Pest Management Regulatory Agency
Agence de réglementation de la lutte antiparasitaire

Re-evaluation Decision Document RRD2006-21

Etridiazole

The purpose of this Re-evaluation Decision Document (RRD) is to notify registrants, pesticide regulatory officials and the Canadian public that Health Canada's Pest Management Regulatory Agency (PMRA) has re-evaluated the active ingredient etridiazole and its associated uses as a fungicide on greenhouse non-food crops, terrestrial food crops, outdoor ornamentals, indoor plants and plantscapes as well as turf.

On 21 March 2006, the Proposed Acceptability for Continuing Registration document PACR2006-03, *Re-evaluation of Etridiazole* was published for consultation. The PMRA did not receive any comments concerning PACR2006-03.

The PMRA has determined that this active ingredient is acceptable for continued registration. Mitigation measures to further protect workers, users, bystanders and the environment are specified in PACR2006-03. The registrants have been informed by letter of the specific requirements, including confirmatory data requirements as defined in this RRD (Appendix I), affecting their product registrations and the regulatory options available to comply with this decision.

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Appendix I Revised Data Requirements

The registrants of etridiazole will be required to submit the following within 24 months of finalization of the re-evaluation decision:

- a submission to register a source of the technical grade active ingredient; and
- efficacy data to support a maximum rate of 4.26 kg a.i./ha etridiazole on golf course greens or a rationale for this rate that includes relevance to the Canadian use pattern.

Registrants should note that specific data, selected from the data package that was submitted to the United States Environmental Protection Agency to support reregistration of this active ingredient, may be required by the PMRA in the future with respect to use expansions, special reviews or minor uses, or to establish maximum residue limits (MRLs).