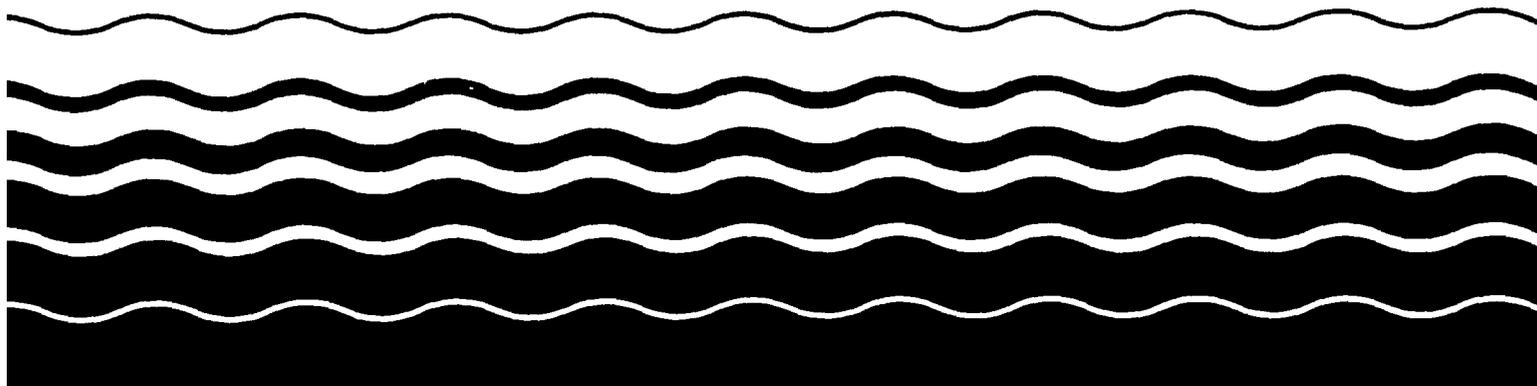




# Technical Support Document For Water Quality-based Toxics Control



# **TECHNICAL SUPPORT DOCUMENT FOR WATER QUALITY-BASED TOXICS CONTROL**

*This copy represents the second printing of this document.*

*Changes made to this document reflect corrections of typographical errors and the following update of the interim guidance on criteria for metals: The Agency has issued "Interim Guidance Interpretation and Implementation Aquatic Life Criteria for Metals." The interim guidance supersedes criteria document statements expressing criteria in terms of a acid soluble analytical method and also the metals discussion of Section 5.7.3. The availability of this document appeared in the June 5, 1992 Federal Register (Vol. 57, No. 109, pg. 24401).*

**March 1991  
Office of Water Enforcement and Permits  
Office of Water Regulations and Standards  
U.S. Environmental Protection Agency  
Washington, DC 20460**

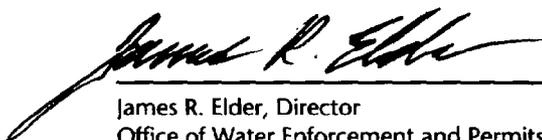
---

## FOREWORD

The U.S. Environmental Protection Agency (EPA) and the State pollution control agencies have been charged with enforcing the laws regarding pollution of the natural environment. Environmental pollution is an urgent and continuing problem and, consequently, the laws grant considerable discretion to the control authorities to define environmental goals and develop the means to attain them. Establishing environmentally protective levels and incorporating them in a decisionmaking process entails a considerable amount of scientific knowledge and judgment. One area where scientific knowledge is rapidly changing concerns the discharge of toxic pollutants to the Nation's surface waters.

This document provides technical guidance for assessing and regulating the discharge of toxic substances to the waters of the United States. It was issued in support of EPA regulations and policy initiatives involving the application of biological and chemical assessment techniques to control toxic pollution to surface waters. This document is agency guidance only. It does not establish or affect legal rights or obligations. It does not establish a binding norm and is not finally determinative of the issues addressed. Agency decisions in any particular case will be made applying the law and regulations on the basis of specific facts when permits are issued or regulations promulgated.

This document is expected to be revised periodically to reflect advances in this rapidly evolving area. Comments from users will be welcomed. Send comments to U.S. EPA, Office of Water Enforcement and Permits, 401 M Street, SW, Mailcode EN366, Washington, DC 20460.



---

James R. Elder, Director  
Office of Water Enforcement and Permits



---

Martha G. Prothro, Director  
Office of Water Regulations and Standards

**TABLE OF CONTENTS**

Section	Page
Foreword . . . . .	iii
Acknowledgment . . . . .	xiii
Executive Summary . . . . .	xiv
List of Abbreviations . . . . .	xvii
Glossary . . . . .	xix
Introduction . . . . .	xxiii
1. APPROACHES TO WATER QUALITY-BASED TOXICS CONTROL . . . . .	1
1.1 INTRODUCTION . . . . .	1
1.2 CHEMICAL-SPECIFIC APPROACH FOR AQUATIC LIFE PROTECTION . . . . .	1
1.2.1 Correlation of Chemical-specific Measurements to Actual Receiving Water Impacts . . . . .	2
1.2.2 Chemical-Specific Analytical Method Precision . . . . .	2
1.3 WHOLE EFFLUENT APPROACH FOR AQUATIC LIFE PROTECTION . . . . .	4
1.3.1 Toxic Units . . . . .	6
1.3.2 Correlation of Whole Effluent Toxicity Measurements to Actual Receiving Water Impact . . . . .	6
1.3.3 Toxicity Test Method Precision . . . . .	11
1.3.4 Considerations Involved When Implementing the Whole Effluent Toxicity Approach . . . . .	11
1.4 BIOLOGICAL CRITERIA/BIOASSESSMENT AND BIOSURVEY APPROACH FOR AQUATIC LIFE PROTECTION . . . . .	18
1.4.1 Use of Biosurveys and Bioassessments in Water Quality-based Toxics Control . . . . .	18
1.4.2 Conducting Biosurveys . . . . .	19
1.5 INTEGRATION OF THE WHOLE EFFLUENT, CHEMICAL-SPECIFIC, AND BIOASSESSMENT APPROACHES . . . . .	20
1.5.1 Capabilities and Limitations of the Chemical-Specific Approach . . . . .	20
1.5.2 Capabilities and Limitations of the Whole Effluent Approach . . . . .	21
1.5.3 Capabilities and Limitations of the Bioassessment Approach . . . . .	22

**Table of Contents (Continued)**

Section	Page
1.6 OTHER FACTORS INFLUENCING WATER QUALITY-BASED TOXICS CONTROL .....	23
1.6.1 Persistence .....	23
1.6.2 Additivity, Antagonism, and Synergism .....	24
1.6.3 Test Interferences .....	24
1.7 HUMAN HEALTH PROTECTION .....	24
1.7.1 Types of Health Effects .....	25
REFERENCES .....	26
2. WATER QUALITY CRITERIA AND STANDARDS .....	29
2.1 INTRODUCTION .....	29
2.1.1 Overview of Water Quality Standards .....	29
2.1.2 Water Quality Standards and State Toxics Control Programs .....	30
2.2 GENERAL CONSIDERATIONS .....	31
2.2.1 Magnitude, Duration, and Frequency .....	31
2.2.2 Mixing Zones .....	33
2.3 WATER QUALITY CRITERIA FOR AQUATIC LIFE PROTECTION .....	34
2.3.1 Development Process for Criteria .....	34
2.3.2 Magnitude for Single Chemicals .....	34
2.3.3 Magnitude for Whole Effluent Toxicity .....	35
2.3.4 Duration for Single Chemicals and Whole Effluent Toxicity .....	35
2.3.5 Frequency for Single Chemicals and Whole Effluent Toxicity .....	36
2.4 WATER QUALITY CRITERIA FOR HUMAN HEALTH PROTECTION .....	36
2.4.1 Overview .....	36
2.4.2 Magnitude and Duration .....	37
2.4.3 Human Exposure Considerations .....	37
2.4.4 Fish Consumption Values .....	37
2.4.5 Bioaccumulation Considerations for Reference Ambient Concentration Development .....	38
2.4.6 Updating Human Health Criteria and Generating RACs Using IRIS .....	38
2.4.7 Calculating RACs for Non-carcinogens .....	39
2.4.8 Calculating RACs for Carcinogens .....	40
2.4.9 Deriving Quantitative Risk Assessments in the Absence of IRIS Values .....	40
2.4.10 Deriving Reference Tissue Concentrations for Monitoring Fish Tissue .....	41

**Table of Contents (Continued)**

Section	Page
2.5. BIOLOGICAL CRITERIA . . . . .	41
2.5.1 Regulatory Bases for Biocriteria . . . . .	41
2.5.2 Development and Implementation of Biocriteria . . . . .	41
2.6. SEDIMENT CRITERIA . . . . .	42
2.6.1 Current Developments in Sediment Criteria . . . . .	42
2.6.2 Approach to Sediment Criteria Development . . . . .	42
2.6.3 Application of Sediment Criteria . . . . .	43
2.6.4 Sediment Criteria Status . . . . .	43
REFERENCES . . . . .	45
3. EFFLUENT CHARACTERIZATION . . . . .	47
3.1 INTRODUCTION . . . . .	47
3.1.1 NPDES Regulation Requirements . . . . .	47
3.1.2 Background for Toxic Effects Assessments on Aquatic Life and Human Health . . . . .	48
3.1.3 General Considerations in Effluent Characterization . . . . .	49
3.2 DETERMINING THE NEED FOR PERMIT LIMITS WITHOUT EFFLUENT MONITORING DATA FOR A SPECIFIC FACILITY . . . . .	50
3.3 DETERMINING THE NEED FOR PERMIT LIMITS WITH EFFLUENT MONITORING DATA . . . . .	51
3.3.1 General Considerations . . . . .	51
3.3.2 Addressing Uncertainty in Effluent Characterization by Generating Effluent Monitoring Data . . . . .	52
3.3.3 Effluent Characterization for Whole Effluent Toxicity . . . . .	53
3.3.4 Use of Toxicity Testing in Multiple-source Discharge Situations . . . . .	59
3.3.5 Ambient Toxicity Testing . . . . .	61
3.3.6 Special Considerations for Discharges to Marine and Estuarine Environments . . . . .	61
3.3.7 Using a Chemical-specific Limit to Control Toxicity . . . . .	61
3.3.8 Effluent Characterization for Specific Chemicals . . . . .	62
3.3.9 Effluent Characterization for Bioconcentratable Pollutants . . . . .	64
3.3.10 Analytical Considerations for Chemical . . . . .	65
REFERENCES . . . . .	66

## Table of Contents (Continued)

Section	Page
4. EXPOSURE AND WASTELOAD ALLOCATION .....	67
4.1 INTRODUCTION .....	67
4.2 TOTAL MAXIMUM DAILY LOADS AND WASTELOAD ALLOCATIONS .....	67
4.2.1 Total Maximum Daily Loads .....	67
4.2.2 Wasteload Allocation Schemes .....	69
4.3 INCOMPLETELY MIXED, DISCHARGE RECEIVING WATER SITUATIONS .....	69
4.3.1 Determination of Mixing Zone Boundaries .....	70
4.3.2 Mimimizing the Size of Mixing Zones .....	71
4.3.3 Prevention of Lethality to Passing Organisms .....	71
4.3.4 Prevention of Bioaccumulation Problems for Human Health .....	72
4.4 MIXING ZONE ANALYSES .....	72
4.4.1 General Recommendations for Outfall Design .....	73
4.4.2 Critical Design Periods for Waterbodies .....	73
4.4.3 General Recommendations for Tracer Studies .....	74
4.4.4 Discharge-induced Mixing .....	75
4.4.5 Ambient-induced Mixing .....	77
4.5 COMPLETELY MIXED DISCHARGE-RECEIVING WATER SITUATIONS .....	78
4.5.1 Wasteload Modeling Techniques .....	78
4.5.2 Calculating the Allowable Effluent Concentration Distribution and the Return Period .....	82
4.5.3 General Recommendations for Model Selection .....	83
4.5.4 Specific Model Recommendations .....	83
4.5.5 Effluent Toxicity Modeling .....	85
4.6 HUMAN HEALTH .....	87
4.6.1 Human Health Considerations .....	87
4.6.2 Determining the TMDL Based on Human Health Toxicants .....	87
REFERENCES .....	90
5. PERMIT REQUIREMENTS .....	93
5.1 INTRODUCTION .....	93
5.1.1 Regulatory Requirements .....	93

---

---

**Table of Contents (Continued)**

Section	Page
5.2	BASIC PRINCIPLES OF EFFLUENT VARIABILITY . . . . . 93
5.2.1	Variations in Effluent Quality . . . . . 93
5.2.2	Statistical Parameters and Relationship to Permit Limits . . . . . 95
5.2.3	Expression of Permit Limits . . . . . 96
5.3	ENSURING CONSISTENCY WITH THE WASTELOAD ALLOCATION . . . . . 96
5.3.1	Statistical Considerations of WLAs . . . . . 96
5.3.2	Types of Water Quality Models and Model Outputs . . . . . 96
5.4	PERMIT LIMIT DERIVATION . . . . . 98
5.4.1	EPA Recommendations for Permitting for Aquatic Life Protection . . . . . 98
5.4.2	Other Approaches to Permitting for Aquatic Life . . . . . 103
5.4.3	Special Permitting Requirements . . . . . 104
5.4.4	EPA Recommendations for Permitting for Human Health Protection . . . . . 104
5.5	SPECIAL CONSIDERATIONS IN USE OF STATISTICAL PERMIT LIMIT DERIVATION TECHNIQUES . . . . . 105
5.5.1	Effect of Changes of Statistical Parameters on Permit Limits . . . . . 105
5.5.2	Coefficient of Variation . . . . . 106
5.5.3	Number of Samples . . . . . 107
5.5.4	Probability Basis . . . . . 110
5.6	PERMIT DOCUMENTATION . . . . . 110
5.7	EXPRESSING LIMITS AND DEVELOPING MONITORING REQUIREMENTS . . . . . 110
5.7.1	Mass-based Effluent Limits . . . . . 110
5.7.2	Energy Conservation . . . . . 111
5.7.3	Considerations in the Use of Chemical specific Limits . . . . . 111
5.7.4	Considerations in the Use of Whole Effluent Toxicity Limits . . . . . 112
5.7.5	Selection of Monitoring Frequencies . . . . . 113
5.7.6	Analytical Variability . . . . . 113
5.7.7	Antibacksliding . . . . . 113
5.8	TOXICITY REDUCTION EVALUATIONS . . . . . 114
5.8.1	TRE Guidance Documents . . . . . 114
5.8.2	Recommended Approach for Conducting TREs . . . . . 114
5.8.3	Circumstances Warranting a TRE . . . . . 117
5.8.4	Mechanisms for Receiving TREs . . . . . 118
	REFERENCES . . . . . 121

---



---

## Table of Contents (Continued)

Section	Page
6. COMPLIANCE MONITORING AND ENFORCEMENT .....	123
6.1 INTRODUCTION .....	123
6.2 PERMIT REQUIREMENTS .....	123
6.3 COMPLIANCE MONITORING .....	123
6.3.1 Self-monitoring Reports .....	123
6.3.2 Discharge Monitoring Reports/Quality Assurance .....	124
6.3.3 Inspections .....	124
6.4 VIOLATION REVIEW .....	124
6.5 ENFORCEMENT .....	125
6.6 REPORTING OF VIOLATIONS .....	126
REFERENCES .....	127
7. CASE EXAMPLES .....	129
7.1 INTRODUCTION .....	129
7.2 CASE 1: INDUSTRIAL DISCHARGE .....	129
7.2.1 General Site Description and Information .....	129
7.2.2 Effluent Characterization for Specific Chemicals .....	129
7.2.3 Effluent Characterization for Whole Effluent Toxicity .....	131
7.2.4 Determine Wasteload Allocations .....	132
7.2.5 Develop Permit Limits .....	132
7.2.6 Determining and Expressing the Controlling Effluent Limit .....	133
7.2.7 Comparing Different Limit Development Methods .....	133
7.3 CASE 2: POTW DISCHARGE .....	134
7.3.1 General Site Description and Information .....	134
7.3.2 Effluent Characterization for Specific Chemicals .....	134
7.3.3 Effluent Characterization for Whole Effluent Toxicity .....	136
7.3.4 Determine Wasteload Allocations .....	136
7.3.5 Develop Permit Limits .....	136
7.3.6 Determining and Expressing the Controlling Effluent Limits .....	137
7.3.7 Comparing Different Limit Development Methods .....	137
7.4 CASE 3: MULTIPLE DISCHARGERS INTO THE SAME REACH .....	137
7.4.1 Effluent Characterization .....	137
7.4.2 TMDLs and WLAs .....	138
7.3.3 Permit Limit Development .....	139
INDEX .....	140

---

---

**APPENDICES**

<b>Section</b>	<b>Page</b>
Appendix A-1: Toxicity Test Precision Data . . . . .	A-1
Appendix A-2: Effluent Variability Data . . . . .	A-2-1
Appendix A-3: Acute to Chronic Ratio Data . . . . .	A-3-1
Appendix B-1: Summary of Clean Water Act Provisions . . . . .	B-1-1
Appendix B-2: Policies for Toxics Control . . . . .	B-2-1
Appendix B-3: Regulations for Toxics Control . . . . .	B-3-1
Appendix B-4: Whole-Effluent Toxicity Permitting Principles and Enforcement Strategy . . . . .	B-4-1
Appendix B-5: Quality Control Fact Sheets . . . . .	B-5-1
Appendix B-6: Case Decisions on Whole-Effluent Toxicity . . . . .	B-6-1
Appendix C: Ambient Toxicity Testing and Data Analysis . . . . .	C-1
Appendix D: Duration and Frequency . . . . .	D-1
Appendix E: Lognormal Distribution and Permit Limit Derivations . . . . .	E-1
Appendix F: Sampling . . . . .	F-1
Appendix G: The Development of a Biological Indicator Approach to Water Quality-based Human Health Toxics Control . . . . .	G-1
Appendix H: Reference Dose (RfD): Description and Use in Health Risk Assessments . . . . .	H-1
Appendix I: Chemicals Available in IRIS . . . . .	I-1

## **ACKNOWLEDGMENT**

The preparation of the revised *Technical Support Document for Water Quality-based Toxic Control* began with a 3-day conference held in Williamsburg, Virginia, in December 1988. Representatives of EPA Headquarters and Regions, States, private industry, municipalities, academia, and various interest groups attended this meeting and provided valuable input. The principal authors of this document were Bill Swietlik, James Taft, Jacqueline Romney, Kathryn Smith, John Cannell, Robert Wood, James Pendergast, and Rick Brandes of the Permits Division; Sheila Frace and Margarete Heber of the Enforcement Division, Elizabeth Southerland and Richard Healy of the Assessment and Watershed Protection Division; and Charles Delos, Warren Banks, and Robert April of the Criteria and Standards Division. Listed below are the contributors to specific chapters of this document.

### ***Approaches to Water Quality-Based Toxics Control***

Margaret Heber, U.S. EPA, Enforcement Division  
Kathryn Smith, U.S. EPA, Permits Division

### ***Water Quality Criteria and Standards***

Charles Delos, U.S. EPA, Criteria and Standards Division  
Warren Banks, U.S. EPA, Criteria and Standards Division  
Kathy Barylski, U.S. EPA, Criteria and Standards Division  
Robert April, U.S. EPA, Criteria and Standards Division  
David Moon, U.S. EPA, Criteria and Standards Division  
Jacqueline Romney, U.S. EPA, Permits Division

### ***Effluent Characterization***

Robert Wood, U.S. EPA, Permits Division  
Bill Swietlik, U.S. EPA, Permits Division  
James Pendergast, U.S. EPA, Permits Division

### ***Exposure and Wasteload Allocation***

Elizabeth Southerland, U.S. EPA, Assessment and Watershed Protection  
Richard Healy, U.S. EPA, Assessment and Watershed Protection

### ***Permit Requirements***

Bill Swietlik, U.S. EPA, Permits Division  
James Pendergast, U.S. EPA, Permits Division  
John Cannell, U.S. EPA, Permits Division

### ***Compliance Monitoring and Enforcement***

Sheila Frace, U.S. EPA, Enforcement Division  
Theodore Coopwood, U.S. EPA, Enforcement Division

### ***Human Health Component of All Chapters***

John Cannell, U.S. EPA, Permits Division  
Katherine Dowell, U.S. EPA, Permits Division  
William Morrow, U.S. EPA, Permits Division

### ***Case Examples Workgroup***

Bill Swietlik, U.S. EPA, Permits Division  
James Pendergast, U.S. EPA, Permits Division  
Charles Delos, U.S. EPA, Criteria and Standards Division  
Jacqueline Romney, U.S. EPA, Permits Division

### ***Appendices***

Appendix A: Margaret Heber, U.S. EPA, Enforcement Division  
Appendix B: U.S. EPA, Permits Division  
Appendix C: U.S. EPA, Permits Division  
Appendix D: Nelson Thomas, U.S. EPA, ERL/ORD, Duluth, MN  
Appendix E: Henry Kahn and Marla Smith, U.S. EPA, Analysis and Evaluation Division  
Appendix F: U.S. EPA, Permits Division  
Appendix G: U.S. EPA, Permits Division  
Appendix H: U.S. EPA, RfD Workgroup

### 3.2 DETERMINING THE NEED FOR PERMIT LIMITS WITHOUT EFFLUENT MONITORING DATA FOR A SPECIFIC FACILITY

If the regulatory authority so chooses, or if the circumstances dictate, the authority may decide to develop and impose a permit limit for whole effluent toxicity or for individual toxicants without facility-specific effluent monitoring data, or prior to the generation of effluent data. Water quality-based permit limits can be set for a single toxicant or for whole effluent toxicity based on the available dilution and the water quality criterion or the State standard in the absence of facility specific effluent monitoring data. However, in doing so, the regulatory authority must satisfy all the requirements of 40 *CFR* 122.44(d)(1)(ii).

When determining whether or not a discharge causes, has the reasonable potential to cause, or contributes to an excursion of a numeric or narrative water quality criterion for individual toxicants or for toxicity, the regulatory authority can use a variety of factors and information where facility-specific effluent monitoring data are unavailable. These factors also should be considered with available effluent monitoring data. Some of these factors are the following:

- **Dilution**—Toxic impact is directly related to available dilution for the effluent. Dilution is related to the receiving stream flow and the size of the discharge. The lower the available dilution, the higher the potential for toxic effect. If an effluent's concentration at the edge of a mixing zone in a receiving water is expected to reach 1 percent or higher during critical or worst-case design periods, then such an effluent may require a toxicity limit (see discussion in Section 3.3.3). Assessment of the amount of stream dilution available should be made at the conditions required by the water quality standards or, if not specified in the standards, at the harmonic mean flow and the 7Q10 flow. Figure 3-3 (Pg. 57) shows that, whereas a majority of NPDES permittees nationwide discharge to areas during annual mean flow ranging in dilution from 100 to 1,000, the majority of dischargers fall into the 1 to 10 dilution range during low-flow conditions.
- **Type of industry**—Although dischargers should be individually characterized because toxicity problems are site-specific, the primary industrial categories should be of principal toxicity concern. EPA's treatment technology data base generally suggests that secondary industrial categories may have less potential for toxicity than primary industries. However, based on experience, it is virtually impossible to generalize the toxicity of effluents with any certainty. If two plants produce the same type of product, one effluent may be toxic while the other may not be toxic due to the type and efficiency of the treatment applied, general materials handling practices, and the functional target of the compound(s) being produced.
- **Type of POTW**—POTWs with loadings from indirect dischargers (particularly primary industries) may be candidates for toxicity limits. However, absence of industrial input does not guarantee an absence of POTW discharge toxicity problems. For example, commercial pesticide applicators often discharge to POTWs, resulting in pesticide concentrations in the POTW's effluent. Household disposal of pesticides, detergents, or other toxics may have a similar effect. The types of industrial users, their product lines, their raw materials, their potential and actual discharges, and their control equipment should be evaluated. POTWs should also be characterized for the possibility of chlorine and ammonia problems.
- **Existing data on toxic pollutants**—Discharge monitoring reports (DMRs) and data from NPDES permit application forms 2C and 2A may provide some indication of the presence of toxicants. The presence or absence of the 126 "priority pollutants" may or may not be an indication of the presence or absence of toxicity. There are thousands of "nonpriority" toxicants that may cause effluent toxicity. Also, combinations of several toxicants can produce ambient toxicity where the individual toxicants would not. EPA regulations at 40 *CFR* 122.21(j) require POTWs with design flows equal to or greater than 1 MGD and POTWs with approved pretreatment programs, or POTWs required to develop a pretreatment program, to submit the results of whole effluent toxicity tests with their permit applications. These regulations also provide discretion to the permitting authority to request such data from other POTWs at the time of permit application.
- **History of compliance problems and toxic impact**—Regulatory authorities may consider particular dischargers that have had difficulty complying with limits on toxicants or that have a history of known toxicity impacts as probable priority candidates for effluent toxicity limits.
- **Type of receiving water and designated use**—Regulatory authorities may compile data on water quality. Examples of available data include fish advisories or bans, reports of fish kills, State lists of priority waterbodies, and State lists of waters that are not meeting water quality standards. Regulatory authorities should use this information as a means of identifying point sources that discharge to impaired waterbodies and that thus may be contributing to this impairment. One source of this information is the lists of waters generated by states to comply with Section 304(l) regulations at 40 *CFR* 130.10(d)(6); 50 *FR* 23897-98, June 2, 1989:
  - 1) Waters where fishing or shellfish bans and/or advisories are currently in effect or are anticipated;
  - 2) Waters where there have been repeated fish kills or where abnormalities (cancers, lesions, tumors, etc.) have been observed in fish or other aquatic life during the last ten years;
  - 3) Waters where there are restrictions on water sports or recreational contact;
  - 4) Waters identified by the state in its most recent state section 305(b) report as either "partially achieving" or "not achieving" designated uses;

- 5) Waters identified by the states under section 303(d) of the Clean Water Act as waters needing water quality-based controls;
- 6) Waters identified by the state as priority water bodies;
- 7) Waters where ambient data indicate potential or actual excursions of water quality criteria due to toxic pollutants from an industry classified as a primary industry in Appendix A of 40 CFR Part 122;
- 8) Waters for which effluent toxicity test results indicate possible or actual excursions of state water quality standards, including narrative "free from" water quality criteria or EPA water quality criteria where state criteria are not available;
- 9) Waters with primary industrial major dischargers where dilution analyses indicate exceedances of state narrative or numeric water quality criteria (or EPA water quality criteria where state standards are not available) for toxic pollutants, ammonia, or chlorine;
- 10) Waters with POTW dischargers requiring local pretreatment programs where dilution analyses indicate exceedances of state water quality criteria (or EPA water quality criteria where state water quality criteria are not available) for toxic pollutants, ammonia, or chlorine;
- 11) Waters with facilities not included in the previous two categories such as major POTWs, and industrial minor dischargers where dilution analyses indicate exceedances of numeric or narrative state water quality criteria (or EPA water quality criteria where state water quality criteria are not available) for toxic pollutants, ammonia, or chlorine;
- 12) Water classified for uses that will not support the "fishable/swimmable" goals of the Clean Water Act;
- 13) Waters where ambient toxicity or adverse water quality conditions have been reported by local, state, EPA or other Federal Agencies, the private sector, public interest groups, or universities;
- 14) Waters identified by the state as impaired in its most recent Clean Lake Assessments conducted under 314 of the Clean Water Act; and
- 15) Surface waters impaired by pollutants from hazardous waste sites on the National Priority List prepared under section 105(8)(A) of CERCLA.
- 16) Waters judged to be impaired as a result of a bioassessment/biosurvey.

The presence of a combination of these factors, such as low available dilution, high-quality receiving water, poor compliance record, and clustered industrial and municipal discharges, could constitute a high priority for effluent limits.

Regardless, the regulatory authority, if it chooses to impose an effluent limit after conducting an effluent assessment without facility-specific monitoring data, will need to provide adequate justification for the limit in its permit development rationale or in its permit fact sheet. A clear and logical rationale for the need for the limit covering all of the regulatory points will be necessary to defend the limit should it be challenged. In justification of a limit, **EPA recommends that the more information the authority can acquire to support the limit, the better a position the authority will be in to defend the limit if necessary.** In such a case, the regulatory authority may well benefit from the collection of effluent monitoring data prior to establishing the limit.

If the regulatory authority, after evaluating all available information on the effluent, in the absence of effluent monitoring data, is not able to decide whether the discharge causes, has the reasonable potential to cause, or contributes to, an excursion above a numeric or narrative criterion for whole effluent toxicity or for individual toxicants, the authority should require whole effluent toxicity or chemical-specific testing to gather further evidence. In such a case, the regulatory authority can require the monitoring prior to permit issuance, if sufficient time exists, or it may require the testing as a condition of the issued/reissued permit.

Under these circumstances, the regulatory authority may find it protective of water quality to include a permit reopener for the imposition of an effluent limit should the effluent testing establish that the discharge causes, has the reasonable potential to cause, or contributes to excursion above a water quality criteria. A discussion of these options is provided later in this chapter.

### **3.3 DETERMINING THE NEED FOR PERMIT LIMITS WITH EFFLUENT MONITORING DATA**

#### **3.3.1 General Considerations**

When characterizing an effluent for the need for a whole effluent toxicity limit, and/or an individual toxicant limit, the regulatory authority should use any available effluent monitoring data, together with any information like that discussed under Section 3.2 above, as the basis for a decision. The regulatory authority may already have effluent toxicity data available from previous monitoring, or it may decide to require the permittee to generate effluent monitoring data prior to permit issuance or as a condition of the issued permit. EPA regulations at 40 CFR 122.21(j) require POTWs with design flows equal to or greater than 1 MGD and POTWs with approved pretreatment programs, or POTWs required to develop a pretreatment program, to submit the results of whole effluent toxicity tests with their permit applications. These regulations also provide discretion to the permitting authority to request such data from additional POTWs at the time of permit application.

In the instance where the permittee is required to generate data in advance, data collection should begin 12 to 18 months in advance of permit development to allow adequate time for conducting toxicity tests and chemical analyses. The type of data, including toxicity testing data, should be specified by the regulatory authority at the outset so that decisions on permit actions will not be delayed. **EPA recommends monitoring data be generated on effluent toxicity prior to permit limit development for the following reasons: (1) the presence or absence of effluent toxicity can be more clearly established or refuted and (2) where toxicity is shown, effluent variability can be more clearly defined.** Several basic factors that should be considered in generating effluent monitoring data are discussed below.

### 3.3.2 Addressing Uncertainty in Effluent Characterization by Generating Effluent Monitoring Data

All toxic effects testing and exposure assessment parameters, for both effluent toxicity and individual chemicals, have some degree of uncertainty associated with them. The more limited the amount of test data available, the larger the uncertainty. The least amount of uncertainty of an effluent's impact on the receiving water exists where (1) a complete data base is available on the effects of acute and chronic toxicity on many indigenous species, (2) there is a clear understanding of ecosystem species composition and functional processes, and (3) actual measured exposure concentrations are available for all chemicals during seasonal changes and dilution situations. The uncertainty associated with such an ideal situation would be minimal. However, generation of these data can be very resource intensive.

An example of uncertainty that results from limited monitoring data is if a regulatory authority has only one piece of effluent data (e.g., an LC<sub>50</sub> of 50 percent) for a facility. Effluent variability in such a case, given the range of effluent toxicity variability seen in other effluents, may range between 20 percent and 100 percent (see Appendix A). It is impossible to determine from one piece of monitoring data where in this range the effluent variability really falls. More monitoring data would need to be generated to determine the actual variability of this effluent and reduce this source of uncertainty.

To better characterize the effects of effluent variability and reduce uncertainty in the process of deciding whether to require an effluent limit, EPA has developed the statistical approach described below and in Box 3-2. This approach combines knowledge of effluent variability as estimated by a coefficient of variation with the uncertainty due to a limited number of data to project an estimated maximum concentration for the effluent. The estimated maximum concentration is calculated as the upper bound of the expected lognormal distribution of effluent concentrations at a high confidence level. The projected effluent concentration after consideration of dilution can then be compared to an appropriate water quality criterion to determine the potential for exceeding that criterion and the need for an effluent limit.

The statistical approach has two parts. The first is a characterization of the highest measured effluent concentration based on the desired confidence level. The relationship that describes this is the following:

$$p_n = (1 - \text{confidence level})^{1/n}$$

where  $p_n$  is the percentile represented by the highest concentration in the data and  $n$  is the number of samples. The following are some examples of this relationship at a 99 percent confidence level:

- The largest value of 5 samples is greater than the 40 percentile
- The largest value of 10 samples is greater than the 63 percentile
- The largest value of 20 samples is greater than the 79 percentile
- The largest value of 100 samples is greater than the 96 percentile.

The second part of the statistical approach is a relationship between the percentile described above and the selected upper bound of the lognormal effluent distribution. EPA's effluent data base suggests that the lognormal distribution well characterizes effluent concentrations (see Appendix E). For example, if five samples were collected (which represents a 40th percentile), the coefficient of variation is 0.6, and the desired upper bound of the effluent distribution is the 99th percentile, then the two percentiles can be related using the coefficient of variation (CV) as shown below:

$$\frac{C_{99}}{C_{40}} = \frac{\exp(2.326\sigma - 0.5\sigma^2)}{\exp(-0.258\sigma - 0.5\sigma^2)} = 4.2$$

where  $\sigma^2 = \ln(CV^2 + 1)$  and 2.326 and -0.258 are the normal distribution values for the 99th and 40th percentiles, respectively. The use of the 99th percentile is for illustrative purposes here. Although it does represent a measure of the upper bound of an effluent distribution, other percentiles could be selected by a regulatory agency. The relationship shown above can be calculated for other percentiles and CVs by replacing the values in the equation.

Tables 3-1 and 3-2 show the combined effects of both parts for a 99-percent confidence level and upper bounds of the 99th and 95th percentiles, respectively. The factors shown in the tables are multiplied by the highest concentration in an effluent sample to estimate the maximum expected concentration.

This procedure can be used for both single and multiple discharges to the same receiving waterbody. This is accomplished for multiple dischargers by summing the projected RWCs for the pollutant or pollutant parameter of concern from each individual discharger, and comparing it to the water quality standard. This involves an assumption of conservative additivity of the pollutant after discharge, which may not accurately reflect the true behavior of the toxicant. To overcome this, and to further refine the proportional contribution of each discharger and the resultant limits, the permitting authority should supplement this evaluation with multiple source WLA modeling and/or ambient water concentration monitoring.

### Box 3-2. Determining "Reasonable Potential" for Excursions Above Ambient Criteria Using Effluent Data Only

EPA recommends finding that a permittee has "reasonable potential" to exceed a receiving water quality standard if it cannot be demonstrated with a high confidence level that the upper bound of the lognormal distribution of effluent concentrations is below the receiving water criteria at specified low-flow conditions.

- Step 1** Determine the number of total observations (" $n$ ") for a particular set of effluent data (concentrations or toxic units [TUs]), and determine the highest value from that data set.
- Step 2** Determine the coefficient of variation for the data set. For a data set where  $n < 10$ , the coefficient of variation (CV) is estimated to equal 0.6, or the CV is calculated from data obtained from a discharger. For a data set where  $n > 10$ , the CV is calculated as standard deviation/mean (see Figure 3-1). For less than 10 items of data, the uncertainty in the CV is too large to calculate a standard deviation or mean with sufficient confidence.
- Step 3** Determine the appropriate ratio from Table 3-1 or 3-2.
- Step 4** Multiply the highest value from a data set by the value from Table 3-1 or 3-2. Use this value with the appropriate dilution to project a maximum receiving water concentration (RWC).
- Step 5** Compare the projected maximum RWC to the applicable standard (criteria maximum concentration, criteria continuous concentration [CCC], or reference ambient concentration). EPA recommends that permitting authorities find reasonable potential when the projected RWC is greater than an ambient criterion.

#### Example

Consider the following results of toxicity measurements of an effluent that is being characterized: 5  $TU_C$ , 2  $TU_C$ , 9  $TU_C$ , and 6  $TU_C$ . Assume that the effluent is diluted to 2 percent at the edge of the mixing zone. Further assume that the CV is 0.6, the upper bound of the effluent distribution is the 99th percentile, and the confidence level is 99 percent.

- Step 1** There are four samples, and the maximum value of the sample results is 9  $TU_C$ .
- Step 2** The value of the CV is 0.6.
- Step 3** The value of the ratio for four pieces of data and a CV of 0.6 is 4.7.
- Step 4** The value that exceeds the 99th percentile of the distribution (ratio times  $x_{max}$ ) after dilution is calculated as:
- $$[9 TU_C \times 4.7 \times 0.02] = 0.85 TU_C.$$
- Step 5** 0.85  $TU_C$  is less than the ambient criteria concentration of 1.0  $TU_C$ . There is no reasonable potential for this effluent to cause an excursion above the CCC.

### 3.3.3 Effluent Characterization for Whole Effluent Toxicity

Once an effluent has been selected for whole effluent toxicity characterization after consideration of the factors discussed above, the regulatory authority should require toxicity testing in accordance with appropriate site-specific considerations and the recommendations discussed below. In the past 5 years, significant additional experience has been gained in generating effluent toxicity data upon which to make decisions as to whether or not an effluent will cause toxic effects in the receiving water in both freshwater and marine environments.

### General Considerations and Assumptions

EPA has revised its initial effluent toxicity data generation recommendations based on three observations made over the last 5 years:

- 1) Only rarely have effluents discharged by NPDES permittees been observed to have  $LC_{50}$ s less than 1.0 percent or no observed effect concentrations (NOECs) less than 0.1 percent. However, there is always a chance that an effluent could be toxic at such low effluent concentrations.

**Table 3-1. Reasonable Potential Multiplying Factors: 99% Confidence Level and 99% Probability Basis**

Number of Samples	Coefficient of Variation																			
	0.1	0.2	0.3	0.4	0.5	0.6	0.7	0.8	0.9	1.0	1.1	1.2	1.3	1.4	1.5	1.6	1.7	1.8	1.9	2.0
1	1.6	2.5	3.9	6.0	9.0	13.2	18.9	26.5	36.2	48.3	63.3	81.4	102.8	128.0	157.1	190.3	227.8	269.9	316.7	368.3
2	1.4	2.0	2.9	4.0	5.5	7.4	9.8	12.7	16.1	20.2	24.9	30.3	36.3	43.0	50.4	58.4	67.2	76.6	86.7	97.5
3	1.4	1.9	2.5	3.3	4.4	5.6	7.2	8.9	11.0	13.4	16.0	19.0	22.2	25.7	29.4	33.5	37.7	42.3	47.0	52.0
4	1.3	1.7	2.3	2.9	3.8	4.7	5.9	7.2	8.7	10.3	12.2	14.2	16.3	18.6	21.0	23.6	26.3	29.1	32.1	35.1
5	1.3	1.7	2.1	2.7	3.4	4.2	5.1	6.2	7.3	8.6	10.0	11.5	13.1	14.8	16.6	18.4	20.4	22.4	24.5	26.6
6	1.3	1.6	2.0	2.5	3.1	3.8	4.6	5.5	6.4	7.5	8.6	9.8	11.1	12.4	13.8	15.3	16.8	18.3	19.9	21.5
7	1.3	1.6	2.0	2.4	2.9	3.6	4.2	5.0	5.8	6.7	7.7	8.7	9.7	10.8	12.0	13.1	14.4	15.6	16.9	18.2
8	1.2	1.5	1.9	2.3	2.8	3.3	3.9	4.6	5.3	6.1	6.9	7.8	8.7	9.6	10.6	11.6	12.6	13.6	14.7	15.8
9	1.2	1.5	1.8	2.2	2.7	3.2	3.7	4.3	5.0	5.7	6.4	7.1	7.9	8.7	9.6	10.4	11.3	12.2	13.1	14.0
10	1.2	1.5	1.8	2.2	2.6	3.0	3.5	4.1	4.7	5.3	5.9	6.6	7.3	8.0	8.8	9.5	10.3	11.0	11.8	12.6
11	1.2	1.5	1.8	2.1	2.5	2.9	3.4	3.9	4.4	5.0	5.6	6.2	6.8	7.4	8.1	8.8	9.4	10.1	10.8	11.5
12	1.2	1.4	1.7	2.0	2.4	2.8	3.2	3.7	4.2	4.7	5.2	5.8	6.4	7.0	7.5	8.1	8.8	9.4	10.0	10.6
13	1.2	1.4	1.7	2.0	2.3	2.7	3.1	3.6	4.0	4.5	5.0	5.5	6.0	6.5	7.1	7.6	8.2	8.7	9.3	9.9
14	1.2	1.4	1.7	2.0	2.3	2.6	3.0	3.4	3.9	4.3	4.8	5.2	5.7	6.2	6.7	7.2	7.7	8.2	8.7	9.2
15	1.2	1.4	1.6	1.9	2.2	2.6	2.9	3.3	3.7	4.1	4.6	5.0	5.4	5.9	6.4	6.8	7.3	7.7	8.2	8.7
16	1.2	1.4	1.6	1.9	2.2	2.5	2.9	3.2	3.6	4.0	4.4	4.8	5.2	5.6	6.1	6.5	6.9	7.3	7.8	8.2
17	1.2	1.4	1.6	1.9	2.1	2.5	2.8	3.1	3.5	3.8	4.2	4.6	5.0	5.4	5.8	6.2	6.6	7.0	7.4	7.8
18	1.2	1.4	1.6	1.8	2.1	2.4	2.7	3.0	3.4	3.7	4.1	4.4	4.8	5.2	5.6	5.9	6.3	6.7	7.0	7.4
19	1.2	1.4	1.6	1.8	2.1	2.4	2.7	3.0	3.3	3.6	4.0	4.3	4.6	5.0	5.3	5.7	6.0	6.4	6.7	7.1
20	1.2	1.3	1.6	1.8	2.0	2.3	2.6	2.9	3.2	3.5	3.8	4.2	4.5	4.8	5.2	5.5	5.8	6.1	6.5	6.8

**Table 3-2. Reasonable Potential Multiplying Factors: 95% Confidence Level and 95% Probability Basis**

Number of Samples	Coefficient of Variation																			
	0.1	0.2	0.3	0.4	0.5	0.6	0.7	0.8	0.9	1.0	1.1	1.2	1.3	1.4	1.5	1.6	1.7	1.8	1.9	2.0
1	1.4	1.9	2.6	3.6	4.7	6.2	8.0	10.1	12.6	15.5	18.7	22.3	26.4	30.8	35.6	40.7	46.2	52.1	58.4	64.9
2	1.3	1.6	2.0	2.5	3.1	3.8	4.6	5.4	6.4	7.4	8.5	9.7	10.9	12.2	13.6	15.0	16.4	17.9	19.5	21.1
3	1.2	1.5	1.8	2.1	2.5	3.0	3.5	4.0	4.6	5.2	5.8	6.5	7.2	7.9	8.6	9.3	10.0	10.8	11.5	12.3
4	1.2	1.4	1.7	1.9	2.2	2.6	2.9	3.3	3.7	4.2	4.6	5.0	5.5	6.0	6.4	6.9	7.4	7.8	8.3	8.8
5	1.2	1.4	1.6	1.8	2.1	2.3	2.6	2.9	3.2	3.6	3.9	4.2	4.5	4.9	5.2	5.6	5.9	6.2	6.6	6.9
6	1.1	1.3	1.5	1.7	1.9	2.1	2.4	2.6	2.9	3.1	3.4	3.7	3.9	4.2	4.5	4.7	5.0	5.2	5.5	5.7
7	1.1	1.3	1.4	1.6	1.8	2.0	2.2	2.4	2.6	2.8	3.1	3.3	3.5	3.7	3.9	4.1	4.3	4.5	4.7	4.9
8	1.1	1.3	1.4	1.6	1.7	1.9	2.1	2.3	2.4	2.6	2.8	3.0	3.2	3.3	3.5	3.7	3.9	4.0	4.2	4.3
9	1.1	1.2	1.4	1.5	1.7	1.8	2.0	2.1	2.3	2.4	2.6	2.8	2.9	3.1	3.2	3.4	3.5	3.6	3.8	3.9
10	1.1	1.2	1.3	1.5	1.6	1.7	1.9	2.0	2.2	2.3	2.4	2.6	2.7	2.8	3.0	3.1	3.2	3.3	3.4	3.6
11	1.1	1.2	1.3	1.4	1.6	1.7	1.8	1.9	2.1	2.2	2.3	2.4	2.5	2.7	2.8	2.9	3.0	3.1	3.2	3.3
12	1.1	1.2	1.3	1.4	1.5	1.6	1.7	1.9	2.0	2.1	2.2	2.3	2.4	2.5	2.6	2.7	2.8	2.9	3.0	3.0
13	1.1	1.2	1.3	1.4	1.5	1.6	1.7	1.8	1.9	2.0	2.1	2.2	2.3	2.4	2.5	2.5	2.6	2.7	2.8	2.9
14	1.1	1.2	1.3	1.4	1.4	1.5	1.6	1.7	1.8	1.9	2.0	2.1	2.2	2.3	2.3	2.4	2.5	2.6	2.6	2.7
15	1.1	1.2	1.2	1.3	1.4	1.5	1.6	1.7	1.8	1.8	1.9	2.0	2.1	2.2	2.2	2.3	2.4	2.4	2.5	2.5
16	1.1	1.1	1.2	1.3	1.4	1.5	1.6	1.6	1.7	1.8	1.9	1.9	2.0	2.1	2.1	2.2	2.3	2.3	2.4	2.4
17	1.1	1.1	1.2	1.3	1.4	1.4	1.5	1.6	1.7	1.7	1.8	1.9	1.9	2.0	2.0	2.1	2.2	2.2	2.3	2.3
18	1.1	1.1	1.2	1.3	1.3	1.4	1.5	1.6	1.6	1.7	1.7	1.8	1.9	1.9	2.0	2.0	2.1	2.1	2.2	2.2
19	1.1	1.1	1.2	1.3	1.3	1.4	1.5	1.5	1.6	1.6	1.7	1.8	1.8	1.9	1.9	2.0	2.0	2.0	2.1	2.1
20	1.1	1.1	1.2	1.2	1.3	1.4	1.4	1.5	1.5	1.6	1.7	1.7	1.8	1.8	1.8	1.9	1.9	2.0	2.0	2.0

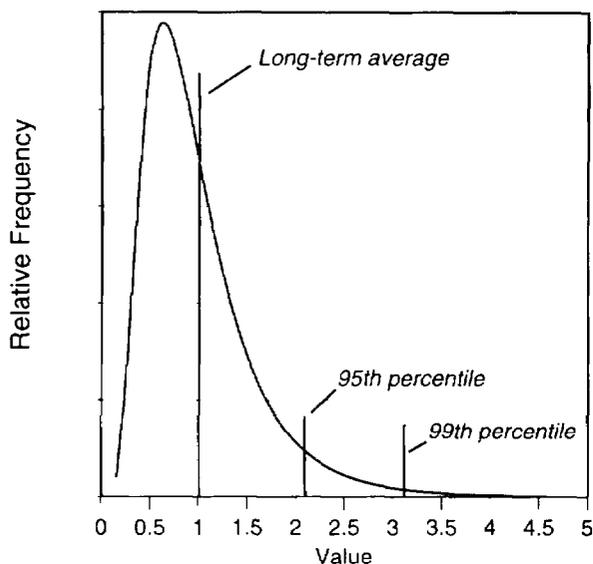


Figure 3-1a. Frequency Distribution of Values for a Lognormal Distribution with a Mean of 1.0 and a Coefficient of Variation of 0.6

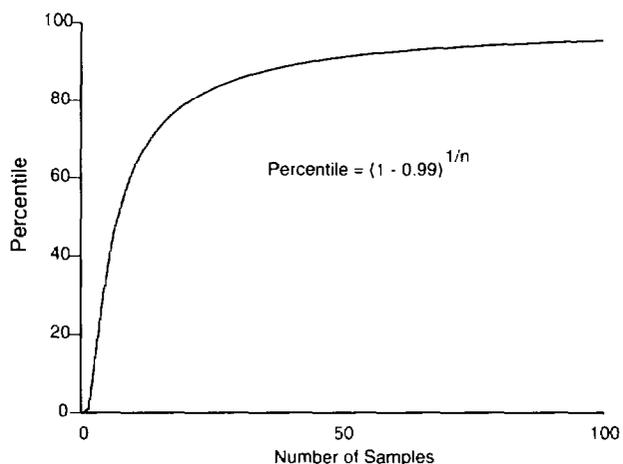


Figure 3-1c. Relationship Between the Largest Value of n Samples and the Percentile It Exceeds with 99 Percent Confidence

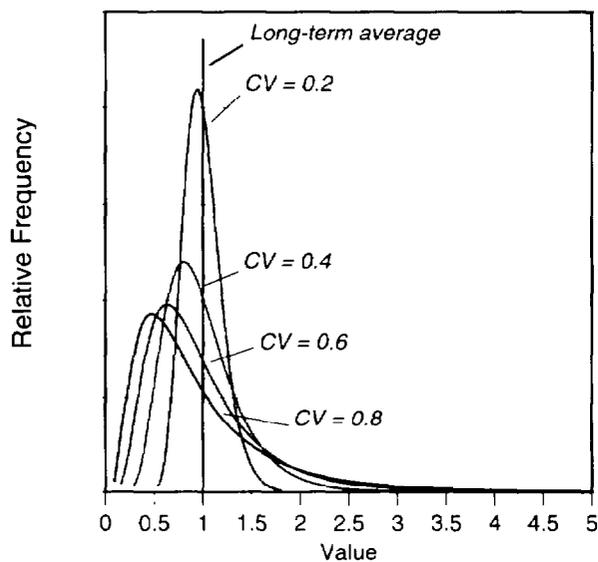


Figure 3-1b. Comparison of Relative Frequencies of Lognormal Distributions with a Mean of 1.0 for Different Coefficients of Variation

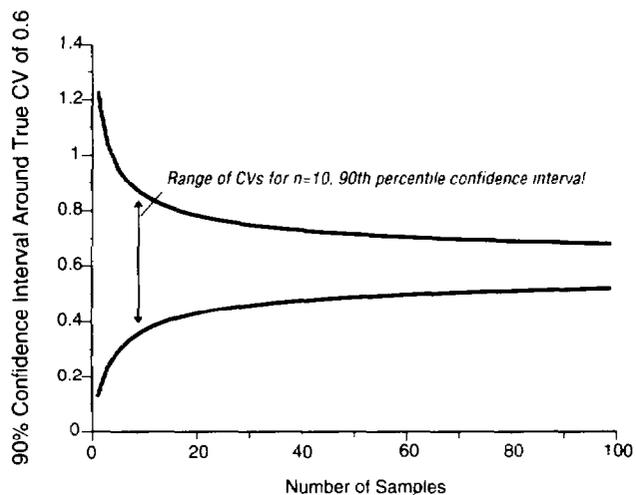
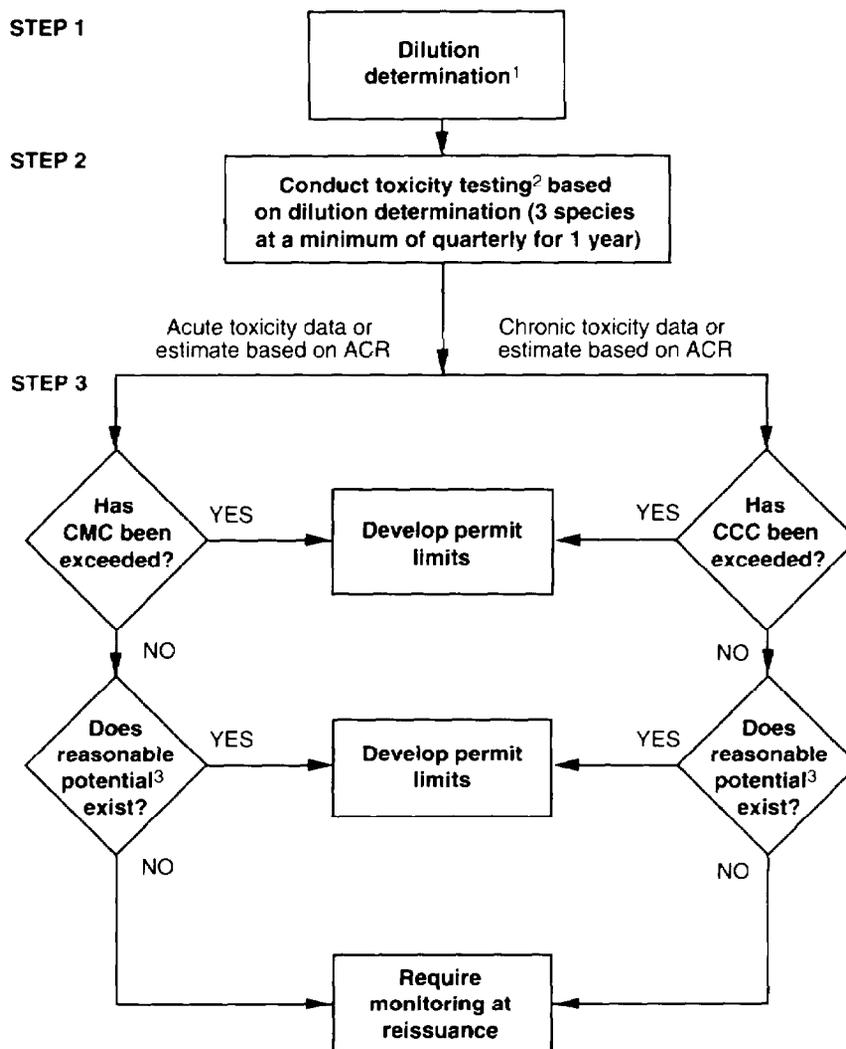


Figure 3-1d. Example of 90 Percent Confidence Intervals Around Coefficient of Variation Estimates for Numbers of Samples

- 2) With the exception of a small number of "outliers" for which confirmation is not possible, acute-to-chronic ratios (ACRs) above 20 for effluents discharged by NPDES permittees have not been observed by EPA. The majority of observed ACRs are very seldom above 10. However, higher ACRs may be found for selected facilities.
- 3) The use of the three commonly used freshwater species and of three of the five commonly used marine organisms has generally been sufficient to measure any effluent's toxicity for the purposes of projecting effluent toxicity impact and making regulatory decisions.

Figure 3-2 is a flow chart of EPA's recommendations for data generation for three different dilution scenarios. It is divided into three basic steps: determining initial dilution, developing toxicity testing procedures, and developing decision criteria for permit limit. There are certain basic assumptions built into this flow chart. The basic principle used in making decisions is to compare available dilution to known or projected toxic effect concentrations in order to place an effluent into one of three categories:



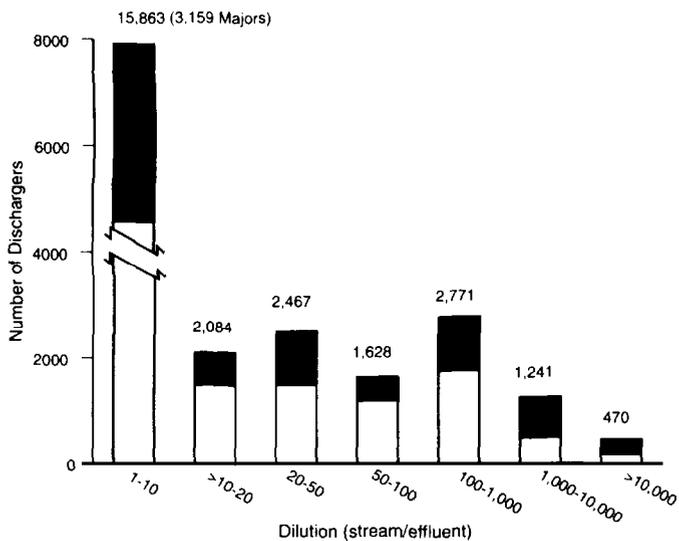
Notes:

- <sup>1</sup>Dilution determinations should be performed for critical flows and any applicable mixing zones.
- <sup>2</sup>Toxicity testing recommendations
  - a. Dilution > 1000:1: acute testing, check CMC only.
  - b. 100:1 < Dilution < 1000:1: acute or chronic testing, check CMC and CCC with data or ACR.
  - c. Dilution < 100:1: conduct chronic testing, check CCC with data and CMC using acute data or ACR.
- <sup>3</sup>Reasonable potential: Use procedures in Box 3-3.

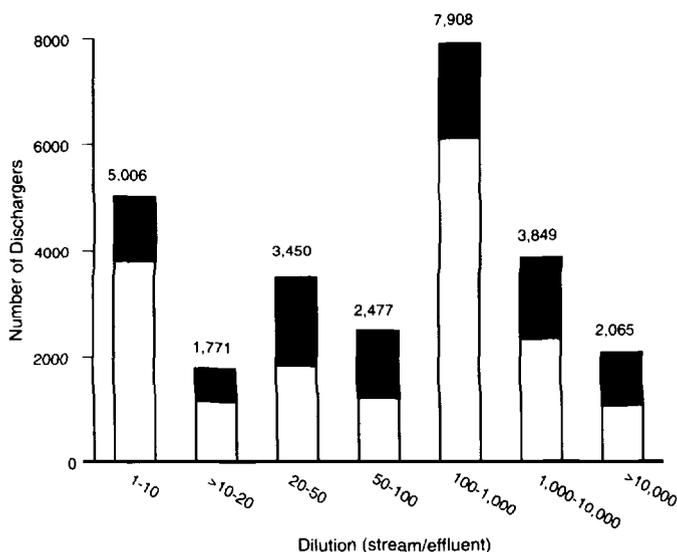
Figure 3-2. Effluent Characterization for Whole Effluent Toxicity

- 1) The effluent causes or contributes to an excursion of a numeric or narrative water quality criterion and the permit requires a limit on toxicity.
- 2) The effluent has a reasonable potential of causing or contributing to an excursion of a numeric or narrative water quality criterion and a limit is required.
- 3) The effluent has a very low probability of causing or contributing to an excursion of a water quality standard and no limit is required.

This categorization is accomplished by using dilution estimates in the first step and the results of the toxicity tests in the next steps. In addition, all these impact estimates assume discharge at critical conditions and imposition of any applicable mixing zone requirements. Therefore, a conservative assumption is used to determine whether or not an impact is projected to occur. Estimates of possible toxic impact are made assuming that the effluent is most toxic to the most sensitive species or lifestage at the time of lowest available dilution.



(a) At Low Flow (7Q10)



(b) At Annual Mean Flow

**Figure 3-3. National Distribution of NPDES Dilution Conditions at 7Q10 and at Annual Mean Flow**

The changes to the EPA's data generation recommendations eliminate the application of multiple sets of safety margins that was proposed in the 1985 version of this document. Rather, general observations on effluent toxicity described above now allow regulatory authorities to tighten the bounds of the initial dilution categorization, eliminate the species sensitivity uncertainty factor and target LC<sub>50</sub>s of 1 percent and NOECs of 0.1 percent as the most extreme toxicity measurements that can normally be expected for the vast majority of effluents discharged by NPDES permittees for acute and chronic toxicity, respectively. The observation of toxicity was based on multiple dilution tests. The same observation may not hold for toxicity measured with single dilution tests (pass/fail). As reflected in Chapter 1, single

dilution toxicity tests are much more variable than multiple dilution tests. **Therefore, the use of single concentration toxicity tests is strongly discouraged for this data generation process.**

Since the new data generation requirements are much less expensive than the previous requirements, tiered testing (less expensive, single-concentration, initial screening followed by increasingly expensive definitive data generation, using multiconcentration tests, as described in the September 1985 version of the technical support document) is unnecessary. However, **elimination of the requirement to conduct toxicity testing on the basis of projections using dilution alone is not recommended.** Although EPA's data review suggests that an LC<sub>50</sub> of 1 percent and an NOEC of 0.1 percent are the lower bounds on effluent toxicity, there may be other effluents that are presently unmeasured that are more toxic. Testing data are always desirable for fully characterizing discharges of concern.

**Steps in Whole Effluent Characterization Process**

The following is a detailed description of the major steps presented in Figure 3-2 and the rationale behind each.

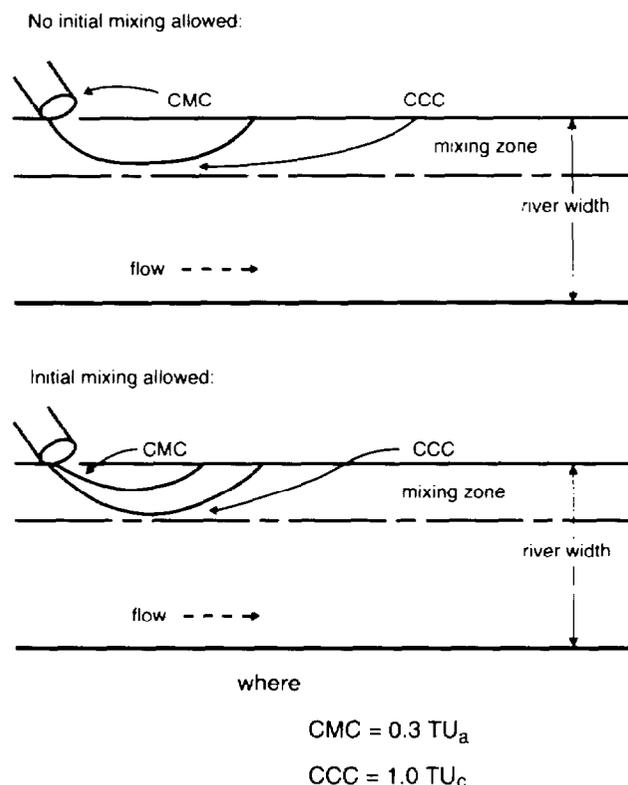
**Step 1: Dilution Determination**

The initial step is to determine the dilution of the effluent at the edge of the mixing zone, assuming the State allows mixing zones. Figure 3-4 shows a schematic representation of typical mixing zone requirements for both acute and chronic toxicity. Calculating the dilution at the edges of mixing zones for site-specific situations can be complicated. Modeling can be employed using either steady-state or dynamic approaches to calculate the dilution (see Chapter 4). However, for complex situations, such as marine and estuarine waters or lakes, dye studies (or other techniques used to assess mixing zones) may still be required.

Some State water quality standards do not allow the use of mixing in the control of acute toxicity. For these States, acute toxicity is often limited at the end of the pipe. Permit limits derived to enforce such requirements would be considered "water quality-based" because they would be based upon an ambient criterion (as opposed to an arbitrary test endpoint). Regardless, both chronic and acute toxicity must be assessed in these situations.

**Step 2: Toxicity Testing Procedures**

Where toxicity tests are required in order to make decisions regarding appropriate next steps in a screening protocol, **EPA recommends as a minimum that three species (for example, a vertebrate, an invertebrate, and a plant) be tested quarterly for a minimum of 1 year.** As discussed in Chapter 1, the use of three species is strongly recommended. Experience indicates that marine algae can be a highly sensitive test species for some effluents. Using a surrogate species of the plant kingdom adds another trophic level to the testing regimen. For both freshwater and marine situations, the use of three species is more protective than two species since a wider range of species sensitivity can be measured. EPA is continuing to develop toxicity test methods using additional organisms including plants. In addition, EPA has revised the test for *Selenastnum*, which has improved the test precision.



**Figure 3-4. Schematic Representation of Mixing Zone Areas Where the CMC and CCC Apply**

**EPA recommends against selecting a “most sensitive” species for toxicity testing.** For one organism to consistently be the most sensitive in a battery of toxicity tests, two conditions must occur: (1) the toxicants causing toxicity must remain the same, and (2) the ratios of the toxicants in the effluent (if more than one) must remain the same. Based on EPA’s experience at the Duluth research laboratory, neither of these conditions is likely to occur. For example, the causes of effluent toxicity in POTWs can vary on a seasonal basis. Toxicity in the summer can be caused by pesticides to which invertebrates are most sensitive. However, the winter toxicity could be caused by ammonia to which fathead minnows will respond most sensitively. The most sensitive species for an effluent actually may not exist and at best is difficult to identify.

**Conducting toxicity tests using three species quarterly for 1 year is recommended to adequately assess the variability of toxicity observed in effluents.** Below this minimum, the chances of missing toxic events increase. The toxicity test result for the most sensitive of the tested species is considered to be the measured toxicity for a particular effluent sample.

The data generation recommendations in Figure 3-2 represent minimum testing requirements. Since uncertainty regarding whether or not an effluent causes toxic impact is reduced with more data, **EPA recommends that this test frequency be increased where necessary to adequately assess effluent vari-**

**ability.** If less frequent testing is required in the permit, it is preferable to use three species tested less frequently than to test the effluent more frequently with only a single species whose sensitivity to the effluent is not well characterized.

**EPA recommends that a discharger conduct acute toxicity testing if the dilution of the effluent is greater than 1000:1 at the edge of the mixing zone [3].** Such a discharger would be considered a low priority for chronic toxicity testing. The rationale for this is that the effluent concentration would be below 0.1 percent at the edge of the mixing zone and thus incapable of causing an excursion above the CCC. A worst case NOEC of 0.1 percent translates into 1,000  $\text{TU}_c$ , which would result in a concentration of less than 1.0  $\text{TU}_c$  at the edge of the mixing zone for this dilution category. The test results would be compared to the CMC after consideration of any allowable mixing.

**EPA recommends that a discharger conduct either acute or chronic toxicity testing if the dilution of the effluent falls between 100:1 and 1,000:1 at the edge of the mixing zone.** Effluents have been shown to be both acutely and chronically toxic within this range of receiving water dilution. Under worst-case scenarios,  $\text{LC}_{50}$ s of 1.0 percent and ACRs of 10 will result in excursions above both the CCC and CMC at the edge of the regulatory mixing zone.

Although either acute or chronic testing can be required within this dilution range, acute testing would be more appropriate at the higher end of this dilution range (1,000:1 or 0.1 percent). At the lower end of this dilution range (100:1 or 1.0 percent), chronic tests may be more appropriate. Where other factors are equal, chronic testing may be preferable since the interim results in a chronic test gives data on acute toxicity as well. The acute endpoint data can then be used to compare directly to the CMC without the need for an ACR.

Whichever type of toxicity test (either acute or chronic) is specified, the results from that test should be compared to the criterion associated with that type of test. For example, a chronic test would be compared to the CCC. Comparisons to the other criteria can be made by using the ACR or additional data generated to convert a chronic test result to an acute endpoint and vice versa. For example, a chronic NOEC of 5 percent effluent (or 20  $\text{TU}_c$ ) represents an acute  $\text{LC}_{50}$  of 50 percent (or 2  $\text{TU}_a$ ) at an ACR of 10.

**EPA recommends that a discharger conduct chronic toxicity testing if the dilution of the effluent falls below 100:1 at the edge of the mixing zone.** The rationale for this recommendation is that chronic toxicity has been observed in some effluents down to the 1.0 percent effect concentration. Therefore, chronic toxicity tests, although somewhat more expensive to conduct, should be used directly in order to make decisions about toxic impact.

There is a potential for acute toxicity within this dilution range, although this is less likely as the 100:1 dilution level is approached. Thus, the recommended screening protocol shown in Figure 3-2 includes a determination of whether excursions above the CMC are projected [4]. This analysis may be performed by assuming an ACR, applying this value to the chronic toxicity testing data, and allowing for any allowable initial mixing. Alternatively, the regulatory authority may use the interim results in the chronic test to calculate the acute toxicity.

Both the chronic and acute toxicity test data would be compared to their respective criterion. The chronic test results would be compared to the CCC, and the acute results, regardless of how calculated, would be compared to the CMC.

### Step 3: Decision Criteria for Permit Limit Development

Once the toxicity data have been generated for a discharger, the regulatory authority must decide whether or not the results show that the permittee causes, has the reasonable potential to cause, or contributes to an excursion of an applicable numeric or narrative water quality criterion and therefore needs to limit effluent toxicity. To do this, these data should be used to project receiving water concentrations, which are then compared to the CCC and CMC. One of four outcomes will be reached when following the screening protocol shown in Figure 3-2:

- 1) **Excursion Above CMC or CCC**—Where any one data point shows an excursion above the State’s numeric or narrative criterion for the parameter toxicity, EPA regulations require a permit limit be set for whole effluent toxicity (40 *CFR* 122.44(d)(1)(iv or v)), unless limits on a specific chemical will allow the narrative water quality criterion to be attained or maintained. In the absence of a State numeric criterion for the parameter toxicity, **EPA recommends that 1.0 TU<sub>c</sub> and 0.3 TU<sub>a</sub> be used as the CCC and CMC, respectively.** The decision to develop permit limits based upon an excursion above either the CMC or CCC will lead to protection against both acute and chronic toxicity if the permit derivation procedures in Chapter 5 are used to set effluent limits.
- 2) **Reasonable Potential for Excursion Above CMC or CCC**—EPA believes that “reasonable potential” is shown where an effluent is projected to cause an excursion above the CCC or CMC. This projection is based upon a statistical analysis of available data that accounts for limited sample size and effluent variability. EPA’s detailed recommendations for making a statistical determination based upon effluent monitoring data alone are shown in Box 3-2. Where a regulatory authority finds that test results alone indicate a “reasonable potential” to cause an excursion above a State water quality criterion in accordance with 40 *CFR* 122.44(d)(1)(ii), a permit limit must be developed.

A regulatory authority may select an alternative approach for assessing reasonable potential. For example, an authority may opt to use a stochastic dilution model that incorporates both ambient dilution and effluent variability for determining reasonable potential. Such an approach is analogous to the statistical approach shown in Box 3-2. Whatever approach selected by the authority, it must use all the factors that account for all the factors listed in 40 *CFR* 122.44(d)(1)(ii).

In some cases the statistical analysis of the effluent data may not actually project an excursion above the CMC or CCC but may be close. Under such conditions, reasonable potential determinations will include an element of judgment on the part of the regulatory authority. Other factors will need to be considered and given appropriate weight in the decisionmaking process, including value of waterbody (e.g., high-use fishery), relative proximity to the CCC or CMC, existing controls on point and nonpoint sources, informa-

tion on effluent variability, compliance history of the facility, and type of treatment facility. These factors are summarized in Box 3-2 and are discussed in detail in Section 3.1. **EPA recommends regulatory authorities establish a written policy and procedure for making determinations of “reasonable potential” under these circumstances.**

- 3) **No Reasonable Potential for Excursions Above CMC or CCC**—In these situations, **EPA recommends that the toxicity tests recommended above be repeated at a frequency of at least once every 5 years as a part of the permit application.** Such testing is required for certain POTWs under 40 *CFR* 122.21(j).
- 4) **Inadequate Information**—Where a regulatory authority has inadequate information to determine reasonable potential for an excursion of a numeric or narrative water quality criterion, there may still be a basis for concern on the part of the authority. The permit should contain whole effluent toxicity monitoring requirements and a reopener clause. This clause would require reopening of the permit and establishment of a limit based upon any test results, or other new factors, which substantiate that the effluent causes, has the reasonable potential of causing, or contributes to an excursion above the CCC or CMC.

### 3.3.4 Use of Toxicity Testing in Multiple-source Discharge Situations

Where more than one discharge to the same receiving waterbody contributes, or has the reasonable potential to contribute to an excursion of water quality standards, permit limits must be developed for each individual discharger on that waterbody. For the regulatory authority to make this assessment, additional testing may be needed to provide the authority with the information necessary to assess the relative impact of each source. For purposes of this discussion, a multiple-source discharge situation is defined as a situation where impact zones overlap, or where ambient receiving water concentrations of a pollutant are elevated due to upstream discharges. In multiple-source discharge situations, additivity, antagonism, and persistence of toxicity can be of concern. To collect additional data, the permit authority should employ the toxicity testing procedures for multiple dischargers described in Box 3-3. In addition, ambient toxicity testing, as described below, could be used.

Assuming that screening has been conducted that reveals the need for permit limits, two options for controlling the discharges exist. The first option is for the permit authority to regulate each source separately using the procedures for individual point sources. In this option, the permitting authority would require use of upstream ambient water as a diluent in the toxicity test so as to be able to evaluate the contributions of upstream sources of toxicity. A second option is to treat each discharge as an interactive component of a whole system. In this option, the permit writer would determine a total maximum daily load for the receiving waterbody and develop individual wasteload allocations for each discharger using the procedures discussed in Chapter 4.



United States  
Environmental Protection Agency  
EN-336  
Washington, DC 20460

Official Business  
Penalty for Private Use  
\$300