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Value Assessment of Pest Control Products

PMRA Guidance Document



*Protecting the health and
environment of Canadians*



*Protéger la santé des Canadiens
et l'environnement*

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Updated	Update/Rationale:
September 2013	Initial publication of DIR2013-03 <i>Value Assessment of Pest Control Products</i>
March 2022	Revisions based on current documentation standards. Re-issued as PMRA Guidance Document.

Disclaimer

This document does not constitute part of the Pest Control Products Act or its regulations and in the event of any inconsistency or conflict between the Act or regulations and this document, the Act or the regulations take precedence. This document is an administrative document that is intended to facilitate compliance by the regulated party with the Act, the regulations and the applicable administrative policies.

Related documents

This guideline should be read in conjunction with the accompanying notice and the relevant sections of the following guidance documents:

- Value Guidelines for New Plant Protection Products and Label Amendments
- Value Guidelines for New Personal Insect Repellent Products and Label Amendments
- Pesticide Resistance Management Labelling Based on Target Site/Mode of Action
- A framework for Risk Assessment and Risk Management of Pest Control Products
- Value Guidelines for New Antimicrobial Pest Control Products and Label Amendments

This guideline does not identify or establish requirements that are outside or in addition to current legislation. See the [Pest control products \(pesticides\) acts and regulations](#) section of the Canada.ca website for a list of all regulatory requirements.

Please note that this guidance document is in effect as of March 2022 and should be used for applications submitted on or after 1 April 2022.

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Executive Summary

This document describes the Pest Management Regulatory Agency's (PMRA) approach to the value assessment of pesticides. It outlines the considerations and key concepts of value assessments and describes information needed to support an application for proposed pesticide uses from a value perspective.

Previously, the PMRA relied primarily on efficacy information to establish the value of proposed uses of pesticides. The current approach is based on the weight of evidence from information on various components, which includes efficacy and benefits of the proposed use. In addition, the type of information that can be provided can come from a variety of sources.

The PMRA's approach to value assessment was developed over a number of years and was subject to informal consultation by industry and grower groups, as well as the Canadian public through a regulatory proposal that preceded DIR2013-03, *Value Assessment of Pest Control Products*. The consideration of the definition of value as stated in the *Pest Control Products Act* provides flexibility in fulfilling information requirements, which will help reduce regulatory burden for applicants. It will also provide opportunities for alignment with member countries of the Organisation for Economic Cooperation and Development (OECD) and an increase in predictability and efficiency of value assessments.

1.0 Purpose

1.1 Overview

In accordance with the *Pest Control Products Act*, only pest control products that are determined to be of acceptable value are approved for use in Canada. The value of a pest control product, as defined by the *Pest Control Products Act* (paragraph 2(1)), refers to the product's actual or potential contribution to pest management, taking into account its conditions or proposed conditions of registration. This includes the product's efficacy, effect on host organisms, health, safety and environmental benefits derived from use of the product and social or economic impacts.

The value assessment is an integral part of the evaluation conducted on all pesticides before they are registered in Canada. To determine acceptable value, a weight of evidence approach is taken that considers all the factors that may contribute to a product's value.

This document explains the PMRA's approach to the value assessment of pesticides. As the specific details regarding the use of a pest control product can vary, there is flexibility in interpretation and application of the approach with regards to the manner in which value requirements can be addressed. The type of information that can be provided for the value assessment can include one or a combination of the following:

- Experimental data generated from research trials,
- Use history information from other jurisdictions,
- Rationales based on accepted scientific principles,
- Existing scientific literature, and
- Benefits information.

Trial data are required for uses that relate to control of pests having a direct effect on human health, such as swimming pool and spa products and disease vector control products.

1.2 Definitions and regulations

Test Data as defined by the *Pest Control Products Act*, are: scientific or technical information respecting the health or environmental risks or the value of a pest control product.

Value, in respect of a pest control product, means the product's actual or potential contribution to pest management, taking into account its conditions or proposed conditions of registration, and includes the product's:

- (a) efficacy;
- (b) effect on host organisms in connection with which it is intended to be used; and
- (c) health, safety and environmental benefits and social and economic impact.

An application to register or amend the registration of a pest control product must include any characteristics that are relevant to its health or environmental risks or value (refer to Pest Control Products Regulations section 6). Per section 7(3) (a) of the *Pest Control Products*

Act, the applicant must conduct any evaluations that the PMRA considers necessary with respect to the health or environmental risks or the value of the pest control product. Moreover, the applicant has the burden of persuading the Minister that the value of the product is acceptable (refer to *Pest Control Products Act* section 7(6)).

In addition to the information required as per *Pest Control Products Act* section 7, the applicant must provide any other information required to evaluate the health and environmental risks and the value of the pest control product.

2.0 Value assessment: approach and considerations

2.1 Efficacy

Pesticide products should have a level of efficacy that significantly contributes to pest management in order to be registered in Canada. Different types of information may be used to address product efficacy. These include use history information, published papers, scientific rationales, and/or trial data. The information provided to support efficacy should permit an assessment of the level, duration and consistency of control provided by the product. It should demonstrate the product's efficacy (in other words, the resulting level of control or intended effect) under the proposed conditions of use in Canada.

When use history information is provided, it should include a comparison of the Canadian-proposed and foreign-registered use patterns, an indication of the performance and experience associated with the use of the product in the foreign jurisdiction, and a validation of the use history information, where appropriate. Published scientific papers may also be used to address efficacy provided that it is clearly related to the proposed use. Arguments based on established scientific principles and precedent registrations may be used to address efficacy. When a rationale is provided, applicants should indicate its scientific basis and/or any specific details related to precedent registrations that apply.

When trial data are used as supporting information, a summary table in a spreadsheet format should be provided; the findings should be discussed in relation to the proposed label claims in the value summary document. Information on field trials generated outside of Canada may be submitted provided that it can be shown to be relevant to Canadian conditions. Information from greenhouse trials conducted outside of Canada is acceptable and does not require additional information regarding trial conditions unless there are differences with respect to crop production practices.

Consistent with the goal of sustainable pest management, applicants should demonstrate that the proposed application rate provides an acceptable level of control under representative conditions without being excessive.

2.2 Effects on host organisms

The effects of the pesticide on host organisms or use sites should be explained in the value summary report. Information in the form of use history, published studies, scientific rationales or research trials may be provided. For field trials, observations on host crop tolerance may be made within efficacy trials unless phytotoxic reactions occur that necessitate further

study. Applicants should describe the basis on which they have determined that an adequate margin of crop safety exists for each host and rotational crop. If the potential for crop injury exists then warning statements to this effect should be included on the product label.

2.3 Health, safety and environmental benefits

Information concerning the actual or potential benefits associated with the proposed pesticide use may be considered during the value assessment. Information provided in this regard should demonstrate how and to what extent the proposed registration would benefit Canadian users and other direct stakeholders (for example, consumers or applicators) from a health /safety and/or environment perspective. For example, a product may be needed to control a poisonous plant, a pest that impacts public health (for example, mosquitos or ticks), an invasive species (for example, zebra mussels), or a plant disease with harmful effects on humans/livestock (for example, ergot). Actual or potential contribution to risk reduction of the proposed pesticide use could also be explained, such as a product that is considered a replacement for a pesticide that is being phased out following re-evaluation.

Information on the compatibility of the proposed use with an integrated pest management strategy may also be considered during the value assessment. For example, spot application of products with narrow pest spectrum or limited or no negative impacts on biological control agents or beneficial insects. Consistent with the PMRA's goal of sustainable pest management, an explanation of how the proposed use contributes to resistance management in consideration of other registered alternatives should be provided, when applicable.

2.4 Social and economic impact

This information on social or economic impacts associated with the proposed use can also contribute to the value assessment process. This includes effects on:

- The sustainability of the sector or trade implications especially the impact on competitiveness of Canadian growers,
- Crop value (farm-gate, market value),
- Acreage devoted to crop,
- Influence of the pest on crop quality and marketability,
- Additional costs associated with the pest presence (for example, drying costs for grain), and/or
- Indirect effects of the pest on the crop (for example, alternate host for a crop disease).

The projected economic benefits should be described in relation to the pest problem and the system in which the pesticide is to be used. Quantitative estimates are preferable, but qualitative information is also useful. Additional information such as an attribute that contributes to a product's value could also be included. For example, if a product is stable for longer periods without the need for refrigeration, this attribute affects product cost.

2.5 Weight of evidence approach to value assessment

Applicants should consider how best to support the value of proposed pesticide uses considering that a combination of various types of information may be submitted. If the product is registered in a foreign jurisdiction and the use pattern there is similar to that proposed in Canada, supporting information for efficacy could mainly consist of use history information. In cases where the active ingredient is new and the product is not registered elsewhere, efficacy trials may be the main source of value information. Scientific rationales to extend support for other uses based on information on certain uses or published information may also be provided.

In all cases, it is useful to provide the PMRA with the benefits and the social and economic impact of the proposed use. This is particularly important when the scientific information to support the proposed use is limited. The PMRA's value assessment of pesticides is based on a weight of evidence approach that considers all the value information provided to support the application.

3.0 Joint Health Canada/Agriculture and Agri-Food Canada (HC/AAFC) Minor Use Pesticide Program

The joint HC/AAFC Minor Use Pesticide Program is aimed at adding minor uses to the labels of registered pest control products. A minor use is defined as a necessary use of a pest control product for which the anticipated volume of sales is not sufficient to persuade a manufacturer to register and sell the product in Canada (for further details, please refer to the document [User Requested Minor Use Label Expansion](#) in the pesticides portion of Canada.ca). In order for a minor use of a pesticide to be registered, as with all uses of pest control products, it must be shown to have acceptable value.

Within this program, if a minor use is prioritized by growers at the Annual Canadian Minor Use Priority Setting Workshop as a project for which Agriculture and Agri-Food Canada's Pest Management Center will generate supporting information (in other words, an 'A priority'), and is submitted for registration under the joint HC/AAFC Minor Use Program, then the PMRA will not require value information to support the minor use registration. The priority setting exercise itself establishes the primary pest management needs for minor crops and satisfies the need to demonstrate acceptable value. A label review will be conducted to determine the extent to which the proposed use pattern is consistent with the registered use pattern and that any differences between them are justified.

4.0 Structuring the value information package

The value information package consists of a value summary report (see Appendix I) and associated documents such as use history information, or individual trial reports, if applicable. The value summary report provides a description of the overall value associated with the proposed use. It is an executive summary of the entire value information package. Key elements related to efficacy, host and rotational crop tolerance, health, safety and

environmental benefits and social and economic impact should be integrated and discussed as they apply.

Note: The PMRA will also accept applicant generated supporting value information submitted to other regulatory agencies (for example, public interest finding documents submitted to the United States Environmental Protection Agency and biological dossiers submitted to OECD).

Supporting guidance documents are available separately on the [Policies and Guidelines page in the pesticides section of the Canada.ca website](#) to stakeholders to assist in the preparation of the value information package. These documents are not intended to be prescriptive, but rather are an additional resource to provide more specific guidance related to the preparation of value information drawing on established international protocols such as OECD and European and Mediterranean Plant Protection Organization guidelines.

In addition to the guidance documents, templates to assist in the preparation of the value information package are also available regarding summarizing efficacy data and use-history information. Additional templates may be developed in the future. Furthermore, a pre-submission consultation can be requested in which the PMRA provides advice regarding the preparation of the value information package.

Appendix I Template for the value assessment of pesticides

This template outlines the information considered in an application to register a pesticide or to add a proposed use to a currently registered product. It is intended to assist applicants in the preparation of the Value Summary upon which the value assessment is based. This template should be used in conjunction with relevant guidance documents, summary tables for efficacy or crop tolerance trials, and use history information templates, as appropriate.

Value summary

1.0 Introduction

1.1 Product description

Provide a description of the formulated product and the active ingredient.

1.2 Use pattern

1.2.1 Registered use pattern

For products that are already registered in Canada, provide a summary description of the use pattern currently registered on the Canadian label. Include information that is relevant to any new uses that are being proposed below.

1.2.2 Proposed use pattern/Amendments to registered use pattern

Provide a description of the proposed use pattern (for unregistered products) or proposed amendments to the registered use pattern (for registered products), for example, crop(s), site(s), pest(s), application rate, etc.

1.3 Description of the pest problem

Provide a description of the pest(s) proposed to be added to the label, including common and Latin binomial name(s), the nature and severity of the damage to the crop(s) associated with the pest(s).

2.0 Efficacy

This portion of the value summary presents all information related to efficacy. This may include experimental results from research trials, published scientific literature, scientific rationales, and use history information.

2.1 General factors affecting efficacy

Describe any general factors that may influence product efficacy.

2.2 Supporting information from earlier formulations of the product or similar products

If efficacy information is available for product formulations tested during earlier stages of development, or similar products, rationales and bridging data should be presented in this section to demonstrate equivalence between the products.

2.3 Requirements for adjuvants

If an adjuvant is proposed to be used with the product, demonstrate why the adjuvant is required, or under what circumstances it is required.

2.4 Support for proposed claims

Each claim should be identified (each application timing, method of application, pest, tank mix, etc.), and the approach and information used to support each of these claims, with respect to efficacy, should be clearly indicated.

3.0 Effects on host organisms

This portion of the value package presents all information related to non-safety adverse effects (for example, phytotoxicity to the host or rotational crop, damage to the site of application, etc.).

3.1 General factors influencing effects on host organisms

Describe any general factors that may influence effects on host organisms.

3.2 Supporting information from earlier formulations of the product or similar products

If information is available for product formulations tested during earlier stages of development, or similar products, rationales and bridging data should be presented in this section to demonstrate equivalence between the products.

3.3 Support for proposed claims

3.3.1 Host crop claims

Each claim should be identified (in other words, each application timing, method of application, pest, tank mix, etc.), and the approach and information used to support each of these claims should be clearly indicated.

3.3.2 Rotational crop claims

Each claim should be identified (in other words, each rotational crop and its proposed replanting interval relative to application), and the approach and information used to support each of these claims should be clearly indicated.

4.0 Consideration of benefits

4.1 Alternatives

An overview of products registered in Canada for the same uses currently being proposed. If the current alternatives do not address grower/user needs, an explanation should be provided.

4.2 Compatibility with current management practices including integrated pest management

A description of how the proposed use can be integrated into the production system, including its contribution to integrated pest management.

4.3 Resistance management

A description of how the proposed use contributes to resistance management, in consideration of other registered alternatives.

4.4 Contribution to risk reduction

A description of how the proposed use contributes to risk reduction, in consideration of other registered alternatives, should be provided.

4.5 Social and economic impacts

A description of any social or economic impacts associated with the proposed use such as effects on the sustainability of the sector or trade implications. Information explaining why the product is needed as well as how and to what extent product registration would benefit Canadian users should be provided.

4.6 Health, safety and environmental benefits

A summary of any potential health, safety or environmental benefits that could result from the proposed use of the pesticide should be provided. It is not a summary of the information provided to support the human health or environmental risk assessment.

5.0 Summary and conclusions

An integrated summary of all information provided to support the value of the proposed use(s).

6.0 References

A list of reference materials or documents cited in the value summary report.