Applicators and Residents Applying Chlorpyrifos in the Residential/Recreational Environment**

| Application Scenario                | Clothing          | Method of Evaluation | MOE    |            |       | Risk Characterization/<br>Uncertainties |
|-------------------------------------|-------------------|----------------------|--------|------------|-------|---|
|                                     |                   |                      | Dermal | Inhalation | Total |   |
| Residential Applicator (short-term) | SS, SP, no gloves | Residential SOPs     | 120    | 1150       | 110   |   |

**Table 10. Estimates of Risks to Commercial Applicators and Residents Applying Chlorpyrifos in the Residential/Recreational Environment**

| Application Scenario   | Clothing  | Method of Evaluation   | MOE      |            |                                 | Risk Characterization/<br>Uncertainties  |
|--|---|--|----------|------------|---------------------------------|--|
|  |   |  | Dermal   | Inhalation | Total                           |  |
| <b>Termiticide Treatments</b>  |   |  |          |            |                                 |  |
| <b>(9) Pre-Construction (1.44% chlorpyrifos as Dursban TC) (EPA Reg. 62719-47) (long-term)</b> |   |  |          |            |                                 |  |
| Mixer/Loader/<br>Applicator (3 hour<br>average exposure)                                       | label-specified PPE:<br>single layer clothes and<br>forearm-length<br>chemically-resistant<br>gloves (forearm length<br>gloves not required by<br>label)  | Dosimetry and air<br>monitoring from<br>Registrant Study<br>MRID No. 44589001                | 19       | 67         | 15                              | Low-end risk estimates for workers that wore double layer of clothing and forearm length gloves not required by the label; Central-tendency risk estimates for workers that wore a single layer of clothing and forearm length gloves; assumes 3 hour exposure, which could underestimate risks to workers exposed > 3 hrs/day, or that use 2% ai to treat utility poles or fences |
|  | double layer clothes<br>(LS,LP, coveralls, rubber<br>boots, and forearm-length<br>gloves) (forearm-length<br>gloves not required by<br>label)   |  | 63       | 67         | 33                              |  |
| Tarp puller  | with forearm-length<br>gloves (LS,LP, leather<br>and/or rubber boots and<br>hat)  | Dosimetry and air<br>monitoring from<br>Registrant Study (1-8<br>tarps)<br>MRID No. 44589001 | 170-1300 | 180-1400   | 87 (8 tarps)<br>690<br>(1 tarp) | Central-tendency risk estimates; assumes workers pull 1-8 tarps/day (7 min/tarp), could underestimate risks to workers who pull > 8 tarps/day (i.e., >1 hr exposure/day). All total MOEs < 100 for 8 tarp/day. Also, workers wore forearm length gloves not required by the label which reduce estimated exposure.   |
|  | without gloves (LS,LP,<br>leather and/or rubber<br>boots and hat)   |  | 47-370   | 240-2000   | 39 (8 tarps)<br>310<br>(1 tarp) |  |
| <b>(10) Post-Construction (1% chlorpyrifos as Dursban TC) (EPA Reg. 62719-47) (long-term)</b>  |   |  |          |            |                                 |  |
| Mixer/Loader/<br>Applicator  | Label-specified PPE: LS,<br>LP, chemically resistant<br>gloves, hat, eye protection<br>and half face piece<br>respirator in confined<br>spaces;<br>During M/L: 2 layers<br>clothes and chemically-<br>resistant shoes | Biomonitoring: 4.3<br>MRID No. 44729402<br>(n=5)   | 7        |            | 7                               | Central-tendency risk estimate, could underestimate risks for workers that apply 2% ai to treat utility poles or fences  |
|  |   | Dosimetry and air<br>monitoring<br>MRID No. 44729402<br>(n=14)                               | 12       | 33         | 9                               | Central-tendency risk estimate; excludes worker with higher exposure (10X greater than mean) due to a broken hose  |

**Table 10. Estimates of Risks to Commercial Applicators and Residents Applying Chlorpyrifos in the Residential/Recreational Environment**

| Application Scenario  | Clothing  | Method of Evaluation   | MOE                      |                            |                          | Risk Characterization/<br>Uncertainties   |
|---|---|--|--------------------------|----------------------------|--------------------------|---|
|   |   |  | Dermal                   | Inhalation                 | Total                    |   |
| <b>(11) Paint Brush (Short-term) (Dursban 1-12 Insecticide, EPA Reg. 62719-56)</b>  |   |  |                          |                            |                          |   |
| Residential Applicator  | SS, SP, no gloves   | Residential SOPs; 1 gallon for worst case and 1 quart for typical case | 37 (1 gal)<br>148 (1 qt) | 590 (1 gal)<br>2300 (1 qt) | 35 (1 gal)<br>140 (1 qt) | Central-tendency risk estimates for typical case and high end risk estimates for worst case; low to medium confidence in dermal exposure estimates and medium confidence in inhalation exposure estimates; Assumes resident applies 1 gallon or 1 quart of diluted product in a day |
| <b>(12) Ornamental Application (Short-term) (Dursban 1-12 Insecticide, EPA Reg. 62719-56)</b>                                   |   |  |                          |                            |                          |   |
| Residential Mixer/Loader/ Applicator<br>Low pressure Handwand   | SS, SP, no gloves   | Residential SOPs (minimum : 1 oz/3gal H2O)                             | 270                      | 18,000                     | 270                      | Central-tendency to high-end risk estimates; low and medium confidence in the dermal and inhalation exposure estimates, respectively. Assumes resident applies 5 gallons of diluted product/day.  |
|   |   | Residential SOPs (typical 4 oz/3 gal H2O)                              | 70                       | 4,700                      | 69                       |   |
|   |   | Residential SOPs (max. 1 qt/3 gal H2O)                                 | 8                        | 560                        | 8                        |   |
| Residential Mixer/Loader/ Applicator<br>Hose End Sprayer  | SS, SP, no gloves   | Residential SOPs (minimum : 1 oz/3gal H2O)                             | 900                      | 57,000                     | 880                      | Central-tendency to high-end risk estimates; low confidence in the dermal and inhalation exposure estimates. Assumes resident applies 5 gallons of diluted product/day.   |
|   |   | Residential SOPs (typical 4 oz/3 gal H2O)                              | 230                      | 15,000                     | 230                      |   |
|   |   | Residential SOPs (max. 1 qt/3 gal H2O)                                 | 28                       | 1,800                      | 28                       |   |
| <b>(13) Mosquitocide Mixer/Loader/Applicator (PHED V1.1) (Short- and intermediate-term) (Mosquitomist One EPA Reg. 8329-24)</b> |   |  |                          |                            |                          |   |
| Mixer/Loader--Aerial  | PPE double layer clothes and gloves                                     | PHED V1.1  | 120 (ST)<br>24 (IT)      | 34 (ST&IT)                 | 26 (ST)<br>14 (IT)       | High end risk estimates. Application rate of 0.023 lb ai/acre for 7500 acres  |
|   | Engineering Controls (enclosed cockpit) single layer clothes and gloves |  | 236 (ST) 47 (IT)         | 490 (ST&IT)                | 160 (ST) 43 (IT)         |   |

**Table 10. Estimates of Risks to Commercial Applicators and Residents Applying Chlorpyrifos in the Residential/Recreational Environment**

| Application Scenario                   | Clothing   | Method of Evaluation | MOE                   |               |                      | Risk Characterization/<br>Uncertainties  |
|--|--|----------------------|-----------------------|---------------|----------------------|--|
|  |  |                      | Dermal                | Inhalation    | Total                |  |
| Mixer/Loader--<br>Ground-based fogger_ | PPE, single layer clothes and gloves   |                      | 1010 (ST)<br>200 (IT) | 390 (ST&IT)   | 280 (ST)<br>133 (IT) | High end risk estimates. Application rates of 0.005 and 0.01 lb ai//acre for 3000 acres. Surrogate ground-based fogger exposure data are not available, and therefore, it was necessary to extrapolate from airblast exposure data |
|  | engineering controls (enclosed cab) and single layer clothes and gloves        |                      | 270 (IT)              | 2800 (IT)     | 250 (IT)             |  |
| Aerial Applicator                      | engineering controls (enclosed cockpit) and single layer clothes and no gloves |                      | 400 (ST)<br>81 (IT)   | 600 (ST&IT)   | 240 (ST)<br>71 (IT)  | High end risk estimates. Application rate of 0.023/acre for 7500 acres   |
| Ground-based fogger Applicator         | engineering controls (enclosed cab) and single layer clothes and no gloves     |                      | 610-1230 (ST)         | 520-1040 (ST) | 280-560 (ST)         | High end risk estimates. Application rates of 0.005 and 0.01 lb ai/acre for 3000 acres. Surrogate ground-based fogger exposure data are not available, and therefore, it was necessary to extrapolate from airblast exposure data  |
|  |  |                      | 120-250 (IT)          | 520-1040 (IT) | 100-200 (IT)         |  |

LS=Long sleeves; LP = Long pants; SS = short sleeves; SP = short pants  
H2O = water; ST = short-term (1- 30 days); IT = intermediate term (30 days to 6 months) LT = long term (> 6 months)  
NE = Not evaluated



| TABLE 11<br>Crop Grouping Matrix by Potential for Dermal Contact |  |   |  |
|--|--|---|--|
| Potential for Dermal Contact                                     | Transfer Coefficient (cm <sup>2</sup> /hr) | Activities                                  | Crops  |
| Low  | 2,500                                      | Harvest                                     | Alfalfa, asparagus, small grains (wheat, sorghum, milo), soybeans, cole crops, mint  |
|  |  | Sort/Pack                                   | Sugar beets, radishes, rutabagas   |
| Medium   | 4,000                                      | Harvest, stake/tie, scout, irrigate         | Cranberries, strawberries  |
|  |  | Irrigate                                    | Christmas trees  |
|  |  | Late season scouting                        | Cotton   |
| High   | 10,000                                     | Harvest                                     | Sunflowers, sugar beets, corn (up to 1.5 lb ai/A as a foliar treatment), sweet potatoes, radishes, rutabagas, turfgrass (sodfarm) for fire ants, almond harvesting |
|  |  | Cut/harvest, prune, transplant, ball/burlap | Christmas trees  |

| TABLE 12<br>Restricted Entry Intervals (REIs) for Chlorpyrifos: General |                        |           |                               |           |
|---|------------------------|-----------|-------------------------------|-----------|
| Potential for Dermal Contact  | Short-Term REIs (days) |           | Intermediate-Term REIs (days) |           |
|   | 1 lb ai/A              | 2 lb ai/A | 1 lb ai/A                     | 2 lb ai/A |
| LOW   | 1                      | 1         | 1                             | 1         |
| MEDIUM  | 1                      | No Crops  | 1                             | No Crops  |
| HIGH  | 1                      | 1         | 1                             | 2         |
| Scouting (Various Crops)  | 0                      | 1         | 1                             | 1         |

| TABLE 13<br>Restricted Entry Intervals (REIs) for Chlorpyrifos:<br>Cauliflower, Citrus and Tree Nuts & Fruit |                        |        |        |             |        |                               |        |        |             |        |
|--|------------------------|--------|--------|-------------|--------|-------------------------------|--------|--------|-------------|--------|
| Activity   | Short-Term REIs (days) |        |        |             |        | Intermediate-Term REIs (days) |        |        |             |        |
|  | Almonds                | Apples | Pecans | Cauliflower | Citrus | Almonds                       | Apples | Pecans | Cauliflower | Citrus |
| Scouts   | 2                      | 1      | 0      | 1 to 3      | 2      | 2                             | 1      | 0      | 1 to 3      | 2      |
| Harvesting   | 5                      | 3      | 1      | 5 to 8      | 5      | 7                             | 4      | 2      | 7 to 10     | 5      |
| Pruning (wet cond.)  | NE                     | NE     | NE     | NA          | 4      | NE                            | NE     | NE     | NA          | 5      |
| Pruning (dry cond.)  | NE                     | NE     | NE     | NA          | 2      | NE                            | NE     | NE     | NA          | 2      |

NE = Not Evaluated

| Table 14<br>Chlorpyrifos Surrogate Occupational Postapplication Assessment for Golf Course Turf Treatment |                  |         |                                       |  |                    |  |                    |
|---|------------------|---------|---------------------------------------|--|--------------------|--|--------------------|
| Crop  | Application Rate | DAT (a) | TTR from WP (Fg/cm <sup>2</sup> ) (b) | Mow/Maintain Transfer coefficient =500 cm <sup>2</sup> /hr |                    | Mow/Maintain Transfer coefficient =1,000 cm <sup>2</sup> /hr |                    |
|   |                  |         |                                       | Potential Dermal Dose (mg/kg/day) (c)                      | Short-term MOE (d) | Potential Dermal Dose (mg/kg/day) (c)                        | Short-term MOE (d) |
| Golf Course Turf  | 4.0              | 0       | 0.414                                 | 0.024  | 210                | 0.047  | 110                |

- (a) DAT is "days after treatment."
- (b) Turf Transferable residues (TTR) from MRID 448296-01 based on average of CA, IN and MS sites following application of 4 lb ai/ Acre of Dursban 50W.
- (g) Dermal Dose = TTR (Fg/cm<sup>2</sup>) x Transfer coefficient (cm<sup>2</sup>/hr) x conversion factor (1 mg/1,000) x 8 hr/day duration x dermal absorption x 1/70 kg body weight. The target MOE of 100 is based on 10x interspecies and 10x intraspecies.
- (d) Short-term MOE = NOAEL of 5 mg/kg/day / Potential dermal dose (mg/kg/day).

**Table 15. Estimates of Post-Application Risks to Residents/Recreational Users**

| Reentry Scenario  | Method of Evaluation  | Central-tendency MOE       |                            | Risk Characterization/<br>Uncertainties   |
|---|---|----------------------------|----------------------------|---|
|   |   | Adult                      | Child                      |   |
| <b>(1) Crack &amp; Crevice Treatment of Kitchen and Bathroom (0.5% Dursban Pro diluted spray, EPA Reg. 62719-166) (Short and Intermediate Term)</b> |   |                            |                            |   |
| Maximum 1-Day Inhalation Exposure:  | Biomonitoring Study, with environmental measurements  | 560                        | 130                        | Central-tendency to High-end risk estimates; assumes exposure exclusively through inhalation and that children spend 21 hours/day (50th percentile for 1-4 yr old at home) in a treated room (i.e., home, schools, day care centers, etc). This could over-or under-estimate risk because it is compared to a 90 day inhalation NOAEL for rats exposed 6 hours/day.                 |
| 10-Day TWA Inhalation Exposure  |   | 670                        | 360                        |   |
| <b>(2) Crack &amp; Crevice Treatment Using Residential SOPs (0.5% Dursban Pro diluted spray, EPA Reg. 62719-166) (Short-term)</b>                   |   |                            |                            |   |
| Dermal Exposure From Carpets  | Highest deposition from <u>untreated</u> family room in biomonitoring study (room adjacent to treatment) and Residential SOPs | 1950                       | 1360                       | Low-end risk estimates; highest deposition from <u>untreated room</u> used in conjunction with updated SOP assumptions (i.e., 5% of residues are dislodgeable, 50% extracted in saliva, transfer coefficients of 6,000 and 16,700 cm <sup>2</sup> for children and adults, respectively). Inadequate deposition data collected in treated rooms in registrant study.                |
| Dermal Exposure From Surfaces   |   | 3900                       | 2700                       |   |
| Oral Exposure   |   | NE                         | 4100                       |   |
| <b>Total Crack &amp; Crevice (Sum of 1 and 2) Inhalation, Dermal and Oral</b>   |   | 390 (1 day)<br>440 (10day) | 110 (1 day)<br>240 (10day) | Central-tendency risk estimates. Inhalation estimates are central-tendency to high end, but dermal and oral exposure estimates are low end.   |
| <b>(3) Pet Collar Uses (11 month efficiency) (Long-term)</b>  |   |                            |                            |   |
| Dog Collar ( EPA No. 45087-49; 3.44 g ai); Cat Collar (EPA No. 4306-16; 0.93 g chlorpyrifos)  |   |                            |                            |   |
| Total Exposure  | Residential SOPs  | 670 (dog)<br>2500 (cat)    | 140 (dog)<br>530 (cat)     | Central-tendency to high-end risk estimates; assume that a total of 1% ai is available from collar over 11 months only from dermal exposure. Assumes incidental ingestion and inhalation are negligible. Based on preliminary data, equivalent to approximately 2 , 3 <u>or</u> 105 min per day of vigorous dermal contact with collar, neck fur <u>or</u> back fur over 11 months. |

**Table 15. Estimates of Post-Application Risks to Residents/Recreational Users**

| Reentry Scenario   | Method of Evaluation  | Central-tendency MOE      |                        | Risk Characterization/<br>Uncertainties  |
|--|---|---------------------------|------------------------|--|
|  |   | Adult                     | Child                  |  |
| <b>(4) Termiticide Treatment Includes Risk Mitigation (adjustment to 0.5% ai as Dursban TC) (Intermediate and Long-term) (See Table A-1, Appendix A)</b> |   |                           |                        |  |
| <b>Basement Construction</b>   |   |                           |                        |  |
| 90-Day Incremental Time-weighted- average (TWA)  | Registrant study that collected air measurements in 7 homes from 7 days to 1 year post-treatment. | 13,000<br>(2,100-30,000)  | 3800<br>(600-8700)     | Median MOE with range of MOEs presented in parentheses. Values adjusted from 1% ai (typical rate) to 0.5% ai (minimum rate). Assumes a child spend 20 hours in a treated residence.  |
| 1-Year Incremental TWA   |   | 3,800<br>(930-8,800)      | 1,100<br>(270-2,500)   |  |
| <b>Crawl-Space-type Construction</b>   |   |                           |                        |  |
| 90-Day Incremental Time-weighted- average (TWA)  | See comments under basement construction.   | 7,300<br>(3,300-25,000)   | 2,100<br>(950-7,200)   | See comments under basement construction.  |
| 1-Year Incremental TWA   |   | 1,800<br>(1,200-7,400)    | 530<br>(340-2,100)     |  |
| <b>Slab Type Construction</b>  |   |                           |                        |  |
| 90-Day Incremental Time-weighted- average (TWA)  | See comments under basement construction.   | 6,600<br>(1,500-20,000)   | 1,900<br>(440-5,800)   | See comments under basement construction.  |
| 1-Year Incremental TWA   |   | 2,100<br>(960-7,600)      | 600<br>(280-2,200)     |  |
| <b>Plenum-Type Construction</b>  |   |                           |                        |  |
| 90-Day Incremental Time-weighted- average (TWA)  | See comments under basement construction.   | 6,600<br>(1,600 - 22,000) | 1,900<br>(460 - 6,400) | See comments under basement construction. 1-Year incremental TWA based on five houses, due to insufficient sampling for two houses. Sampling not conducted beyond days 30 and 7 for houses P-6 and P-7, respectively. Based on available data, these houses had higher air concentrations than the other houses. |
| 1-Year Incremental TWA   |   | 2,600<br>(940-9,500)      | 760<br>(270-2,700)     |  |

**Table 15. Estimates of Post-Application Risks to Residents/Recreational Users**

| Reentry Scenario  | Method of Evaluation  | Central-tendency MOE                      |                                  | Risk Characterization/<br>Uncertainties   |
|---|---|---|----------------------------------|---|
|   |   | Adult                                     | Child                            |   |
| <b>(5) Insecticidal Dust Products (Insufficient data to evaluate; see text)</b>                             |   |   |                                  |   |
| <b>Broadcast Turf Application (Residential/Recreational) (Short-term)</b>                                   |   |   |                                  |   |
| <b>(6) Chlorpyrifos Spray (Dursban Turf Insecticide)</b>  |   |   |                                  |   |
| Inhalation  | Biomonitoring Study, with environmental measurements. Application of 0.29% chlorpyrifos spray at 4 lb ai/acre | 170                                       | 20                               | Average represents central-tendency risk estimates based on arithmetic mean exposure from biomonitoring study in adults, where chlorpyrifos applied at the maximum label rate of 4 lb ai/acre. Based on 2 hour dermal contact with lawn the day of treatment. Maximum represents the highest exposed individual in the study. Study does not adequately address frequent hand to mouth activity of children, or incidental ingestion of soil or residues on treated grass by children. Application at typical rate of 1 lb ai/acre would potentially result in lower exposures (see below). |
| Dermal  |   | 10  | 12                               |   |
| Oral  |   | NE  | 400                              |   |
| Total Absorbed Dose   |   | Average: 9 -24<br>Maximum: 5.6-15         | Average: 7.5-15<br>Maximum: 6-12 |   |
| Total Absorbed Dose   | Biomonitoring Study with adjustment for 1 lb ai/acre  | Average: 36-96                            | Average: 30-60                   | Low to Central-tendency risk estimates, based on typical application rate of 1 lb ai/acre.  |
| <b>(7) Granular Formulation of 0.5% Chlorpyrifos (Dursban Insecticide) (1.8 lb ai/acre)</b>                 |   |   |                                  |   |
| Inhalation  | Biomonitoring Study, with environmental measurements  | 330                                       | 400                              | Average represents central-tendency risk estimates based on arithmetic mean exposure from biomonitoring study in adults. Based on 2 hour dermal contact with lawn the day of treatment; does not adequately address frequent hand to mouth activity of children, or incidental ingestion of soil or granules by children. Maximum MOE is for the highest exposed individual in the study.   |
| Dermal  |   | 190                                       | 90                               |   |
| Oral  |   | NE  | 6000                             |   |
| Total Absorbed Dose   |   | Average: 110-120<br>Maximum: 42-45        | Average: 73-75<br>Maximum: 29    |   |
| <b>(8) Golf Course Treatment (Dursban Turf Insecticide; EPA Reg 62719-35) (1-4 lb ai/acre) (Short-term)</b> |   |   |                                  |   |
| Adolescent Golfer (12 yrs; 44kg)  | Residential SOPs and surrogate residue data from flurprimidol study the day of treatment                      | 360 (4 lb ai/acre)<br>1500 (1 lb ai/acre) |                                  | High-end risk estimates. Assumes exclusively dermal exposure the day of turf treatment. Assumes a 4 hour exposure for a 18 hole round of golf.  |

**Table 15. Estimates of Post-Application Risks to Residents/Recreational Users**

| Reentry Scenario | Method of Evaluation | Central-tendency MOE                      |       | Risk Characterization/<br>Uncertainties |
|------------------|----------------------|---|-------|---|
|                  |                      | Adult                                     | Child |   |
| Adult Golfer     |                      | 600 (4 lb ai/acre)<br>2400 (1 lb ai/acre) |       |   |

**Table 15. Estimates of Post-Application Risks to Residents/Recreational Users**

| Reentry Scenario   | Method of Evaluation   | Central-tendency MOE |   | Risk Characterization/<br>Uncertainties   |
|--|--|----------------------|---|---|
|  |  | Adult                | Child   |   |
| <b>(9) Aerial and Ground-Based Fogger Mosquitocide Application (Mosquitomist One, EPA Reg. 8329-24) (0.01 lb ai/acre) (Short-term)</b> |  |                      |   |   |
| Dermal   | Literature studies, the AgDrift Model and the updated Residential SOPs | 42,000               | 26,000  | High-end risk estimates based on the updated Residential SOPs. Assumes long-term inhalation exposure is negligible based on low application rate and infinite dilution.   |
| Oral (hand to mouth)   |  | NE                   | 13,000  |   |
| Oral (Turfgrass Ingestion)   |  | NE                   | 54,000  |   |
| Oral (Soil Ingestion)  |  | NE                   | 20,000,000  |   |
| Total Exposure   |  | 42,000               | 15,000  |   |
| <b>(10) Yard and Ornamental Sprays (Evaluated based on analogy to Lawn Products; see text)</b>   |  |                      |   |   |
| <b>(11) Perimeter Treatment of Residence (Dursban Pro, EPA Reg. 62719-166) (4.35 lb ai/acre) (Short-term)</b>                          |  |                      |   |   |
| Dermal   | Updated Residential SOPs Residential                                   | NE                   | 8 minutes of play is equivalent to a MOE of 1000      | High-end risk estimates based on the updated Residential SOPs. Assumes a child plays on treated turf the day of treatment. The most critical items are the probability that a child would play within 6 to 10 feet of a residence and for what duration a child would be in the treatment zone. |
| Oral (hand to mouth)   |  | NE                   | 7 hand to mouth events is equivalent to a MOE of 1000 |   |
| Oral (Soil Ingestion)  |  | NE                   | MOE = 2300  |   |

#### 4.4.4.4 Incident Reports

Chlorpyrifos is one of the most widely used insecticides in the home both by consumers and PCOs or exterminators. In a 1990 EPA-sponsored survey of pesticide use in households, chlorpyrifos was the fourth most commonly used insecticide, present in 18% of all households. A 1993 EPA survey of PCOs found it was the number one insecticide in use and accounted for a quarter of the poundage used in residential settings. Consequently, there have been many reports of human exposure and poisonings due to the widespread use of chlorpyrifos. The human poisoning incidents associated with chlorpyrifos exposure have been evaluated and summarized in the attached memorandum from J. Blondell to D. Smegal, April 20, 2000. HED notes that approximately 98% of chlorpyrifos exposures discussed below are due to products removed under the risk mitigation plan.

Data from the Nation's Poison Control Centers in 1996 reported approximately 116,000 unintentional exposures to all pesticides, of which, 16% were due to organophosphate (OP) pesticides, and 5,188 or 4.5% were attributed to chlorpyrifos. These numbers are based on exposures to single products, a small proportion of which may contain additional active ingredients besides chlorpyrifos. Given that 30% of the organophosphate poisonings were not specifically identified by active ingredient, the actual number of chlorpyrifos cases is probably close to 7,000 or 6% of all pesticide-related exposures. Many of these exposures involve small children who were exposed but never developed symptoms. In 1996 there were 1,109 symptomatic cases related to chlorpyrifos that were judged to have effects related to the exposure, although most (83%) had only minor symptoms (e.g., headache, nausea, vomiting, dizziness and diarrhea) that could be treated at home. From 1993 through 1996, there were an average of 116 unintentional chlorpyrifos cases per year with moderate to severe outcomes (including one fatality) reported in residential settings.

The possibility of risk from chlorpyrifos exposure is very similar to the other OP pesticides (e.g., diazinon, malathion, dichlorvos) that have significant residential uses for both children and adults. The one exception is the percent of cases with fatal or life-threatening outcome (not including suicide attempts), where chlorpyrifos had the highest percentage (0.46% based on 18 cases) of any of the other 13 OP pesticides, that was 50% higher than any of the non-OP pesticides. Between 1993 and 1996, there was one fatality and 34 life-threatening cases attributed to chlorpyrifos exposure. The fatality was a 22 month old boy who accidentally ingested chlorpyrifos that had



been placed in a cup. Measures called for in the 1997 Chlorpyrifos Risk Reduction Plan, in part, were aimed at preventing such poisoning incidents.

Chlorpyrifos ranked third of the 13 OPs for serious outcomes resulting from exposure to environmental residues left after application or use. Environmental residues accounted for 15% of the chlorpyrifos exposures and 30% of the cases with serious outcomes (moderate or life-threatening), which was double the incidence for non-OP pesticides.

A particular concern with chlorpyrifos are reports of exposures and poisonings related to use by PCOs. A review of the Poison Control Center data for four years (1993-1996) found over 1000 reports of exposure (250 per year) to chlorpyrifos products that would most commonly be used by PCOs in residential settings. A total of 325 of these cases were symptomatic, 241 cases were seen in a health care facility, 35 were hospitalized and 16 were admitted to an intensive care unit (ICU). Chlorpyrifos PCO products accounted for 9% of the exposures, but 21-24% of the life-threatening/fatal cases, hospitalized cases and cases seen in an ICU. Note that the number of cases involving PCO products is relatively small compared to the exposure and symptomatic cases involving consumer products. Just 4% of the product-identified chlorpyrifos exposures in children under age six involved PCO products, and for adults and children over age six the figure was 15%. Also, some of the more serious cases, both for PCO and homeowner products, were due to broadcast carpet treatment, fogger and pet uses that were voluntarily canceled in 1997.

Another source of concern with all the OP pesticides, including chlorpyrifos, are the frequent anecdotal reports of chronic neurobehavioral effects and multiple chemical sensitivity. Kilburn (1999) documented neurobehavioral effects (including signs consistent with peripheral neuropathy in 11 cases) among 22 patients reporting exposure to chlorpyrifos, 10 of which were self-referred and 12 referred by attorneys. In addition to these reports, there were 14 self-reported but unconfirmed cases (without medical documentation) of chronic neurobehavioral effects submitted by Dow AgroSciences during 1998-1999. Another 73 cases were reported to EPA during the public comment period (October-December 1999) for chlorpyrifos. A few of these cases may have overlapped the reports from Kilburn and Dow AgroSciences. Twelve of the 73 cases provided some, often very limited, medical documentation of their effects. Out of all of the cases reported by Kilburn, Dow AgroSciences or directly to EPA there were only about 3-4 with laboratory confirmation (e.g., reduced cholinesterase) of their

exposures. Neurobehavioral effects reported include persistent headaches, blurred vision, muscle weakness, fatigue, and problems with mental function including memory, concentration, depression, and irritability.

HED suspects that these chronic neurobehavioral effects are caused by the acute poisoning, partly from a case-control study in California partly from case-control (cross sectional) studies of other OP pesticides similar to chlorpyrifos, and most recently from a NIOSH study. With EPA support, NIOSH completed a study of 191 current and former PCOs that apply chlorpyrifos as a termiticide in North Carolina. An extensive battery of neurological and neurobehavioral tests was administered. The study (Steenland et al. 2000), concluded "this cross-sectional study of workers exposed to chlorpyrifos . . . found few exposure related effects for most tests, including a clinical exam. However, the exposed did not perform as well as the non-exposed on pegboard turning tests and some postural sway tests. Furthermore, exposed subjects reported more symptoms than non-exposed subjects; this is a cause for concern because previous studies lend some support to this finding." Among acutely poisoned subjects the study stated, "Eight men who reported past chlorpyrifos poisoning had a pattern of low performance on a number of tests, which is consistent with prior reports of chronic effects of organophosphate poisoning." Finally, the study noted the following reservation, partly due to the relatively heavy exposure experienced by study participants, "Although this was a relatively large study based on a well-defined target population, the workers we studied may not be representative of all exposed workers and caution should be exercised in generalizing our results." (Steenland et al. 2000). These findings are consistent with an earlier review that suggested chlorpyrifos may be a cause of chronic neurobehavioral effects in some subsets of sensitive people who have been poisoned (Blondell and Dobozy 1997). In addition to the studies described above, DAS has agreed to undertake an epidemiologic study of manufacturing workers.

As noted previously, four uses of chlorpyrifos have been voluntarily canceled and removed from the market: paint additives; shampoos, sprays and dips used on pets; indoor broadcast flea control products; and household foggers. Poison Control Center data for 1993-1996 suggest that as many as 20-25% of symptomatic exposures in residential settings were related to these uses. All of these residential uses involve either concentrates or widespread applications that involve greater potential for exposure to consumers than do other forms and uses of chlorpyrifos. Therefore, substantially less exposures and hazards are expected when additional years of

poisoning surveillance data become available. DAS is continuing its' efforts to monitor poisoning incidents through its agreement with a Poison Control Center that takes telephone contacts from the public and the health care community concerning chlorpyrifos. Follow up information to determine the circumstances that lead to exposure and poisoning should be useful.

#### 4.4.5 Pet Incident Reports

A review and analysis of the poisoning incident reports on domestic animals for chlorpyrifos was conducted in 1995 (attached memo from V. Dobozy to B. Kitchens, January 23, 1995) and was updated in 1999 (attached memo from V. Dobozy to D. Smegal, April 26, 1999, D255514). In the 1995 analysis, poisoning incidents in dogs and cats were categorized as exposure by direct applications (flea and tick dips, sprays, collars, etc) or by premise applications (household and lawn treatments). The analysis found that the majority of the incidents in domestic animals involved cats, although the chemical is registered only for use in flea collars for this species. Cats that were exposed to products registered only for use on dogs, mainly dips, experienced a high incidence of death (30%). There was also evidence of misuse of treatment products, including practices such as applying these products directly to animals and not removing pets from premises during applications.

In 1996, PR Notice 96-6 was finalized, which requires the revision of labels for all products administered directly to animals to ensure adequate directions for use and warning information. In 1997, the registrant voluntarily agreed to cancel chlorpyrifos registrations for indoor broadcast flea control and direct application pet products (sprays, shampoos, and dips), except flea collars, to establish specific protection measures for pets during and immediately after application, and to expedite implementation of PR Notice 96-6 on pet products.

An evaluation of incident reports for domestic animals for the years 1996 through 1998 (memo from V. Dobozy to D. Smegal, April 26, 1999, D255514) revealed that there has been a decrease in the percentage of incidents resulting from exposure to products registered for direct use on animals, but an increase in the percentage of incidents resulting from premise exposure. In addition, deaths are still being reported, especially for cats. The cancellation of indoor broadcast flea control applications and products for direct application to dogs and cats should reduce the risk of serious adverse reactions and deaths, however time is required to eliminate all chlorpyrifos products from store shelves. Therefore, it may be premature to review the Incident Data System (IDS) for evidence that these actions were effective.

#### 4.5 Chlorpyrifos Exposure Estimates in the U.S. Population

Because of chlorpyrifos' extensive use on food and in homes and the workplace, the majority of the U.S. population is exposed to this pesticide. Literature studies, in addition to several of the registrant-submitted biomonitoring studies, have estimated typical or baseline exposure to chlorpyrifos by measuring the urinary excretion of 3,5,6-TCP, the primary metabolite of chlorpyrifos. TCP has a biological half-life of approximately 27 hours, therefore, the urinary TCP levels reflect recent exposure. It should be noted however, that exposure to chlorpyrifos-methyl, 3,5,6-TCP (the animal, and plant metabolite and environmental degradate of chlorpyrifos and chlorpyrifos-methyl), and trichlorpyr (a herbicide) also contribute to an unknown degree to 3,5,6-TCP urinary concentrations, thus the chlorpyrifos exposure estimates presented in this section represent an upper-bound estimate. Chlorpyrifos contributes significantly more to urinary TCP than chlorpyrifos-methyl and trichlorpyr based on relative annual U.S. usage of approximately 21 to 24 million pounds of chlorpyrifos (of which approximately 11 million are used in residential and recreational settings) versus 92,000 pounds of chlorpyrifos-methyl and 700,000 pounds of trichlorpyr.

HED has conducted a preliminary risk assessment for TCP, which is in the attached memorandum from S. Knizner to D. Smegal, D265035 June 5, 2000.

Table 16 summarizes the typical upper-bound baseline exposure to chlorpyrifos estimated from the registrant submitted biomonitoring studies of TCP measurements, and the scientific literature. These values represent worst case estimates because all of the TCP was attributed to chlorpyrifos.

##### Registrant Residential Biomonitoring Studies

DAS recently conducted four biomonitoring studies to quantify exposures to residential populations following the use of chlorpyrifos products in the home. Volunteers were typically adults of both sexes between the ages of 25 and 65. Other details were not provided (i.e., ethnicity). For all of these studies, baseline chlorpyrifos exposures of the volunteers were quantified by analysis of urinary 3,5,6-TCP prior to commencement of the study. Quantification of baseline chlorpyrifos exposure for each volunteer was necessary in order to determine actual exposure associated with a product's use. For each of these studies, baseline TCP measurements were subtracted from total TCP measurements to quantify chlorpyrifos exposure in the biomonitoring study. In addition, residents were instructed to avoid chlorpyrifos exposure for several days (typically one week to 10 days) prior to the measurement of baseline levels. Therefore, the baseline exposures are most likely attributed to dietary exposure of chlorpyrifos, chlorpyrifos-methyl and TCP.

In August 1999, DAS submitted a TCP Biomonitoring study that assesses children's potential household exposure to chlorpyrifos and its environmental degradate, TCP (MRID 44889501). The study evaluated urinary TCP concentrations of 416 children 0-6 years of age in North and South Carolina; 120 children were from households treated with a termiticide containing chlorpyrifos, and 296 children were from households identified from the general population sample. TCP was detected in 100% of the children's urine. The 24 hour TCP excretion ranged from 0.09 to 75.79 Fg TCP/g creatinine/kg body weight, with a mean value of 1.19 Fg TCP/g creatinine/kg body weight. These values correlate to approximately 0.045 to 38 Fg chlorpyrifos /kg/day, with a mean value of 0.6 Fg/kg/day. It should be noted that 73% (303/413) and 11% (47/413) of the children in this survey lived in homes that had been treated with a chlorpyrifos-containing insecticide indoors or with a termiticide, respectively within the past year. In addition, 64% of the children (264/412) also were from homes that had a lawn treatment within the past year. HED is currently reviewing this study.

### **Scientific Literature**

The study published by Hill et al. (1995) measured the biomarker 3,5,6-TCP in 993 adults (20-59 years old) participating in the National Health and Nutrition Examination Survey III, known as NHANES III from 1988 - 1994. The individuals were selected from a broad spectrum of the U.S. population reflecting both sexes and different age groups, races/ethnicities, urban/rural residences and regions of the country. 3,5,6-TCP was detected in 82% of the individuals evaluated. The average TCP concentration was 4.5 Fg/L or 3.1 Fg TCP/g creatinine. The results of NHANES III differ significantly from the NHANES II survey collected between 1976 and 1980, where only 5.8% of the 6990 people evaluated had concentrations of 3,5,6-TCP greater than the detection limit of 5 Fg/L. In the NHANES III survey, 31% of the 993 people had 3,5,6-TCP concentrations greater than 5 Fg/L. It should be noted however, that the lower detection limit of 1 Fg/L in the NHANES III study could partially account for the increased frequency of detection of 82%. The results of this study are presented below in Table 14. It is possible that the registration of chlorpyrifos-methyl for use on stored grains in 1985 contributes to the increased frequency and concentration of TCP measurements between the NHANES II and III results. In addition, chlorpyrifos-methyl was detected at greater frequencies than chlorpyrifos in the 1991-1997 Total Diet Study (FDA 1999). In this study, 100% of samples for several commodities containing flour (i.e., whole wheat bread, tortilla flour, rye bread, cracked wheat bread, english muffin, teething biscuits, pretzels, fish sticks, white roll, and butter type crackers) contained measurable chlorpyrifos-methyl residues.

A recent study of 65 recently-exposed termiticide applicators (Steenland et al. 2000) reported an average urinary TCP level of 629.5 Fg/L, compared to the 4.5 Fg/L for the general U.S. population from Hill et al. (1995).

The Minnesota Children's Pesticide Exposure Study, which is one of the National Human Exposure Assessment Surveys (NHEXAS), evaluated 102 children ages 3-12 (mean  $7.6 \pm 2.9$  yrs), stratified by those with more frequent residential insecticide usage (personal communication with James Quackenboss, March 1, 1999). This study was initiated to assess children's actual exposures to pesticides. The study examined the relationship between environmental concentrations and urinary biomarker levels of 3,5,6-TCP from a population-based study of total exposure in urban and non-urban children. Tap water, personal, indoor, and outdoor air, house dust, and soil were monitored over 6 days while food and beverage monitoring was conducted over 4 days. Urine samples were obtained for 87% (89) of the study subjects. Preliminary data were presented at the International Society for Environmental Epidemiology (ISEA) conference in Boston in August 1998 (Adgate et al. 1998), where 92% of the 89 children had measurable levels of 3,5,6-TCP in their urine. It should be noted, however, that the study over sampled homes that frequently used pesticides, and 30% of the households had used chlorpyrifos. The results from the metabolite analysis suggest that these children have higher concentrations of 3,5,6-TCP than was reported for the NHANES-III adult population (medians of 8 and 2 Fg/L TCP, respectively) (Quackenboss et al. 1998). The final study results are anticipated to be available in 2000.

Macintosh et al. (1999) evaluated urinary TCP levels in 80 individuals in Maryland during 1995-1996. Up to six samples were collected from each individual over a period of a year. TCP was detected in 96% of the 346 samples at a median concentration of 5.3 Fg/L and 4.6 Fg/g creatinine. The geometric mean concentrations of TCP were significantly greater in samples collected during the spring and summer of 1996 than in the preceding fall and winter. In addition, the geometric mean TCP concentrations differed significantly between Caucasian (GM = 5.7 Fg/g creatinine) and African-American (GM = 4 Fg/g creatinine) participants and among education levels but were not significantly different among groups classified by gender, age, or household income. The mean and median TCP concentrations in this study (5.8 and 4.6 Fg/g creatinine) are approximately twofold greater than those measured in the NHANES III (3.1 and 2.2 Fg/g creatinine, respectively) (Hill et al. 1995), however the upper end of the distributions are approximately equal. Individual urinary TCP levels varied over time and were highly variable, indicating that a single measure of urinary TCP levels is not sufficient to adequately characterize the relative magnitude of a person's typical exposure to chlorpyrifos.

Buckley et al. (1997) evaluated 18 nonsmoking adults from nine homes in the Lower Rio Grande Valley (LRGV) in Texas during the spring and summer 1993. Urinary TCP was significantly higher in the summer relative to the spring, and was correlated with air and dust concentrations. TCP was detected in 77% (13/17) and 92% (11/12) of the spring and summer samples, respectively at median concentrations of 1.9 and 3.2 Fg/L, respectively.

Table 16 summarizes the typical upper-bound baseline exposure to chlorpyrifos estimated from the Hill et al. (1995) and DAS biomonitoring studies of TCP measurements. These values represent worst case estimates because all of the TCP was attributed to chlorpyrifos. All exposure estimates have been normalized for creatinine excretion. The assumptions and equations are presented in the footnotes.

**Table 16  
Upper Bound Chlorpyrifos Exposure Estimates Based on  
Biomonitoring of Urinary TCP**

| Source/Study   | Sample Size             | Percent with TCP in urine  | Mean Chlorpyrifos Dose Fg/kg/day | 95 <sup>th</sup> Percentile Fg/kg/day | Range of Chlorpyrifos Dose Fg/kg/day |
|--|-------------------------|----------------------------|----------------------------------|---------------------------------------|--------------------------------------|
| <b>Residential Biomonitoring Studies</b>   |                         |                            |                                  |                                       |                                      |
| Child TCP Biomonitoring study (0-6 yrs old, North and South Carolina, 1998) (a)                          | 416                     | 100%                       | 0.6                              | 1.32                                  | 0.045-4.7                            |
| Residential exposures from Lawn treated with Chlorpyrifos Spray (MRID 43013501) (Adults) (b)             | 8                       | 100%                       | 0.3                              | NE                                    | 0.09 - 0.6                           |
| Residential Exposures from Lawn treated with Granular Chlorpyrifos (MRID 44167101) (Adults) (b)          | 9                       | 100%                       | 0.5                              | NE                                    | 0.21 - 1.47                          |
| Residential Exposure from Crack and Crevice Application (MRID 44458201) (Adults) (b)                     | 6                       | 100%                       | 0.4                              | NE                                    | 0.1-0.86                             |
| Residential Exposures from Application of a Ready-to-Use Formulated Product (MRID 44739301) (Adults) (b) | 15                      | 100%                       | 0.12                             | NE                                    | 0.05-0.3                             |
| <b>Literature Studies</b>  |                         |                            |                                  |                                       |                                      |
| Hill et al. 1995 (NHANES III) (Adults, 1988-1994) (c)  | 993                     | 82%                        | 0.2 (b)                          | 0.52                                  | ND - 2                               |
| MacIntosh et al. 1999 (Adults, Maryland, 1995-1996) (d)  | 80 people (329 samples) | 96%                        | 0.37                             | 1                                     | 0.013-2.2                            |
| Buckley et al. (1997) (Adults, Texas, 1993) (e)  | 18                      | Spring: 77%<br>Summer: 92% |                                  |                                       |                                      |

ND = not detected

NE = not estimated



- (a) Creatinine adjusted concentrations for 24 hour TCP excretion ranged from 0.09 to 15.8 Fg TCP/g creatinine/kg body weight, with a mean value of 1.19 Fg TCP/g creatinine/kg. In the initial study, the highest child was 75.79 Fg TCP/g creatinine/kg, which is equal to approximately 38 Fg/kg/day chlorpyrifos. A more recent submission, March 2000, reported lower levels of TCP in this child of 15.8 Fg TCP/g creatinine/kg, which is equivalent to approximately 4.7 Fg/kg/day chlorpyrifos. The 95<sup>th</sup> percentile was 2.63 Fg TCP/g creatinine/kg. Assumes child specific body weight, and average creatinine excretion of 0.2 g/day from 416 children. Assumes steady-state between exposure and excretion.
- (b) Based on pre-study 3,5,6-TCP results in urine. See HED study reviews for details
- (c) Creatinine adjusted concentrations of mean 3.1 and maximum of 34 Fg TCP/g creatinine, respectively that assumes an average creatinine excretion rate of 1.8 g/day (Tietz 1982), a body weight of 70 kg, and that 72% of chlorpyrifos is excreted in the urine. A molecular weight adjustment was also made 350.6 chlorpyrifos/ 198 TCP. Assumes steady-state between exposure and excretion. Example calculation: Dose (Fg/kg/day) =  $[(3.1 \text{ Fg TCP/g creatinine} * 350.6/198 * 1.8 \text{ g/day}) / (70 \text{ kg} * 0.72 \text{ (fraction chlorpyrifos excreted as TCP)})]$ .
- (d) creatinine adjusted concentrations of <0.2, 5.8, 16 and 35 Fg TCP/g creatinine for minimum, mean, 95<sup>th</sup> percentile and maximum, respectively. Assumes an average creatinine excretion rate of 1.8 g/day (Tietz 1982), a body weight of 70 kg, and that 72% of chlorpyrifos is excreted in the urine. A molecular weight adjustment was also made 350.6 chlorpyrifos/ 198 TCP. Example calculation: Dose (Fg/kg/day) =  $[(35 \text{ Fg TCP/g creatinine} * 350.6/198 * 1.8 \text{ g/day}) / (70 \text{ kg} * 0.72 \text{ (fraction chlorpyrifos excreted as TCP)})]$ .
- (e) Creatinine adjusted concentrations not presented. Median TCP concentrations of 1.9 and 3.2 Fg/L and maximum concentrations of 6.4 and 11 Fg/L for spring and summer, respectively.

## 5.0 Aggregate Risk Assessments and Risk Characterization

The Food Quality Protection Act amendments to the Federal Food, Drug, and Cosmetic Act (FFDCA, Section 408(b)(2)(A)(ii)) require that for establishing a pesticide tolerance "that there is reasonable certainty that no harm will result from aggregate exposure to pesticide chemical residue, including all anticipated dietary exposures and other exposures for which there are reliable information." Aggregate exposure is the total exposure to a single chemical (or its residues) that may occur from dietary (i.e., food, and drinking water), residential and other non-occupational sources, and from all known or plausible exposure routes (oral, dermal and inhalation). Aggregate risk assessments are typically conducted for acute (1 day), short-term (1-30 days), intermediate-term (30 days to several months), and chronic (several months to lifetime) exposure.

DAS has submitted a probabilistic Integrated Exposure Assessment (MRID No. 44104001, September 1996). This submission is in internal HED review, because the Agency policy on aggregate probabilistic risk assessment is still in development. This submission, however, has been used by the Agency in developing policy and will be evaluated once this policy is finalized and has undergone peer review.

The total residential MOEs (dermal, inhalation, and inadvertent oral exposures) for all the residential post-application exposure scenarios, except mosquitocide use, and golf course use alone exceed HED's level of concern. In addition the acute dietary exposure and risk estimates exceed HED's level of concern. However, HED conducted acute, short-term and chronic aggregate assessments assuming the mitigation plan is adopted. As noted previously, the mitigation plan would reduce potential chlorpyrifos exposures on apples, grapes and tomatoes, and mitigate the residential/recreational exposures.

### 5.1 Acute Aggregate Risk

The acute aggregate risk estimate to chlorpyrifos addresses exposures from food and drinking water. For the highly refined acute probabilistic dietary exposure analysis, PDP, FDA and NFS monitoring data were used to the greatest extent possible, along with field trial data, and cooking and processing factors to assess dietary exposures. This aggregate assessment incorporates the mitigation plan (i.e., reduction of apple tolerance to 0.01 ppm based on dormant application, reduction of grape tolerance to 0.01 ppm based on domestic use pattern and deletion of the use on tomatoes).

With the mitigation measures, the chlorpyrifos acute dietary risk estimates range from 4.1% to 82% of the aPAD, with children (1-6 yrs) being the highest exposed population subgroup. Thus, the mitigated acute dietary (food) risk estimate associated with chlorpyrifos exposure is below the Agency's level of concern. Using conservative screening-level models, the acute estimated concentrations (EECs) of chlorpyrifos in groundwater (SCI-GROW) range from 0.007 to 0.103 Fg/L. The acute surface water EECs, based on upper-bound

monitoring data results, are 0.026 to 0.4 Fg/L, respectively. As shown previously on Table 7, and on Table 17 below, the EECs are less than the DWLOCs for all populations (highest EEC of 0.4 Fg/L is less than the lowest DWLOC of 0.9 Fg/L), indicating that acute food and drinking water exposures (except possible well contamination) do not exceed HED's level of concern. It should be noted that neither the SCI-GROW model nor the monitoring data reflect concentrations after dilution (from source to treatment to tap) or drinking water treatment. HED concludes that **acute aggregate chlorpyrifos exposure in food and water does not exceed HED's level of concern.**

| <b>Table 17<br/>                     Summary of Acute Aggregate Exposure<br/>                     Includes Risk Mitigation</b> |                       |                                      |                                     |  |  |                            |
|--|-----------------------|--------------------------------------|-------------------------------------|--|--|----------------------------|
| Population Subgroup (a)  | Acute PAD (Fg/kg/day) | Food Exposure 99.9th (Fg/kg/day) (b) | Max. Water Exposure (Fg/kg/day) (c) | Surface Water (Monitoring Data) (Fg/L) | Ground Water SCI-GROW, (excluding well contamination) (Fg/L) | Acute DWLOC (Fg/L) (d,e,f) |
| U.S. Population  | 5                     | 0.237                                | 4.76                                | 0.026 to 0.4                           | 0.007 to 0.103   | 166                        |
| All Infants (< 1 Year)   | 0.5                   | 0.258                                | 0.242                               |  |  | 2.4                        |
| Children (1-6 years)   | 0.5                   | 0.410                                | 0.09                                |  |  | 0.9                        |
| Females (13-50 years)  | 0.5                   | 0.201                                | 0.299                               |  |  | 9                          |

- (a) In addition to the U.S. population (all seasons), the most highly exposed subgroup within each of the infants, children, female groups is listed.
- (b) 99.9th percentile exposure. Values are from Table 3 (and rounded).
- (c) Maximum Water Exposure (Fg/kg/day) = Acute PAD (Fg/kg/day) - [Acute Food Exposure (Fg/kg/day)].
- (d) DWLOC (Fg/L) = Maximum water exposure (Fg/kg/day) x body wt (kg) ÷ water consumed daily (L/day)]
- (e) HED default body weights are: general U.S. population, 70 kg; adult females, 60 kg; and infants/children, 10 kg.
- (f) HED default daily drinking water rates are 2 L/day for adults and 1 L/day for children.

Acute exposure to chlorpyrifos in groundwater as a result of well contamination from termiticide use could potentially result in exposures of concern. However, as noted previously, the groundwater exposures from well contamination resulting from termiticide use are highly localized. The implementation of PR 96-7 for termiticides has reduced the reported incidents of groundwater contamination resulting from termiticide treatments. For example, incidents associated with termiticide use were 28.2 per 100,000 homes in 1997 (pre PR-96-7), and were 8.3

per 100,000 homes in 1998 (post PR-96-7).

## 5.2 Short-Term Aggregate Risk

The short-term aggregate risk estimate includes chronic dietary (food and water) from chlorpyrifos uses, and short-term non-occupational exposures (i.e., residential/recreational uses). As noted previously, this aggregate assessment is based on the mitigation plan that would reduce potential chlorpyrifos exposures in food (apples, grapes and tomatoes) and in the residential/recreational environment. This assessment evaluates potential exposures resulting from continued chlorpyrifos use on golf courses at a reduced rate of 1 lb ai/acre (i.e., risks to golfers), in addition to potential exposures as a result of mosquito abatement activities.

Table 18 presents the aggregate exposure estimates for chlorpyrifos from diet and residential/non-occupational uses (golfing and mosquitocide abatement activities). Based on the mitigation plan, it was assumed that children (1-6 years) could be exposed to chlorpyrifos residues on turf as a result of ground-based fogger applications of a chlorpyrifos-containing mosquitocide, and through dietary exposures. Children 7-12 years were assumed to be dermally exposed to chlorpyrifos residues while playing golf (the day of treatment), and to ingest chlorpyrifos residues in the diet. Female residents were assumed to be concurrently exposed to chlorpyrifos via mosquito abatement activities (i.e., dermal contact with residues on turf), golfing (dermal contact turf residues the day of treatment), in addition through dietary exposures. The results of the exposure analysis for the individual scenarios are presented in detail in the Occupational /Residential Exposure Chapter for the RED for Chlorpyrifos (D266562, June 2000).

As shown on Table 18, aggregate MOEs are greater than 1000 for children 1-6 years, children 7-12 years and females 13-50 years, and therefore do not exceed HED's level of concern. Therefore, short-term DWLOCs were estimated to account for potential drinking water exposures.

**Table 18**  
**Summary of Aggregate Short-Term Exposure**  
**Chronic Diet and Short-Term Residential Use**  
**(Excludes Water)**  
**Includes Risk Mitigation**

| Population Subgroup   | Dietary Exposure with Risk Mitigation                  | Short-Term Residential/Recreational Exposure (Fg/kg/day)/ MOE Risk Mitigation |                         |   | Total Aggregate MOE Estimate (b)           |
|-----------------------|--|---|-------------------------|---|--|
|                       |  | Mosquitocide Postapplication  |                         | Golf Course Postapplication Exposure (1 lb ai/acre) | Diet and Residential/Recreational Exposure |
|                       | Chronic Diet Exposure with FHE (Fg/kg BW/day) (a)/ MOE | Oral  | Dermal                  | Dermal  | Oral and Dermal                            |
| Children (1-6 years)  | 0.008<br>MOE = 62,500                                  | 0.013<br>MOE = 38,500   | 0.19<br>MOE = 26,000    | NE  | 12,000                                     |
| Children (7-12 years) | 0.015<br>MOE = 33,000                                  | NE  | NE                      | 3.4<br>MOE = 1,500                                  | 1,400                                      |
| Females 13-50         | 0.006<br>MOE = 83,000                                  | NE  | 0.14 (c)<br>MOE= 36,000 | 2.45 (c)<br>MOE = 2,000                             | 1,900                                      |

NE = not evaluated.

FHE = Food Handling Establishment Use

- (a) MOE calculated based on acute oral NOAEL of 500 Fg/kg/day, and short-term dermal NOAEL of 5000 Fg/kg/day for dermal exposures. No dermal absorption is necessary because dermal NOAEL is based on a dermal rat study.
- (b) Oral and dermal exposures were combined because the oral and dermal endpoints are both based on plasma and RBC ChE inhibition.
- (c) Adjusted from 70 kg to 60 kg for aggregate exposure.

The short-term DWLOC values are presented in Table 19. For each population subgroup listed, the acute PAD and the chronic dietary (food) exposure (from Table 4) for that subgroup were used to calculate the short-term DWLOC for the subgroup, using the formulas in footnotes of Table 19. The EECs are less than the DWLOCs for all populations (highest EEC of 0.1 Fg/L is less than the lowest DWLOC of 1.4 Fg/L), indicating that chronic food and drinking water exposures (except possible well contamination), in addition to exposures from mosquitoicide abatement and golfing activities do not exceed HED's level of concern. In conclusion, potential **short-term aggregate exposure to chlorpyrifos resulting from food, water and residential/recreational use, assuming the mitigation plan is adopted, does not exceed HED's level of concern.** This analysis is considered conservative because, HED assumed that there could be concurrent residential and recreational exposures to chlorpyrifos (i.e., golfing and mosquitoicide abatement activities on the same day). In addition, neither the SCI-GROW model nor the monitoring data reflect concentrations after dilution (from source to treatment to tap) or drinking water treatment.

**Table 19**  
**Summary of Short-Term Aggregate Exposure DWLOCs**  
**Chronic Diet and Short-Term Residential Use**  
**Includes Risk Mitigation**

| Population Subgroup (a) | Acute oral NOAEL (Fg/kg/day) | Short-Term MOE (Food and Residential) (Fg/kg/day) (a) | MOE Water (b) | Max. Water Exposure (Fg/kg/day) c) | Surface Water (Monitoring Data) (Fg/L) | Ground Water SCI-GROW, (excluding well contamination) (Fg/L) | Short-Term DWLOC (Fg/L) (d,e,f) |
|-------------------------|------------------------------|---|---------------|------------------------------------|--|--|---------------------------------|
| Children (1-6 years)    | 500                          | 1,200   | 1,090         | 0.4587                             | 0.026                                  | 0.007 to 0.103   | 4.5                             |
| Children (7-12 years)   |                              | 1,400   | 3,450         | 0.14                               |  |  | 1.4                             |
| Females (13-50 years)   |                              | 1,900   | 2,100         | 0.238                              |  |  | 7.1                             |

- (a) Values are from Table 18.
- (b)  $MOE_{WATER} = 1 / [(1/MOE_{AGG} - [1/MOE_{FOOD} + 1/MOE_{DERMAL} + 1/MOE_{ORAL}])]$ , where  $MOE_{AGG}$  is 1000.
- (c) Maximum Water Exposure (Fg/kg/day) = Acute NOAEL of 500 (Fg/kg/day) ÷  $MOE_{WATER}$
- (d) DWLOC (Fg/L) = Maximum water exposure (Fg/kg/day) x body wt (kg) ÷ water consumed daily (L/day)]
- (e) HED default body weights are: adult females, 60 kg; and infants/children, 10 kg.
- (f) HED default daily drinking water rates are 2 L/day for adults and 1 L/day for children.



### 5.3 Intermediate-Term Aggregate Risk

Based on the mitigation plan, there are no residential/recreational uses that result in exclusively intermediate-term exposures (i.e., > 30 days but less than 6 months). Therefore, an intermediate-term aggregate risk estimate was not evaluated.

### 5.4 Chronic Aggregate Risk

The chronic aggregate risk estimate to chlorpyrifos addresses exposures from food and drinking water. For the highly refined chronic dietary exposure analysis, PDP, FDA and NFS monitoring data were used to the greatest extent possible, along with field trial data, and cooking and processing factors to assess dietary exposures. This aggregate assessment incorporates the mitigation plan (i.e., reduction of apple tolerance to 0.01 ppm based on dormant application, reduction of grape tolerance to 0.01 ppm based on domestic use pattern and deletion of the use on tomatoes), and assumes there are no chronic exposures from termiticide treatments.

The chlorpyrifos chronic noncancer dietary risk estimates range from 2.5 to 51% of the cPAD, with children (1-6 yrs) being the highest exposed population subgroup. Thus, the chronic dietary (food) risk estimate associated with chlorpyrifos exposure is below the Agency's level of concern.

Using conservative screening-level models the groundwater EECs range from 0.007 to 0.103 Fg/L. The upper-bound surface water EEC, based on monitoring data, is 0.026 Fg/L. As noted previously, DWLOCs were calculated based on food (including food handling establishment uses) and water exposure alone to account for the mitigation options. The chronic non-cancer DWLOC values were presented previously in Table 8, and are shown below on Table 20. For each population subgroup listed, the chronic PAD and the chronic dietary (food) exposure (from Table 4) for that subgroup were used to calculate the chronic DWLOC for the subgroup, using the formulas in footnotes of Table 20. As shown, the upper-bound EEC of 0.103 Fg/L is less than the DWLOCs, and therefore does not exceed HED's level of concern. It should be noted that neither the SCIGROW model nor the monitoring data reflect actual drinking water concentrations after dilution (from source to tap) or drinking water treatment.

| <b>Table 20</b><br><b>Summary of Short-Term Aggregate Exposure DWLOCs</b><br><b>Includes Risk Mitigation</b> |                         |  |                                     |                                      |   |                              |
|--|-------------------------|--|-------------------------------------|--------------------------------------|---|------------------------------|
| Population Subgroup (a)  | Chronic PAD (Fg/kg/day) | Chronic Food Exposure with FHE (Fg/kg/day) (b) | Max. Water Exposure (Fg/kg/day) (c) | Surface Water Monitoring Data (Fg/L) | Ground Water SCI-GROW (excluding well contamination) (Fg/L) | Chronic DWLOC (Fg/L) (d,e,f) |
| U.S. Population  | 0.3                     | 0.008  | 0.292                               | 0.026                                | 0.007 to 0.103  | 10                           |
| All Infants (< 1 Year)   | 0.03                    | 0.01   | 0.02                                |                                      |   | 0.2                          |
| Children (1-6 years)   | 0.03                    | 0.015  | 0.015                               |                                      |   | 0.15                         |
| Females (13-50 years)  | 0.03                    | 0.006  | 0.024                               |                                      |   | 0.72                         |

- (a) In addition to the U.S. population (all seasons), the most highly exposed subgroup within each of the infants, children, female groups is listed.
- (b) Values are from Table 4 (and rounded).
- (c) Maximum Water Exposure (Fg/kg/day) = Chronic PAD (Fg/kg/day) - [Chronic Food Exposure + Chronic Residential Exposure (Fg/kg/day) (if applicable)]. Chronic residential uses were not considered based on mitigation options.
- (d) DWLOC (Fg/L) = Maximum water exposure (Fg/kg/day) x body wt (kg) ÷ water consumed daily(L/day)]
- (e) HED default body weights are: general U.S. population, 70 kg; adult females, 60 kg; and infants/children, 10 kg.
- (f) HED default daily drinking water rates are 2 L/day for adults and 1 L/day for children.

As noted previously, long-term exposure to chlorpyrifos as a result of well contamination from termiticide use could potentially result in exposures of concern. However, the groundwater risk estimates from well contamination resulting from termiticide use are highly localized. The implementation of PR 96-7 for termiticides has reduced the reported incidence of groundwater contamination resulting from termiticide treatments.

Although not all of the risk estimates for termiticide use achieve a margin of exposure of 1000, the Agency believes that individuals are unlikely to experience adverse health effects from the termiticide use of chlorpyrifos. This conclusion is based on: the public health protective assumptions; the 1000 fold safety factor; and the additional 3 to 10 fold cushion between the NOAEL and the LOAEL. Mitigation measures will further reduce exposures and risk associated with the termiticide use. For example, the removal of whole house barrier treatment addressed the exposures of most concern. It is expected that the limited spot and localized treatment, and pre-construction treatments would represent less exposure and risk. In conclusion, based on the mitigation plan, and best professional and scientific judgement, the Agency concludes that the chronic aggregate risk including termiticide use, does not raise a concern.

## 6.0 Cumulative Exposure and Risks

The Food Quality Protection Act (1996) stipulates that when determining the safety of a pesticide chemical, EPA shall base its assessment of the risk posed by the chemical on, among other things, available information concerning the cumulative effects to human health that may result from dietary, residential, or other non-occupational exposure to other substances that have a common mechanism of toxicity. The reason for consideration of other substances is due to the possibility that low-level exposures to multiple chemical substances that cause a common toxic effect by a common mechanism could lead to the same adverse health effect as would a higher level of exposure to any of the other substances individually. A person exposed to a pesticide at a level that is considered safe may in fact experience harm if that person is also exposed to other substances that cause a common toxic effect by a mechanism common with that of the subject pesticide, even if the individual exposure levels to the other substances are also considered safe.

Chlorpyrifos is a member of the organophosphate (OP) class of pesticides. All pesticides of this class contain phosphorus and other members of this class of pesticides are numerous and include azinphos methyl, chlorpyrifos-methyl, diazinon, dichlorvos, dicrotophos, dimethoate, disulfoton, methamidophos, methidathion, monocrotophos, oxydemeton methyl, phorate, phosmet, and pirimiphos-methyl to name a few. EPA considers organophosphates to express toxicity through a common biochemical interaction with cholinesterase which may lead to a myriad of cholinergic effects and, consequently the organophosphate pesticides should be considered as a group when performing cumulative risk assessments. HED recently published the final guidance that it now uses for identifying substances that have a common mechanism of toxicity (FR 64(24) 5796-5799, February 5, 1999).

HED has recently developed a framework that it proposes to use for conducting cumulative risk assessments on substances that have a common mechanism of toxicity. This framework was presented to the SAP. The SAP was in general agreement with the framework, and made recommendations for improving it. HED plans to release the proposed framework for public comment in March 2000. The framework is available from the Internet at: <http://www.epa.gov/scipoly/>. In the framework it is stated that a cumulative risk assessment of substances that cause a common toxic effect by a common mechanism will not be conducted until an aggregate exposure assessment of each substance has been completed. The framework is expected to be finalized by the fall of 2000. When the methods are completed and peer reviewed, EPA will proceed with a cumulative assessment of the organophosphates. The current assessment addressed only the risks posed by chlorpyrifos.

## **7.0 Confirmatory Data**

Additional data requirements have been identified in the attached Science Chapters and are summarized here.

### **7.1 Toxicology Data for OPPTS Guidelines**

HED has recommended and the registrant has developed a protocol for a Repeated Exposure Neurotoxicity Study of Sensory Electrophysiology. This study will also include measurement of neurotoxic esterase (NTE). It is expected that this would be a 28 day 2 dose, oral exposure study. In addition to the neurophysiological and neurochemical measures, neuropathological assessment focused on central/peripheral axonopathic changes associated with OPIDN (organophosphate-induced delayed neuropathy should also be performed). This is special study for which no single EPA guideline provides complete guidance. EPA has a guideline for 28 day hen studies of organophosphates that may cause OPIDN that includes guidance for neuropathology and NTE measurements (US EPA 1998; 870.6100). EPA has a guideline for examining peripheral nerve function (US EPA 85-SS1998; 870.6850) and a guideline for sensory evoked potentials (US EPA 1998; 870.6855). The current protocol for this special study has been developed by the registrant working voluntarily in conjunction with EPA. While EPA has not required this study, EPA maintains the right to require further study, based on concerns for potential health effects, consistent with its obligations under FIFRA.

### **7.2 Product and Residue Chemistry Data for OPPTS Guidelines**

#### **7.2.1 Product Chemistry**

Forty (40) MP's have been identified. Guideline 830.6314 data requirements remain outstanding for the DAS 99% T. Data remain outstanding for all other chlorpyrifos MP's; for many MP's no product chemistry data have been submitted. The reregistration guidelines for

product chemistry data requirements are complete, provided that the registrants submit the data required in the attached summary tables for the chlorpyrifos MPs, and either certify that the suppliers of starting materials and the manufacturing processes for the chlorpyrifos technicals and manufacturing-use products have not changed since the last comprehensive product chemistry review or submit complete updated product chemistry data packages.

### 7.2.2 Residue Chemistry

The following confirmatory data requirements and/or label revisions for magnitude of the residue in plants (Guideline 860.1500) remain outstanding or are now required:

- For asparagus, no additional residue data are required. However, a label revision is needed. The maximum equivalent rate of 1.9 lb ai/A specified by a homeowner-use label (EPA Reg. No. 62719-56) should be adjusted to reflect the maximum registered rate of 1.0 lb ai/A for which adequate residue data are available. In a letter to the Agency dated 5/8/95 the registrant committed to correcting the label directions to 1.0 lb ai/A at the next label printing.
- For corn, label restrictions prohibiting feeding of silage, forage, or fodder to meat or dairy animals are not practical and must be removed from SLN DE930004 and FL940003 labels. Additional data must be submitted to determine if established tolerances on corn forage and fodder are adequate for these uses. Alternatively, these SLN uses may be canceled.
- For cotton, feeding restrictions for gin trash (gin by-products) are not practical and must be removed from product labels. Appropriate tolerances for cotton gin by-products must be proposed. The proposal must be supported by adequate residue data conducted according to the maximum use patterns.
- For crops grown solely for seed (clover, and grasses), tolerance proposals and adequate field residue data are required to support SLN (Section 24-c) uses. The Oregon Clover Association has indicated that it will support chlorpyrifos SLN (OR850032) use on clover grown for seed. The requirements specified in the Addendum to the Chlorpyrifos SRR remain outstanding. For grasses grown for seed, appropriate tolerances for residues of chlorpyrifos *per se* in/on grass forage and hay must be proposed. The proposal must be supported by adequate residue data conducted according to the maximum use patterns specified by NV940002, and OR94032.

Alternatively, these SLN uses may be canceled.

- For mint, Table 1 (OPPTS Test Guidelines 860, August 1996) requires data for peppermint and spearmint tops (leaves and stems). Mint hay is no longer considered a RAC. Additional data are required for peppermint and spearmint tops (leaves and stems).

- For peppers, the requirements specified by the Addendum to the Chlorpyrifos SRR to submit English translations of labels for all products that permit use of chlorpyrifos on peppers imported to the U.S. have not been fulfilled. Chlorpyrifos use on peppers was approved at the issuance of the SRR, SLN (FL920007, FL920009, GA930003, and GA930004).
- For sorghum, data are required for aspirated grain fractions.
- For the tree nuts group (almonds, filberts, pecans, and walnuts), the Addendum to the Chlorpyrifos SRR did not require additional data to support the established crop group tolerance. However, an examination of the recently amended labels for the 4 lb/gal EC formulation (EPA Reg. Nos. 62719-23 and 62719-220) indicated that a maximum seasonal rate of 10 lb ai/A was inadvertently approved for pecans. The available residue data, reflecting combined residues of chlorpyrifos and TCP in/on pecans and other representative members of this crop group, only support a maximum seasonal rate of 5 lb ai/A. If the registrant wishes to support a seasonal rate of 10 lb ai/A, then additional data are required. Alternatively, the labels for pecans may be revised to reflect a maximum seasonal rate of 5 lb ai/A. In a letter to the Agency dated 5/8/95, DAS stated that they would modify labels to reflect a maximal seasonal use rate of 5 lb ai/A for pecans at the next label printing. The latest approved label for Lorsban 4E (EPA Reg. No. 62719-220), dated 4/8/96 did not include this modification. The labels should be revised or appropriate residue data supplied.
- For wheat, data are required for aspirated grain fractions.

[Note: The field trial data submitted for asparagus, apples, sugar beets, and tree nuts depict combined residues of chlorpyrifos and TCP. In the absence of adequate data depicting chlorpyrifos *per se* on the commodities of these crops, the established tolerances, for tolerance reassessment purposes, should remain at the existing levels. It is the registrant's prerogative to petition the Agency and submit additional field residue data depicting chlorpyrifos *per se* in/on these crops if tolerance-level reductions or lower anticipated residue calculations are desired.]

#### GLN 860.1520: Magnitude of the Residue in Processed Food/Feed

According to Table 1 (August 1996) OPPTS 860.1000 Test Guidelines residue data for sorghum flour are not needed at this time because it is used exclusively as a component of drywall, and not as a food or animal feed item, in the US. However, because 50% of the worldwide

sorghum production is used for human consumption, data may be needed at a later time.

The requirements for processing data on alfalfa meal are waived because residue data indicate that levels of chlorpyrifos *per se* are not likely to exceed the established tolerance in alfalfa hay following tests conducted according to registered uses. In addition, no sweet corn processing data are required since adequate corn forage data are available.

The available processing data for apples and sugar beets depict combined residues of chlorpyrifos and TCP. In the absence of adequate data depicting chlorpyrifos *per se* on the processed commodities of these crops, the established feed additive tolerances, for tolerance reassessment purposes, should remain at the existing levels. It is the registrant's prerogative to petition the Agency and submit additional processing data depicting chlorpyrifos *per se* in/on these commodities if tolerance-level reductions or lower anticipated residue calculations are desired.

#### GLNs 860.1850 and 860.1900: Confined/Field Rotational Crops

Provided that DAS modifies all labels for its chlorpyrifos containing products to limit application to 5 lb ai/A/season on those crops where rotation to another crop could occur (as was stated in their letter to the Agency dated 8/12/94), HED will not require field rotational crop studies. Furthermore, a 30 day plant back interval for rotational crops would then be appropriate.

### **7.3 Occupational Exposure Data for OPPTS Guidelines**

HED has insufficient data for the following agricultural handler scenarios:

- seed treatment uses
- dip applications (e.g., preplant peaches)
- dry bulk fertilizer applications to citrus orchard floors

These scenarios are of concern given the results from the other scenarios assessed.

For postapplication agricultural worker exposures, there is insufficient information (e.g., timing of applications -- dormant/bark versus foliar treatments) and exposure data to assess postapplication activities for ornamental and soil incorporated uses. The data needed to assess these uses include ornamental dislodgeable foliar residues in greenhouses and biological monitoring data for reentry into treated areas with soil directed applications.



In addition, HED could not evaluate the postapplication exposures and risks associated with use of insecticidal dust products due to an absence of chemical-specific data or recommended procedures in the Residential SOPs. Nevertheless, HED has concerns about the use of these products based on the low MOEs calculated using the surrogate data from the scientific literature for residents or workers that could apply these products. HED recommends that the registrant provide additional information on the potential post-application residential exposures associated with these products.

HED requests additional data for indoor crack, crevice and spot uses of chlorpyrifos. Specifically, HED requests treated room residue data for floors, furniture and other surfaces available for contact by children for both chlorpyrifos, and its primary degradation metabolite, 3,5,6-TCP following multiple treatments. Additionally, HED requests chlorpyrifos air measurements in treated rooms following multiple treatments (i.e., at a minimum 3 treatments 7 days apart). Residue data for 3,5,6-TCP are important due to the potential for accumulation and persistence of this environmental degradate.

HED requests confirmatory air monitoring data immediately following ground-based fogger application due to potential concern for short-term inhalation exposures.

In addition, HED requests exposure and/or environmental data for all registered products and/or uses that are not assessed in this risk assessment.

## 8.0 References

Adgate, J., Quackenboss, J, Needham, L., Pellizari, P., Liroy, P, Shubat, P., and Sexton, K . 1998. Comparison of Urban versus Rural Pesticide Exposure in Minnesota Children. Annual Conference of International Society for Environmental Epidemiology (ISEE) and International Conference for Society of Exposure Analysis (ISEE). July 1998, Volume 9 No. 4. Supplement. Abstract 92 O.

Blondell, J., and Dobozy, 1997. Memorandum to Linda Propst: Review of Chlorpyrifos Poisoning Data. January 14, 1997. U.S. Environmental Protection Agency, Washington, D.C.

Buckley T.J., Liddle J., Ashley D.L., Paschal D.C., Burse V.W., Needham L.L., and Akland G. 1997. Environmental and Biomarker Measurements in Nine Homes in the Lower Rio Grande Valley: Multimedia Results for Pesticides, Metals, PAHs and VOCs. Environmental International. 23(5):705-732.

Campbell, C.G., Seidler, F.J, and Slotkin, T.A. (1997). Chlorpyrifos interferes with cell development in rat brain regions (Brain Res. Bull 43(2):179-189.

Capodicasa, E., Scapellato, M.L., Moretto, A., Caroldi S., and Lotti, M. 1991.

Chlorpyrifos-induced delayed polyneuropathy. *Arch Toxicol.* 65:150-155.

Chanda, S.M., Mortensen, S.R., Barone, S., Moser, V.C., and Padilla, S. 1997. Developmental Profiles of two organophosphate detoxifying enzymes: carboxylesterase and A-esterase [abstract 1757]. *Toxicologist* 36(1):346.

Costa LG, Li WF, Richter RJ, Shih DM, Lulis A, and Furlong CE. 1999. The role of paraoxonase (PON1) in the detoxication of organophosphates and its human polymorphism. *Chemico-Biological Interactions* 119-120: 429-438

Coulston, F., Golberg, L. and Griffin, T. 1972. Safety Evaluation of DOWCO 179 in Human Volunteers. Institute of Experimental Pathology and Toxicology, Albany Medical College, Albany, New York. MRID No. 95175. HED Doc No. 000179, 03822, 04363.

Dam K, Garcia SJ, Seidler FJ, Slotkin TA (1999a) Neonatal chlorpyrifos exposure alters synaptic development and neuronal activity in cholinergic and catecholaminergic pathways. *Developmental Brain Res.* 116:9-20.

Dam K; Seidler FJ; Slotkin TA (1999b) Chlorpyrifos releases norepinephrine from adult and neonatal rat brain synaptosomes. *Brain Res Dev Brain Res*, 118(1-2):129-33.

Das KP, Barone S (1999) Neuronal differentiation in PC 12 cells is inhibited by chlorpyrifos and its metabolites: Is acetylcholinesterase inhibition the site of action? *Toxicol. Applied Pharmacol.* 160:217-230.

Davies HG, Richter RJ, Keifer M, Broomfield CA, Sowalla J, and Furlong CE. 1996. The effect of the human serum paraoxonase polymorphism is reverse with diazoxon, soman and sarin. *Nat Genet.* Nov 14(3):334-6.

Dittenber, D.A 1997. Chlorpyrifos: Evaluation of Single Oral Doses on Cholinesterase and Neurotoxic Esterase Inhibition in F344 Rats. Toxicology Laboratory, Dow Chemical Co. Study No. 960036. March 13, 1997. MRID No. 44273901.

Dow AgroSciences. 1998. Chlorpyrifos Technical Bulletin: Toxicity. Urban Exposure Considerations. Dow AgroSciences LLC. Indianapolis, IN. July.

EPA 1992. National Study of Chemical Residues in Fish. Office of Science and Technology (WH-551), Washington, D.C. Office of Water. EPA 823-R-92-008a. September 1992.

EPA 822-R-96-001; Drinking Water Regulations and Health Advisories; Office of Water; February 1996.

Food and Drug Administration (FDA). 1999. Total Diet Study. Summary of Residues Found Ordered by Pesticide Market Baskets 91-3 - 97-1. June, 1999.

Furlong, CE., Li WF., Costa, LG., Richter RJ., Shih DM, and Lulis AJ. 1998 Genetically determined susceptibility to organophosphorus insecticides and nerve agents: developing a mouse model for the human PON1 polymorphism. *Neurotoxicology*. Aug-Oct: 19(4-5):645-60

Hill R.H., Head, S.L., Baker, S., Gregg, M., Shealy, D.B., Bailey, S.L., Williams, C.C., Sampson, E.J., and Needham, L.L. 1995. Pesticide Residues in Urine of Adults Living in the United States: Reference Range Concentrations. *Environmental Research*. 71:99-108.

Hoberman A.M. 1998a,b. Developmental neurotoxicity study of chlorpyrifos administered orally via gavage to CrI:CD®BR VAF/Plus® presumed pregnant rats. Argus Research Laboratories, Inc., Horsham, Pennsylvania, laboratory study No. 304-001, sponsor study No. K-044793-109, May 1, 1998: MRID 44556901, MRID 44661001.

Jefferson Davis Associates, Inc. 1999. Lawn Care Applicator Exposure to Dursban. A Study of Typical Treatment Practices. A Quantitative Study. Prepared for Dow AgroSciences. December 1999.

Jett D.A., Navoa, R.V., Lyons, M.A. 1999. Additive inhibitory action of chlorpyrifos and polycyclic aromatic hydrocarbons on acetylcholinesterase activity in vitro. *Toxicology Letters*. 105:223-229.

Johnson, D.E., Seidler F.J., and Slotkin, T.A. 1998. Early Biochemical Detection of Delayed Neurotoxicity Resulting from Developmental Exposure to Chlorpyrifos. *Brain Research Bulletin*. 45(2):143-147.

Kilburn KH. 1999. Evidence for chronic neurobehavioral impairment from chlorpyrifos an organophosphate insecticide (Dursban) used indoors. *Environmental Epidemiology and Toxicology* 1:153-162.

Kisicki J.S., Seip, C.W., and Combs M.L. 1999. A Rising Dose Toxicology Study to Determine the No-Observable-Effect-Levels (NOEL) for Erythrocyte Acetylcholinesterase (AChE) Inhibition and Cholinergic Signs and Symptoms of Chlorpyrifos at Three Dose Levels. MDC Harris Laboratory, Lincoln Nebraska, Study No. 21438 (for the Harris Project) and DR K-0044793-284 (for Dow AgroSciences), April 19, 1999, MRID No. 44811002.

Lassiter TL, Padilla S, Mortensen SR, Chanda SM, Moser VC, Barone S (1998) Gestational exposure to chlorpyrifos: Apparent protection of the fetus? *Toxicol. Applied Pharmacol.* 152: 56-65.

Li WF, Costa LG, Furlong CE. 1993. Serum paraoxonase status: a major factor in determining resistance to organophosphates. *J Toxicol Environ Health*. Oct-Nov: 40(2-3):337-46.

MacIntosh D.L., Needham L.L., Hammerstrom K.A., and Ryan P.B. 1999. A longitudinal investigation of selected pesticide metabolites in urine. *J. of Exposure Analysis and Environ Epidemiol.* 9:494-501.

Mattsson J.L., Maurissen J.P., Spencer, P.J., Brzak K.A., and Zablony C.L. 1998. Effects of Chlorpyrifos administered via gavage to CD rats during gestation and lactation on plasma, erythrocyte, heart and brain cholinesterase and analytical determination of chlorpyrifos and metabolites. Health and Environmental Research Laboratories, The Dow Chemical Co. for Dow AgroSciences, August 31, 1998. Unpublished Study. MRID 44648101.

Maurissen J.P., Shankar, M.R., Mattsson J.L. 1996. Chlorpyrifos: cognitive study in adult Long-Evans rats. The Toxicology Research Laboratory, Health and Environmental Studies, The Dow Chemical Co. Midland, MI. Laboratory Project Study ID K-044793-096. April 29, 1996. MRID No. 44020901. Unpublished.

Mendrala A.L., and Brzak K.A. 1998. Chlorpyrifos: Part A-concentration-time course of chlorpyrifos and chlorpyrifos-oxon in blood. Health and Environmental Research Laboratories. The Dow Chemical Co. Midland MI. Laboratory Project Study ID: 971187A. August 31, 1998. MRID No. 44648102. Unpublished.

Mortensen, S.R., Hooper M.J. S. Padilla. 1998. Rat brain acetylcholinesterase activity: developmental profile and maturational sensitivity to carbamate and organophosphorus inhibitors. *Toxicology.* 125:13-19.

Moser, V.C. and S. Padilla. 1998. Age- and gender-related differences in the time-course of behavioral and biochemical effects produced by oral chlorpyrifos in rats. *Toxicology and Applied Pharmacology.* 149:107-119.

Moser, V.C., Chanda, S.M., Mortensen S.R., and Padilla, S. 1998. Age- and Gender-Related Differences in Sensitivity to Chlorpyrifos in the Rat Reflect Developmental Profiles of Esterase Activities. *Toxicological Sciences.* 46:211-222.

Nolan R.J., Rick D.L., Freshour M.L., and Saunders J.H. 1982. Chlorpyrifos: Pharmacokinetics in human volunteers following single oral and dermal doses. The Dow Chemical Co. Biomedical Medical Research Lab. Toxicology Research Lab. Midland MI. Accession No. 249203.

Pope, C.N., Chakraborti, T.K., Chapman, M.L., Farrar, J.D., and Arthur, D.(1991). Comparison of in vivo Cholinesterase Inhibition in Neonatal and Adult Rats by Three Organophosphorothioate Insecticides. *Toxicology*

Pope, C.N and Liu, J (1997). Age-Related Differences in Sensitivity to Organophosphorous Pesticides. *Environmental Toxicol. And Pharmacol.* 4;309-314.

Quackenboss, J.J., Pellizari, E., Freeman, N., Head, S., Whitmore, R., Zelon, H., Stroebel, C. 1998. Use of Screening Questionnaires to Identify Exposed and Sensitive Population Groups in the Region V NHEXAS Children's Pesticide Study. Annual Conference of International Society for Environmental Epidemiology (ISEE) and International Conference for Society of Exposure Analysis (ISEE). July 1998, Volume 9 No. 4. Supplement. Abstract 440 O.

Roy TS, Andrews JE, Seidler FJ, Slotkin TA (1998) Chlorpyrifos elicits mitotic abnormalities and apoptosis in neuroepithelium of cultured rat embryos. *Teratology* 58:62-68.

Shih DM, Gu L, Xia YR, Navab M, Li WF, Hama S, Castellani LW, Furlong CE, Costa LG, Fogelman AM and Lusic AJ. 1998. Mice Lacking serum paraoxonase are susceptible to organophosphate toxicity and atherosclerosis. *Nature*. Jul 16: 394 (6690):284-7

Slokin T.A. 1999. Developmental Cholinotoxicants: Nicotine and Chlorpyrifos. *Environmental Health Perspectives*. 107, Supplement 1, 71-80.

Song, X., Seidler, F.J., Saleh, J.L., Zhang, J. Padilla, S., Slotkin T.A. 1997. Cellular mechanisms for developmental toxicity of chlorpyrifos: targeting the adenylyl cyclase signaling cascade. *Toxicol Appl. Pharmacol*. 145:158-174.

Steenland K., Dick RB., Howell RJ., Chrislip DW., Hines CJ., Reid TM., Lehman E., Laber P., Krieg EF. Jr., Knott C. 2000. Neurologic function among termiticide applicators exposed to chlorpyrifos. *Environmental Health Perspectives*. 108(4):293-300. February.

Tang J, Carr RL, Chambers JE (1999) Changes in rat brain cholinesterase activity and muscarinic receptor density during and after repeated oral exposure to chlorpyrifos in early postnatal development. *Toxicological Sciences* 51:265-272.

Tietz, N.W. 1982. *Fundamentals of Clinical Chemistry*, 3rd Edition, W.B., Saunders Company, Philadelphia, PA. pg. 1950.

TruGreen/ChemLawn. 1999. Comments Submitted during Phase 2 Public Comment Period. November 29, 1999. Includes letter from R.A. Yeary, and studies of Pesticide Exposure of ChemLawn Employees 1975-1983.

U.S. Environmental Protection Agency. 1992. National Study of Chemical Residues in Fish. Volume 1. Office of Science and Technology. WH-551. Washington DC. EPA 823-R-92-008a. .

U.S. Environmental Protection Agency. 1997. Exposure Factors Handbook. Volume 1. General Factors. Office of Research and Development. Washington, D.C. Page 1-7. EPA/600/P-95/002Fa.

Whitney, K.D., Seidler, F.J., and Slotkin, T.A (1995). Developmental Neurotoxicity of

Chlorpyrifos Cellular Mechanism. *Toxicol. And Pharmacol*134:53-62

Wright, C.G., Leidy, R.B., and Dupree, H.E., Jr. 1988. Chlorpyrifos in the Ambient Air of Houses Treated for Termites. *Bull. Environ. Contam. Toxicol.* 40:561-568.

Wright, C.G., Leidy, R.B., and Dupree, H.E., Jr. 1994. Chlorpyrifos in the Air and Soil of Houses Treated Eight Years after its Application for Termite Control. *Bull. Environ. Contam. Toxicol.* 52:131-134.

Zheng, Q., Olivier K., Won Y., and Pope C. 1999. Comparative Cholinergic Neurotoxicity of Oral Chlorpyrifos Exposures in Neonatal and Adult Rats. Abstract and Poster Presentation. presented at the 38<sup>th</sup> Annual Society of Toxicology Meeting in New Orleans, March 14-18. *The Toxicologist* Vol 48, No.1-S, #874. March 1999.

Zheng Q, Olivier K, Won YK, Pope CN (2000) Comparative cholinergic neurotoxicity of oral chlorpyrifos exposures in preweanling and adult rats. *Toxicological Sciences.* 55:124-132.

## APPENDIX A: Sensitivity/Susceptibility of the Young

The following summary has been extracted from the following report: "Chlorpyrifos Children's Hazard: Sensitivity and Susceptibility" HED Doc No. 014074, March 28, 2000. The entire document is also an appendix to the April 6, 2000 HIARC report (which is an attachment to the risk assessment).

The weight of evidence provides appreciable support for the increased sensitivity of the young compared to adult rats to the neurotoxic effects of chlorpyrifos and for the susceptibility of the developing brain to chlorpyrifos. A number of different rat studies clearly demonstrate that at a given oral dose the young rat will respond more to the anticholinesterase effects of chlorpyrifos (as defined biochemically and behaviorally) than adult animals. The differential found between pups and adult animals is a function of the treatment dose, duration of treatment, timing of treatment (*i.e.*, developmental stage) and of measurements (*i.e.*, time to peak effect), and the toxicological endpoint examined. At high acute doses, chlorpyrifos is fatal to the rat pup, but produces no lethality and little to no behavioral changes in the adult rat (*e.g.*, LD<sub>10</sub> and MTD doses = neonate-15 mg/kg; adult-136 and 100 mg/kg, respectively). At the LD<sub>10</sub> or MTD doses neonates are up to ~5-fold more sensitive than adult rats to ChEI (brain and blood) and clinical/behavioral effects. Furthermore, at a single treatment of 15 mg/kg, the down-regulation of the cholinergic (muscarinic) receptors was more extensive in the pups than in adults treated with 80 mg/kg. The magnitude of change, the effective time points, and the brain regions involved were different in pups versus adult rats. This suggests that the cholinergic receptors are more readily altered in the pup following chlorpyrifos treatment. Although the consequence of this is unknown, cholinergic receptors play an important role in normal brain development.

The increase in sensitivity between young and adult animals appears to occur at acute doses below 15 mg/kg. The study by Zheng *et al.* (2000) using lower dose levels (ranging from 0.15 mg/kg to 15 mg/day) provides cholinesterase inhibition (ChEI) data in 7-day old animals and adult male rats showing a greater sensitivity (up to ~3-fold for RBC and plasma, and perhaps at least 5-fold for brain) of pups compared with adult males. In the Zheng *et al.* study, the adult did not respond at the high dose of 15 mg/kg for brain ChEI. Thus, a difference in response greater than 5-fold can not be ruled out. Because of the lack of data, the extent of differences in brain ChEI between pups and the pregnant female rat remains uncertain. Although the young animal appears to recover at least two times faster than the adult animal from the ChEI induced by acute chlorpyrifos treatment, other toxicities (*e.g.*, delays in brain development, behavioral effects) may persist or appear at later times.

Repeated dosing with chlorpyrifos does not appear to result in an increase in brain or blood ChEI in neonates relative to adults with one exception. Based on ED<sub>50</sub>'s, there is a 1.5-fold difference in the response of PND 7 pups to brain ChEI compared to adult males (Zheng *et al.*, 2000). In contrast to the rapid recovery from ChEI observed with acute chlorpyrifos treatments of neonates (Pope and Liu, 1997), repeated dosing with

chlorpyrifos (every other day, 11 treatments during PND 1 to PND 21) indicates ChEI persists for ~9 to >19 days depending on the dose administered (Tang *et al.*, 1999). Body weight changes and behavioral effects occur at ~3-fold lower doses in neonates versus adult rats with repeated treatments of chlorpyrifos doses equal to or above 3 mg/kg/day.

It is apparent that cholinesterase activity is inhibited in the fetus if the dam is treated with a chlorpyrifos dose which can be absorbed by the fetus. The magnitude of brain, plasma, and RBC ChEI in the fetus is less or equal to that observed in dams with acute or repeated treatments of dams with chlorpyrifos. The lack of an apparent differential response of the fetus (or neonate with repeated dosing) versus the maternal system to treatment of dams with chlorpyrifos may be due to the increased new synthesis or more rapid turnover of inhibited molecules of cholinesterases in the fetal brain than in the adult (Lassiter *et al.*, 1998; Mortensen *et al.*, 1998).

Differences in detoxification between the young and adults may explain the increased sensitivity of exposed pups to chlorpyrifos toxicity. Chlorpyrifos and its oxon (*i.e.*, the anticholinesterase metabolite) are detoxified by binding to carboxylesterases and hydrolysis by A-esterases. The young animal has minimal activity of these detoxification enzymes compared to adult animals. The precise influence of these enzymes on sensitivity to chlorpyrifos treatment has not been established. Because detoxification enzyme activities increase with age, the enzymatic profile of newborn rats raises concern that the newborn may be even more sensitive than older neonates to an acute chlorpyrifos treatment. There is some evidence (albeit at high doses) that suggests that the magnitude of the differential sensitivity between young and adult animals depends on the age of the animal. Based on the LD<sub>10</sub> data in Zheng *et al.* and from the ChEI data in Zheng *et al.* and Moser and Padilla (1998), the order of sensitivity is PND 7 > PND 17 > PND 27 > adult female > adult male. Therefore, given that 7-day old rats are the youngest animals evaluated to date, it is uncertain whether the magnitude of differential sensitivity would be greater with pups exposed earlier than 7 days.

The developmental neurotoxicity study, which involved treatment of dams with 5, 1, or 0.3 mg/kg/day chlorpyrifos from GD 6 through lactation day 11 (Hoberman, 1998a,b), offspring were observed to have alterations in brain structure that are suggestive of a developmental defect that may predispose the neonate to unique adverse consequences. In this study, morphometric measurements in PND 11 pups of the high dose included, decreases in anterior to posterior measurements of the cerebellum, reduced height of the cerebellum, decreased thickness of the parietal cortex, and decreased thickness of the hippocampal gyrus. These effects at the high dose occurred in the presence of maternal toxicity (*e.g.*, maximum brain, RBC and plasma ChEI) but in the absence of effects on body weights, food consumption, pregnancy parameters, or deaths among the dams. In mid- and high-dose PND 66 offspring, effects on brain structure included marginal but statistically significant decreases in the thickness of the parietal cortex and non-significant decreases in the thickness of the hippocampal gyrus. This difference in the qualitative severity of the findings seen in adult and neonatal animals is indicative of susceptibility of the offspring. It is also important to note that morphometric evaluation of the low-dose



brains was not conducted. So it is not known whether alterations are occurring at lower doses.

Additionally, a number of the treatment-related findings in the offspring appear to be delayed in expression of perturbations in earlier neurological development, because functional and morphological changes are observed at study termination (~PND 61 - 66), approximately 50 - 55 days after cessation of maternal dosing. At the high dose, these findings included increased motor activity in females at PND 61, alterations in auditory startle measurements (increased latency to peak response and decreased peak response amplitudes) at PND 62, and morphometric alterations in the parietal cortex and hippocampal gyrus on PND 66.

A variety of *in vitro* and *in vivo* studies published in the peer reviewed literature show that chlorpyrifos can alter macromolecular synthesis, neuronal activity, neurotransmitter levels, neurite outgrowth and branching, and cell signaling in the developing rat brain (reviewed by Slotkin, 1999). Although these studies did not include accompanying measures of direct adverse effects (*e.g.*, functional effects) but rather used biomarkers, they nevertheless raise concern that chlorpyrifos potentially can affect processes occurring in both early and late developmental periods of brain growth that influence cell replication and differentiation needed for normal function. Although the data primarily come from one laboratory, multiple studies from this group have shown a consistency in the different responses measured. Furthermore, several of the key responses observed are highly significant and robust (*e.g.*, effects on norepinephrine turnover, DNA synthesis, adenylyl cyclase transduction). Also, the responses reported tend to have little variability in the data. Finally, effects on the developing brain reported in the literature are consistent with the morphometric changes observed in the guideline developmental neurotoxicity study by Hoberman (1998) even though a direct linkage of effects can not be made. The available data suggest a selective action of chlorpyrifos on the developing brain, given the regional and temporal pattern of responses. Thus, it seems unlikely that the observed effects are due to nonspecific toxicity.

Although there are strengths of these studies, there are also some limitations and questions raised which are not addressed by the results. As discussed above, the mechanism of action for chlorpyrifos in the developing brain is unclear. Also, the *in vivo* studies using macromolecular biomarkers have primarily been conducted using the subcutaneous injection (SC) route of exposure and DMSO as the vehicle. It should be noted that DMSO controls were conducted in all the studies. DMSO would result in a rapid uptake and full absorption of the compound. Compounds administered via SC injection enter directly into the general circulation and bypass hepatic metabolism once, thus bypassing hepatic activation of chlorpyrifos to its active metabolite chlorpyrifos-oxon. The SC route of exposure can not be reliably compared to the oral route given the lack of pharmacokinetic data on this dosing regime. Also, this is not a pathway of human exposure. Thus the DMSO-SC dosing regime makes quantitative interpretation and extrapolation of the results problematic. Nevertheless, these studies still provide important qualitative information on the potential for chlorpyrifos to affect neurodevelopmental processes. Cholinesterase inhibition was not measured in most of these studies except for Song *et al.* (1997). In that study, no extreme cholinesterase inhibition is found in the brainstem at the low dose used in the study: approximately 20-25% cholinesterase

inhibition is found when 1 mg/kg of chlorpyrifos is administered during PND 1-4 and cholinesterase activity (measured 24 hours after the last dose) is almost completely recovered by 10 days of age (Song *et al.*, 1997). Given that key effects in the postnatal brain are found at the low dose, the concern of a rapid delivery of a toxic dose with this standard dosing regime is reduced. Also, no significant changes in body or brain weight and no mortality occurs with this dosing regime (1 mg/kg at PND 1-4 or 5 mg/kg at PND 11-14). Additionally, it should be noted that chlorpyrifos is rapidly absorbed and transported to the brain with oral dosing (Mendrala and Brzak, 1998). Thus, the findings derived from the SC/DMSO dosing regime can not be discounted as an artifact of the vehicle and route of exposure and raise concerns for the unique susceptibility of the young.

The mechanism(s) of action for the chlorpyrifos-induced changes (*e.g.*, macromolecular synthesis, cell signaling) is/are unclear. However, given that these effects can be found after intracisternal injection of chlorpyrifos, with *in vitro* TCP treatment, and *in vitro* PC12 cell cultures with limited capability to activate chlorpyrifos to its ChE-inhibiting oxon, raises the issue of whether these effects can occur independent of cholinesterase inhibition. Although it is not possible to link each effect reported with another effect or with a functional outcome, the data show a consistent pattern of the potential for chlorpyrifos to produce qualitatively different effects in the central nervous system (CNS) of young versus adult animals. Potential implications of the effects include alteration of synaptic responses that are programmed by neural input, disruption of cell replication and differentiation, and temporary or persistent delays in the development of CNS structures.

In conclusion, the weight of the evidence raises concern for an increase in both the sensitivity and susceptibility of the fetus or young animal to adverse biochemical, morphological, or behavioral alterations from chlorpyrifos treatment during brain development. With respect to cholinesterase inhibition, an increase in sensitivity of the young compared to adults was seen all along the dose response curve, even at relatively low doses. There is a clear differential response (2- to ~5-fold) in the young compared to the adult animal after an acute treatment to a relatively low dose of chlorpyrifos. There is also increased sensitivity found after repeated dosing (up to 9-fold), but at the LD<sub>10</sub> and MTD. It is important to point out that an uncertainty remains concerning the magnitude of the differential response, given that newborn animals (less than PND 7) have not been characterized for sensitivity. Results of multiple studies have consistently shown that the developing brain is susceptible to chlorpyrifos treatment. Effects on the developing CNS that are indicative of the unique susceptibility to the young animal include changes in macromolecular synthesis, altered cell signaling and muscarinic receptor down-regulation, as well as morphological alterations in brain development. An uncertainty remains regarding the NOAELs for the susceptibility effects. The effects observed raise a high degree of concern that the fetus or young animal is particularly susceptible to adverse outcome if exposed to chlorpyrifos.