

URMULE Update

User Requested Minor Use Label Expansion Program Registrations for April 1, 2002 – April 30, 2002

The User Requested Minor Use Label Expansion (URMULE) Update provides timely information on the latest proposals for minor use label expansion of pesticides that have been evaluated and accepted for registration.

For the following pesticides, a Certificate of Registration has been issued based on a Supplemental Label, making the use legal. Registrants are required to add the new use to the full product label at the next printing. Provincial authorities, in publishing recommendations on the use of agricultural chemicals, may incorporate such acceptable uses in the knowledge that they meet the requirements of the *Pest Control Products Act*. Users must consult the product label as the ultimate authority on whether or not the product can be used to control a specific pest on a specific site, as a product's uses are subject to change.

NOTE: The URMULE Updates are available on the Pest Management Regulatory Agency's Internet site, address provided below. If Internet access is not available, arrangements can be made to receive this publication via facsimile or electronic mail. Please contact the Publications Coordinator at the location provided below.

(publié aussi en français)

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\mathbf{MRL}^1

Vegetables

Rimsulfuron (Prism 25DF - DuPont)		
registration number 23938 for the control		
of hairy nightshade on fresh market field		
tomatoes.	proposed 0.05 ppm	
Quizalofop-p-ethyl (Assure II - DuPont)		
registration number 25462 for the control of		
listed weeds in rutabaga in Ontario and		
Quebec only.	proposed 0.1 ppm	
Quizalofop-p-ethyl (Assure II - DuPont)		
registration number 25462 for the control of		
listed weeds in pinto and navy beans in		
Western Canada only.	proposed 0.15 ppm ³	
Quizalofop-p-ethyl (Assure II - DuPont)		
registration number 25462 for the control of		
listed weeds in beans - white, white and		
red kidney, cranberry, black, brown and		
yellow-eye in Southern Ontario only.	proposed 0.15 ppm ³	
Quizalofop-p-ethyl (Assure II - DuPont)		
registration number 25462 for the control of		
listed weeds in lima, mung and adzuki beans		
in Southern Ontario only.	proposed 0.15 ppm ³	
	proposed one ppm	
Fruit		
Tebufenozide (Confirm 240F - Rohm & Haas)		
registration number 24503 extension of temporary	cranberries	1 ppm ²
registration for insect control on cranberry.	dried cranberries	5 ppm^2
Fenhexamid (Elevate 50 WDG - Arvesta)		
registration number 25900 for the control of		
Botrytis on red and black raspberries, loganberries		
and blackberries.	proposed 20 ppm ³	

Field crops

Rimsulfuron+**nicosulfuron** (Ultim 75DF - DuPont)

registration number 24736 for weed control in field corn in Manitoba.

Azoystrobin (Quadris - Syngenta)

registration number 26153 for the control of rust on seed corn in Ontario only.

nicosulfuron 0.1 ppm rimsulfuron 0.1 ppm

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Footnotes

- ¹ Maximum Residue Limits (MRLs) have been added to this document for the user's information and convenience. Under the *Food and Drugs Act*, it is prohibited to sell food containing residues of pest control products at a level greater than 0.1 ppm unless a higher MRL has been established in Table II, Division 15, of the Food and Drug Regulations or an Interim Marketing Authorization (IMA) has been issued. In case of discrepancy between this document and the Food and Drug Regulations, information in the latter will prevail. For more information on MRLs and IMAs, please refer to the PMRA URMULE Regulatory Directive (DIR2001-01, Sections 4.2 and 5 (step 5)) or on the Internet at <u>http://www.hc-sc.gc.ca/pmraarla/english/legis/maxres-e.html.</u>
- ² An IMA was issued on August 4, 2001 to permit the sale of dried cranberries and cranberries containing residues of tebufenozide with MRLs of 5 ppm and 1 ppm respectively while the regulatory process to amend the Food and Drug Regulations is completed.
- ³ A request for an IMA is being processed to permit the sale of the food containing residues of the active ingredient up to the proposed MRL listed above while the regulatory process to amend the Food and Drug Regulations is completed.