

**ONTARIO'S WATER QUALITY
OBJECTIVE DEVELOPMENT
PROCESS**

MARCH 1992



Environment
Environnement

ISBN 0-7729-7532-9

ONTARIO'S WATER QUALITY OBJECTIVE DEVELOPMENT PROCESS

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MARCH 1992

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PIBS 1885
log 90-2309-042

Notice

The Ministry of the Environment sets Provincial Water Quality Objectives and Guidelines for the protection of aquatic life and recreation. Initiatives to develop multimedia criteria to ensure protection of all compartments of the environment are currently under way. Although PWQO/Gs provide full protection for aquatic life, development of multimedia criteria may result in the lowering of some PWQO/Gs to afford adequate protection for other compartments or receptors. As a result, the present PWQO/G should be regarded as *provisional* pending development of multimedia criteria.

ABSTRACT

As part of its program to ensure that the surface waters of Ontario are satisfactory for aquatic life and recreation, the Ontario Ministry of the Environment develops Provincial Water Quality Objectives (PWQOs) and Guidelines (PWQGs). This report details the procedures used to develop PWQOs and PWQGs. PWQOs are developed from the best available knowledge on the effects of substances on aquatic life. Data are required for acute and chronic toxicity, bioaccumulation and mutagenicity. Also considered is available information on environmental fate, physical-chemical properties, taste and odour in water and tainting of fish tissues. If adequate information is unavailable for setting a PWQO, a PWQG is developed using uncertainty factors which depend on the quantity and quality of the toxicological data.

RÉSUMÉ

Le ministère de l'Environnement de l'Ontario élabore des objectifs et des lignes directrices pour la qualité de l'eau de la province dans le cadre de son programme visant à rendre les eaux de surface propices à la vie aquatique et aux activités récréatives. Ce rapport explique les méthodes utilisées pour l'élaboration de ces objectifs et lignes directrices. Les objectifs provinciaux de qualité de l'eau sont établis à partir des données les plus récentes sur les effets de certaines substances sur la vie aquatique. Il reste cependant des lacunes à combler en ce qui concerne les seuils de toxicité aiguë et chronique, le facteur de bioaccumulation et la mutagénicité. Le rapport tient compte des renseignements existants sur le destin d'une substance dans l'environnement, les propriétés physico-chimiques, le goût et l'odeur de l'eau ainsi que l'altération de la chair des poissons. S'il est impossible d'obtenir suffisamment de renseignements pertinents pour établir des objectifs provinciaux de qualité de l'eau, des lignes directrices seront élaborées à partir de coefficients d'incertitude qui sont fonction de la quantité et de la qualité des données toxicologiques.

ACKNOWLEDGEMENTS

This document is the product of the Aquatic Criteria Development Committee. What started out as an attempt to formalize existing procedures for developing PWQOs turned into a reassessment of the process itself. The following people made significant contributions to the document through discussions within the committee and in writing drafts:

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The final draft report was widely circulated across Canada and the United States for scientific peer review. Many comments and suggestions were received, all were carefully evaluated and appropriate changes were subsequently made to the document. The final draft report was also reviewed by the Ministry of the Environment's Advisory Committee on Environmental Standards. The Aquatic Criteria Development Committee appreciates the assistance of all those people who contributed their time and knowledge to the completion of this document.

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1 INTRODUCTION

1.1 The Basis for Provincial Water Quality Objectives

The mandate for water quality management in Ontario is legislated under several statutes, most notably, the Water Resources Act. In addition, the Ministry of the Environment (MOE) has specific policies (MOE, 1984) with the goal:

"to ensure that the surface waters of the province are of a quality which is satisfactory for aquatic life and recreation."

To support this goal, the Ministry's policy further states that:

"... the Objectives are set at such values as to protect all forms of aquatic life and all aspects of the aquatic life cycles. The clear intention is to protect all life stages during indefinite exposure to the water."

In order to achieve these water management goals, a comprehensive understanding of, and sound management program for the control of physical, chemical and bacteriological pollutants is required. Since 1978, Ontario has relied on Provincial Water Quality Objectives and Guidelines as principal tools for surface water quality management.

Objectives are numerical and narrative values designed to protect all stages of aquatic life cycles during indefinite exposure to the water. They are not legal standards *per se*, but they are employed in various types of water quality models to establish acceptable wastewater loading limits on a

site-specific basis. The purpose of this document is to set forth detailed procedures for developing Provincial Water Quality Objectives (PWQOs). The PWQO development process closely adheres to the water management policy of the Ministry by requiring that PWQO setting considers a wide variety of representative aquatic organisms, critical life stages and toxic effects, including mutagenicity. All the following types of information are used in setting a PWQO (* indicates mandatory factors):

- * Toxicity to aquatic life (acute, chronic, lethal, sublethal)
- * Bioaccumulation (animals consuming aquatic organisms and humans consuming sport fish)
- * Mutagenicity to aquatic life
- * Fate and physical-chemical factors
- * Sources and ambient levels
- * Taste and odour in water and tainting of fish tissues
- * Impacts on wildlife
- * Recreation
- * Sediment quality
- * Objectives of other agencies (for comparison only).

PWQO setting is not influenced by analytical limitations, economics, social factors, or waste treatment availability. All of these are important in water management, and are considered in the site-specific application of the Objectives. PWQOs are designed to take into account both cold-water and warm-water environments in a single Objective, with the exception of the PWQO for dissolved oxygen. An assessment of toxicity to both cold and warm-water species

is a requirement of the PWQO setting process. Consequently, Objectives are designed to be applied independent of the nature of the possible resident species (e.g. cold-water or warm-water species), the type of waterbody (lake or stream) and its trophic status.

The influence of water temperature and other specific physical and chemical factors such as pH and hardness are, however, considered when they affect the toxicity of the candidate substances. The PWQO for ammonia, for example, is related to both pH and water temperature and the Objectives for some of the metals are related to water hardness.

The process for developing PWQOs has evolved with experience to effectively take into account crucial aspects of the hazard of substances in aquatic environments. The process is also efficient at developing Objectives and Guidelines with limited resources. The purpose of this document is to describe this process. By doing so, the applicability as well as the limitations of the Objectives will better be understood by water quality managers, by the public, as well as by those who assist in the process.

1.2 The Basis for Provincial Water Quality Guidelines

It is recognized that, for many substances of concern, there are not sufficient toxicological or other data to prepare a PWQO. In an effort to achieve a consistent approach in establishing environmental protection values for substances with limited information, a process for setting Provincial Water Quality Guidelines (PWQG) was established.

The PWQG setting process was developed to parallel, as closely as possible, the Objective setting process by employing toxicological data as well as considering bioaccumulation, mutagenicity and other factors, yet permitting the development of substance-specific environmental protection values with even a small amount of data. The PWQG process sets a margin of uncertainty whose size is dependent on the quantity and quality of available data. The process encourages the publication of new aquatic toxicity data to further refine the PWQG or allow it to become a PWQO.

1.3 Application of PWQOs and PWQGs

This document is not intended to describe in detail the application of Objectives and Guidelines in Ontario's water quality management programs. Objectives and Guidelines are designed for application to all surface waters (rivers, streams, lakes, etc.) in Ontario regardless of their existing use or condition. With specific respect to the use of PWQOs and PWQGs, the Ministry's surface water quality management goal is supported by a series of policies and application procedures. Key policies state:

"In areas which have water quality better than the Provincial Water Quality Objective [or Guideline], water quality shall be maintained at or above the Objectives [or Guidelines]

and

"Water quality which presently does not meet the Provincial Water Quality Objectives [or Guidelines] shall not be degraded further and all practical measures shall be taken to upgrade the water quality to the Objectives [or Guidelines]."

PWQOs and PWQGs are, therefore, used in combination with water quality monitoring data to assess ambient conditions in Ontario and to identify areas with existing or potential water quality problems.

The PWQO development process follows the policy of the Ministry to treat all surface waters in the province alike, except for distinguishing between waters that are of acceptable quality and those that are degraded. It is essential therefore to set Objectives that are stringent enough to protect the most sensitive environments.

Although Objectives and Guidelines are not legal standards *per se*, they are employed in various water quality models to establish acceptable wastewater loading limits on a site-specific basis. These loading limits, contained in Control Orders or other similar instruments, are legally enforceable.

The Ministry's water management policy lists several substances with "zero tolerance limits" which "... should be completely eliminated [from discharges]". The Municipal and Industrial Strategy for Abatement (MISA) is based on the goal of the virtual elimination of persistent toxic substances from point source discharges. Both of these policies relate to effluent control and in regulatory instruments they may supersede waste loading limits derived from a PWQO or PWQG. Objectives and Guidelines are needed, however, to assess the aquatic environmental significance of the materials that may be found in waterbodies as the result of spills and uncontrolled sources, and to protect local water quality in situations where technology based requirements may not be stringent enough to protect aquatic life.

PWQOs are set for individual chemicals or homologous groups and, for the most part, do not account for joint toxicity (i.e. additive, synergistic or antagonistic effects) resulting when a mixture of hazardous substances is present in a waterbody. In application, models addressing combined toxic effects should be employed wherever possible. At the very least, water quality managers should consider that waters with a quality at or near the Objectives for several parameters may not afford adequate protection to aquatic life. In these situations, the empirical approach of monitoring the toxicity of whole effluent or receiving water samples may be useful in assessing the joint toxicity of complex mixtures.

1.4 Other Water Quality Objectives and Guidelines

PWQO's and PWQG's are designed primarily to meet the Ministry's goal of protecting aquatic life and recreational uses. There are other uses and criteria that should be considered. The Canadian Water Quality Guidelines (CCREM, 1987) for agricultural uses (irrigation and livestock watering) and industrial water supply provide protection for these uses and should be considered where applicable.

PWQOs are not Drinking Water Objectives and thus do not apply to drinking water. The Ministry has set Drinking Water Objectives which apply specifically to finished drinking water and not raw water. These Objectives are developed independently of the PWQOs and are listed with supporting documentation in the MOE publication "Ontario Drinking Water Objectives" (MOE, 1983).

1.5 Water Quality Objective Setting Administrative Structure

The setting of PWQOs is primarily the responsibility of the Water Resources Branch of the Ontario Ministry of the Environment. Currently, the Aquatic Criteria Development Committee is responsible for all activities related to the development of Provincial Water Quality Objectives and Guidelines. This committee is comprised of staff of the Water Resources Branch, Hazardous Contaminants Coordination Branch, Laboratory Services Branch, Investigations and Enforcement Branch, the MOE Regions and Environment Canada.

1.6 What are Objectives and Guidelines? - A Definition of Terms

Throughout this report, references are made to *Objectives* and *Guidelines*. There are many government agencies using these terms throughout the world and they each have their own definitions. For water quality management in Ontario, the terms are defined as:

1.6.1 Objective

A Provincial Water Quality Objective (PWQO) is a numerical and/or narrative limit recommended to protect all forms of aquatic life and all aspects of the aquatic life cycles during indefinite exposure and to protect recreational water uses. Objectives are established when a defined minimum information base is available. They are used to assess ambient surface water quality, identify areas with degraded conditions, assess impacts of wastewater discharges

and spills, and provide a basis for establishing industrial and municipal wastewater discharge limits.

1.6.2 Guideline

For many substances there are some scientifically sound aquatic toxicological data available with which to assess potential aquatic environmental impacts, but there is not enough information to meet the minimum requirements for setting a PWQO. In these cases, the available data are assessed and, using a prescribed method of applying uncertainty factors determined by the quality and quantity of data, a Guideline value is calculated. This PWQG represents the best available value and is employed the same way as a PWQO with the understanding that it is more likely to change as additional information becomes available. Guidelines are recommended with the intention of upgrading them to PWQO status as sufficient information becomes available.

1.6.3 Interim Guideline

An Interim Guideline is a numerical water quality limit set using the Provincial Water Quality Guideline procedures, normally, to meet an emergency need for the Ministry of the Environment. It is based on a search of the best information at hand. An Interim Guideline is set by qualified Ministry staff and then checked and approved by other senior Ministry scientists. Unlike the PWQO or PWQG, an Interim Guideline is not subject to a formal peer review or further Ministry approval and publication.

1.7 Priority of Candidate Substances for PWQOs

The availability of toxicological data is limited for many of the substances now being detected. With the rapidly expanding capabilities of analytical chemists to measure contaminants in the environment, those responsible for the setting of water quality Objectives face a major challenge to keep pace. In addition, there is an increasing need to develop PWQOs for the constituents of complex effluents.

In order to make the most efficient use of resources and scientific data, most agencies develop priority substance lists. Ontario has lists in the areas of air, waste and water resources management. Resource limitations may also preclude the possibility of setting environmental protection values for all substances in the short term. Consequently, the absence of a published PWQO or PWQG for a substance does not necessarily mean that it is harmless or that future action will not be taken.

The development of a water quality Objective takes a wide variety of effects into consideration. In order to achieve consistency in the process and to efficiently employ available resources, a comprehensive and systematic approach has been developed.

The Ministry developed a screening process to identify a list of contaminants of greatest concern for Ontario. This list, the Effluent Monitoring Priority Pollutants List (EMPPPL), is supported by a published document (MOE, 1987). The screening process consisted of a chemical identification stage and a preliminary hazard assessment stage. A

number of factors were considered for each chemical. These include: potential presence in effluent, persistence, potential to bioaccumulate, and acute and chronic toxicity to organisms, including humans. The substances on the list have been derived from an ongoing assessment process which so far has considered in excess of 1500 chemicals. Substances currently not on the EMPPPL but identified through complete characterization of effluent samples will be candidates for inclusion on the EMPPPL. Chemicals may also be added to reflect new information on environmental fate or toxicity.

This list is used to establish priorities for Objective setting. Data on environmental persistence, bioaccumulation, fate and aquatic toxicity data are assessed and the substances are categorized in terms of their potential impact on the aquatic communities and uses such as fish consumption. Objectives are being set for EMPPPL substances with the highest potential for impairment and the intention is to establish Objectives or Guidelines for all of the EMPPPL substances as rapidly as resources and the availability of data permit.

Requests for Objectives are received from MOE Regional and Branch staff. Such requests are ranked in order of priority and assessed as soon as possible (see Section 1.8.1).

1.8 Summary of PWQO and PWQG Setting

The following briefly describes the key components of the process and provides reference to detailed discussions in subsequent chapters of this document (Figure 1).

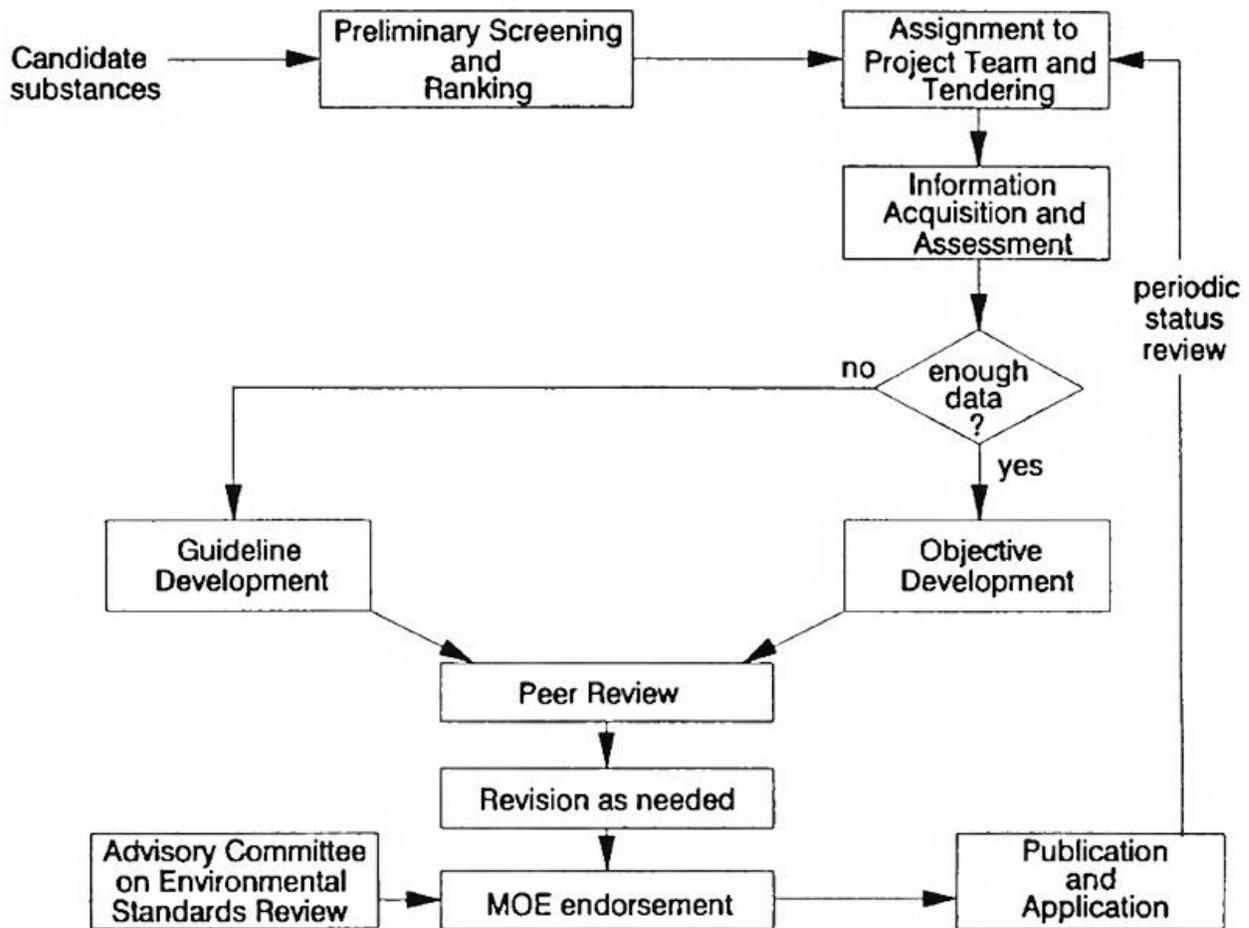


Figure 1. The process for developing PWQOs and PWQGs.

1.8.1 Selection of Candidate Substances

Requests for new Objectives originate in various pollution control activities of the Ministry of the Environment and from other Ministries. Candidate substances are screened for existing or potential hazard and occurrence in the aquatic environment. They are then ranked for immediate or future action.

To satisfy the immediate need for an Objective, an Interim Guideline may be set. Based on the results of the Interim Guideline setting, Interim Guidelines may

be recommended for PWQO or PWQG development.

1.8.2 Assignment

An Objective development document is prepared by a Ministry project team, normally with the assistance of a consultant.

1.8.3 Information Acquisition and Assessment

The world literature is searched for information that will be needed for setting a PWQO. The reviewer is seeking information on aquatic toxicity, bioaccumulation, taste

and odour in water or tainting of fish tissues, mutagenicity, fish consumption limits, impacts on wildlife, environmental fate, sources, objectives of other agencies, and, any other appropriate information found in the literature.

The information so obtained is then assessed. Rules for the adequacy and admissibility of data on aquatic toxicity, bioaccumulation, mutagenicity and tainting are described later in this document. All candidate information is screened for acceptability. Refereed publications are preferred. The fact that information is published is not sufficient, however, to justify its use in PWQO development.

1.8.4 Setting a Provincial Water Quality Objective

Provincial Water Quality Objectives are set to protect aquatic life and recreational water uses. The assessment of aquatic toxicity, bioaccumulation and mutagenicity data is fundamental to the process but other factors such as taste and odour in water, tainting of fish tissues, wildlife protection, recreation and aesthetics are also considered. Further details are provided in Chapter 2 of this document. Information on the aquatic environmental fate is included for reference.

The final draft report of the Objective Development Document and the critical papers used to derive the PWQO are reviewed by staff of the Ministry of the Environment and forwarded to recognized experts in Canada and other countries. The comments of the reviewers are assessed and appropriate revisions or additions are made to the draft report. The final report is submitted for approval and

publication by the Ministry, and this may include a public review coordinated by the MOE's Advisory Committee on Environmental Standards.

1.8.5 Setting a Provincial Water Quality Guideline

Once all of the information is obtained and assessed, a decision is made as to whether a PWQO can be set. If sufficient toxicological information (which meets MOE rules for admissibility) is available then a PWQO can be set.

When some aquatic toxicological data are available, but the minimum data set requirements for a PWQO cannot be met, a PWQG can usually be calculated. As in the Objective setting process, the Guideline process begins by selecting a value based on the lowest toxic effect concentration for an appropriate aquatic species and dividing that value by an uncertainty factor. The magnitude of the uncertainty factor is determined by the quantity and quality of available toxicity data and the physical-chemical properties of the substance. The Guideline setting process has a screening mechanism to accommodate concerns about bioaccumulation. Like the PWQO setting process, the Guideline setting process also considers mutagenicity. A numeric Guideline based on mutagenicity will only be set for substances with demonstrated mutagenic properties and where data for aquatic organisms are available.

The development of a Guideline can also consider factors such as taste and odour in water and fish tissue tainting. Further details are provided in Chapter 3 of this document.

The final draft of the Guideline Development Document is reviewed by staff of the Ministry of the Environment and forwarded to recognized experts in Canada and other countries. The Guideline is then submitted

for approval and publication by the Ministry, and this may include a public review coordinated by the MOE's Advisory Committee on Environmental Standards.

2 SETTING PROVINCIAL WATER QUALITY OBJECTIVES

2.1 Substance Physical-Chemical Properties

The fate and effects of organic contaminants in aquatic systems are largely determined by their inherent physical-chemical properties. Hence a knowledge of the fundamental substance properties (e.g. octanol-water partition coefficient) is essential in any assessment of potential environmental effects (Kobayashi, 1981; Klein *et al.*, 1988; Kollig, 1988; Kollig and Kitchens, 1990; U.S.E.P.A., 1991a;). Where this information is available, it is included in the PWQO development document.

2.2 Acute and Chronic Toxicity

PWQOs are primarily developed using the available knowledge of aquatic toxicity of the substances to which they refer. Many people relate aquatic toxicity with the survival of fish but the development of PWQOs also considers other impacts on aquatic life. A substance is generally considered toxic if it reduces longevity, fecundity, growth, reproductive success, produces atypical behaviour, physiological effects, histological effects or deformities, including tumours. Ontario's Water Quality Objectives are intended to provide limits for substances in fresh water environments. If these limits are achieved, observable deleterious impacts are unlikely. Since the PWQOs are intended to protect Ontario's fresh water environments, data relating to fresh water aquatic life resident in the province must be included.

2.2.1 Acute Versus Chronic Tests

Since it makes little sense to protect aquatic

life only from short term exposures to toxic substances, PWQOs are primarily based on the results of chronic test exposures. Decisions regarding classification of exposures as acute or chronic are complex and are assessed on a case-by-case basis.

An acute effect comes rapidly to an endpoint while a chronic effect is long lasting or continued. Acute effects are not synonymous with lethal effects. The words acute and chronic in connection with toxicity tests refer to short (acute) and long (chronic) exposure. Such terms require a specific organism as a reference. For a *Daphnia* with a life cycle of weeks, acute usually means two days or less. However, for trout with a life cycle of years, acute usually means four days or less.

When classifying a test results for a substance as either acute or chronic based on the duration of exposure, the nature of the substance and how rapidly it acts must be considered. Lethality caused by chronic exposure is a valid observation for PWQO development. For some substances, an acute endpoint may be used to meet the minimum data requirements for Objective development if it is known that the toxicant exerts all of its effects before the end of the exposure period. For example, if lethality due to chronic exposures can easily be predicted from a toxicity curve expressing LC50 values as a function of duration of exposure, then these data can and should be used to develop PWQOs. Justification for the use of acute data in meeting minimum data set requirements must be clearly documented.

Also important in Objective development,

are sublethal effects which occur at lower concentrations. Some judgement must be made as to whether these effects are harmful to the organism.

2.2.2 Primary and Secondary Data

All candidate toxicological information is screened for acceptability. All information that meets the following requirements is considered primary data:

- Toxicity tests must employ accepted laboratory practices of exposure and environmental controls. While all tests must be evaluated on a case by case basis, those tests following published protocols of government agencies or standard setting associations are generally acceptable.
- Any tests may be acceptable, including static tests if it can be shown that concentrations of the toxicant are not changing (significantly) throughout the test and adequate environmental conditions for the test species are maintained with respect to such factors as dissolved oxygen and removal of metabolic wastes. Generally, continuous flow exposures, and renewal tests (i.e. static tests with replacement) are acceptable if appropriate rates of renewal of toxicant are maintained. Static tests are acceptable if concentrations of the toxicant are measured in the exposure vessel at the beginning and end of the test and no more than 10% of the toxicant is lost during the test. The use of chemical carriers is acceptable as long as the concentration of the toxicant does not exceed water solubility in the absence of the carrier. Appropriate chemical carrier controls must also be included.

- Dissolved concentrations of toxicant in the exposure vessels must be constant and verified by measurements rather than calculated or measured only in stock solutions. Tests will generally be considered unacceptable if more than 10% of the toxicant is lost during the test.
- Test end points and lengths of exposure must be appropriate to the life stage of the species tested and the characteristics of the substance. Although the definitive bench mark for chronic toxicity is a whole life cycle test, partial life cycle and short term or early life stage tests are acceptable as chronic data (McKim, 1977). An example of this type of test is the fathead minnow, 7 day growth test (Norberg and Mount, 1985).
- Relevant environmental parameters such as temperature, pH and hardness must have been recorded.
- Responses and survival of controls must be appropriate for the species and test used.

Data on vertebrates and invertebrates not meeting all of the above are denoted as secondary in objective development documents. Secondary data are inadmissible in the derivation of an Objective, but are listed and discussed in the PWQO development documents. Most tests using aquatic plants will also be classified as secondary due to the frequent use of artificial media or a lack of standardized protocols; however, plant data may be used as the critical endpoint for Objective development subject to best scientific judgement.

2.2.3 Minimum Data Requirements

The goal in setting PWQOs is to protect all forms of aquatic life. It is essential, therefore, to ensure that the Objectives are based on toxicity tests representative of a wide variety of life forms. The data must include vertebrates, invertebrates and plants and must consider the most sensitive lifestages.

PWQOs must be derived from literature conforming to the following rules for minimum data:

FISH

- Toxicological data from at least three different species including at least one cold-water fish species (e.g. Rainbow trout) and one warm-water fish species (e.g. Fathead minnow) must be present.
- At least one fish species must be resident in Ontario.
- Marine or brackish water species must not be used.
- The data must include two different chronic whole organism responses (e.g. mortality, growth, reproduction), and at least one of these must involve an early life stage.
- Acute lethal responses are generally not used to meet minimum data requirements for an Objective except where a convincing case can be made that they are indicative of chronic lethality (See Section 2.2.1).

INVERTEBRATES

- At least two different orders of invertebrates must be represented, one of

which must be from the class Crustacea.

- Marine or brackish water species must not be used.
- Data from no more than one tropical invertebrate species may be used.
- Data must include two different responses and at least one of these must involve an early life stage.
- Acute lethality data are generally not used except when a convincing case can be made that acute lethal responses are indicative of chronic lethality (See Section 2.2.1).

ALGAE/AQUATIC PLANTS

- At least one algae or aquatic plant species must be represented.
- Algae/plants must be freshwater species resident in temperate North America.

When assessing the aquatic impacts of phytotoxic substances (e.g. herbicides), special data requirements may be considered for algae and aquatic plants.

Additional minimum data requirements are also found in Sections 2.3 and 2.4 dealing with bioaccumulation and mutagenicity, respectively.

For any given substance, if the toxicological data available do not meet the minimum requirements, an Objective cannot be set and the process reverts to Guideline setting as described in Chapter 3.

2.2.4 Modifying Factors

Frequently the available toxicological literature indicates that factors such as water

hardness, pH or temperature clearly modify the effects of a substance. Where the data for such a relationship are sufficiently clear, Objectives will be developed as a function of these modifying parameters. In these cases, the PWQO is expressed as a table of numbers and/or an equation defining the Objective at various levels of modifying parameters (i.e. ranges expected to occur in Ontario).

When modifying factors must be considered, it becomes more complicated to define minimum data requirements. Generally, the available data should support the Objective in the ranges of the modifying parameters that are included in the table.

2.2.5 Safety Factors

A preliminary Objective is calculated based on the aquatic toxicity data. As long as the minimum data required to establish an Objective exist for a given substance, the lowest water concentration (including primary chronic and acute endpoints) causing significant responses is divided by a safety factor to obtain a preliminary Objective value.

The "safe" concentration may be different to protect against growth effects than it would be to protect against other toxic effects. Therefore, since all types of undesirable effects need to be prevented, the safety factor should account for relationships among concentrations causing various types of responses that could be observed for a substance. The minimum data requirements for establishing a PWQO cover this concern in part since there is a requirement to include at least one vertebrate and one invertebrate early life stage test. Early life stages are known to generally exhibit the most sensitive endpoints. Even with this consideration,

however, there needs to be some safety factor to account for variation due to endpoints and for those cases where the early life stage is not the most sensitive.

The safety factor should also protect sensitive species that are not feasible to use as bioassay test organisms. Bioassay test species are not necessarily the most sensitive in aquatic environments. They are chosen partly because they are robust enough or are well enough known to be cultured in the laboratory. Inter-species differences in sensitivity to a toxicant are generally greater than are intra-species differences.

A margin of safety is also needed to account for the fact that laboratory animals are generally in better condition than are those in the field since they are fed daily under optimum conditions. Field animals are often under the stresses of winter fasting, parasites, predators etc. It is true that there is greater genetic diversity in the field allowing survival of the population in changing environments. However, since the Objectives are intended to protect individuals as well as populations, a safety factor is needed to account for the relative sensitivities of field versus laboratory-reared individual animals.

Safety factors are not intended to account for joint toxicity (i.e. additive or synergistic or antagonistic effects) resulting when a mixture of hazardous substances is present in a waterbody (See Section 1.3). These phenomena are still poorly understood and adjustment of the safety factor to account for these effects is not possible at this time.

Few comparative studies exist that would assist in estimating these differences in sensitivity to a substance due to differences in species, endpoints etc. (e.g. Benoit *et al.*, 1982; Mayer *et al.*, 1986). The existing water

quality criteria development documents are additional sources of comparative information. **A safety factor of 10 applied to the lowest effect concentration reported in a primary study is used for setting the preliminary Objective value based on toxicity.**

2.2.6 Summary

These initial steps in developing an Objective, based on acute and chronic toxicity, are summarized as follows:

A review of the world literature on acute and chronic toxicity is prepared for the substance.

The available information is critically assessed to determine whether it is acute or chronic, and primary or secondary data.

If an adequate amount of primary information exists, the preliminary Objective is recommended at the lowest observed effect concentration, reported in a primary study, divided by 10.

2.3 Bioaccumulation

Potential effects that may occur at the upper trophic levels in the food chain, from the consumption of contaminated biota, are not considered in the initial assessment involving toxicity. A separate assessment of bioaccumulation in the PWQO development process is therefore required. Regardless of whether bioaccumulation is the determining factor in setting the final PWQO value for a substance, a minimum amount of information is required for the assessment (Figure 2).

If both a reliable laboratory determination of

residues in fish exposed to a known water concentration [i.e. a bioconcentration factor (BCF)] and information on the potential health effects in predators and/or human consumers of contaminated biota (generally these values are fish consumption limits endorsed by a recognized environmental or human health protection agency) are available, then a preliminary Objective based on bioaccumulation can be determined. This calculation is described in Sections 2.3.1 through 2.3.4.

If information on consumption limits is not available, a preliminary Objective based on bioaccumulation cannot be set.

However, if reliable information suggests that BCF is less than 1000, bioaccumulation is not considered significant and the calculation of a preliminary Objective based on bioaccumulation is not required.

If BCF is greater than or equal to 1000, the whole assessment defaults to the Guideline setting process described in Chapter 3.

If no information on BCF is available, an octanol-water partition coefficient (K_{ow}) measured by a recognized method or, if reliable, a calculated K , will be considered. For nonpolar organic compounds, the low solubility in water is the primary physical-chemical property influencing the bioconcentration potential of a compound. The octanol-water partition coefficient is the most widely used quantitative measure of relative chemical solubility properties. Octanol is also considered a surrogate for naturally-occurring lipids of aquatic organisms - lipids in which these hydrophobic chemicals concentrate (see Section 3.1 and Appendix A for a more detailed discussion). It is assumed that substances with an octanol-water partition

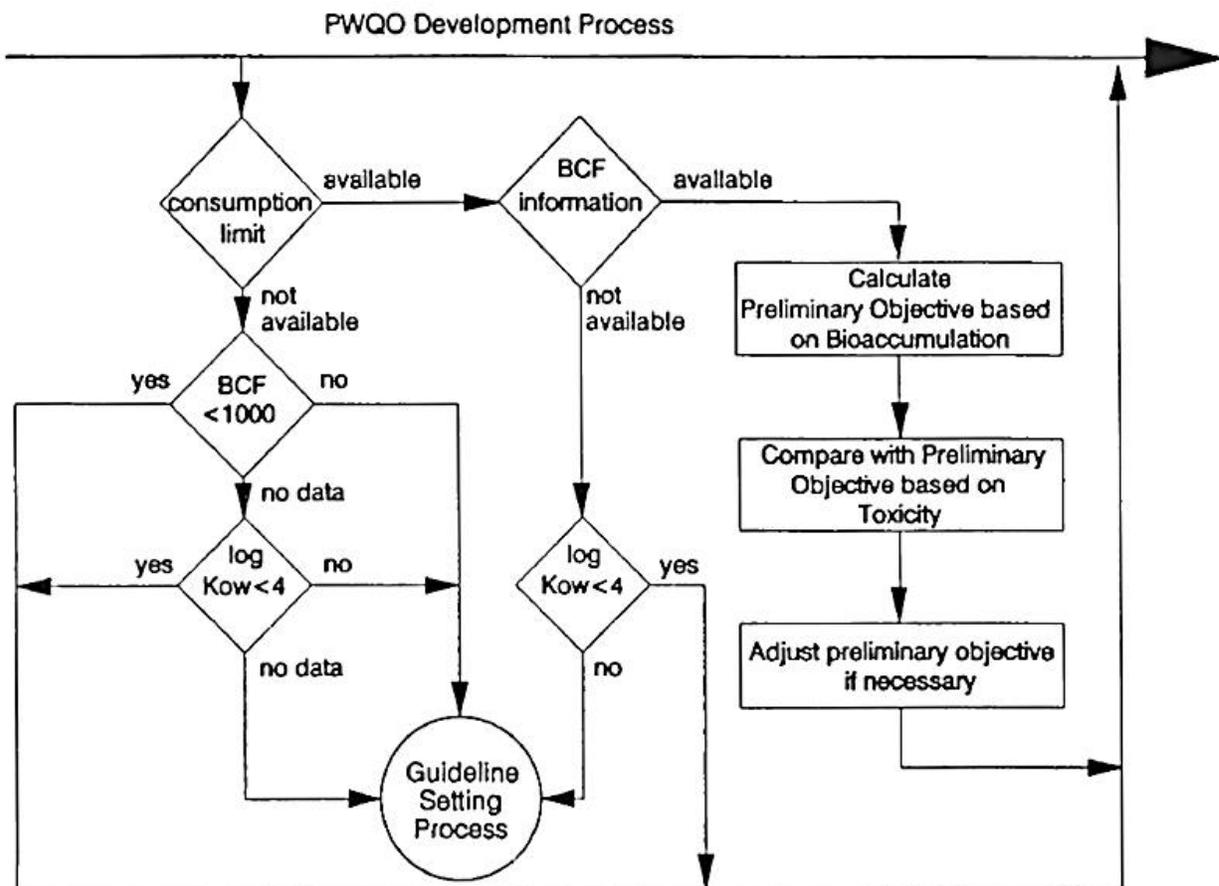


Figure 2. Bioaccumulation assessment for the PWQO development process.

coefficient ($\log K_{ow}$) of less than four do not bioaccumulate significantly and the calculation of a preliminary Objective based on bioaccumulation is not required.

If the $\log KOW$ is greater than or equal to four, or there are no data on K_{ow} , the whole assessment defaults to the Guideline setting process described in Chapter 3.

For metals, special considerations are needed on a case-by-case basis. Mechanisms of uptake and storage of metals in aquatic life are diverse and strongly influenced by chemical speciation and interactions with complexing ligands.

2.3.1 Consumption Limits

The BCF must be based on a controlled laboratory exposure of fish in which the water concentration was measured, the substance was bioavailable and present at less than known toxic levels. The BCF must be determined on a wet weight basis at steady-state equilibrium. Since lipid is the critical storage compartment for highly bioaccumulated nonpolar organics, the BCF must also be normalized to a lipid content of 10% as follows:

$$\text{BCF} = \frac{C_t}{C_w} \times \frac{10}{\% \text{ lipid}}$$

where C_t is the concentration in whole fish and C_w is the water concentration.

The relatively high lipid concentration of 10% was chosen to be conservative. If more than one reliable BCF value is available, the highest value is used.

Fish consumption limits designed to protect human health or wildlife from the effects of eating contaminated fish are employed to define acceptable fish concentrations.

Fish consumption limits are developed by Health and Welfare Canada or similar health protection agencies (e.g. U.S. Food and Drug Administration, World Health Organization). If more than one consumption limit is available, the value endorsed by Health and Welfare Canada is given preference and is used to calculate the concentration for determining the preliminary Objective based on bioaccumulation.

2.3.2 Fish Equivalency Concentration

Unfortunately, fish consumption limits are not available for many substances. The following may assist in developing surrogate limits from related information. Caution must be exercised, particularly if the bioaccumulation calculation determines the final PWQO.

If an allowable daily intake (ADI) value is available, a fish equivalency concentration (FEC) can be calculated. Fish equivalency concentrations are not consumption limits. They are used in this process as a gross surrogate to address bioaccumulation concerns. Fish equivalency concentrations

for organic chemicals are calculated to protect a 70 kg human consumer of sport fish. This is done by calculating a total allowable daily intake for the chemical in question. This allowable daily intake is converted to a chemical concentration in the edible portion (muscle fillet) of fish by allotting 50% of the ADI to fish and dividing this value by 25 grams, the average daily fish consumption by an angler in Ontario, (Cox *et al.*, 1989). For example, with an ADI value for a chemical pollutant of concern of 1 $\mu\text{g}/\text{kg}$ body weight per day, a fish equivalency concentration of 1.4 $\mu\text{g}/\text{g}$ is derived by substituting the appropriate values into the following equation:

$$\text{FEC} = \frac{(\text{ADI} \times 70 \text{ kg} \times 0.5)}{25 \text{ g fish/day}}$$

where FEC is the fish equivalency concentration.

2.3.3 Determining the Acceptable Water Concentration

There is currently a lack of information on which to base ambient water concentrations to protect wildlife from the deleterious effects caused from consuming contaminated fish. The Aquatic Criteria Development Committee is currently developing procedures to establish fish contaminant residue limits for the protection of predacious fish and wildlife. Until these limits are developed, the contaminant residue limits of the International Joint Commission, the non-carcinogenic based limits of the New York State Department of Environmental Conservation (Newell *et al.*, 1987), or other agencies, as deemed appropriate by the Aquatic Criteria Development Committee of the MOE, may be directly substituted into the equation as the whole fish concentration to calculate an acceptable water concentration.

It has been observed that the concentrations of organic chemicals in whole fish are approximately 2.5 times higher than those in the edible portions (Niimi and Oliver, 1983). If bioaccumulation factors represent the ratio between the concentration of a chemical contaminant in whole fish and the concentration in the exposure water, consumption limits must be expressed on a whole fish basis. Therefore, the fish consumption limit or the FEC is multiplied by 2.5 to convert this value to an acceptable concentration in whole fish. A water concentration designed to limit bioaccumulation in fish to meet the fish consumption limits is calculated by dividing the whole fish concentration by the lipid-normalized bioconcentration factor (BCF).

$$C_w = \frac{\text{whole fish concentration}}{\text{lipid normalized BCF}}$$

where C_w is the water concentration.

2.3.4 Calculating the Bioaccumulation-Based Preliminary Objective

Maximum allowable residues in fish would theoretically result if the calculated ambient water concentrations to protect against effects caused by the accumulation of chemicals in aquatic biota were permitted. **Since it is not desirable to reach a maximum acceptable residue concentration of a substance in fish on a consistent basis, a safety factor of 10 is applied to the calculated water concentration.**

The final water concentration based on the BCF is compared to the preliminary Objective based on toxicity and if it is more stringent, the value based on bioac-

cumulation becomes the preliminary Objective.

In some parts of Ontario, subsistence fishing is an important use of lakes and rivers. Available ADI values may not offer adequate protection to year-round consumers of large quantities of fish. In applying PWQOs to waterbodies where subsistence fishing occurs, additional safety factors, determined on a site-specific basis, may be warranted.

2.3.5 Summary

In summary, the steps for setting a preliminary Objective based on bioaccumulation are as follows:

- Information is sought on laboratory derived BCF and consumption limits designed to protect consumers of fish. If these data are available, a water concentration can be determined.
- A preliminary Objective is set by dividing the lowest consumption limit by the highest BCF value found in the literature and then dividing that value by a safety factor of ten (10).
- This preliminary Objective value is compared to the preliminary Objective calculated using the aquatic toxicological information, the more stringent of these two values is carried forward to the next step in PWQO setting.
- If consumption limits are not available or cannot be derived, a preliminary Objective based on bioaccumulation cannot be set. The next step, then, is to assess the available BCF data or, in its absence, the octanol-water partition coefficient ($\log K_{ow}$).
- If the BCF is less than 1000 or, in the absence of BCF data, the $\log K_{ow}$ is less

than 4.0, bioaccumulation is not considered significant and the calculation of an Objective based on bioaccumulation is not required.

- If the BCF is greater than or equal to 1000 or, in the absence of BCF data, the $\log K_{ow}$, is greater than or equal to 4.0, bioaccumulation is considered a significant factor to be addressed and the whole assessment then defaults to the PWQG setting process described in Chapter 3.
- If, by lack of data, BCF and the $\log Kow$ data are not known, the substance could, potentially, be of concern for bioaccumulation. Therefore, a PWQO can not be calculated. The whole assessment then defaults to the PWQG setting process described in Chapter 3.

2.4 Mutagenicity

The induction of mutagenic events and mutation related diseases in organisms as a result of their exposure to pollutants is a serious and undesirable effect. Genotoxic damage, mutations and mutation related diseases, which are measurable in microbial, plant, insect and mammalian systems, can also be measured in aquatic organisms. The potential effects of such exposure are mutagenicity related diseases, such as heritable mutations, birth defects or cancer. If sufficient information is available, the preliminary Objective for mutagenicity will be set to protect aquatic organisms from the effects of mutagenic or genotoxic pollutants. The preliminary Objective for mutagenicity will be compared to the preliminary Objective based on toxicity or bioaccumulation. The most stringent of these values will be adopted for consideration as the final PWQO.

A mutagenic pollutant is defined as a chemical which possesses the ability to induce permanent and heritable genetic change at the cellular or whole organism level. A genotoxic pollutant is defined as a chemical with the ability to induce damage to DNA, to trigger DNA repair systems, to form DNA adducts, to induce damage and/or sister chromatid exchanges in chromosomes, or to induce mutagenic events.

The data used to set these Objectives will be derived from experiments using aquatic species exposed under appropriate conditions. These mutagenicity Objectives are not designed to protect the consumers of these aquatic species, for example humans, terrestrial mammals or fish-eating birds.

Such concerns are addressed in other areas of the PWQO development document dealing with bioaccumulation and the consumption of fish (Section 3.2).

Setting Objectives for mutagenicity assumes that a no effect concentration for mutagens can be identified (Schaeffer, 1981). Justification for such a threshold model arises from the fact that for most organisms mutations occur naturally at a given background rate, as the result of errors in DNA synthesis and/or repair. When organisms are exposed to a mutagenic pollutant, the lowest observable effect concentration (LOEC) would be that concentration inducing an increase in mutagenic events to a frequency which is significantly greater than the background mutation rate. In setting mutagenicity Objectives, a practical goal is to reduce the mutagenic risk for aquatic organisms to this background mutation rate and not to some theoretical and perhaps unachievable lower level.

Consistent with the PWQO setting process for aquatic toxicity (Section 2.2), the LOEC and not the no observed effect concentration (NOEC) will be used to set the preliminary Objective based on mutagenicity concerns.

Objective setting for mutagenic chemicals will be based on data from acceptable tests employing aquatic organisms as test species. The lowest concentration of a mutagenic compound which induces mutagenic events exceeding the background rate, in the most sensitive aquatic species, will be used as the critical value. This value will be adjusted by a safety factor to establish a mutagenicity Objective for that compound.

In the initial stage of Objective development, the mutagenic potential of the chemical is determined using all available mutagenicity data. Only chemicals which are identified as mutagenic or genotoxic will continue to the stage of setting preliminary Objectives or Guidelines for mutagenicity. If a substance is shown to be non-mutagenic, this component of the PWQO process will be terminated.

Mutagenicity Objectives will be set based on the data from a minimum of three primary mutagenicity studies conducted on whole aquatic organisms. The critical concentration, that is the lowest concentration inducing a mutagenic effect in the most sensitive aquatic species, adjusted by a safety factor, will be considered the mutagenicity Objective. If the mutagenicity data available for a chemical are insufficient to establish an Objective or, in the case where data are not available to determine the mutagenic potential of a chemical, then a Guideline for mutagenicity will be set. That process is described in Chapter 3.

2.4.1 Acceptable Data

All mutagenicity data from tests employing acceptable protocols may be used in determining the mutagenic potential of a chemical. Objectives for mutagenicity, however, will be set only on data derived from aquatic organisms. Since PWQO's are expressed in units of concentration in water (e.g. mg/L), the Objectives for mutagenic chemicals will be derived from experiments using suitable species exposed in aqueous solutions. This includes data derived from tests using aquatic vertebrates, such as fish and amphibians, aquatic invertebrates, such as crustaceans, as well as aquatic plants. Ideally data on the incidence of mutagenicity related diseases will be used, such as cancer or heritable mutations induced in aquatic organisms by an identified mutagenic compound. Data on the incidence of cellular mutations, DNA damage or chromosome damage in whole aquatic organisms are also acceptable. Data from tests using measured aquatic concentrations of the chemical would be most appropriate. Data from tests reporting only nominal chemical concentrations are also acceptable but these data are secondary. In addition, data derived from bacterial, yeast or algal tests, exposed in the absence of endogenous activation systems, are acceptable. Such microbial data are representative of *in vivo* (intact) organisms in the aquatic environment.

Data derived from non-aquatic organisms, such as, rodents (mammals), are unacceptable for setting Objectives and Guidelines. Similarly, results from tests using cell tissue-culture or microbial tests requiring endogenous activation systems (such as S-9) are not acceptable. These two data types are excluded because neither measures an effect in intact aquatic organisms.

Only data from tests which expose organisms for a period of time permitting adequate cell division (two cellular generations) or DNA replication are acceptable. The process of DNA replication profoundly influences the development and expression of mutagenic events.

Only tests using fresh water organisms will be acceptable for setting mutagenicity Objectives. Data from tests using marine organisms may be considered in Guidelines. Examples of acceptable tests and acceptable species are given in Appendix B.

2.4.2 Minimum Test Protocol Requirements

Only data from properly conducted tests will be used. Each test must contain adequate quality control checks including negative controls and appropriate positive controls. Each test must provide data at a minimum of five concentrations, preferably including one approaching the acutely lethal concentration. Tests must have sufficient replication of control and treatment groups to assure credible statistical analysis. Tests must be conducted over a sufficient dose range to permit definition of a concentration-effect relationship as well as demonstration of a no-effect and lowest-effect concentration. Data from tests which do not meet minimum data requirements will not be used in the setting of Objectives but may be considered in the development of Guidelines.

A chemical will be considered mutagenic if it induces a significant, (tested statistically, ANOVA or other such test), concentration-related increase in mutagenic events. In addition the substance is considered mutagenic if two or more concentrations induce a significantly greater

number of these events than the control. A chemical which does not induce a concentration-related increase in mutagenic events will be considered non-mutagenic.

2.4.3 Primary and Secondary Data

Tests on aquatic species which meet minimum test protocol requirements and which utilize constant, measured, and continuous (flow-through) concentrations of chemical in aqueous solution, will be considered primary. Static and renewal tests (i.e. static tests with replacement) will also be considered primary if it can be shown that chemical concentrations did not change appreciably (i.e. < 10%) over the duration of the exposure. The use of chemical carriers is acceptable as long as the concentration of the toxicant does not exceed its water solubility in the absence of the carrier. Appropriate solvent controls must also be included.

Ideally, primary data should also define the chemical's mode of action. Such data should identify the most prominent genotoxic endpoint, the organ or tissue most affected, the physiological activation and detoxification pathways as well as the formation of DNA adducts with the chemical directly or its metabolically activated products.

Tests which utilize any of the following will be considered secondary and unsuitable for the setting of Objectives:

- Tests not employing accepted laboratory practices of exposure and environmental controls. While all tests must be evaluated on a case by case basis, those tests following published protocols of government agencies or standard setting associations are generally acceptable.
- Tests using nominal (i.e. unmeasured)

chemical concentrations.

- Tests exposures with appreciably variable (decreasing) chemical concentrations (i.e. > 10%).

Only data classified as primary will be utilized to set Objectives. Guidelines may be set using data classified as secondary or primary.

2.4.4 Complicating Factors

If test conditions confound the outcome of the experiment, then the data should be considered secondary. Factors should be identified which may affect the concentration or integrity of the test chemical, such as the presence of light when testing a photosensitive chemical. Factors which may alter the sensitivity of the test organism, such as the use of a second chemical to induce detoxification or activation metabolic pathways, should be noted. In addition, use of a second chemical as a promoter should also be noted.

If however, mutagenicity is a function of environmental variability, then the Objective may be a function of that variable. Modifying parameters which may influence the expression of mutagenic events or which may influence the condition of the test organism could include temperature, dissolved oxygen, pH, or conductivity of the aqueous medium.

If specific information is available for effects such as those noted above, it is to be noted in the mutagenicity Objective development document.

2.4.5 Process for Setting Objectives Based on Mutagenicity

The process for setting Objectives based on mutagenicity consists of first determining the mutagenic potential of the substance and then calculating the Objective based on mutagenicity.

2.4.5.1 Determining Mutagenic Potential

A minimal set of data is necessary to set mutagenicity Objectives. The process for setting such Objectives is outlined in Figure 3.

At the initial assessment level, data from experiments utilizing non-aquatic as well as aquatic species are considered in order to screen for mutagenic properties. At this assessment level, relevant data would include the results of short-term bacterial tests, mammalian tissue-culture assays, mammalian whole organism tests, as well as tests with aquatic species.

If data from a minimum of two test systems, including tests for mutagenic as well as chromosomal damage endpoints, clearly demonstrate a chemical to be nonmutagenic, mutagenicity Objectives will not be set. Objectives for that chemical will be based on other types of toxicological data.

Data demonstrating a chemical's ability to induce a mutagenic or genotoxic response in at least two test systems, are necessary to consider a chemical mutagenic and to initiate the setting of a mutagenicity Objective.

If both positive and negative mutagenicity data are available, the chemical will be regarded as either mutagenic or non-mutagenic based on weight of evidence and

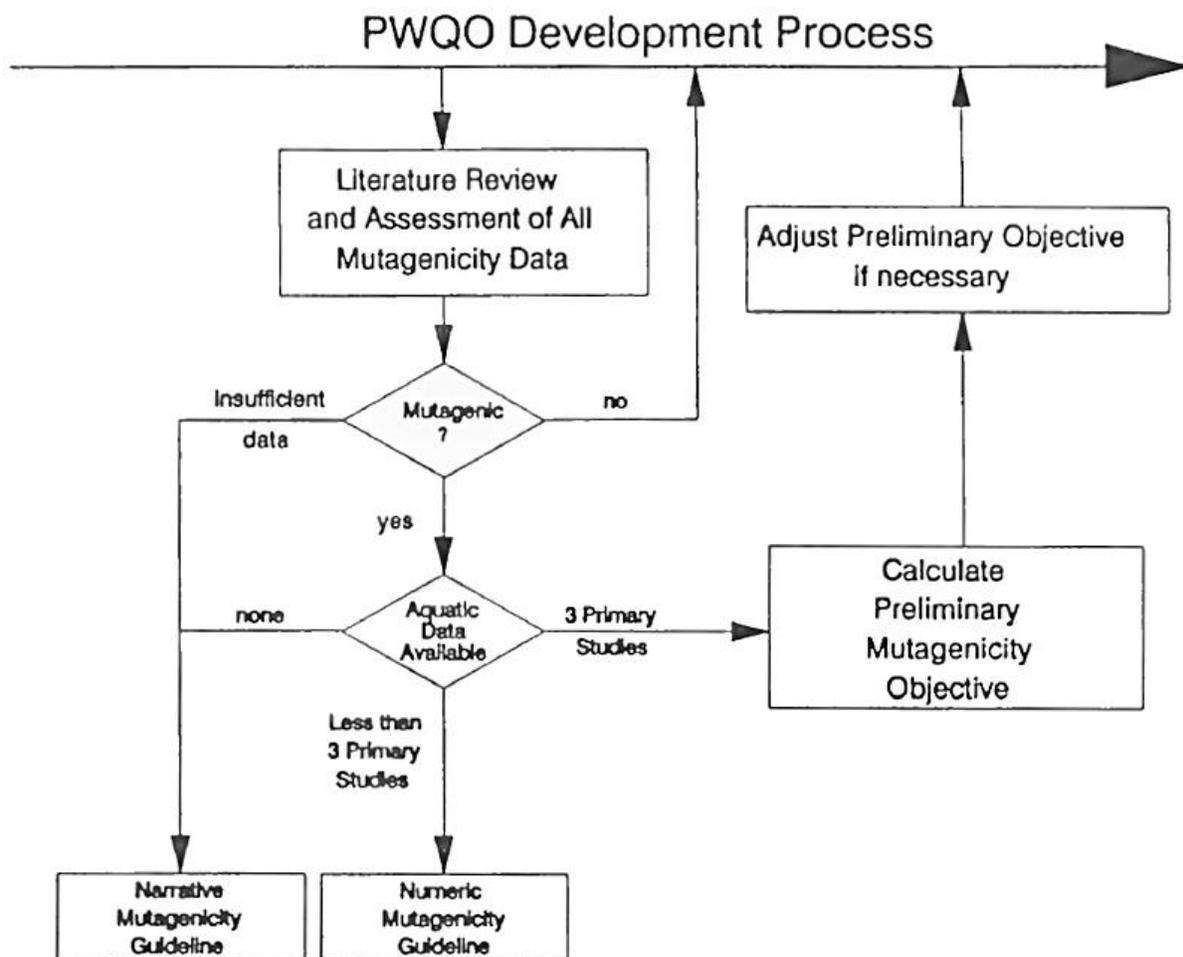


Figure 3. Mutagenicity Assessment for the PWQO Development Process.

through consultation with MOE authorities.

If mutagenicity data are not available, or if confirmatory data from two test systems are not available, the chemical would be directed into the Guideline process.

2.4.5.2 Setting Mutagenicity Objectives

To set a preliminary Objective based on mutagenicity, the chemical must be shown to be mutagenic or induce mutagenicity related diseases in aquatic organisms. With many chemicals, sufficient data may not be available to assess its mutagenicity in aquatic

species. If data from experiments using aquatic organisms are not available, the chemical will be directed into the Guideline setting process.

In order to set a preliminary Objective based on mutagenicity, data must be available for at least three primary studies demonstrating mutagenic events or mutagenicity related diseases in aquatic bacteria, plant, invertebrate or vertebrate species. Data need not be available for each level of organization (for example, three fish studies are acceptable for setting an Objective). However, the minimal data

set must include results from at least one vertebrate study and must not include results from more than one plant or bacterial study. For each primary study, the lowest concentration inducing a measurable effect (LOEC) exceeding background would then be used in setting the mutagenicity Objective. The LOEC for the most sensitive aquatic species will be the critical value. This value is reduced by a safety factor to set a mutagenicity Objective.

2.4.6 Safety Factors

A safety factor will be used in the setting of mutagenicity Objectives. The use of this factor corrects for the possibility that the most sensitive mutagenic or genotoxic endpoint may not be measured in the available data. For example, tests measuring chromosomal damage may not detect additional genetic damage (such as mutations) which may also be induced. In addition, the safety factor would add assurance in those instances where data from the most sensitive species are not available. Moreover, the sensitivity in detection of a mutagenic effect is dependent on the size of the population examined. The safety factor, accounts for the possibility that effects may be measured at lower concentrations of the chemical if larger numbers of organisms had been used in experiments. Finally a factor must be applied to account for variability in exposure-effect data experience within and between laboratories employing similar test procedures. To account for all uncertainties a factor of 10 is considered appropriate. **The lowest concentration of a mutagenic compound inducing a significant effect in the most sensitive aquatic species reported in a primary study, is divided by a safety factor of 10 to establish a preliminary Objective for mutagenicity.**

2.4.7 Selecting a Preliminary Objective Value

The preliminary Objective based on mutagenicity is compared with the preliminary Objective based on toxicity and bioaccumulation, as described in Sections 2.2 and 2.3. The most stringent concentration of all the preliminary Objectives is then selected as the candidate PWQO.

2.4.8 Summary

In summary, the steps for setting a preliminary Objective for mutagenicity for the protection of aquatic organisms are as follows:

- A thorough search of the available mutagenicity test data is conducted.
- Only data from acceptable tests, employing acceptable test protocols, will be used.
- The substance is initially assessed for its ability to induce mutagenic or genotoxic damage in tests employing both aquatic and non-aquatic species.
- If a minimum of two data sets, which include tests for both mutagenic and chromosomal damage endpoints clearly demonstrate (no positive data) the chemical is non-mutagenic, the mutagenicity Objective setting exercise is terminated.
- If data from two different test systems confirm the mutagenic potential of the chemical, data from aquatic organisms only will be assessed for the purpose of setting a mutagenicity Objective.
-
- If no data are available to assess its

mutagenic properties, or if data from a single test demonstrates a chemical mutagenic but a second data set demonstrating mutagenic or genotoxic properties is not available, the chemical is directed to the Guideline setting process.

- In some cases where data are conflicting, indicating both mutagenic and non-mutagenic results, the chemical will be considered either mutagenic or non-mutagenic based on weight of evidence and in consultation with MOE authorities.
- If data from acceptable tests employing aquatic organisms are not available, or if fewer than three primary aquatic organism data sets are available, the chemical is directed into the Guideline setting process.
- If a minimum of three primary aquatic organism data sets are available the LOEC (lowest concentration inducing a measurable mutagenic effect) for each data set is summarized. The LOEC for the most sensitive aquatic species is selected as the critical value.
- The critical LOEC value is divided by a safety factor of 10 to set the mutagenicity Objective.
- This preliminary Objective for mutagenicity is compared with the preliminary Objective based on toxicity and bioaccumulation. If the mutagenicity Objective is the most stringent concentration, then this value is carried forward to the next step in PWQO development.

2.5 Impacts on the Taste and Odour of Water and Tainting of Fish Tissues

To adequately protect the recreational uses of surface waters of the province, PWQOs must ensure the natural quality of these waters are not impaired by the imparting of taste and odour in water and/or fish by chemical pollutants.

Odour in water, taste in water, and tainting (taste or odour) of fish tissue which may affect its palatability are effects that can clearly influence the quality and thereby impact on the recreational use of surface water (i.e. fishing, swimming, bathing etc.).

The data required for Objective setting are normally obtained through a review of pertinent literature. These data should be derived from properly designed experiments with measured concentrations of, at least, the stock solutions of chemical (Amoore and Hautala, 1983). Taste and odour effects of substances in water should be obtained with water temperatures of 30°C or less.

Threshold concentrations eliciting an odour or taste in water or an off-flavour or odour in fish-tissue may be used to set a preliminary Objective. While the MOE is not requiring strict adherence to any specific protocol for the admissibility of such data, there are some fundamental conditions that must be met if the data are to be used (MOE 1981):

Fish Tainting

- Fish used in the test should be exposed to the substance for a minimum exposure time of 48 hours before being sacrificed and prepared for the taste panel.
- Tainting values should be expressed as

water concentrations that result in taste or odour in the fish tissue.

- The dilution series should elicit a range of responses from strong to no response.
- Panels should consist of at least six people, preferably non-smokers.
- Taste testing should be carried out in an odour-free room (preferably with filtered air).
- Precautions should be taken to ensure no loss of chemical during testing of volatile chemicals.

Taste and Odour of Water

- Dilution series should range from strong response to no response.
- A panel size of 20, preferably non-smokers, is desirable although a minimum panel size of six is acceptable.
- Dilutions of test compounds must be prepared using odour-free water (preferably obtained by passing water through an activated carbon filter).
- Taste and/or odour testing should be carried out in an odour free room (preferably with filtered air).
- Since warm water may taste unpleasant, the samples should be presented at room temperature for judgement of taste.
- When judging the odour of a chemical in water, the temperature of the solution should not exceed 30°C.

If an effect concentration is below or just above (i.e. within an order of magnitude) the

preliminary Objective, based on aquatic toxicity, bioaccumulation or mutagenicity, special attention should be given to identifying the most accurate value of effect concentration.

Because fish tainting and taste or odour of water are expressed as threshold water concentrations that can be perceived by people with good taste and odour acuity and because these effects are not health related, a safety factor of two (2) was selected to calculate a "no-effect" concentration or taste/odour protection value.

If the value calculated from the assessment of impacts on taste and odour in water and tainting of fish tissues is more stringent than the preliminary Objective value calculated from toxicity, bioaccumulation or mutagenicity, then this value is carried forward to the next step in the PWQO setting.

2.6 Recreation

Traditionally, the Objectives for the protection of recreational water uses have been based on public health concerns (e.g. microbiological contamination, eye irritation) and aesthetic considerations (e.g. colour, clarity). Procedures for the development of such Objectives tend to be issue specific.

In setting PWQOs for recreational use, the principal concern relates to the impact of chemical substances on humans during recreational uses of water through direct dermal exposure or accidental ingestion of water while swimming or during similar recreational activities. At this time, the Ministry is unable to make conclusive assessments on the risks to bathers

through such pathways but will continue to evaluate procedures for directly assessing recreational use requirements in PWQO setting activities.

Should information on recreational uses of water be found during the literature search, it will be documented and assessed.

2.7 Wildlife Protection

The protection of wildlife is also a consideration when setting a PWQO. Some wildlife occupy niches that can potentially expose them to elevated levels of contaminants in the aquatic environment. Those that rely on aquatic environments and particularly those birds (e.g. osprey, gulls, etc.) and mammals (e.g. mink) that consume large quantities of fish and other aquatic organisms are at greatest risk.

The health significance of residue concentrations within aquatic organisms is currently under study by MOE. Substance-specific guidelines will be set for permissible residue concentrations. Procedures will then be established for relating these guidelines to PWQO values.

Where appropriate data on BCF and acceptable intake levels are available, the protection of fish-eating birds or mammals is addressed in the preliminary bioaccumulation Objective (Chapter 2.3).

No specific procedures are yet in place for evaluating proposed PWQO values in relation to the health of wildlife drinking the water. Any relevant information found during the literature review phase is assessed on a case by case basis for possible use in adjusting the PWQO.

2.8 Sediment Quality

Contaminants in sediment are a concern because of their potential effect on overlying water quality and water use, their impact on abundance and community composition of the benthic community and their effect, through bioaccumulation, on the organisms higher in the food chain which feed on benthos.

Work is underway to develop a set of guidelines for sediment quality. They would represent a desirable level of sediment quality the Ministry strives to maintain in sediments of the Province.

Because of the links between sediment quality and water quality, proposed PWQOs should be evaluated to determine their potential effect on sediment quality. As sediment quality guidelines become available, these evaluations can be made.

2.9 Toxicant Interactions

The Objective setting process does not specifically address interactions with other contaminants (e.g. synergistic, additive or antagonistic toxicity) or ecological effects. When the effects of mixtures of contaminants are better understood, a re-evaluation may be undertaken.

2.10 Recommending the Final PWQO

The most stringent of the preliminary Objectives developed in the preceding sections of the document is recommended as the PWQO.

2.11 Other Uses and Objectives of Other Agencies

The use of conservative safety factors normally results in an Objective that affords adequate protection for most other beneficial uses of surface waters for individual contaminants. For example, most PWQOs are more stringent than the Ministry's Drinking Water Objectives (MOE, 1983) and the Canadian Water Quality Guidelines for Agricultural Uses and Livestock Watering (CCREM, 1987).

Where available, aquatic environmental criteria of other agencies (e.g. CCREM, 1987) are noted and discussed in the PWQO development document. Since the criteria of other agencies have different development procedures and management strategies, they are not used to alter the PWQO, but do provide a useful comparison.

2.12 Research Needs

Even though the existing data are of sufficient quality and quantity to permit Objective development, this Section of the PWQO development document should focus research toward providing new insights into the toxicity and environmental behaviour of contaminants which would aid in future management decisions. This differs somewhat from the aims of Section 3.10 (Research Needs) where the PWQG development document should focus on procurement of additional data which could be used to reduce the uncertainty associated with the Guideline.

A compilation of current areas of research interests - *Strategic Directions* - is published by the Ministry of the Environment's Research and Technology Branch.

3 . SETTING PROVINCIAL WATER QUALITY GUIDELINES

The Ontario Ministry of the Environment recognizes the need for a procedure to derive water quality guidelines for environmental contaminants where data on aquatic toxicity, mutagenicity and bioaccumulation are insufficient for the development of Provincial Water Quality Objectives. The procedure described in this chapter provides a rational method for consistent development of Provincial Water Quality Guidelines (PWQGs) for organic and inorganic pollutants.

A preliminary Guideline value is calculated by applying a safety factor (called a final uncertainty factor) to the lowest water concentration shown to cause a toxic effect. Since the available database on aquatic toxicity varies from substance to substance, so does the level of assurance in the final product (i.e. the PWQG). In order to account for this and ensure environmental protection, a specific uncertainty factor is calculated for each substance. The size of the uncertainty factor is determined by the physical-chemical properties and the quality and quantity of the aquatic toxicity database.

Although Guidelines and Objectives serve the same purposes, the following points must be kept in mind:

- Guidelines are not as scientifically supportable as PWQOs because they lack the comprehensive data base required for the derivation of Objectives.
- The intention of the Guideline derivation process is to ensure consistency in the interpretation and use of data for setting numerical values and to develop the best

possible recommendation based on available data.

- As new data become available, Guidelines will be periodically reassessed and, if warranted, modified using the Ministry's PWQO setting process. The Guideline setting process requires identification of the scientific research needed to support this step.

3.1 Substance Physical-Chemical Properties

The fate and effects of organic contaminants in aquatic systems are largely determined by their inherent physical-chemical properties. Hence a knowledge of fundamental substance properties (e.g. molecular weight) is essential in any assessment of potential environmental effects (Kobayashi, 1981; Klein *et al.*, 1988; Kollig, 1988; Kollig and Kitchens, 1990; U.S.E.P.A., 1991a;). Where this information is available, it is recorded on the Physical-Chemical Properties sheet (Table 1 - Appendix C).

The most useful chemical properties are those which describe the equilibrium tendencies of a chemical to partition between the various phases (water, biota and sediment) present in an aquatic system. These properties are usually solubilities, vapour pressures or partition coefficients. They may include information about dissociation or speciation of the chemical.

A key property in this regard is octanol-water partition coefficient (K_{ow}). K_{ow} is the ratio of chemical solubility in octanol (a nonpolar phase) to solubility in water (a polar phase). Nonpolar organic chemicals with a high K_{ow}

(log K_{ow} >4) accumulate and reside in nonpolar phases of the environment such as the fatty or lipid-rich tissues of aquatic organisms. Since chemical affinity for octanol is reasonably well correlated with affinity for lipids, octanol has become, for environmental applications, a useful surrogate or model nonpolar organic phase.

It is generally accepted that substances with high K_{ow} 's have significant potential to persist and accumulate in aquatic biota, reaching concentrations which could pose a direct threat to aquatic life and to wildlife and humans consuming aquatic species (Alexander, 1981; Niemi *et al.*, 1987; Veith *et al.*, 1988; Connell, 1990).

Well-known contaminants such as mirex and DDT are examples of substances with high octanol-water partition coefficient values which persist in the environment and accumulate in biota.

Values for K_{ow} are commonly obtained by measurements using equilibrium shake-flask methods or chromatographic retention times (Veith *et al.*, 1979; Doucette and Andren, 1988; De Bruijn *et al.*, 1989). Where a measured value is unavailable, a K_{ow} value can be estimated using correlations with chemical structure or equilibrium properties (Lyman *et al.*, 1982; Miller *et al.*, 1985). Other procedures for *a priori* calculation of an unknown K_{ow} value are outlined in Hansch and Leo (1979) and Lyman *et al.* (1982). It should be noted that the calculation of K_{ow} can be quite complex and measured values are preferred for setting Guidelines.

K_{ow} values are available in the literature for most of the substances which are candidates for Guideline setting (Leo *et al.*, 1971; Hansch & Leo, 1979; Veith *et al.*

1979; Lyman *et al.*, 1982; Doucette and Andren, 1988; De Bruijn *et al.*, 1989). In many cases, the variation between reported values is substantial, especially in the high K_{ow} range where measurement of K_{ow} is difficult and calculated values are often overestimates.

3.2 Selection Of Baseline Uncertainty Factor

If available information indicates that a BCF has been reliably determined for a substance in an aquatic animal, a baseline uncertainty factor of 1000 is chosen for a BCF of less than 1000 (whole body basis) or a baseline uncertainty factor of 10,000 is chosen for a BCF greater than 1000.

If BCF information is unavailable, K_{ow} values are used in Guideline development to select an initial uncertainty factor, as follows:

If log K_{ow} <4.0, the baseline uncertainty factor =1000

If log K_{ow} > 4.0, the baseline uncertainty factor = 10,000

In the development of a preliminary Guideline, the baseline uncertainty factor is adjusted with calibration factors (see Table 4), based on the quality and quantity of available toxicological data, to derive a final uncertainty factor.

While these baseline uncertainty factors and the calibration factors were arbitrarily selected values, their derivation was calibrated so that the PWQG for a substance with a log K_{ow} <4.0 and a complete toxicity data set (Table 4) would have a final uncertainty factor slightly greater than 10. This value approximates the safety factor used to derive a PWQO (See Section 2.2.5).

Where both a measured and estimated or calculated K_{OW} value is available, the measured value is selected as the final value on Table 4 (Appendix C). Although an accurate value for K_{OW} is desirable, selection of the baseline uncertainty factor depends only on a determination that $\log K_{OW}$ is less than or greater than four. If $\log K_{OW}$ is unknown, or if a substance has a range of values spanning $\log K_{OW} = 4.0$, select 10,000 as the baseline uncertainty factor.

Inorganic forms of metals are assigned a baseline uncertainty factor of 1000.

3.3 Evaluation of Toxicological Data

A scientific assessment of the toxicity data must be documented in the Guideline report, with special attention given to the studies from which the lowest observed effect concentrations were obtained. All relevant data, including test conditions, obtained from the literature review, must be listed on the Aquatic Toxicity Data Table (Table 2). Toxicity values (e.g. LC50s) are also plotted on the Guideline Derivation Graph (Figure C1). The toxicity value which is the lowest observed effect concentration must be highlighted on both the table and the graph. When necessary, the toxicity values may be plotted as ranges to conserve space on the graph.

3.3.1 Primary Or Secondary Data

Guidance for categorizing data as acute or chronic and primary or secondary is provided in PWQO Sections 2.2.1 and 2.2.2, respectively. Standard test procedures or protocols do not always generate primary data. Therefore, best scientific judgement must be used to justify a primary or secondary designation for toxicity data, on a case-by-case basis.

3.3.2 Optimal Data Requirements

All available aquatic toxicity data on a substance are reviewed to determine the lowest water concentration associated with an adverse biological effect. In addition, the available data, to a maximum of 11 data within the categories of the optimal data requirements (see below and Table 4), are used in the determination of the final uncertainty factor (see Section 3.3.4). These categories are designed to represent a range of acute and chronic biological effects in a variety of aquatic species.

The following list indicates the optimal data requirements for use in setting Guidelines (Table 4):

CHRONIC DATA ON:

- Three different species of fish.
- Two invertebrate species from different orders.
- One algal or aquatic plant species.

ACUTE DATA ON:

- Three different fish species.
- Two invertebrate species representing different orders.

Toxicological data on marine or brackish water species may not be used in the Guideline setting process.

Data on the aquatic life stage of amphibians can be substituted, appropriately, for one of the six possible fish data listed above.

Data on the aquatic life stage of protozoa can be substituted, appropriately, for one of the

four possible invertebrate data listed above.

3.3.3 Simulated Data

Toxicity data simulated by quantitative structure-activity relationships (QSARs) and acute-chronic ratios (ACRs), which meet the requirements, for different species may be used to a limited extent where measured toxicity data are not available. Information on these data simulation approaches is provided in Könemann, 1981; Kenaga, 1982; Richter *et al.*, 1983; Veith *et al.*, 1983; Bobra *et al.*, 1985; Call *et al.*, 1985; McCarty *et al.*, 1985 and Appendix A.

In Using Simulated Data the Following Rules Apply:

- A QSAR equation may be used once in Guideline setting to interpolate a species-specific simulated acute toxicity value for a compound within the chemical class to which the QSAR applies.
- An ACR, derived from a study designed to establish this ratio, may be applied once to an acute toxicity value for a second fish species to calculate a simulated chronic toxicity value.
- Simulated toxicity values may be used in the absence of measured data to contribute to a decrease in the final uncertainty factor and are plotted on the Guideline Derivation Graph. In the case of ACR's, the acute value used to set the chronic value from the ACR is not used again at later stages in the Guideline setting process.
- Simulated data cannot be employed as the critical data point to which the final uncertainty factor is applied.

3.3.4 Application of Toxicological Data

Each toxicity data point, up to a maximum of 11, which qualifies as a different type of information is now assigned a calibration factor (listed in Table 4) according to its appropriate data category; i.e. acute or chronic and primary or secondary. Calibration factors are multipliers applied to the baseline uncertainty factor to calculate a final uncertainty factor. They represent the weight or influence of the information in reducing baseline uncertainty and provide the link between the quality and quantity of available toxicity information and the size of the final uncertainty factor.

Specific values for the calibration factors were selected so that a complete data set of primary acute and chronic test results for fish, invertebrates and aquatic plants would produce a final uncertainty factor compatible with the existing safety factor used in PWQO development. The calibration factors assigned for plant data and secondary data account for lesser adjustments to the uncertainty factor.

In order to obtain consistency between chemicals, the Guideline setting process must be strictly followed with very little deviation from the prescribed methods. The calculation of a preliminary Guideline is documented on the Toxicity Guideline Uncertainty Factor Worksheet (Table 4). The steps are as follows:

1. Using the bioconcentration factor (BCF), if available, or the octanol-water partition coefficient, K_{ow} (use the log K_{ow} value), select the baseline uncertainty factor indicated on Table 1. Re-enter the baseline uncertainty factor at the bottom of Table 4.

2. Enter in the boxes on Table 4 the key data on each toxicity test.

If more than one test result has been categorized as the same type of information (e.g. if there are three 96h LC50s for rainbow trout), preference for inclusion on the form should be given to the test results which have been assigned the lowest calibration factors (i.e. primary over secondary data and measured over simulated data).

The data which yield the lowest final uncertainty factor should be used on the uncertainty factor worksheet (Table 4). **The lowest observed toxic effect concentration (i.e. the critical value) need not be entered in one of the boxes on the uncertainty factor worksheet (Table 4), even though it is used in the calculation of the preliminary Guideline.**

If there are unfilled acute data boxes, unused chronic data may be used in these boxes provided other requirements (i.e. species diversity) are met. The extra data are considered, for the purpose of reducing uncertainty, as useful as acute information and are entered and assigned a calibration factor for acute information.

If all data boxes cannot be filled, one simulated acute and/or one simulated chronic toxicity value may be used.

3. Re-enter the calibration factor associated with each test result at the bottom of Table 4. Multiply the baseline factor by all the calibration factors to determine the final uncertainty factor. The final uncertainty factor is not to be less than a value of 13.

4. To calculate a preliminary Guideline value, divide the lowest observed toxic effect concentration by the final uncertainty factor.

The minimum data requirement to set a Guideline is one piece of toxicity data (e.g. an LC50 value), acute or chronic, primary or secondary, for a fish or invertebrate. Guidelines cannot be based only on plant, amphibian, protozoan and/or simulated data alone.

A preliminary Guideline based on aquatic toxicity data may be superseded by consideration of other effects described in the following sections.

3.4 Bioaccumulation

Information on both bioaccumulation and consequent health effects is unlikely to be available for the vast majority of industrial organic contaminants. The Guideline setting process considers this limitation and permits the calculation of a Guideline which acknowledges concerns about bioaccumulation. This is done initially through the selection of the baseline uncertainty factor based on a reliably determined BCF or the octanol-water partition coefficient (See Section 3.2 and Figure 2).

If scientifically sound data on bioconcentration and health effects are available, the information is used to calculate a preliminary Guideline as outlined for PWQOs (Section 2.3) and compared to the toxicity-derived Guideline. The lower value would then be recommended as the final PWQG. If bioaccumulation data determine the Guideline, the study used to derive the BCF must be thoroughly investigated and documented in the final report.

3.5 Mutagenicity

A mutagenicity Guideline will be set when sufficient primary information is not available to set a PWQO based on mutagenicity concerns or when there is insufficient data to assess the mutagenic potential of the chemical. The data requirements to set a mutagenicity Objective or to consider a chemical non-mutagenic are presented in Section 2.4.5.1.

A chemical clearly shown to be non-mutagenic need not be assessed further in either the mutagenicity Objective or Guideline process.

Only acceptable data, as outlined in Mutagenicity Objective Section 2.4.1, with the minimum test protocol requirements, as outlined in Section 2.4.2, will be used in setting these Guidelines. However, this Guideline process will consider both primary and secondary data as defined in Section 2.4.3. In addition data from tests using marine organisms may be considered as secondary data in setting mutagenicity Guidelines but not in Objectives.

In many instances primary studies will not be available. In such cases, the available primary information, in combination with any available secondary information will provide the data for setting a mutagenicity Guideline. However, Guidelines will not be set based solely on the results of microbiological or algal studies.

In the case of mutagenicity Guidelines, simulated data derived from QSAR studies will not be used.

The circumstances which require a chemical to be considered in the Guideline process for mutagenicity are identified in Section 2.4.5.

Depending on the availability of acceptable data, a narrative or numeric mutagenicity Guideline may be set. A narrative Guideline will only be set when there are insufficient data to either assess mutagenic potential or to set a numeric Guideline for aquatic organisms.

3.5.1 Numeric Mutagenicity Guidelines

Numeric Guidelines may be set when acceptable data, either primary or secondary are available from tests using aquatic organisms. Only data meeting the minimum requirements will be considered. Available mutagenicity data are summarized, and the pertinent information entered into Table 5 (Appendix C).

If available information indicates that a BCF has been reliably determined for a substance in a species of fish, the preliminary Guideline based on mutagenicity is calculated using a baseline uncertainty factor of 1000 for a BCF of less than 1000 (whole body basis) or a baseline uncertainty factor of 10,000 for a BCF greater than 1000.

If this information is unavailable, a baseline uncertainty factor is chosen based on the log K_{ow} as follows:

If $\log K_{ow} < 4.0$, the baseline uncertainty factor = 1000

If $\log K_{ow} > 4.0$, the baseline uncertainty factor = 10,000

This value is entered into Table 5.

The use of an uncertainty factor of 10,000 for mutagenic chemicals with a $BCF \geq 1000$ or a $\log K_{ow} > 4$ is based on potential accumulation of the chemical in the aquatic food chain by organisms incapable of

metabolizing or excreting the chemical. Predators having the ability to metabolize these chemicals could thus receive a much larger dose through the diet than through waterborne exposure alone.

3.5.1.1 Optimal Guideline Data Requirements

The available acceptable data are sorted as primary or secondary studies, and aquatic vertebrate, invertebrate, plant or microbial organism categories. The suitable studies fall into the following categories:

Primary data from either combination:

2 vertebrate species, and 1 plant species or microbial species

OR

1 vertebrate and 1 invertebrate species, and 1 plant species or microbial species.

Secondary data from:

2 vertebrate species
2 invertebrate species
1 plant species
1 microbial species

The range of studies should include data demonstrating mutagenic and/or genotoxic endpoints as well as whole organism health effects such as heritable mutations or carcinogenicity (for mutagenic carcinogens). In addition, attempts should be made to obtain data from organisms which can and cannot metabolize the chemical.

For each selected study the LOEC values are summarized on Table 5 (Appendix C). In

addition, each study is assigned a calibration factor, as outlined in the Legend, on the basis of primary or secondary data classification. Within organism categories, data from the most sensitive species will be used.

The baseline uncertainty factor is multiplied by all the calibration factors to determine the final uncertainty factor.

The preliminary mutagenicity Guideline is calculated. The mutagenicity LOEC value for the most sensitive species from all organism categories is selected as the critical value. This LOEC value is divided by the final uncertainty factor to determine the preliminary Guideline based on mutagenicity. If this value is more stringent than the preliminary Guideline determined for toxicity and for bioaccumulation, then this value is carried forward to the next step in PWQO setting.

3.5.2 Narrative Mutagenicity Guidelines

A chemical which has not been tested for genotoxic properties or which has not been tested in aquatic organisms may be given a narrative Guideline. Under these conditions the chemical will remain in the Guideline process until data become available and will include one of the following cautions:

- a) No mutagenicity data available.
 - for chemicals which have not been tested for mutagenic or genotoxic activity in any system.
- b) Insufficient data to confirm mutagenicity.

- for chemicals shown to be mutagenic or genotoxic in a single acceptable test but which have been shown to be non-mutagenic or non-genotoxic in all other available tests, or where no additional data are available.

c) Mutagenic - no aquatic data available.

- for chemicals which have been confirmed mutagenic in two tests with acceptable protocols, but where data with aquatic organisms are not available.

3.6 Impacts on the Taste and Odour of Water and Tainting of Fish Tissues

The acquisition, evaluation and use of data follows the same steps as outlined for PWQOs.

3.7 Toxicant Interactions

The Guideline setting process does not specifically address interactions with other contaminants (e.g. synergistic, additive or antagonistic toxicity) or ecological effects. When the effects of mixtures of contaminants are better understood, a re-evaluation may be undertaken.

3.8 Recommending the Final PWQG

The most stringent of the preliminary Guidelines developed in the preceding sections of the document is recommended as the PWQG.

3.9 Other Uses and Objectives of Other Agencies

The use of conservative safety factors normally results in an Guideline that affords adequate protection for most other beneficial uses of surface waters for individual contaminants. For example, most PWQGs are more stringent than the Ministry's Drinking Water Objectives (MOE, 1983) and the Canadian Water Quality Guidelines for Agricultural Uses and Livestock Watering (CCREM, 1987).

Where available, aquatic environmental criteria of other agencies (e.g. CCREM, 1987) are noted and discussed in the PWQO development document. Since the criteria of other agencies have different development procedures and management strategies, they are not used to alter the PWQO, but do provide a useful comparison.

3.10 Research Needs

The intent of this section is to identify any data which would reduce the uncertainty associated with the Guideline developed above. A second aim is to highlight data required to promote a Guideline to an Objective. Specifically, any data gaps identified in Table 6 - Data Requirements for Provincial Water Quality Objectives - are obvious areas for further work. In addition this section should focus research toward providing new insights into the toxicity and environmental behaviour of contaminants which would aid in future management decisions.

A compilation of current areas of research interests - *Strategic Directions* - is published by the Ministry of the Environment's Research and Technology Branch.

4 GLOSSARY OF TERMS

Acceptable Daily Intake (ADI) - an estimated (i.e. theoretically calculated) amount that would result in a "negligible" risk if intaken daily by humans for a lifetime (after HWC, 1990). Use of the term ADI in the current document also covers Tolerated Daily Intake (TDI).

Acute-Chronic Ratio (ACR) - the ratio of the acute toxicity of a toxicant to its chronic toxicity, determined in a single toxicity test. In PWQG setting, it can be used for estimating chronic toxicity from acute toxicity data (after USEPA, 1991a).

Acute - "a stimulus severe enough to rapidly induce an effect; in aquatic toxicity tests, an effect observed in 96 hours or less typically is considered acute" (USEPA, 1991a). An acute effect need not be lethal. "Acute" in connection with toxicity tests refers to short exposures and requires a specific organism as a reference (see Section 2.2 of the current document for examples).

Additive toxicity - "the toxicity of a mixture of chemicals which is approximately equivalent to that expected from a simple summation of the known toxicities of the individual chemicals present in the mixture" (Rand and Petrocelli, 1985).

Antagonistic toxicity - "a phenomenon in which the toxicity of a mixture is less than that which would be expected from a simple summation of the toxicities of the individual chemicals present in the mixture" - i.e. less than additive (Rand and Petrocelli, 1985).

Baseline Uncertainty Factor - in PWQG derivation, the uncertainty factor (1000 or 10000) assigned to a substance based on its potential to bioconcentrate.

Bioaccumulation - the "process by which chemicals are taken up by aquatic organisms from water directly or through consumption of food containing the chemicals" (Rand and Petrocelli, 1985).

Bioaccumulation Factor (BAF) - the ratio of a substance's concentration in tissue or whole body versus its concentration in ambient water, in situations where contaminant uptake is from both food and water (after USEPA, 1991a).

Bioconcentration - the "process by which there is a net accumulation of a chemical directly from water into aquatic organisms resulting from simultaneous uptake (e.g. by gill or epithelial tissue) and elimination" (Rand and Petrocelli, 1985).

Bioconcentration Factor (BCF) - "the ratio of a substance's concentration in tissue versus its concentration in water, in situations where the food chain is not exposed or contaminated. For non-metabolized substances, it represents the equilibrium partitioning between water and organisms" (USEPA, 1991a).

Biomagnification - "result of the processes of bioconcentration and bioaccumulation by which tissue concentrations of bioaccumulated chemicals increase as the chemical passes up through two or more trophic levels. The term implies an efficient transfer of chemical from food to consumer, so that residue concentrations increase from one trophic level to the next." (Rand and Petrocelli, 1985).

Carcinogen - any substance that incites development of a malignant tumour or any other sort of malignancy (after Lapedes, 1974).

Calibration factor - in the PWQG setting process, a weighted multiplication factor assigned to a piece of toxicity data according to its quality. The baseline uncertainty factor (qv) is multiplied by the calibration factors to calculate a substance-specific final uncertainty factor.

Chronic - "a stimulus that lingers or continues for a relatively long period of time, often one-tenth of the life span or more". Endpoint responses to chronic exposures may be lethality, reduced growth, reduced reproduction, etc.. (USEPA, 1991a). The term requires a specific organism and life stage as a reference.

Consumption limit - a concentration of chemical contaminant in the edible portion of fish, below which is considered to be protective of public health or a concentration of chemical contaminant in whole fish, below which is considered to be protective of wildlife.

Cold-water fish species - "those fish species whose populations thrive in relatively cold water, including but not limited to trout, salmon, whitefish and cisco" (MDNR, 1986)

Endogenous activation system - an enzyme complex, normally obtained from the livers of mammals, which have the ability to convert some contaminants into an active mutagenic form.

Final uncertainty factor - in the PWQG setting process, this value is the product of the baseline uncertainty factor (qv) and the calibration factors (qv). The lowest significant effect concentration is divided by this value to calculate the preliminary guideline.

Fish equivalency concentration (FEC) - in PWQO setting, a surrogate for fish consumption limits (qv) when these are unavailable. It is calculated from the ADI (qv) and information on fish consumption by the average Ontario angler.

Genotoxicity - the ability of a substance to damage the integrity of an organism's genetic material.

LC50 - median lethal concentration - the concentration of a material in water to which test organisms are exposed that is estimated to be lethal to 50 percent of a test organisms. The LC50 is usually expressed as a time-dependent value." (Rand and Petrocelli, 1985).

Life-cycle test - with fish and other aquatic organisms, "a study in which all the significant stages of an organism are exposed to a test material. Generally, a life-cycle involves an entire reproductive cycle of the organism" (Rand and Petrocelli, 1985).

Lowest observed effect concentration (LOEC) - "the lowest concentration of a material used in a toxicity test that has a statistically significant adverse effect on the exposed population of test organisms as compared with the controls." (Rand and Petrocelli, 1985).

Mutagenicity - a measure of a chemicals ability to cause heritable change (a mutation) in genetic material in either somatic or reproductive cells.

Narrative mutagenicity guideline - a qualitative caution applied to a suspected mutagenic chemical for which there is insufficient information to set a numeric value.

No observed effect concentration (NOEC) - "the highest concentration of a material in a toxicity test that has no statistically significant effect on the exposed population of test organisms as compared with the controls" (Rand and Petrocelli, 1985).

Octanol-water partition coefficient - "the ratio at equilibrium, in a two-phase system, of a chemical in the n-octanol phase to that in the water phase" (USEPA, 1991b).

Partial life-cycle test - with fish and other aquatic organisms, a test in which the most sensitive life stage of an organism (based on previous testing) is exposed to the toxicant. For most contaminants this represents a good surrogate for full life-cycle toxicity tests.

Persistence - a chemical characteristic which renders it resistant to decay, degradation, transformation, volatilization, hydrolysis, or photolysis (after USEPA, 1991a).

Primary data - in PWQO and PWQG setting, data which meets certain minimum requirements in the areas of laboratory practice, test conditions, toxicant concentrations, test endpoints, length of exposure, measurement of environmental parameters, responses and survival controls (see Section 2.2.2).

QSAR - quantitative structure-activity relationship "in which some property of a series of molecules (such as solubility or toxicity) is correlated with molecular structure". This enables toxicity and physical-chemical properties to be estimated rather than measured (Suntio *et al.*, 1988).

Safety factor - in PWQO setting, a factor

applied to the LOEC for the most sensitive organism. It is intended to provide a margin of safety to account for the fact that more sensitive species may not have been tested, and the generally stressed condition of field animals in comparison to the laboratory animals used to derive the LOEC.

Secondary data - Data not meeting all of the requirements for primary data but which meet certain minimum requirements for use in PWQG development documents.

Simulated data - data simulated by either quantitative structure-activity relationships (*qv*) or acute-chronic ratios (*qv*). Used in PWQG setting, to a limited extent, where measured data are unavailable.

Sublethal - "below the concentration that directly causes death. Exposure to sublethal concentrations of material may produce less obvious effects on behaviour, biochemical and/or physiological function, and histology of organisms" (Rand and Petrocelli, 1985).

Synergistic (toxicity) - "a phenomenon in which the toxicity of a mixture of chemicals is greater than that which would be expected from a simple summation of the toxicities of the individual chemicals present in the mixture" (Rand and Petrocelli, 1985).

Tainting - imparting a taste or odour in fish tissues.

Teratogenic - capable of causing alteration in the normal development of cells, tissues, or organs of the embryo.

Tertiary data - data which do not qualify as either primary or secondary in quality and are not used in either PWQO or PWQG development documents. This data may be discussed in the text.

Uncertainty factor - in PWQG setting, a factor weighted by the quality and quantity of available toxicity or mutagenicity information, applied to the LOEC for the most sensitive organism. It is intended to provide a margin of safety to account for the fact that more sensitive species may not have been tested, the generally stressed condition of field animals in comparison to the laboratory animals used to derive the

LOEC, and the paucity of toxicity information.

Warm-water fish species - "those fish species whose populations thrive in relatively warm water, including but not limited to bass, pike, walleye and panfish" (MDNR, 1986).

5 REFERENCES

- Alexander, M. 1981. Biodegradation of chemicals of environmental concern. *Science*, 211: 132-139.
- Amoore, J. E., and E. Hautala. 1983. Odour as an aid to chemical safety: odour thresholds compared with threshold limit values and volatilities for 214 industrial chemicals in air and water dilution. *J. Appl. Toxicol.*, 3: 272-290.
- Benoit D. A., F. A. Puglisi and D. L. Olson. 1982. A fathead minnow *Pimephales promelas* early life stage toxicity test method evaluation and exposure to four organic chemicals. *Environ. Poll. (series A).*, 28: 189-197.
- Bobra, A., W. Y. Shiu and D. Mackay. 1985. Quantitative structure-activity relationships for the acute toxicity of chlorobenzenes to *Daphnia magna*. *Environ. Toxicol. Chem.*, 4: 297-305.
- Call, D., L. Brooke, M. Knuth, S. Poitier and M. Hoglund. 1985. Fish subchronic toxicity prediction model for industrial organic chemicals that produce simple narcosis. *Environ. Toxicol. Chem.*, 4: 335-341.
- CCREM. 1987. Canadian Water Quality Guidelines. Canadian Council of Resource and Environment Ministers, Toronto. .
- Connell, D.W. 1990. Bioaccumulation of xenobiotic compounds. CRC Press, Florida. ISBN 0-8493-4810-2.
- Cox, C., A. Vaillancourt and A. F. Johnson. Feb. 1989. A Method for determining the intake of various contaminants through the consumption of Ontario sport fish. Aquatic Biology Section, Water Resources Branch.
- De Bruijn, J., F. Busser, W. Seinen, and J. Hermens. 1989. Determination of octanol-water partition coefficients for hydrophobic organic chemicals with the "slow-stirring" method. *Environ. Toxicol. Chem.*, 8: 499-512.
- Doucette, W. J., and A. W. Andren. 1988. Estimation of octanol/water partition coefficients: evaluation of six methods for highly hydrophobic aromatic hydrocarbons. *Chemosphere*, 17: 345-359.
- Hansch, C., and A. J. Leo. 1979. Substituent constants for correlation analysis in chemistry and biology. John Wiley and Sons, New York.
- HWC. 1990. Health Risk Determination: The challenge of health protection. Health and Welfare Canada. 36 pp.

- Kenaga, E. E. 1982. Predictability of chronic toxicity from acute toxicity of chemicals in fish and aquatic invertebrates. *Environ. Toxicol. Chem.*, 1: 347-358.
- Klein, A. W., W. Klein, W. Kordel, and M. Weiss. 1988. Structure-activity relationships for selecting and setting priorities for existing chemicals - A computer assisted approach. *Environ. Toxicol. Chem.* 7: 455-467.
- Kobayashi K. 1981. Safety examination of existing chemicals - Selection, testing, evaluation, and regulation in Japan. *Proceedings, Workshop on the control of existing chemicals under the patronage of the Organization for Economic Co-operation and Development, West Berlin, F.R.G. June 10-12, 1981.* pp. 141-163.
- Kollig, H.P. 1988. Criteria for evaluating the reliability of literature data on environmental process constants. *Toxicol. Environ. Chem.*, 28: 95-103.
- Kollig, H.P. and B.E. Kitchens. 1990. Problems associated with published environmental fate data. *Toxicol. Environ. Chem.*, 28: 95-103.
- Könemann, H. 1981. Quantitative structure-activity relationships in fish toxicity studies. Part 1: relationship for 50 industrial pollutants. *Toxicology*, 19: 209-221.
- Lapedes, D. N. (ed.) 1974. *Dictionary of Scientific and Technical Terms.* McGraw Hill Book Co., New York.
- Leo, A., C. Hansch, and D. Elkins. 1971. Partition coefficients. *Chem. Rev.*, 17: 525-616.
- Lyman, W. J., W. F. Reehl and D. H. Rosenblatt (ed.). 1982. *Handbook of chemical property estimation methods: Environmental behavior of organic compounds.* McGraw Hill Book Co., New York. (A computer version of this approach is also available, entitled CHEMEST).
- Mayer, F.L., K.S. Mayer and M.R. Ellersieck. 1986. Relation of survival to other endpoints in chronic toxicity test with fish. *Environ. Toxicol. Chem.* 5: 737-748.
- McCarty, L. S., P. V. Hodson, G. R. Craig and K. L. E. Kaiser. 1985. The use of quantitative structure-activity relationships to predict the acute and chronic toxicities of organic chemicals to fish. *Environ. Toxicol. Chem.*, 4: 595-606.
- McKim, J. M. 1977. Evaluation of tests with early life stages of fish for predicting long-term toxicity. *J. Fish. Res. Board Can.*, 34: 1148-1154.
- MDNR. 1986. *State of Michigan Water Quality Standards.* Michigan Department of Natural Resources. Water Resources Commission.

- Miller, M. M., S. P. Wasik, G. L. Huang, W. Y. Shiu and D. Mackay. 1985. Relationships between octanol-water partition coefficient and aqueous solubility. *Environ. Sci. Technol.*, 19: 522-278.
- MOE. 1981. Fish Tainting. Ontario Ministry of the Environment. 9 pp.
- MOE. 1983. Ontario Drinking Water Objectives. Ontario Ministry of the Environment. 56 pp.
- MOE. 1984. Water Management: Goals, Policies, Objectives and Implementation Procedures of the Ministry of the Environment. Ontario Ministry of the Environment. 70 pp.
- MOE. 1987. Development of an Ontario effluent monitoring priority pollutants list. Ontario Ministry of the Environment. ISBN 0-7729-2784-7
- Newell, A. J., D. W. Johnson and L. K. Allen. 1987. Niagara River Biota Contamination Project: Fish Flesh Criteria for Piscivorous Wildlife. New York State Department of Environmental Conservation.
- Niemi, G. J., G. D. Veith, R. R. Regal and D. D. Vaishnov. 1987. Structural features associated with degradable and persistent chemicals. *Environ. Toxicol. Chem.*, 6: 515-527.
- Niimi, A. and B. Oliver. 1983. Biological half-lives of polychlorinated biphenyl (PCB) congeners in whole fish and muscle of rainbow trout (*Salmo gairdneri*). *Can. J. Fish. Aquat. Sci.*, 40: 1388-1394.
- Norberg, T. J. and D. I. Mount. 1985. A new fathead minnow subchronic toxicity test. *Environ. Toxicol. Chem.*, 4: 711- 718.
- Rand, G. M. and S. R. Petrocelli (ed.). 1985. *Fundamentals of Aquatic Toxicology: Methods and Applications*. Hemisphere Publishing Corporation, Washington.
- Richter, J. E., S. F. Peterson and C. F. Kleiner. 1983. Acute and chronic toxicity of some chlorinated benzenes, chlorinated ethanes and tetrachloroethylene to *Daphnia magna*. *Arch. Environ. Contam. Toxicol.*, 12: 679-684.
- Schaeffer, D. A. 1981. Is "no-threshold" a "non-concept". *Environmental Management* 5 (6), 475-481.
- Suntio, R. S., W. Y. Shiu and D. Mackay. 1988. A review of the nature and properties of chemicals present in pulp mill effluents. *Chemosphere*, 17: 1249-1290.
- USEPA. 1991a. Technical support document for water quality-based toxics control. United States Environmental Protection Agency. Office of Water. PB91-127415.

- USEPA 1991b. (DRAFT) Assessment and control of bioconcentratable contaminants in surface waters. United States Environmental Protection Agency. Office of Water.
- Veith, G. D., N. M. Austin and R. T. Morris. 1979. A rapid method for estimating log P for organic chemicals. *Water Research*, 13: 43-47.
- Veith, G. D., D. J. Call and L. T. Brooke. 1983. Structure- toxicity relationships for the fathead minnow, *Pimephales promelas*: narcotic industrial chemicals. *Can. J. Fish. Aquat. Sci.*, 40: 743-748.
- Veith, G.D., E. Greenwood, R.S. Hunter, G.J. Niemi and R.R. Regal. 1988. On the intrinsic dimensionality of chemical structure space. *Chemosphere* 17(8): 1617-1630.
- Verscheuren, K. 1983. Handbook of environmental data on organic chemicals. Van Nostrand Reinhold Co., Toronto.

APPENDIX A - Bioaccumulation: Theoretical Considerations

Bioconcentration generally refers to the process by which an aquatic organism, when exposed only to water containing a concentration of chemical (C_w), absorbs the chemical from solution and accumulates a higher concentration (C_b) (McKim *et al.*, 1985; Connell, 1988). The ratio C_b/C_w is a bioconcentration factor (BCF). Bioconcentration can only be measured in the laboratory because there is, by definition, no exposure via consumption of contaminated food.

In general, BCF refers to a steady-state condition in which uptake and elimination (clearance) rates have equalized. If an organism cannot readily metabolize or eliminate the chemical compound, the steady-state corresponds to an equilibrium condition in which chemical transport by passive diffusion (exchange partitioning) is the prime determinant of BCF.

BCF may also be defined, for a simple fish-water system, as a ratio of uptake and elimination (first order) rate constants, k_1/k_2 , respectively (Neely, 1979). Each rate constant quantifies the fraction of chemical absorbed or eliminated in unit time. They are determined, for a particular substance and fish, by measuring changes over time in the chemical concentration in biota observed in separate uptake and elimination experiments.

A review of empirical BCF data for a wide range of nonpolar organics suggests that uptake rate constants vary by a much smaller factor than do elimination rate constants and it is the latter which accounts for most of the differences in BCF between compounds (e.g. Könemann and van Leeuwen, 1980; Bruggeman *et al.*, 1981). In addition, the exposure time necessary to reach a steady-state equilibrium condition depends only on the elimination rate constant. Niimi (1987) provides a recent review and compilation of values of this key kinetic parameter.

In the ambient aquatic environment organisms may accumulate chemicals from both water and food; a process termed bioaccumulation. The ratio C_b/C_w is, in this case, a bioaccumulation factor (BAF). If the concentration of chemical in food (C_e) is known, C_b/C_e is sometimes referred to as a biomagnification factor (BMF). The relative importance of food as an exposure route to aquatic biota increases with increasing body size of the organism and increasing hydrophobicity of the chemical.

If experimental data on chemical disposition in aquatic biota were available, a pharmacokinetic model could be derived and used to assess bioconcentration potential. The available information is, however, insufficient to justify such a model. Given the large number of contaminants which have been detected in aquatic systems, it is unlikely that a comprehensive data set for each and every chemical will ever be available.

It is clear that a simple screening procedure is required. One which enables predictions or simulations to be made of relative bioaccumulation potential from a standard set of chemical

benchmark properties; so-called partitioning models based on quantitative structure-activity relationships (QSARs). The QSAR approach is a practical and consistent procedure which provides simple descriptors for:

1. *a priori* estimation of BCF from K_{OW} ,
2. extrapolation of measured BCF values for a given water-biota system to other biota or other structurally similar chemicals
3. evaluation of existing BCF data based on "expected behaviour".

For example, many linear relationships of the general form:

$$\log \text{BCF} = a \log K_{OW} + b$$

have been derived and reported in the literature. The slopes (a) and intercepts (b) of these linear correlations vary greatly for a variety of reasons, including both biological differences in test species (Barber *et al.*, 1988) as well as experimental and statistical artifacts and errors (e.g. short exposure times, unconsidered chemical bioavailability and over-estimated K_{OW} values). Any given set of values for the constants in this general equation are therefore specific to the test system from which the data were obtained.

It is not possible, at present, to easily separate or delineate all the factors which contribute to variations between correlations. One solution is to develop correlations based on compiled data sets, as has been done by Lyman *et al.* (1982). However these correlations tend to underestimate the bioaccumulation potential of persistent hydrophobic chemicals.

The following approach, based on simple fish-water exchange partitioning theory and key biological attributes, is recommended as an assessment tool:

The simplest analysis is based on the assumptions that an organism achieves equilibrium with water and that the lipid portion of the organism is equivalent in properties and amount to octanol. The chemical concentration in organism lipids (C_L) will then be the product $C_w \times K_{OW}$. For example, if the water concentration is 0.1 g/m^3 and the K_{OW} is 1000, then the concentration in the lipid will be 100 g/m^3 . If the organism consists of 5% lipids (i.e. lipid volume fraction (L) is 0.05), then the whole body concentration (C_b) will be $0.1 \times 0.05 \times 1000$ or 5 g/m^3 . In general terms:

$$\text{BCF} = L \times K_{OW}$$

This type of analysis is generally satisfactory for compounds with: a high proportion of C-C (aliphatic), C-C (aromatic), C-H and C-Cl bonds, molecular weight <300 g/mol, molar volume <250 cm^3/mol , $\log K_{OW}$ 1-6, water solubility 0.002-18 mol/m^3 (Connell, 1988). These structural attributes and physical-chemical properties are closely interrelated and generally

describe nonpolar, moderately hydrophobic chemicals which are resistant to biotic and abiotic transformation (e.g. persistent chlorinated hydrocarbons).

There is, however, a loss of linearity in the BCF- K_{OW} correlation at high K_{OW} 's, in both laboratory and field situations. There are at least eight possible reasons (both experimental artifacts and real phenomena) which must be considered when judging the validity and applicability of both measured and estimated BCF values:

1. If $\log K_{OW}$ is low, i.e. less than 1, the substance displays an almost equal tendency to partition between lipids and water. Bioconcentration will be underestimated because a significant fraction of the chemical will be present in the organism's water phase (Gobas, *et al.*, 1986). A convenient "correction", for the purpose of estimating BCF, is to add 1.0 to K_{OW} (not $\log K_{OW}$) to account for hydrophilic behaviour.
2. If $\log K_{OW}$ is very large, i.e. greater than 6, estimated BCFs tend to be higher than measured BCFs. This is a particular problem when $\log K_{OW}$ reaches 7 or 8. The reasons for this are not entirely clear. Gobas *et al.* (1987; 1988) address this issue in detail. A simple procedure is to assume that chemicals with a $\log K_{OW}$ exceeding 6.5 have a ceiling value of 6.5. This first order correction improves the prediction of lab-measured BCFs, although the measured value itself may still be an underestimate of bioaccumulation potential.
3. Many compounds (e.g. polynuclear aromatic hydrocarbons) are metabolized and eliminated in some species at a rate which is a significant determinant of the tissue concentration. The higher the rate, the lower the measured BCF compared to an estimated value. It is very difficult, at present, to predict the extent of this decrease. If evidence suggests that metabolism (biotransformation) is operative, the person developing or using the BCFs should search for a reported "field" bioaccumulation factor. In some species, metabolism of a chemical causes a drastic increase in its toxicity. Therefore, metabolism does not always provide assurance of safety.
4. There may be a food chain effect. BCFs derived from laboratory data may underestimate, by a factor of about 10 to 15, the body burdens of hydrophobic ($\log K_{OW}$ 5-7) chemicals in field-caught fish. This is an indication that consumption of contaminated food is the major route of exposure. Residue levels increase with fish size and age and a fish-water equilibrium may never be reached for compounds with a $\log K_{OW} > 5$.

If specific information on food chain effects is not available, a BAF for a top level predacious fish can be roughly estimated as the BCF multiplied by a factor of 3 for each trophic level below the fish. For example, if one views a food chain as an alga-zooplankton-minnow-predacious fish system and the BCF for the fish is 1000, then the BAF = $1000 \times (3 + 3 + 3) = 9000$.

5. Exposure time in a "steady-state" experiment was too short and a fish-water

steady-state equilibrium was not reached such that the organism has a lower concentration than expected. For a series of related chemicals of increasing K_{ow} , elimination rate constants generally decrease (e.g. Bruggeman *et al.*, 1981). In other words, the exposure time required to reach a steady-state concentration in biota (t_{eq}) increases with K_{ow} . The t_{eq} also tends to increase with increasing size and lipid content of the exposed organism.

The following correlating equation, proposed by Gobas and Mackay (1987), can be used to judge the relevance of this phenomenon for a given data set:

$$1/k_2 = (V_L/Q_W) \times K_{ow} + (V_L/Q_L)$$

where V is the volume (litres) of lipid in the fish, Q is a transport parameter (litres per day), the subscripts L and W refer to lipid and water compartments within the fish and k_2 is the elimination (first order) rate constant.

Q_W and Q_L are organism-specific parameters, primarily dependent on fish body weight according to:

$$Q_W = 1.3 * M^{0.64}$$

$$Q_L = 0.014 * M^{0.6}$$

M = fish body weight between 0.1 and 1000 grams

It is thus possible to estimate: (1) k_2 (2) half-life time = $t_{1/2} = 0.693/k_2$ (3) time to steady-state equilibrium = $5 \times t_{1/2}$ for "every" chemical in "all" fish species if data are available for: (1) fish body weight (2) lipid content (3) chemical K_{ow} .

Growth dilution of tissue concentrations (C_b) may also be a significant determinant of the chemical concentration in biota during long term exposures. Estimates of biological half-lives based on body burden (i.e. $C_b \times M$) provide a sounder basis for half-life comparisons of the same chemical among studies and between species (Niimi, 1987).

There may be reduced bioavailability of the chemical in water. Sorption of the chemical to organic carbon-containing matter suspended in water may significantly reduce the concentration of truly dissolved, bioavailable chemical. If octanol is assumed to be an adequate surrogate for organic carbon, then the fraction of chemical which is in true solution is often approximated by:

$$\text{fraction dissolved} = 1 / (1 + K_{ow} C_o)$$

where C_o is the organic carbon content of water in g/g (not mg/L). If $\log K_{ow} = 4$ and $C_o = 10 \text{ mg/L}$ or 10^{-5} g/g , then the fraction dissolved is 0.91, 9% being sorbed and "unavailable".

Experimental work to test the validity of this equation is lacking.

7. The chemical may display unusual or specific properties, for example, it may ionize (e.g. Saarikoski *et al.*, 1986), exhibit surface active (surfactant) properties, chelate, or vary in nature between metallic and organo-metallic forms. In such situations, case by case assessment is required.

While partitioning tendency of organic compounds and organo-metals can be related to a single parameter, octanol-water partition coefficient, no such simple generalization is possible for inorganic forms of metals. Mechanisms of uptake and storage of metals in aquatic biota are diverse and strongly influenced by factors such as chemical speciation and interactions with complexing ligands. Classical concepts of bioaccumulation typically do not hold for most metals.

8. If an organism is benthic in nature, tissue concentrations may be more indicative of those prevailing in the sediments than in the water column. In such situations, case by case assessment is required.

Standard test procedures or protocols do not always consider the potential problems described above. Therefore, best scientific judgement can and must be used to assess data on a case-by-case basis. Unless there are compelling arguments supporting an exception, the final BCF, standardized for fresh fish tissue having a lipid content (L) of 10%, can be determined as follows in order of preference:

- (a) measured steady-state value from a laboratory test.
- (b) projected steady-state value calculated from test data
- (c) measured value from other types of laboratory tests
- (d) measured bioaccumulation factor from a field study if the water concentration was relatively constant, "bioavailable" and less than known toxic levels.
- (e) simulated value calculated as $BCF = L \times K_{ow}$ (described above).
- (f) simulated value calculated using correlations with other physical-chemical properties (see Lyman *et al.*, 1982).

Caution must be exercised because erroneous conclusions may result from a comparison of BCFs expressed in different concentration units. Chemical concentration in whole fish or selected tissues are reported on the basis of wet weight, dry weight or lipid content (lipid normalized). While whole fish values can be used to assess aquatic environmental behaviour, muscle values may be more useful to evaluate consumable fish products for human health concerns. Reported water concentrations should reflect the truly dissolved (bioavailable) fraction of the chemical in water.

REFERENCES

Barber, M. C., L. A. Suarez, and R. R. Lassiter. 1988. Modelling bioconcentration of nonpolar organic pollutants by fish. *Environ. Toxicol. Chem.*, 7: 545-558.

- Bruggeman, W. A., L. B. J.M. Martron, D. Kooiman, and O. Hutzinger. 1981. Accumulation and elimination kinetics of PCBs by goldfish after dietary and aqueous exposure. *Chemosphere*, 10: 811-832.
- Connell, D. W. 1988. Bioaccumulation behavior of persistent organic chemicals with aquatic organisms. *Rev. Environ. Contam. Toxicol.*, 4: 117-154.
- Gobas, F., K. Clark, W. Y. Shiu and D. Mackay. 1988. Bioconcentration of polybrominated benzenes and biphenyls and related super-hydrophobic chemicals in fish: role of bioavailability and elimination in the faeces. *Environ. Toxicol. Chem.*, 8: 231-245.
- Gobas, F., D. Mackay and W. Y. Shiu. 1987. Bioconcentration of highly hydrophobic chemicals. Chapter in *QSAR in Environmental Toxicology II*: 107-123. K.L.E. Kaiser [ed.] D. Reidel Publ. Co.: Dordrecht.
- Gobas, F., and D. Mackay. 1987. Dynamics of hydrophobic chemical bioconcentration in fish. *Environ. Toxicol. Chem.*, 6: 495-504.
- Gobas, F., A. Opperhuizen, and O. Hutzinger. 1986. Bioconcentration of hydrophobic chemicals in fish: relationship with membrane permeation. *Environ. Toxicol. Chem.*, 5: 637- 646.
- Könemann, H., and K. van Leeuwen. 1980. Toxicokinetics in fish: accumulation and elimination of six chlorobenzenes by guppies. *Chemosphere*, 9: 3-19.
- Lyman, W. J., W. F. Reehl, and D. H. Rosenblatt. 1982. Handbook of chemical property estimation methods: Environmental behavior of organic compounds. McGraw Hill Book Co., New York. (A computer version of this approach is also available, entitled CHEMEST).
- McKim, J. M., P. Schmieder, and G. Veith. 1985. Absorption dynamics of organic chemical transport across trout gills as related to Kow. *Toxicol. Appl. Pharmacol.*, 77: 1-10.
- Neely, W. B. 1979. Estimating rate constants for the uptake and clearance of chemicals by fish. *Environ. Sci. Technol.*, 13: 1506-1510.
- Niimi, A. J. 1987. Biological half-lives of chemicals in fish. *Rev. Environ. Contam. Toxicol.*, 99: 1-46.
- Saarikoski, J. R. Lindstrom, M. Tyynela, M. Viluksela 1986. Factors affecting the absorption of phenolics and carboxylic acids in the guppy *Poecilia reticulata*. *Ecotox. Environ. Safety*. 11: 158-173.

APPENDIX B —MUTAGENICITY: ACCEPTABLE TESTS AND SPECIES

ACCEPTABLE TESTS

A. Assessment of Mutagenic Potential

The assessment of mutagenic or genotoxic potential may employ all available mutagenicity data, from tests with acceptable protocols, including data from aquatic as well as mammalian species. In addition, data from microbial or tissue culture test systems, with and without endogenous activation systems may be employed. Although such data are acceptable in this initial assessment, only data from tests employing aquatic organisms are acceptable in setting Objectives or Guidelines for mutagenicity.

B. Acceptable Tests for Objectives and Guidelines.

Only tests using fresh water organisms will be acceptable for setting mutagenicity Objectives. Data from tests using fresh water and marine organisms may be considered in setting Guidelines.

These lists of acceptable tests are examples and are not exclusive.

a. Mutagenicity Related Diseases

Data from these tests are considered when the substance in question is demonstrated to be mutagenic by other assays.

- i. Increased incidence of cancers or tumours in aquatic organisms. Waterborne exposure for significant proportion of life-cycle of aquatic species.
- ii Increased incidence of mutagenicity related diseases in offspring of aquatic organisms including birth defects, heritable mutation or dominant lethal mutations.

b. Mutagenicity (resulting from exposure of whole organisms)

- i Microbial assays such as the Ames test without endogenous activation.
- ii Plant somatic mutation assays such as the *Tradescantia* staminal hair assay.
- iii Somatic cell mutation (for example ouabain resistance) provided the exposure to the chemical was in-vivo (whole animal).
- iv Host-mediated mutation assays.
- v Anaphase or metaphase abnormalities of chromosomes.
- vi Micronuclei.

c. Genotoxicity (resulting from exposure of whole organisms)

- i DNA damage assays (Unscheduled DNA Synthesis)
- ii Microbial DNA damage assays (Chromotest or Induct assays) in the absence of metabolic activation.
- iii Sperm-head abnormality
- iv Sister-chromatid exchanges
- v DNA-adducts

ACCEPTABLE SPECIES

A. Vertebrate

Any freshwater fish species (marine species are acceptable as secondary data for Guidelines)

Amphibians with life-stages resident in fresh water

B. Invertebrate

Freshwater invertebrates, either pelagic or benthic including polychaetes, oligochaetes, crustaceans and molluscs. Data from tests using marine invertebrates, particularly those with freshwater relatives, would not be used in Objectives but may be considered in Guidelines.

C. Plant

Freshwater algae

Freshwater macrophytes, either wholly submergent, or those emergent from water with roots in the bottom substrate.

D. Bacteria

Bacteria shown to be resident in fresh water. Ames test *Salmonella* (without chemical activation) is an acceptable surrogate for freshwater bacteria.

APPENDIX C -WORKSHEETS AND TABLES

TABLE 1: PHYSICAL-CHEMICAL PROPERTIES

COMPOUND:	CHEMICAL FORMULA:	CAS No:
PROPERTIES		
MOLECULAR WEIGHT (MW).		
MELTING POINT:		
BOILING POINT:		
PHYSICAL STATE AT STANDARD TEMPERATURE AND PRESSURE.		
DISSOCIATION CONSTANT:		
LIQUID DENSITY (D):		
MOLAR VOLUME (MW/D):		
VAPOUR PRESSURE (P_s):		
WATER SOLUBILITY (C_s):		
HENRY'S LAW CONST. (P_s/C_s):		
PERSISTENCE		
B.O.D. (mg/L):		
BREAKDOWN PRODUCTS:		
HALF LIFE (DAYS):		
OCTANOL-WATER PARTITION COEFFICIENT (K_{ow})		
RANGE OF AVAILABLE Log K_{ow} VALUES:		
FINAL CHOSEN Log K_{ow} VALUE:		
BASELINE UNCERTAINTY FACTOR FOR GUIDELINE DEVELOPMENT		
IF BCF < 1000, USE 1000		
IF BCF \geq 1000, USE 10000		
IF BCF DATA IS UNAVAILABLE, USE Log K_{ow} :		
IF Log K_{ow} < 4.00, USE 1000		
IF Log K_{ow} \geq 4.00, USE 10000		
BASELINE UNCERTAINTY FACTOR:		
(Enter this value Into Tables 4 & 5)		
<input type="text"/>		

Table 2: Aquatic Toxicity Data Table

COMPOUND:			CAS No.			CONCENTRATION UNITS				
TEST COND.	Species (life stage)	Response	pH	Temp	D.O.	Alk.	Hard.	Effect conc.	Data codes	Ref. no.
ACUTE	VERTEBRATE									
	INVERT.									
CHRONIC	VERTEBRATE									
	INVERT.									
	PLANT									

* - INDICATES VALUES ENTERED INTO TABLE 4

Assign 3 data codes, one from each of the following rows:
 A = acute C = chronic
 S = static R = static/renewal F = flowthrough
 U = unmeasured nominal conc. M = measured conc

TABLE 3: MUTAGENICITY DATA TABLE FOR -

	SPECIES	END-POINT key (1)	TISSUE or ORGAN AFFECTED	LIFE STAGE and TIME or DURATION	TEST CONDITIONS		CONC.'S TESTED (mg/L)	RESPONSE IN CONTROL and at CONC.'S TESTED	LOEL (mg/L)	DATA CODES key (2)	DATA QUALITY key (3)	REF. NO.
					ACTN.	SOLVENT						
VERTEBRATE												
INVERTEBRATE												
PLANT												
MICROBIOLOGICAL												

key (1) END-POINT: AdD = DNA adducts; BsD = DNA strand breaks; CA = chromosomal aberrations; DLM = dominant lethal mutations; dmD = DNA damage/repair HM = heritable mutations; HoM = host-mediated mutation; M = mutagenic in microbial systems; Mn = micronuclei; Neo = neoplasia; SCE = sister chromatid exchange; Sha = spermhead abnormality; SM = somatic mutations; UDS = unscheduled DNA synthesis

key (2) DATA CODES: F = Flowthrough; R = renewed static; S = static; M = measured toxicant concentrations; U = unmeasured toxicant concentrations

key (3) DATA QUALITY: P = primary(1°); S = secondary(2°); Aq = aquatic organism; NAq = non-aquatic organism

Table 4: UNCERTAINTY FACTOR WORKSHEET

CHEMICAL:		CAS No.			CONCENTRATION UNITS mg/L		
Test Conditions	Species (life stage)	Toxicity End Point	Effect conc.	Data Codes	Data Type	Calibration Factor	No. and Reference
ACUTE	VERTEBRATE						
	INVERT.						
CHRONIC	VERTEBRATE						
	INVERT.						
	PLANT						

CALCULATION OF FINAL UNCERTAINTY FACTOR:

Since $BCF < 1000$ (or $\text{Log } K_{ow} < 4$) / $BCF \geq 1000$ (or $\text{Log } K_{ow} \geq 4.00$),

The Baseline Uncertainty Factor = 1000/10000

Baseline Uncertainty Factor X Calibration Factors (maximum number = 11)

x x x x x x x x x x x

= FINAL UNCERTAINTY FACTOR (NOT TO BE LESS THAN A VALUE OF 13)
 CRITICAL VALUE + FINAL UNCERTAINTY FACTOR = PWQG VALUE

= _____ + _____ = _____ mg/L

Assign 2 DATA CODES, one from each of the following rows:
 S - static R - static/renewal F - flowthrough
 U - unmeasured nominal conc. M - measured conc

DATA TYPE:
 1^o = Primary 2^o = Secondary 3^o = Simulated Data
 7 = Unknown (Default Data Quality = 2^o)

TABLE 4 (CONTINUED)

**CALCULATION OF THE FINAL
UNCERTAINTY FACTOR**

CALIBRATION FACTORS FOR TOXICITY DATA

ACUTE DATA	
PRIMARY (1°)	0.8
SECONDARY (2°)	0.9
SIMULATED BY QSAR (3°)	0.9
CHRONIC DATA	
PRIMARY (1°)	0.5
SECONDARY (2°)	0.7
SIMULATED BY ACR (3°)	0.8
PLANTS AND ALGAE	0.9

INSTRUCTIONS FOR COMPLETING THE UNCERTAINTY FACTOR WORKSHEET

1. A maximum of eleven (11) calibration factors may be used in calculating the final uncertainty factor.
2. For the purposes of setting Guidelines, all plant and algal data are considered secondary and chronic.
3. In the absence of a full set of measured acute and chronic toxicity data, simulated data from one ACR and/or one OSAR may be used.
4. The following data may be substituted to reduce the final uncertainty factor:
 - toxicity data on aquatic stages of amphibians may be used once as fish data.
 - aquatic toxicity data on protozoans may be used once as invertebrate data.
5. Guidelines cannot be based on plant, amphibian, protozoan and/or simulated data alone.
6. Unfilled acute data boxes may be filled with unused chronic data, provided other requirements (i.e. species diversity) are met. The acute calibration factor is used.

Table 5: MUTAGENICITY GUIDELINE UNCERTAINTY FACTOR WORKSHEET

CHEMICAL		CAS No.		CONCENTRATION UNITS mg/L		
Test Conditions	Species (life stage)	Toxicity End Point	LOEL (1) conc.	Calibration Factor (2)	No. and Reference	
PRIMARY	VERTEBRATE					
	VERTEBRATE or INVERTEBRATE					
	PLANT or MICROBIAL					
SECONDARY	VERTEBRATE					
	INVERT. VERTEBRATE					
PLANT						
MICROBIAL						

CALCULATION OF FINAL UNCERTAINTY FACTOR:

Since $BCF < 1000$ (or $\text{Log } K_{ow} < 4$) or $BCF \geq 1000$ (or $\text{Log } K_{ow} \geq 4.00$).

The Baseline Uncertainty Factor = 1000/10000

Baseline Uncertainty Factor X Calibration Factors (maximum number = 9)

X X X X X X X X X X

= FINAL UNCERTAINTY FACTOR

CRITICAL VALUE + FINAL UNCERTAINTY FACTOR = PWQG OR INTERIM GUIDELINE VALUE

= _____ + _____ = _____ mg/L

- (1) LOEL = Lowest concentration of chemical Inducing an effect significantly above that of the control.
- (2) Calibration Factor: Primary data = 0.4; Secondary data = 0.8

Table 6: Data Requirements for Provincial Water Quality Objectives

1) Toxicity

All data must be primary, chronic (or equivalent - use an asterisk to denote substitution of acute data to chronic data), and marine or brackish species are not permitted.

Fish

At Least:

	one coldwater species
	one warmwater species
	one other warmwater or coldwater species

With at Least:

	one species resident in Ontario (may be one of above)
	one early lifestage endpoint
	one other whole organism chronic endpoint

Invertebrates

At Least:

	one crustacean
	one non - crustacean

With at Least:

	no more than one tropical species
	one early life stage endpoint
	one other chronic endpoint

Algae/Aquatic Plant

	one algae or aquatic plant resident in temperate North America using scientific procedures and test conditions compatible with recognized algal bioassays
--	---

Table 6 (cont'd): Data Requirements for Provincial Water Quality Objectives

2) Bioaccumulation One of:	
	fish consumption limit (e.g. Health and Welfare Canada Guidelines)
	an acceptable daily Intake limit
	contaminant residue In aquatic biota value

and:

	A Bioconcentration Factor ≥ 1000 (In the absence of consumption limits, bioaccumulation may be significant and the Guideline setting process should be followed.)
--	--

or:

	A Bioconcentration Factor < 1000 (In the absence of consumption information, bioaccumulation is not considered to be significant.)
--	--

If BCF data is unavailable:

	Log $K_{ow} \geq 4.00$, then bioaccumulation is assumed to be significant and the Guideline setting process should be followed
--	---

or:

	Log $K_{ow} < 4.00$, then bioaccumulation is assumed not to be significant
--	---

Table 6 (cont'd): Data Requirements for Provincial Water Quality Objectives

3) Mutagenicity

a) For initial assessment:

	Chemical is considered to be non-mutagenic (i.e. data from a minimum of two test systems, including tests for mutagenic as well as chromosomal damage endpoints, clearly demonstrating the chemical to be non-mutagenic)
	Chemical is considered to be mutagenic in aquatic or mammalian species

b) For setting PWQOs (a total of three studies are required for mutagenicity):

Vertebrates

All data must be primary and measured in whole aquatic organisms. Marine and brackish tests are not permitted.

Data from at least one of the following three categories

	fish - mutagenicity-related diseases
	fish - mutagenicity or chromosomal aberration
	other vertebrate mutagenicity or chromosomal aberration

Invertebrates

Data from a maximum of two of the following three categories

	invertebrate - mutagenicity or chromosomal aberration
	aquatic plant - mutagenicity or chromosomal aberration
	microbial - mutagenicity

FIG. C1: DERIVATION GRAPH -

Toxicity Information Considered		Species and Life Stage	Exposure and Duration	Concentration (mg/L)	Ref. No.
ACUTE	Vert.				
	Invert.				
	Plant				
	other Organisms				
CHRONIC	Vert.				
	Invert.				
	Plant				
	other Organisms				
MUTAGENICITY	Vert.				
	Invert.				
	Plant				
	other Organisms				
taste odour tainting					
BCF					
PWOO/PWOG					
Other Criteria					

o - Critical Value
* - PWOO/PWOG

■ - Toxicity endpoints
Δ - Odour Protection Values
▲ - Taste Protection Values

□ - Criteria of Other Agencies
● - Tainting Protection Values