

Queensland



*Chemical Usage (Agricultural and Veterinary) Control Act 1988*

# **CHEMICAL USAGE (AGRICULTURAL AND VETERINARY) CONTROL REGULATION 1999**

**Reprinted as in force on 14 December 2003  
(includes commenced amendments up to 2003 SL No. 335)**

**Reprint No. 1F**

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The reprint includes a reference to the law by which each amendment was made—see list of legislation and list of annotations in endnotes. Also see list of legislation for any uncommenced amendments.

This page is specific to this reprint. See previous reprints for information about earlier changes made under the Reprints Act 1992. A table of reprints is included in the endnotes.

**Also see endnotes for information about—**

- **when provisions commenced**
- **editorial changes made in earlier reprints.**

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# Queensland



## CHEMICAL USAGE (AGRICULTURAL AND VETERINARY) CONTROL REGULATION 1999

### TABLE OF PROVISIONS

Section		Page
<b>PART 1—PRELIMINARY</b>		
1	Short title . . . . .	5
<b>PART 2—PRESCRIBED AND PROSCRIBED CHEMICALS</b>		
2	Prescribed chemicals—Act, s 4 . . . . .	5
3	Proscribed chemicals—Act, s 11C(2) . . . . .	5
<b>PART 3—PRESCRIBED MAXIMUM RESIDUE LIMITS</b>		
<i>Division 1—Preliminary</i>		
4	Purpose of pt 3 . . . . .	6
5	Definitions for pt 3 . . . . .	6
<i>Division 2—MRLs</i>		
6	MRLs for chemicals for human food—MRL standard . . . . .	7
7	Other MRLs for chemicals for human food . . . . .	7
8	MRLs for chemicals for animal food—sch 2 . . . . .	8
<b>PART 4—USE OF CHEMICAL PRODUCTS</b>		
<i>Division 1—Restricted chemical products</i>		
9	Definitions for div 1 . . . . .	8
10	Restricted chemical products containing bifenthrin or chlorpyrifos . . . . .	9
11	Restricted chemical products containing endosulfan . . . . .	9
11A	Restricted chemical products containing pindone . . . . .	10
12	Other restricted chemical products . . . . .	10
<i>Division 2—Records of chemical product use</i>		
13	Record requirement . . . . .	10
14	Obligation to keep record . . . . .	11

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

---

**PART 5—HORMONAL GROWTH PROMOTANTS**

*Division 1—Preliminary*

15	Definitions for pt 5 . . . . .	11
----	--------------------------------	----

*Division 2—Obligations if HGP treatment given*

16	Obligation to make required earmark . . . . .	13
17	Obligation to record HGP treatment . . . . .	13
18	Obligation to keep HGP treatment record . . . . .	14

*Division 3—Obligations if cattle with HGP free tag are sold*

*Subdivision 1—Saleyard sales by agents*

19	Agent's obligation to give statement . . . . .	14
20	Agent's obligation to keep copy of statement . . . . .	15
21	Buyer's obligation to keep and produce statement . . . . .	15

*Subdivision 2—Other sales*

22	Seller's obligation to give declaration . . . . .	16
23	Requirements for declaration . . . . .	16
24	Seller's obligation to keep copy of declaration . . . . .	17
25	Buyer's obligation to keep and produce declaration . . . . .	17

**PART 6—SUPERVISION FEES AND EXPENSES**

26	Application of pt 6 . . . . .	18
27	Hourly fee . . . . .	18
28	Overnight absence expenses . . . . .	19

**PART 7—MISCELLANEOUS PROVISIONS**

29	Approval of forms . . . . .	19
----	-----------------------------	----

	<b>SCHEDULE 1</b> . . . . .	20
--	-----------------------------	----

**PRESCRIBED AND PROSCRIBED CHEMICALS**

	<b>SCHEDULE 2</b> . . . . .	22
--	-----------------------------	----

**MRLS FOR CHEMICALS FOR ANIMAL FOOD**

**ENDNOTES**

1	Index to endnotes . . . . .	25
2	Date to which amendments incorporated . . . . .	25
3	Key . . . . .	25

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

---

4	Table of reprints . . . . .	26
5	List of legislation . . . . .	26
6	List of annotations . . . . .	27



# **CHEMICAL USAGE (AGRICULTURAL AND VETERINARY) CONTROL REGULATION 1999**

[as amended by all amendments that commenced on or before 14 December 2003]

## **PART 1—PRELIMINARY**

### **1 Short title**

This regulation may be cited as the *Chemical Usage (Agricultural and Veterinary) Control Regulation 1999*.

## **PART 2—PRESCRIBED AND PROSCRIBED CHEMICALS**

### **2 Prescribed chemicals—Act, s 4**

For section 4 of the Act, definition “chemical”, paragraph (b), each substance mentioned in schedule 1 is prescribed to be a chemical.

### **3 Proscribed chemicals—Act, s 11C(2)**

For section 11C(2) of the Act, each chemical mentioned in schedule 1 is a proscribed chemical.<sup>1</sup>

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<sup>1</sup> Section 11C (Governor in Council may proscribe chemicals) of the Act  
For proscribed chemicals, see section 9 (Person not to possess or use proscribed chemical) of the Act.

## **PART 3—PRESCRIBED MAXIMUM RESIDUE LIMITS**

### *Division 1—Preliminary*

#### **4 Purpose of pt 3**

This part prescribes, for the Act section 4, definition “maximum residue limit” and section 38(2)(b), the MRL for certain chemicals for agricultural produce.<sup>2</sup>

#### **5 Definitions for pt 3**

In this part—

“**ERL**” means extraneous residue limit.

“**extraneous residue limit**” means an extraneous residue limit within the meaning of the MRL standard.

“**human food commodity**” means agricultural produce intended or normally used for human consumption.

“**MRL**” means maximum residue limit.

“**MRL standard**” means the National Registration Authority for Agricultural and Veterinary Chemicals, *MRL Standard Maximum Residue Limits in Food and Animal Feedstuffs of Agricultural and Veterinary Chemicals and Associated Substances*, published by the Australian Government Publishing Service, Canberra.<sup>3</sup>

“**prescribed qualification**” means a statement of attainment issued by a registered training organisation stating that an individual has successfully completed each of the following competencies—

- (a) RTC3704—Prepare and apply chemicals;
- (b) RTC3705—Transport, handle and store chemicals.

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2 See also the Food Standards Code, within the meaning of the *Food Act 1981*, standard 1.4.2 (Maximum residue limits (Australia only)).

3 A copy of the MRL standard may be inspected, free of charge, during office hours on business days at the department’s office at 80 Ann Street, Brisbane. The standard may be viewed on the Australian Pesticides and Veterinary Medicines Authority’s website at <http://www.apvma.gov.au/residues>.

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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**“registered training organisation”** means a training organisation registered under the *Training and Employment Act 2000* or under similar legislation of another State.

*Division 2—MRLs*

**6 MRLs for chemicals for human food—MRL standard**

(1) If the MRL Standard fixes an MRL level for a chemical for a human food commodity, that level is the prescribed MRL for the chemical for the commodity as a human food commodity.

(2) If the MRL Standard does not fix an MRL level for the chemical for the commodity but fixes an ERL level for the chemical for the commodity, the ERL level is the prescribed MRL for the chemical for the commodity as a human food commodity.

**7 Other MRLs for chemicals for human food**

(1) If the MRL Standard does not fix an MRL or ERL level for a particular human food commodity, the prescribed MRL for the chemical for the commodity as a human food commodity is zero.

(2) If the MRL Standard does not fix an MRL or ERL level for a chemical for any human food commodity, the prescribed MRL for the chemical for any human food commodity is zero.

(3) However, subsections (1) and (2) do not apply if the use of the chemical in relation to the commodity as a human food commodity is allowed under part 2<sup>4</sup> of the Act.

(4) If subsection (3) applies, no MRL is prescribed for the use of the chemical mentioned in subsection (3).

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4 Part 2 (Use of chemicals and substances having chemical residues) of the Act

## **8 MRLs for chemicals for animal food—sch 2**

(1) The prescribed MRL for a chemical<sup>5</sup> mentioned in schedule 2 for any animal food is the level stated opposite the name of the chemical in schedule 2.

(2) If a chemical is not mentioned in schedule 2, no MRL is prescribed for the chemical for animal food.

(3) In this section—

“**animal food**” means agricultural produce intended or normally used for animal consumption.

“**chemical**” includes a residue of the chemical stated in the MRL Standard, table 3.<sup>6</sup>

## **PART 4—USE OF CHEMICAL PRODUCTS**

### *Division 1—Restricted chemical products*

## **9 Definitions for div 1**

In this division—

“**authorised**”, for a restricted chemical product, means authorised to use the product under—

- (a) an approved label for containers for the product; or
- (b) a permit for the product.

“**prescribed qualification**” means a statement of attainment issued by a registered training organisation stating that an individual has successfully completed each of the following competencies—

- (a) RTC3704—Prepare and apply chemicals;
- (b) RTC3705—Transport, handle and store chemicals.

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5 Under the Act, section 4, definition “chemical”, paragraph (a), the term ‘chemical’ includes a ‘chemical product’.

6 MRL Standard, table 3 (Residue definition)

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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**“registered training organisation”** means a training organisation registered under the *Training and Employment Act 2000* or under similar legislation of another State.

**“restricted chemical product”** means a restricted chemical product under the Agvet Code.

### **10 Restricted chemical products containing bifenthrin or chlorpyrifos**

A person must not use a restricted chemical product containing bifenthrin or chlorpyrifos, unless the person is—

- (a) authorised to use the product; or
- (b) licensed for a pest control activity under the *Pest Management Act 2001* and the licence permits the person to use the product.

*Example of a restricted chemical product containing chlorpyrifos—*

Dursban Pre-Construction Termiticide.

Maximum penalty—40 penalty units.

### **11 Restricted chemical products containing endosulfan**

A person must not use a restricted chemical product containing endosulfan, unless the person—

- (a) is authorised to use the product; or
- (b) holds an unrestricted commercial operator’s licence or a pilot chemical rating licence under the *Agricultural Chemicals Distribution Control Act 1966*; or
- (c) holds an accreditation to use agricultural chemicals from any of the following—
  - (i) a training organisation trading in any State under the name ‘ChemCert’;
  - (ii) Farmcare Australia Farm Chemical User Training Program Incorporated;

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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- (iii) Queensland Agricultural Chemicals Accreditation Council Incorporated, previously trading under the names ‘Chemsafe Training Queensland’ and ‘Chemsmart Training Queensland’; or

(d) holds a prescribed qualification.

Maximum penalty—40 penalty units.

### **11A Restricted chemical products containing pindone**

A person must not use a restricted chemical product containing pindone, unless the person—

- (a) is authorised to use the product; or
- (b) holds an approval granted by the chief executive of the Department of Health under the *Health (Drugs and Poisons) Regulation 1996*, section 18,<sup>7</sup> to obtain, possess and use fluoroacetic acid.

Maximum penalty—40 penalty units.

### **12 Other restricted chemical products**

(1) This section applies only to a restricted chemical product that does not contain bifenthrin, chlorpyrifos, endosulfan or pindone.

(2) A person must not use the product, unless the person is authorised to use the product or holds a prescribed qualification.

Maximum penalty for subsection (2)—40 penalty units.

### ***Division 2—Records of chemical product use***

### **13 Record requirement**

(1) This section applies to a person if—

- (a) the person uses a chemical product; and

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<sup>7</sup> *Health (Drugs and Poisons) Regulation 1996*, section 18 (How chief executive may deal with applications)

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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- (b) either—
    - (i) an approved label for containers for the product contains an instruction; or
    - (ii) a permit for the product that applies to the person is subject to a condition under the Agvet Code;<sup>8</sup> and
  - (c) the instruction or condition requires the person to make a stated record of the use of the product.
- (2) The person must make the record—
- (a) if the instruction or condition states a day by which the record must be made—on or before the stated day; or
  - (b) if paragraph (a) does not apply—as soon as practicable after the chemical product is used.

Maximum penalty for subsection (2)—40 penalty units.

#### **14 Obligation to keep record**

A person who makes a record under section 13 must keep it for at least 2 years after the use to which the record relates, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

## **PART 5—HORMONAL GROWTH PROMOTANTS**

### *Division 1—Preliminary*

#### **15 Definitions for pt 5**

In this part—

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<sup>8</sup> For the instructions or conditions, see the Agvet Code, sections 14 (Grant or refusal of application), 23 (Conditions of approval or registration), 114 (Issue of permit) and 116 (Effect of permit).

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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**“agent”** means a person who is licensed under the *Auctioneers and Agents Act 1971* as an auctioneer or a real estate agent whose licence authorises the person to sell cattle.

**“agent’s statement”** see section 19(2).

**“cattle”** includes bull, calf, cow, heifer, ox and steer.

**“head”** means a head of cattle.

**“HGP”** means hormonal growth promotant.

**“HGP free declaration”** see section 22(2).

**“HGP free tag”** means a HGP free tag in the approved form under the *Stock Act 1915*.

**“HGP treatment”** means implanting a HGP into an animal.

**“HGP treatment record”** see section 17(1).

**“hormonal growth promotant”** means a product that—

- (a) contains an anabolic substance or a hormone; and

*Examples of ‘an anabolic substance or a hormone’—*

- 17 beta oestradiol
- oestradiol benzoate
- progesterone
- testosterone propionate
- trenbolone acetate
- zeranol.

- (b) is used to promote the growth of bovines or bubalines.

**“sell”** includes any of the following—

- (a) supply under an agreement, promise, scheme, transaction (with or without consideration), understanding or undertaking (whether express or implied);
- (b) agree, attempt or offer or agree to sell or supply;
- (c) possess for sale or supply;
- (d) invite or treat or expose for sale or supply;
- (e) cause or permit to be sold or supplied.

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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***Division 2—Obligations if HGP treatment given***

**16 Obligation to make required earmark**

(1) A person must, when giving HGP treatment, permanently mark the animal treated by piercing its right ear with the required earmark so as to leave a space of any size on all sides within the margin of the ear.

Maximum penalty—40 penalty units.

(2) In this section—

**“required earmark”** means—

- (a) for cattle—a mark or cut upon the ear of the head that is approved under the *Brands Act 1915* for the identification of cattle treated with a HGP; or
- (b) for another animal—a mark of an equal sided triangle with sides of 20 mm.

**17 Obligation to record HGP treatment**

(1) A person who has given HGP treatment to an animal must make a written record (a **“HGP treatment record”**) as required by this section—

- (a) identifying the animal treated; and
- (b) stating the following—
  - (i) the HGP with which the animal was treated;
  - (ii) the day the treatment was given (the **“treatment day”**);
  - (iii) any HGP acquired for the treatment that was not used and was disposed of;
  - (iv) the day of the disposal (the **“disposal day”**).

Maximum penalty—40 penalty units.

(2) For subsection (1)(a), the animal may be identified by reference to its sex and breed.

(3) For subsection (1)(b)(i), the HGP may be stated by giving a distinguishing number for, or particulars to identify, the chemical product that contained the HGP.

(4) The information must be entered in the HGP treatment record—

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

---

- (a) for information mentioned in subsection (1)(a) and (b)(i) and (ii)—before the treatment day ends; or
- (b) for information mentioned in subsection (1)(b)(iii) and (iv)—before the disposal day ends.

(5) In this section—

“**disposal**” includes destruction and loss.

### **18 Obligation to keep HGP treatment record**

A person who makes a HGP treatment record must keep it for at least 2 years after the treatment day, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

#### *Division 3—Obligations if cattle with HGP free tag are sold*

##### *Subdivision 1—Saleyard sales by agents*

### **19 Agent’s obligation to give statement**

(1) This section applies if—

- (a) an agent sells a head at a saleyard for someone else; and
- (b) a HGP free tag is attached to the head.

(2) The agent must give a person who buys the head (the “**buyer**”) a written statement (an “**agent’s statement**”)—

- (a) identifying the head; and
- (b) stating that a HGP free tag was attached to the head when it was sold to the buyer.

Maximum penalty—20 penalty units.

(3) For subsection (2)(a), the head may be identified by reference to—

- (a) its sex and breed; or
- (b) a tag number for the animal under the *Stock Act 1915*; or

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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(c) a brand or earmark for the animal under the *Brands Act 1915*.

(4) The agent's statement may be made about more than 1 head.

## **20 Agent's obligation to keep copy of statement**

(1) This section applies to an agent who has given a buyer an agent's statement.

(2) The agent must keep a copy of the statement (the "**agent's copy**") for 2 years after the statement was given, unless the agent has a reasonable excuse.

Maximum penalty—20 penalty units.

(3) If an inspector asks the agent for the agent's copy during the 2 years, the agent must give it to the inspector, unless the agent has a reasonable excuse.

Maximum penalty—20 penalty units.

(4) The inspector may keep the agent's copy to copy it.

(5) However, the inspector must return the agent's copy as soon as practicable after copying it.

## **21 Buyer's obligation to keep and produce statement**

(1) This section applies if a person has been given an agent's statement as a buyer.

(2) The person must keep the agent's statement for 2 years after it was given, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

(3) If an inspector asks the person for the agent's statement during the 2 years, the person must give it to the inspector, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

(4) The inspector may keep the agent's statement copy to copy it.

(5) However, the inspector must return the agent's statement as soon as practicable after copying it.

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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***Subdivision 2—Other sales***

**22 Seller’s obligation to give declaration**

(1) This section applies if—

- (a) a person (the “**seller**”) sells a head; and
- (b) the sale is other than by an agent at a saleyard acting for someone else; and
- (c) a HGP free tag is attached to the head.

(2) The person must, if asked by the person to whom the head is sold (the “**buyer**”), give the buyer a written declaration (a “**HGP free declaration**”) as required under section 23 when the seller delivers the head to the buyer.

Maximum penalty for subsection (2)—20 penalty units.

**23 Requirements for declaration**

(1) A HGP free declaration must—

- (a) identify the animal sold; and
- (b) be made by the seller no more than 7 days before the sale; and
- (c) state the seller has not—
  - (i) given any HGP treatment to the head; or
  - (ii) caused or allowed HGP treatment to be given to the head.

(2) For subsection (1)(a), the animal may be identified by reference to—

- (a) its sex and breed; or
- (b) a tag number for the animal under the *Stock Act 1915*; or
- (c) a brand or earmark for the animal under the *Brands Act 1915*.

(3) Also, if the seller bought the head of cattle from someone else (the “**third person**”), the HGP free declaration must state—

- (a) the third person’s name; and
- (b) either—

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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- (i) that a HGP free tag was attached to the head when it was sold to the seller; or
- (ii) that the seller received a HGP free declaration from the third person.

(4) The HGP free declaration may be made about more than 1 head.

#### **24 Seller's obligation to keep copy of declaration**

(1) This section applies if a person has given a buyer a HGP free declaration.

(2) The person must keep a copy of the HGP free declaration (the “**seller's copy**”) for 2 years after the declaration was given, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

(3) If an inspector asks the person for the seller's copy during the 2 years, the person must give it to the inspector, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

(4) The inspector may keep the seller's copy to copy it.

(5) However, the inspector must return the seller's copy as soon as practicable after copying it.

#### **25 Buyer's obligation to keep and produce declaration**

(1) This section applies if a person has been given a HGP free declaration as a buyer.

(2) The person must keep the declaration for 2 years after it was given, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

(3) If an inspector asks the person for the HGP free declaration during the 2 years, the person must give it to the inspector, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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(4) The inspector may keep the HGP free declaration to copy it.

(5) However, the inspector must return the HGP free declaration as soon as practicable after copying it.

## **PART 6—SUPERVISION FEES AND EXPENSES**

### **26 Application of pt 6**

This part applies to a person if—

- (a) the person has been given a direction under the Act that requires or allows a thing to be done; and
- (b) the direction requires the thing be done under an inspector's supervision.<sup>9</sup>

### **27 Hourly fee**

(1) A fee is payable by the person for each hour or part of an hour of the supervision.

(2) If the supervision, or a part of the supervision, was on a business day, the hourly fee for the supervision or part of the supervision is—

- (a) for working hours—\$30.70; or
- (b) for other than working hours—\$46.05.

(3) If the supervision, or a part of the supervision, was on a day other than a business day, the hourly fee for the supervision or part of the supervision is \$61.40.

(4) In this section—

**“supervision”** includes travelling time for the inspector to travel to and from the place of supervision if the travelling time was for the supervision.

**“working hours”** means the inspector's working hours under any relevant industrial instrument under the *Industrial Relations Act 1999*.

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<sup>9</sup> See section 33 (Supervision by inspector) of the Act.

## **28 Overnight absence expenses**

(1) The person must pay the expense for each overnight absence by the inspector for the supervision.

(2) The expense for each overnight absence is the amount that is, or would be, payable under the *Public Service Act 1996* to the inspector as if the inspector is or were a public service officer travelling on official duty.

## **PART 7—MISCELLANEOUS PROVISIONS**

### **29 Approval of forms**

(1) The chief executive may approve forms for use under the Act.

(2) If a form is approved for a purpose, the approved form is the prescribed form for the purpose.

## SCHEDULE 1

### PRESCRIBED AND PROSCRIBED CHEMICALS

sections 2 and 3

<b>Common name</b>	<b>Chemical name or composition</b>
aldrin	a product containing 95% HHDN
BHC (excluding the gamma isomer)	Mixed isomers of 1,2,3,4,5,6-hexachlorocyclohexane excluding gamma-1, 2,3,4,5,6-hexachlorocyclohexane
chlordane	1,2,4,5,6,7,8,8-octachloro-3a,4,7,7a-tetrahydro-4,7-methanoindane
DDT	Mixed isomers of 1,1,1-trichloro-2,2-bis(chlorophenyl)ethane in which <i>pp'</i> -DDT, 1,1,1-trichloro-2,2-bis(4-chlorophenyl)= ethane, predominates
dieldrin	a product containing 85% HEOD
endrin	1,2,3,4,10,10-hexachloro-6,7-epoxy-1,4,4a,5,6,7,8,8a-octahydro- <i>exo</i> -1,4- <i>exo</i> -5,8-dimethanonaphthalene
HCB	hexachlorobenzene
HEOD	1,2,3,4,10,10-hexachloro-6,7-epoxy-1,4,4a,5,6,7,8,8a-octahydro- <i>endo</i> -1,4- <i>exo</i> -5,8-dimethanonaphthalene
heptachlor	1,4,5,6,7,8,8-heptachloro-3a,4,7,7a-tetrahydro-4,7-methanoindene

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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## SCHEDULE 1 (continued)

<b>Common name</b>	<b>Chemical name or composition</b>
HHDN	1,2,3,4,10,10-hexachloro-1,4,4a,5,8,8a-hexahydro- <i>exo</i> -1,4- <i>endo</i> -5,8-dimethanonaphthalene
TDE	1,1-dichloro-2,2-bis(4-chlorophenyl)ethane

## SCHEDULE 2

### MRLS FOR CHEMICALS FOR ANIMAL FOOD

section 8

#### PART 1—MRLS FOR PRESCRIBED CHEMICALS

<b>Common name</b>	<b>Level (in mg/kg)</b>
aldrin, dieldrin or any total combination of aldrin and dieldrin . . . . .	0.01
BHC (excluding the gamma isomer) . . . . .	0.02
chlordane . . . . .	0.01
DDT . . . . .	0.1
endrin . . . . .	0.03
HCB . . . . .	0.01
heptachlor . . . . .	0.02

#### PART 2—MRLS FOR CHEMICAL PRODUCTS

<b>Chemical product</b>	<b>Level (in mg/kg)</b>
alloydim-sodium . . . . .	0.2
benfluralin . . . . .	0.02

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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## SCHEDULE 2 (continued)

bensulfuron-methyl . . . . .	0.05
bioresmethrin . . . . .	5
bitertanol . . . . .	0.1
carbaryl . . . . .	20
carbofuran . . . . .	2
chlorpyrifos-methyl . . . . .	20
chlorsulfuron . . . . .	10
clopyralid . . . . .	100
cyhalothrin . . . . .	0.01
dichlorvos . . . . .	20
dithiocarbamates (except propineb) . . . . .	30
endosulfan . . . . .	0.3
ethephon . . . . .	10
fenamiphos . . . . .	1
fenitrothion . . . . .	20
fenvalerate . . . . .	10
fluroxypyr . . . . .	25
glyphosate . . . . .	0.3
haloxyfop . . . . .	3
inorganic bromide . . . . .	125
iprodione . . . . .	5
lindane (gamma BHC) . . . . .	0.1
maldison . . . . .	100

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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## SCHEDULE 2 (continued)

methoxychlor . . . . .	1
methyl bromide. . . . .	50
metolachlor. . . . .	5
metribuzin. . . . .	0.2
metsulfuron-methyl . . . . .	0.05
monocrotophos . . . . .	0.2
pirimiphos-methyl . . . . .	20
sethoxydim . . . . .	2
thiodicarb . . . . .	30
tralkoxydim . . . . .	0.02
triadimefon . . . . .	10
triasulfuron . . . . .	5

## ENDNOTES

### 1 Index to endnotes

		Page
2	Date to which amendments incorporated . . . . .	25
3	Key . . . . .	25
4	Table of reprints . . . . .	26
5	List of legislation . . . . .	26
6	List of annotations . . . . .	27

### 2 Date to which amendments incorporated

This is the reprint date mentioned in the Reprints Act 1992, section 5(c). Accordingly, this reprint includes all amendments that commenced operation on or before 14 December 2003. Future amendments of the Chemical Usage (Agricultural and Veterinary) Control Regulation 1999 may be made in accordance with this reprint under the Reprints Act 1992, section 49.

### 3 Key

#### Key to abbreviations in list of legislation and annotations

Key	Explanation	Key	Explanation
AIA	= Acts Interpretation Act 1954	(prev)	= previously
amd	= amended	proc	= proclamation
amdt	= amendment	prov	= provision
ch	= chapter	pt	= part
def	= definition	pubd	= published
div	= division	R[X]	= Reprint No.[X]
exp	= expires/expired	RA	= Reprints Act 1992
gaz	= gazette	reloc	= relocated
hdg	= heading	renum	= renumbered
ins	= inserted	rep	= repealed
lap	= lapsed	(retro)	= retrospectively
notfd	= notified	rv	= revised edition
o in c	= order in council	s	= section
om	= omitted	sch	= schedule
orig	= original	sdiv	= subdivision
p	= page	SIA	= Statutory Instruments Act 1992
para	= paragraph	SIR	= Statutory Instruments Regulation 2002
prec	= preceding	SL	= subordinate legislation
pres	= present	sub	= substituted
prev	= previous	unnum	= unnumbered

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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## 4 Table of reprints

Reprints are issued for both future and past effective dates. For the most up-to-date table of reprints, see the reprint with the latest effective date.

If a reprint number includes a letter of the alphabet, the reprint was released in unauthorised, electronic form only.

### TABLE OF REPRINTS

Reprint No.	Amendments included	Effective	Reprint date
1	none	27 August 1999	2 September 1999
1A	to 2000 SL No. 169	1 July 2000	3 July 2000
1B	to 2001 SL No. 177	28 September 2001	12 October 2001 (Column discontinued) Notes
1C	to 2002 SL No. 351	13 December 2002	
1D	to 2003 SL No. 201	20 September 2003	
1E	to 2003 SL No. 239	4 November 2003	
1F	to 2003 SL No. 335	14 December 2003	

## 5 List of legislation

### **Chemical Usage (Agricultural and Veterinary) Control Regulation 1999 SL No. 203**

made by the Governor in Council on 26 August 1999

notfd gaz 27 August 1999 pp 2224–7

commenced on date of notification

exp 1 September 2009 (see SIA s 54)

Note—The expiry date may have changed since this reprint was published. See the latest reprint of the SIR for any change.

amending legislation—

### **Primary Industries Legislation Amendment Regulation (No. 1) 2000 SL No. 169 pts 1, 4**

notfd gaz 30 June 2000 pp 736–48

ss 1–2 commenced on date of notification

remaining provisions commenced 1 July 2000 (see s 2)

### **Primary Industries Legislation Amendment Regulation (No. 1) 2001 SL No. 177 pts 1, 5**

notfd gaz 28 September 2001 pp 328–30

commenced on date of notification

### **Primary Industries Legislation Amendment Regulation (No. 2) 2002 SL No. 351 pts 1, 5**

notfd gaz 13 December 2002 pp 1266–69

commenced on date of notification

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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**Pest Management Regulation 2003 SL No. 201 ss 1, 2(3), 33 sch 1**

notfd gaz 5 September 2003 pp 57–8

ss 1–2 commenced on date of notification

remaining provisions commenced 20 September 2003 (see s 2(3))

**Primary Industries Legislation Amendment Regulation (No. 1) 2003 SL No. 239  
pts 1, 4**

notfd gaz 3 October 2003 pp 382–5

ss 1–2 commenced on date of notification

remaining provisions commenced 4 November 2003 (see s 2)

**Primary Industries Legislation Amendment Regulation (No. 2) 2003 SL No. 335  
pts 1, 4**

notfd gaz 12 December 2003 pp 1203–7

ss 1–2 commenced on date of notification

remaining provisions commenced 14 December 2003 (see s 2)

## 6 List of annotations

**Proscribed chemicals—Act, s 11C(2)**

**hdg** amd 2003 SL No. 335 s 20(1)

**s 3** amd 2003 SL No. 335 s 20(2)–(3)

**Purpose of pt 3**

**s 4** amd 2003 SL No. 335 s 21

**Definitions for pt 3**

**s 5** amd 2003 SL No. 335 s 22

**Definitions for div 1**

**s 9** def “**prescribed qualification**” ins 2003 SL No. 335 s 23

def “**registered training organisation**” ins 2003 SL No. 335 s 23

**Restricted chemical products containing bifenthrin or chlorpyrifos**

**s 10** amd 2003 SL No. 201 s 33 sch 1

**Restricted chemical products containing endosulfan**

**s 11** amd 2003 SL No. 335 s 24

**Restricted chemical products containing pindone**

**s 11A** ins 2003 SL No. 335 s 25

**Other restricted chemical products**

**s 12** amd 2003 SL No. 335 s 26

**Hourly fee**

**s 27** amd 2000 SL No. 169 s 8; 2001 SL No. 177 s 9; 2002 SL No. 351 s 10; 2003 SL No. 239 s 8

*Chemical Usage (Agricultural and Veterinary)  
Control Regulation 1999*

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**PART 8—REPEAL AND TRANSITIONAL PROVISIONS**

**pt 8 (ss 30–34)** exp 28 August 1999 (see s 34)