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# Gazette

## Agricultural and veterinary chemicals

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Veterinary Medicines Authority**

The *Agricultural and Veterinary Chemical Code Act 1994* (the Act) commenced on 15 March 1995. The Agricultural and Veterinary Chemicals Code (the Agvet Code) scheduled to the Act requires notices to be published in the *Gazette* containing details of the registration of agricultural and veterinary chemical products and other approvals granted by the Australian Pesticides and Veterinary Medicines Authority. The Agvet Code and related legislation also requires certain other notices to be published in the *Gazette*. A reference to Agvet Codes in this publication is a reference to the Agvet Code in each state and territory jurisdiction.

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## **General information**

The APVMA Gazette is published fortnightly and contains details of the registration of agricultural and veterinary chemicals products and other approvals granted by the APVMA, notices as required by the Agricultural and Veterinary Chemicals Code (the Agvet Code) and related legislation and a range of regulatory material issued by the APVMA.

Pursuant to section 8J(1) of the Agvet Code, the APVMA has decided that it is unnecessary to publish details of applications made for the purpose of notifying minor variations to registration details. The APVMA will however report notifications activity in quarterly statistical reports.

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## Agricultural chemical products and approved labels

Pursuant to the Agricultural and Veterinary Chemicals Code scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994*, the APVMA hereby gives notice that it has registered or varied the relevant particulars or conditions of the registration in respect of the following products and has approved the label or varied the relevant particulars or conditions of the approval in respect of the containers for the chemical product, with effect from the dates shown.

**Table 1: Agricultural products based on existing active constituents**

<b>Application no.</b>	150599
<b>Product name</b>	QA MCPA + Picloram SL Herbicide
<b>Active constituents</b>	420 g/L MCPA present as potassium salt, 26 g/L picloram present as potassium salt
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant ACN</b>	680 792 625
<b>Date of registration</b>	2 February 2026
<b>Product registration no.</b>	96758
<b>Label approval no.</b>	96758/150599
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 420 g/L MCPA present as potassium salt, and 26 g/L picloram present as potassium salt, soluble I concentrate (SL) product for the control of climbing buckwheat, common sowthistle, skeleton weed, capeweed, doublegee and other broadleaf weeds in winter cereals and linseed crops

<b>Application no.</b>	145164
<b>Product name</b>	Kilblat Cockroach Killer Spray
<b>Active constituents</b>	2.76 g/kg cyphenothrin, 0.94 g/kg D-tetramethrin
<b>Applicant name</b>	Callington Haven Pty Ltd
<b>Applicant ACN</b>	000 632 404
<b>Date of registration</b>	3 February 2026
<b>Product registration no.</b>	95274
<b>Label approval no.</b>	95274/145164
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 2.76 g/kg cyphenothrin and 0.94 g/kg D-tetramethrin product for control of household crawling insects

<b>Application no.</b>	150620
<b>Product name</b>	Quantum Prosulfocarb 800 EC Herbicide
<b>Active constituent</b>	800 g/L prosulfocarb
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant CAN</b>	680 792 625
<b>Date of registration</b>	3 February 2026
<b>Product registration no.</b>	96764
<b>Label approval no.</b>	96764/150620
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of an 800 g/L prosulfocarb product, formulated as an emulsifiable concentrate (EC) product for the control of annual ryegrass ( <i>Lolium rigidum</i> ) and other grass and broadleaf weeds in barley and wheat as specified in the directions for use table

<b>Application no.</b>	150173
<b>Product name</b>	NovaGuard Pyroxasulfone 850 WG Herbicide
<b>Active constituent</b>	850 g/kg pyroxasulfone
<b>Applicant name</b>	Novaguard Pty Ltd
<b>Applicant ACN</b>	153 121 156
<b>Date of registration</b>	3 February 2026
<b>Product registration no.</b>	96636
<b>Label approval no.</b>	96636/150173
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of an 850 g/kg pyroxasulfone product, formulated as a water dispersible granule (WG) for pre-emergence control of annual ryegrass, barley grass, annual phalaris, silver grass and toad rush and suppression of certain grass weeds in wheat (not durum wheat), triticale and certain winter legume crops

<b>Application no.</b>	148130
<b>Product name</b>	Mozziedot Mosquito Repellent Patch
<b>Active constituent</b>	120 mg/patch citronella oil
<b>Applicant name</b>	Cubitt, Andrea Janine
<b>Applicant ACN</b>	30 923 507 053
<b>Date of registration</b>	4 February 2026
<b>Product registration no.</b>	96083
<b>Label approval no.</b>	96083/148130
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 120 mg/patch citronella oil, formulated as a solid (SO) for use as a household personal insect repellent

<b>Application no.</b>	150174
<b>Product name</b>	EasyFarm Pyroxasulfone 850 WG Herbicide
<b>Active constituent</b>	850 g/kg pyroxasulfone
<b>Applicant name</b>	Novaguard Pty Ltd
<b>Applicant ACN</b>	153 121 156
<b>Date of registration</b>	4 February 2026
<b>Product registration no.</b>	96637
<b>Label approval no.</b>	96637/150174
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 850 g/kg pyroxasulfone product, formulated as a water dispersible granule (WG) for pre-emergence control of annual ryegrass, barley grass, annual phalaris, silver grass and toad rush and suppression of certain grass weeds in wheat (not durum wheat), triticale and certain winter legume crops

<b>Application no.</b>	150597
<b>Product name</b>	QA Hexazinone 250 SL Herbicide
<b>Active constituent</b>	250 g/L hexazinone
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant ACN</b>	680 792 625
<b>Date of registration</b>	6 February 2026
<b>Product registration no.</b>	96756
<b>Label approval no.</b>	96756/150597
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 250 g/L hexazinone product, formulated as a soluble concentrate (SL) for control of certain broadleaf weeds, perennial and annual grasses, woody weeds in pinus radiata plantations, pasture situations and commercial and industrial areas and rights-of-way as per the directions for use

<b>Application no.</b>	150660
<b>Product name</b>	AC Ringstar 920 Herbicide
<b>Active constituents</b>	800 g/L prosulfocarb, 120 g/L S-metolachlor
<b>Applicant name</b>	Axichem Pty Ltd
<b>Applicant ACN</b>	131 628 594
<b>Date of registration</b>	6 February 2026
<b>Product registration no.</b>	96783
<b>Label approval no.</b>	96783/150660
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of an 800 g/L prosulfocarb and 120 g/L S-metolachlor product, formulated as an emulsifiable concentrate (EC) for control of annual ryegrass ( <i>Lolium rigidum</i> ) and other grass and broadleaf weeds in barley, chickpeas, faba beans, field peas, lentils, lupins, potatoes and wheat

<b>Application no.</b>	150598
<b>Product name</b>	QA Effect Wetter Surfactant
<b>Active constituent</b>	1040 g/L octyl phenol ethoxylate
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant ACN</b>	680 792 625
<b>Date of registration</b>	9 February 2026
<b>Product registration no.</b>	96757
<b>Label approval no.</b>	96757/150598
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 1040 g/L octyl phenol ethoxylate product, formulated as a suspension concentrate (SC) for improvement of spray coverage when using agricultural chemicals

<b>Application no.</b>	148160
<b>Product name</b>	Kenso Agcare Amizon Herbicide
<b>Active constituents</b>	300 g/L triclopyr present as the butoxyethyl ester, 100 g/L picloram present as the hexyloxypropylamine salt, 8 g/L aminopyralid present as hexyloxypropylamine salt
<b>Applicant name</b>	Kenso Corporation (M) Sdn. Bhd.
<b>Applicant ACN</b>	N/A
<b>Date of registration</b>	9 February 2026
<b>Product registration no.</b>	96097
<b>Label approval no.</b>	96097/148160
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 300 g/L triclopyr present as butoxyethyl ester, 100 g/L picloram present as hexyloxypropylamine salt and 8 g/L aminopyralid present as hexyloxypropylamine salt in emulsifiable concentrate product for control of a range of environmental and noxious woody and herbaceous weeds

<b>Application no.</b>	150579
<b>Product name</b>	Swan Glufosinate 200 Herbicide
<b>Active constituent</b>	200 g/L glufosinate-ammonium
<b>Applicant name</b>	Swan Chemical Holdings Pty Ltd
<b>Applicant ACN</b>	669 863 067
<b>Date of registration</b>	10 February 2026
<b>Product registration no.</b>	96748
<b>Label approval no.</b>	96748/150579
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 200 g/L glufosinate-ammonium product, formulated as a soluble concentrate (SL) for the non-residual control of broadleaf and grass weeds in various situations

<b>Application no.</b>	150619
<b>Product name</b>	QA Flumioxazin 500 WG Herbicide
<b>Active constituent</b>	500 g/kg flumioxazin
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant ACN</b>	680 792 625
<b>Date of registration</b>	11 February 2026
<b>Product registration no.</b>	96763
<b>Label approval no.</b>	96763/150619
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 500 g/kg flumioxazin product, formulated as a water dispersible granule (WG) for the control of fungal diseases on macadamia nuts, chickpeas, faba beans, lentils and vetch

<b>Application no.</b>	150594
<b>Product name</b>	QA MCPA + Picolinafen EC Herbicide
<b>Active constituents</b>	500 g/L MCPA present as the ethyl hexyl ester, 50 g/L picolinafen
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant CAN</b>	680 792 625
<b>Date of registration</b>	11 February 2026
<b>Product registration no.</b>	96753
<b>Label approval no.</b>	96753/150594
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 500 g/L MCPA present as the ethyl hexyl ester 50 g/L picolinafen emulsifiable concentrate (EC) product for the control of certain broadleaf weeds in winter cereals

<b>Application no.</b>	150617
<b>Product name</b>	QA Propyzamide 500 SC Herbicide
<b>Active constituent</b>	500 g/L propyzamide
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant ACN</b>	680 792 625
<b>Date of registration</b>	11 February 2026
<b>Product registration no.</b>	96762
<b>Label approval no.</b>	96762/150617
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 500 g/L propyzamide product, formulated as a suspension concentrate (SC) for selective control of certain grasses and broadleaf weeds in lettuce, sports turf, home lawns, legumes, seed crops and pastures, as per the direction for use

<b>Application no.</b>	144497
<b>Product name</b>	NAVADET Insecticide
<b>Active constituents</b>	45 g/L chlorantraniliprole, 18 g/L abamectin
<b>Applicant name</b>	United Cropscience PTE., Ltd.
<b>Applicant ACN</b>	N/A
<b>Date of registration</b>	11 February 2026
<b>Product registration no.</b>	95028
<b>Label approval no.</b>	95028/144497
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration and label approval of an 18 g/L abamectin, 45 g/L chlorantraniliprole suspension concentrate product for control of certain mites in pome fruit

<b>Application no.</b>	148658
<b>Product name</b>	NovaGuard Pyriproxyfen 100 Insect Growth Regulator
<b>Active constituent</b>	100 g/L pyriproxyfen
<b>Applicant name</b>	Novaguard Pty Ltd
<b>Applicant ACN</b>	153 121 156
<b>Date of registration</b>	12 February 2026
<b>Product registration no.</b>	96250
<b>Label approval no.</b>	96250/148658
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of 100 g/L pyriproxyfen emulsifiable concentrate product for the control of silverleaf whitefly ( <i>Bemisia tabaci</i> Biotype B) in cotton, rockmelon and capsicum, the control of silverleaf whitefly ( <i>Bemisia tabaci</i> Biotype B) and greenhouse whitefly in tomatoes, and the control of various scale in citrus, mangoes and olives

<b>Application no.</b>	150621
<b>Product name</b>	QA Wetter 600
<b>Active constituent</b>	600 g/L nonyl phenol ethylene oxide condensate
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant ACN</b>	680 792 625
<b>Date of registration</b>	13 February 2026
<b>Product registration no.</b>	96765
<b>Label approval no.</b>	96765/150621
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 600 g/L nonyl phenol ethylene oxide condensate for use with insecticides, fungicides and herbicides

<b>Application no.</b>	149644
<b>Product name</b>	QA Chlorpyrifos 500 EC Insecticide
<b>Active constituent</b>	500 g/L chlorpyrifos (an anti-cholinesterase compound)
<b>Applicant name</b>	Quantum Agrosiences Holdings Pty Ltd
<b>Applicant ACN</b>	680 792 625
<b>Date of registration</b>	13 February 2026
<b>Product registration no.</b>	96520
<b>Label approval no.</b>	96520/149644
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 500 g/L chlorpyrifos product, formulated as an emulsifiable concentrate (EC) for the control of certain insect pests in various situations as per the directions for use

<b>Application no.</b>	150304
<b>Product name</b>	ACP Interact 33-15 Herbicide
<b>Active constituents</b>	33 g/L imazamox as the ammonium salt, 15 g/L imazapyr present as the ammonium salt
<b>Applicant name</b>	Australis Crop Protection Pty Ltd
<b>Applicant ACN</b>	150 711 185
<b>Date of registration</b>	13 February 2026
<b>Product registration no.</b>	96656
<b>Label approval no.</b>	96656/150304
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 33 g/L imazamox as the ammonium salt and 15 g/L imazapyr as the ammonium salt, soluble concentrate product for the early post-emergence control of certain annual grass and broadleaf weeds in imidazolinone-tolerant wheat, barley, and canola

**Table 2: Variations of registration – agricultural chemical products**

<b>Application no.</b>	150568
<b>Product name</b>	Deltanox Duo 10SC Insecticide
<b>Active constituents</b>	80 g/L piperonyl butoxide, 10 g/L deltamethrin, 10 g/L D-tetramethrin (20:80)
<b>Applicant name</b>	Australian Agribusiness (Holdings) Pty Ltd
<b>Applicant ACN</b>	135 355 958
<b>Date of variation</b>	14 October 2025
<b>Product registration no.</b>	68900
<b>Label approval no.</b>	68900/150568
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to the particulars of registration and label approval to change the distinguishing product name and the name that appears on the label from 'Pest Controller's Own Deltamethrin Tetramethrin 10 SC Insecticide' to 'Deltanox Duo 10SC Insecticide'

<b>Application no.</b>	151671
<b>Product name</b>	TITAN Glufosinate 400 Herbicide
<b>Active constituent</b>	400 g/L glufosinate-ammonium
<b>Applicant name</b>	Titan Ag Pty Ltd
<b>Applicant ACN</b>	122 081 574
<b>Date of variation</b>	20 January 2026
<b>Product registration no.</b>	83887
<b>Label approval no.</b>	83887/151671
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to the particulars of registration and label approval to change the distinguishing product name and the name that appears on the label from 'Eureka! Patented Glufosinate 400 Herbicide' to 'TITAN Glufosinate 400 Herbicide'

<b>Application no.</b>	151754
<b>Product name</b>	AC Hussle Spray Adjuvant
<b>Active constituents</b>	700 g/L ethyl and methyl esters of vegetable oil, 116 g/L nonionic surfactants
<b>Applicant name</b>	Axichem Pty Ltd
<b>Applicant ACN</b>	131 628 594
<b>Date of variation</b>	29 January 2026
<b>Product registration no.</b>	69449
<b>Label approval no.</b>	69449/151754
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to the particulars of registration and label approval to change the distinguishing product name and the name that appears on the label from 'AC Hassle Spray Adjuvant' to 'AC Hussle Spray Adjuvant'

<b>Application no.</b>	150484
<b>Product name</b>	Bollgard 3
<b>Active constituents</b>	Bacillus thuringiensis subsp. kurstaki delta endotoxins as produced by the Cry1Ac and Cry2Ab genes and their controlling sequences, bacillus thuringiensis strain AB88 exotoxin as produced by the Vip3A(a) gene and its controlling sequence
<b>Applicant name</b>	Monsanto Australia Pty Ltd
<b>Applicant ACN</b>	006 725 560
<b>Date of variation</b>	2 February 2026
<b>Product registration no.</b>	69656
<b>Label approval no.</b>	69656/106072
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to registration particulars to amend the conditions of registration associated with the resistance management plan (RMP) for northern Australia

<b>Application no.</b>	150665
<b>Product name</b>	Yates Lawn Disease Fungicide Concentrate
<b>Active constituent</b>	125 g/L fludioxonil
<b>Applicant name</b>	Syngenta Australia Pty Ltd
<b>Applicant ACN</b>	002 933 717
<b>Date of variation</b>	3 February 2026
<b>Product registration no.</b>	94228
<b>Label approval no.</b>	94228/150665
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to registration particulars and particulars of label to change product name from 'Munns Professional Lawn Fungicide Concentrate' to 'Yates Lawn Disease Fungicide Concentrate' and to update the statement of claims

<b>Application no.</b>	148206
<b>Product name</b>	BUSHMAN Repellent PERSONAL INSECT REPELLENT
<b>Active constituent</b>	400 g/kg diethyltoluamide
<b>Applicant name</b>	Juno Limited
<b>Applicant ACN</b>	N/A
<b>Date of variation</b>	9 February 2026
<b>Product registration no.</b>	60969
<b>Label approval no.</b>	60969/148206
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to particulars of label, to extend use claims on nuisance flies

<b>Application no.</b>	150622
<b>Product name</b>	Escalate 500 SC Cotton Defoliant
<b>Active constituent</b>	500 g/L thidiazuron
<b>Applicant name</b>	ADAMA Australia Pty Limited
<b>Applicant ACN</b>	050 328 973
<b>Date of variation</b>	10 February 2026
<b>Product registration no.</b>	55126
<b>Label approval no.</b>	55126/150622
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to registration and label particulars of the product 'Escalate 500 SC Cotton Defoliant' (500 g/L thidiazuron) to amend the adjuvant in the directions for use table

<b>Application no.</b>	149962
<b>Product name</b>	Genfarm Panzer 540 Herbicide
<b>Active constituent</b>	540 g/L glyphosate (present as the potassium salt)
<b>Applicant name</b>	Nutrien Ag Solutions Limited
<b>Applicant ACN</b>	008 743 217
<b>Date of variation</b>	11 February 2026
<b>Product registration no.</b>	89795
<b>Label approval no.</b>	89795/149962
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to particulars of registration and label approval to add an alternative soluble concentrate, 540 glyphosate (as the potassium salt) formulation for non-selective herbicide control of many annual and perennial weeds and to decrease the pack size range

**Table 3: Restricted product variation – agricultural chemical products**

<b>Application no.</b>	141062
<b>Product name</b>	Eradicat 1080 Feral Cat Bait
<b>Active constituent</b>	0.18 g/kg sodium fluoroacetate / 4.5 mg sodium fluoroacetate (1080) per bait
<b>Applicant name</b>	Department of Biodiversity, Conservation and Attractions
<b>Applicant ACN</b>	38 052 249 024
<b>Date of variation</b>	9 February 2026
<b>Product registration no.</b>	65272
<b>Label approval no.</b>	65272/141062
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to registration particulars, and particulars of label, to extend use to SA, NT and Qld and set the rate for all use patterns at 50 baits per square km

## Veterinary chemical products and approved labels

Pursuant to the Agricultural and Veterinary Chemicals Code scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994*, the APVMA hereby gives notice that it has registered or varied the relevant particulars or conditions of the registration in respect of the following products and has approved the label or varied the relevant particulars or conditions of the approval in respect of the containers for the chemical product, with effect from the dates shown.

**Table 4: Veterinary products based on existing active constituents**

<b>Application no.</b>	140896
<b>Product name</b>	Varenzin Oral Suspension for Cats
<b>Active constituent</b>	25 mg/mL molidustat sodium
<b>Applicant name</b>	Elanco Australasia Pty Ltd
<b>Applicant ACN</b>	076 745 198
<b>Date of registration</b>	13 February 2026
<b>Product registration no.</b>	93914
<b>Label approval no.</b>	93914/140896
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration and approval of the label for 'Varenzin Oral Suspension for Cats' containing 25 mg/mL of molidustat sodium, as an aid in the control of nonregenerative anaemia associated with chronic kidney disease (CKD) in cats

<b>Application no.</b>	144720
<b>Product name</b>	Abatak Dual Combination Drench for Sheep
<b>Active constituents</b>	50 g/L closantel, 1 g/L abamectin
<b>Applicant name</b>	Abbey Laboratories Pty Ltd
<b>Applicant ACN</b>	156 000 430
<b>Date of registration</b>	3 February 2026
<b>Product registration no.</b>	95111
<b>Label approval no.</b>	95111/144720
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of Abatak Dual Combination Drench for Sheep containing 1 g/L abamectin and 50 g/L closantel for the control and treatment of roundworms, nasal bot, itch mite and liver fluke in sheep

<b>Application no.</b>	150797
<b>Product name</b>	Owtbac Moxidone Long Acting Injection for Cattle
<b>Active constituents</b>	100 mg/mL moxidectin
<b>Applicant name</b>	The United Animal Healthcare (Inner Mongolia) Co Ltd
<b>Applicant ACN</b>	N/A
<b>Date of registration</b>	9 February 2026
<b>Product registration no.</b>	96833
<b>Label approval no.</b>	96833/150797
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration of a 100 mg/mL moxidectin injectable product and label for the treatment and control of moxidectin sensitive internal and external parasites in cattle

<b>Application no.</b>	143998
<b>Product name</b>	Advocate Ultra Chew for Dogs 22.1 – 45 kg
<b>Active constituents</b>	Each chew contains 900 mg lotilaner, 228 mg praziquantel, 228 mg pyrantel (as embonate), 0.9 mg moxidectin
<b>Applicant name</b>	Elanco Australasia Pty Ltd
<b>Applicant ACN</b>	076 745 198
<b>Date of registration</b>	16 February 2026
<b>Product registration no.</b>	94905
<b>Label approval no.</b>	94905/143998
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration and approval of the label of an oral chew containing 900 mg lotilaner, 228 mg pyrantel (as embonate), 228 mg praziquantel, and 0.9 mg moxidectin for the treatment and control of tick species, treatment and prevention of fleas, prevention of heartworm, treatment and control of intestinal worms

<b>Application no.</b>	143999
<b>Product name</b>	Advocate Ultra Chew for Dogs 5.6 – 11 kg
<b>Active constituents</b>	Each chew contains 225 mg lotilaner, 57 mg praziquantel, 57 mg pyrantel (as embonate), 0.225 mg moxidectin
<b>Applicant name</b>	Elanco Australasia Pty Ltd
<b>Applicant ACN</b>	076 745 198
<b>Date of registration</b>	16 February 2026
<b>Product registration no.</b>	94906
<b>Label approval no.</b>	94906/143999
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration and approval of the label of an oral chew containing 225 mg lotilaner, 57 mg pyrantel (as embonate), 57 mg praziquantel, and 0.225 mg moxidectin for the treatment and control of tick species, treatment and prevention of fleas, prevention of heartworm, treatment and control of intestinal worms

<b>Application no.</b>	144000
<b>Product name</b>	Advocate Ultra Chew for Dogs 11.1 – 22 kg
<b>Active constituents</b>	Each chew contains 450 mg lotilaner, 114 mg praziquantel, 114 mg pyrantel (as embonate), 0.45 mg moxidectin
<b>Applicant name</b>	Elanco Australasia Pty Ltd
<b>Applicant ACN</b>	076 745 198
<b>Date of registration</b>	16 February 2026
<b>Product registration no.</b>	94907
<b>Label approval no.</b>	94907/144000
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration and approval of the label of an oral chew containing 450 mg lotilaner, 114 mg pyrantel (as embonate), 114 mg praziquantel, and 0.45 mg moxidectin for the treatment and control of tick species, treatment and prevention of fleas, prevention of heartworm, treatment and control of intestinal worms

<b>Application no.</b>	144001
<b>Product name</b>	Advocate Ultra Chew for Dogs 2.9 – 5.5 kg
<b>Active constituents</b>	Each chew contains 112.5 mg lotilaner, 28.5 mg praziquantel, 28.5 mg pyrantel (as embonate), 0.11 mg moxidectin
<b>Applicant name</b>	Elanco Australasia Pty Ltd
<b>Applicant ACN</b>	076 745 198
<b>Date of registration</b>	16 February 2026
<b>Product registration no.</b>	94908
<b>Label approval no.</b>	94908/144001
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration and approval of the label of an oral chew containing 112.50 mg lotilaner, 28.5 mg pyrantel (as embonate), 28.50 mg praziquantel, and 0.1125 mg moxidectin for the treatment and control of tick species, treatment and prevention of fleas, prevention of heartworm, treatment and control of intestinal worms

<b>Application no.</b>	144002
<b>Product name</b>	Advocate Ultra Chew for Dogs 2.0 – 2.8 kg
<b>Active constituents</b>	Each chew contains 56.25 mg lotilaner, 14.25 mg praziquantel, 14.25 mg pyrantel (as embonate), 0.056 mg moxidectin
<b>Applicant name</b>	Elanco Australasia Pty Ltd
<b>Applicant ACN</b>	076 745 198
<b>Date of registration</b>	16 February 2026
<b>Product registration no.</b>	94909
<b>Label approval no.</b>	94909/144002
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Registration and approval of the label of an oral chew containing 56.25 mg lotilaner, 14.25 mg pyrantel (as embonate), 14.25 mg praziquantel, and 0.05625 mg moxidectin for the treatment and control of tick species, treatment and prevention of fleas, prevention of heartworm, treatment and control of intestinal worms

Table 5: Variations of registration – veterinary chemical products

Application no.	149288
Product name	Equipel Insect Repellent Conditioner
Active constituent	40 mg/mL permethrin (25:75)
Applicant name	Dermcare-Vet Pty Ltd
Applicant ACN	010 280 010
Date of variation	2 February 2026
Product registration no.	93339
Label approval no.	93339/149288
Description of the application and its purpose, including the intended use of the chemical product	Variation to the relevant particulars of the product and label by changing the instructions of use to increase the maximum dosage and align the label with the current Veterinary Labelling Code

Application no.	150766
Product name	Pexion 400 mg Tablets for Dogs
Active constituent	400 mg/tablet imepitoin
Applicant name	Boehringer Ingelheim Animal Health Australia Pty. Ltd.
Applicant ACN	071 187 285
Date of variation	2 February 2026
Product registration no.	67812
Label approval no.	67812/150766
Description of the application and its purpose, including the intended use of the chemical product	Variation to the relevant particulars of the product and label by changing the instructions of use to align the label with the current Veterinary Labelling Code

Application no.	150493
Product name	Simplseal
Active constituent	650 mg/g bismuth subnitrate
Applicant name	Alleeva Animal Health Ltd
Applicant ACN	N/A
Date of variation	5 February 2026
Product registration no.	91411
Label approval no.	91411/150493
Description of the application and its purpose, including the intended use of the chemical product	Variation of the relevant particulars of the product and label approval to vary the instructions for use and net contents

<b>Application no.</b>	151298
<b>Product name</b>	Jurox Domperidone Paste for Horses
<b>Active constituent</b>	600 mg domperidone
<b>Applicant name</b>	Zoetis Australia Pty Ltd
<b>Applicant ACN</b>	156 476 425
<b>Date of variation</b>	5 February 2026
<b>Product registration no.</b>	54030
<b>Label approval no.</b>	54030/151298
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation of relevant particulars of the chemical product and label approval to update the withholding period and align the label with the current Veterinary Labelling Code

<b>Application no.</b>	151146
<b>Product name</b>	Seresto for Kittens and Cats Fleas and Tick Collar
<b>Active constituents</b>	100 mg/g imidacloprid, 45 mg/g flumethrin
<b>Applicant name</b>	Elanco Australasia Pty Ltd
<b>Applicant ACN</b>	076 745 198
<b>Date of variation</b>	6 February 2026
<b>Product registration no.</b>	80642
<b>Label approval no.</b>	80642/151146
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation of the relevant particulars of product registration and label approval to update the general directions of the label

<b>Application no.</b>	147903
<b>Product name</b>	Ovastim Injection for Sheep
<b>Active constituent</b>	600 mg/L polyandroalbumin
<b>Applicant name</b>	Virbac (Australia) Pty Ltd
<b>Applicant ACN</b>	003 268 871
<b>Date of variation</b>	9 February 2026
<b>Product registration no.</b>	51133
<b>Label approval no.</b>	51133/147903
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation of relevant particulars of the product registration and label to update the general direction statements on the label

<b>Application no.</b>	150585
<b>Product name</b>	Quadseal Teat Sealant for Dry Cows
<b>Active constituent</b>	650 mg/g bismuth subnitrate
<b>Applicant name</b>	Alleva Animal Health Ltd
<b>Applicant ACN</b>	N/A
<b>Date of variation</b>	9 February 2026
<b>Product registration no.</b>	92817
<b>Label approval no.</b>	92817/150585
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation of the relevant particulars of the product and label approval to vary the instructions for use and net contents

<b>Application no.</b>	150682
<b>Product name</b>	Biorelease Deslorelin (BRD)
<b>Active constituent</b>	1.25 mg/mL deslorelin (as deslorelin acetate)
<b>Applicant name</b>	Dechra Veterinary Products (Australia) Pty Ltd
<b>Applicant ACN</b>	614 716 700
<b>Date of variation</b>	11 February 2026
<b>Product registration no.</b>	64329
<b>Label approval no.</b>	64329/150682
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to the relevant particulars of a registered chemical product and label approval to update the label statements in accordance with the current Veterinary Labelling Code

<b>Application no.</b>	149350
<b>Product name</b>	Cortavance Cutaneous Spray Solution for Dogs
<b>Active constituent</b>	0.584 mg/mL hydrocortisone aceponate
<b>Applicant name</b>	Virbac (Australia) Pty Ltd
<b>Applicant ACN</b>	003 268 871
<b>Date of variation</b>	11 February 2026
<b>Product registration no.</b>	61129
<b>Label approval no.</b>	61129/149350
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Variation to the relevant particulars of the product and the label particulars and label approval by extending the product claims and the associated use pattern with the new claim

**Table 6: Label approval – veterinary chemical products**

<b>Application no.</b>	151335
<b>Product name</b>	Promin Plus B12 Injection for Cattle
<b>Active constituents</b>	40 g/L zinc (as disodium zinc EDTA), 15 g/L copper (as disodium copper EDTA), 10 g/L manganese (as disodium manganese EDTA), 5 g/L selenium (as sodium selenite), 1.4 g/L cyanocobalamin
<b>Applicant name</b>	Virbac (Australia) Pty Ltd
<b>Applicant ACN</b>	003 268 871
<b>Date of registration</b>	12 February 2026
<b>Product registration no.</b>	96294
<b>Label approval no.</b>	96294/151335
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Approval of a new label for a registered product Promin plus B12 Injection for Cattle with the new label name 'Nutriject with B12 Injection for Cattle'

<b>Application no.</b>	151341
<b>Product name</b>	Promin Plus B12 Injection for Cattle
<b>Active constituents</b>	40 g/L zinc (as disodium zinc EDTA), 15 g/L copper (as disodium copper EDTA), 10 g/L manganese (as disodium manganese EDTA), 5 g/L selenium (as sodium selenite), 1.4 g/L cyanocobalamin
<b>Applicant name</b>	Virbac (Australia) Pty Ltd
<b>Applicant ACN</b>	003 268 871
<b>Date of registration</b>	12 February 2026
<b>Product registration no.</b>	96294
<b>Label approval no.</b>	96294/151341
<b>Description of the application and its purpose, including the intended use of the chemical product</b>	Approval of a new label for a registered product Promin plus B12 Injection for Cattle with the new label name 'Acti-Trace Injection for Cattle with B12'

## Approved active constituents

Pursuant to the Agricultural and Veterinary Chemicals Code scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994*, the APVMA hereby gives notice that it has approved or varied the relevant particulars or conditions of the approval of the following active constituents, with effect from the dates shown.

**Table 7: Approved active constituents**

<b>Application no.</b>	149956
<b>Active constituent</b>	Pergolide mesylate
<b>Applicant name</b>	Abbey Laboratories Pty Ltd
<b>Applicant ACN</b>	156 000 430
<b>Date of approval</b>	3 February 2026
<b>Approval no.</b>	96585
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Approval of the active constituent pergolide mesylate for use in veterinary chemical products

<b>Application no.</b>	147100
<b>Active constituent</b>	Emodepside
<b>Applicant name</b>	Vetoquinol Australia Pty Ltd
<b>Applicant ACN</b>	006 949 480
<b>Date of approval</b>	4 February 2026
<b>Approval no.</b>	95767
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Approval of the active constituent emodepside for use in veterinary chemical products

<b>Application no.</b>	150073
<b>Active constituent</b>	Omeprazole
<b>Applicant name</b>	Bimeda (Australia) Pty. Limited
<b>Applicant ACN</b>	058 196 508
<b>Date of approval</b>	9 February 2026
<b>Approval no.</b>	96613
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Approval of the active constituent omeprazole for use in veterinary chemical products

<b>Application no.</b>	150473
<b>Active constituent</b>	Bismuth subnitrate
<b>Applicant name</b>	Alleva Animal Health Ltd
<b>Applicant ACN</b>	N/A
<b>Date of approval</b>	9 February 2026
<b>Approval no.</b>	96708
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Approval of the active constituent bismuth subnitrate for use in veterinary chemical products

<b>Application no.</b>	150106
<b>Active constituent</b>	Albendazole
<b>Applicant name</b>	Aarambh Life Science
<b>Applicant ACN</b>	N/A
<b>Date of approval</b>	10 February 2026
<b>Approval no.</b>	96615
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Approval of the active constituent albendazole for use in veterinary chemical products

<b>Application no.</b>	149177
<b>Active constituent</b>	Indaziflam
<b>Applicant name</b>	Ningxia Lamtin Agricultural Development Co., Ltd.
<b>Applicant ACN</b>	N/A
<b>Date of approval</b>	13 February 2026
<b>Approval no.</b>	96397
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Approval of the active constituent indaziflam for use in agricultural chemical products

<b>Application no.</b>	143103
<b>Active constituent</b>	Molidustat sodium
<b>Applicant name</b>	Bayer Cropscience Pty Ltd
<b>Applicant ACN</b>	000 226 022
<b>Date of approval</b>	13 February 2026
<b>Approval no.</b>	94656
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Approval of the active constituent molidustat sodium for use in veterinary chemical products

<b>Application no.</b>	142679
<b>Active constituent</b>	Fluopyram
<b>Applicant name</b>	Hebei Lansheng Biotech Co Ltd
<b>Applicant ACN</b>	N/A
<b>Date of approval</b>	2 February 2026
<b>Approval no.</b>	94533
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Approval of the active constituent fluopyram for use in agricultural chemical products

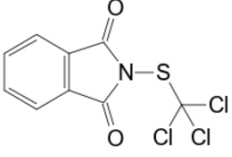
**Table 8: Variations of active constituent**

<b>Application no.</b>	149341
<b>Active constituents</b>	Bacitracin methylene disalicylate
<b>Applicant name</b>	Lifecome Biochemistry Co Ltd
<b>Applicant ACN</b>	N/A
<b>Date of variation</b>	11 February 2026
<b>Approval no.</b>	86765
<b>Description of the application and its purpose, including the intended use of the active constituent</b>	Variation of relevant particulars or conditions of an approved active constituent

## Application for the approval of a new active constituent – Folpet

The APVMA has before it an application for the approval of a new active constituent, folpet, for use in agricultural chemical products.

**Table 9: Particulars of the active constituent**

<b>Common name</b>	Folpet
<b>IUPAC name</b>	2-[(trichloromethyl)sulfanyl]-1 <i>H</i> -isoindole-1,3(2 <i>H</i> )-dione
<b>CAS registry number</b>	133-07-3
<b>Minimum purity</b>	940 g/kg minimum.
<b>Molecular formula</b>	C <sub>9</sub> H <sub>4</sub> Cl <sub>3</sub> NO <sub>2</sub> S
<b>Molecular weight</b>	296.6 g/mol
<b>Structure</b>	
<b>Chemical family</b>	Phthalimides
<b>Mode of action</b>	Folpet's fungicide action relies on its ability to react with thiols, produce the reactive thiophosgene, and potentially induce cytotoxicity and regenerative proliferation, contributing to its broad-spectrum activity and role in disease control.

### Summary of the APVMA's evaluation of folpet active constituent

The APVMA has evaluated the chemistry aspects of folpet (physico-chemical properties, stability, identification, manufacturing process, quality control procedures, batch analysis results and analytical methods) and found them to be acceptable.

The APVMA has considered the toxicological aspects of folpet and concluded that there are no toxicological concerns regarding the approval of this active constituent. An Acceptable Daily Intake (ADI) of 0.09 mg/kg bw/d has been set, based on a No-Observed Adverse Effect Level (NOAEL) of 9 mg/kg bw/d observed in a two-year rat dietary exposure study based on hyperkeratosis and/or acanthosis of the non-glandular gastric mucosa at the 37 mg/kg bw/day dose level (the LOAEL), after applying a safety factor of 100.

An Acute Reference Dose (ARfD) was considered unnecessary due to the low acute toxicity and the lack of evidence of acute neurotoxicity, acute developmental toxicity (in the absence of maternotoxicity) or any other toxicologically relevant effect that might be attributable to a single dose or exposure.

The APVMA's determination is that the impurities in the TGAC folpet above 1 g/kg (0.1%) are not considered to be of toxicological concern. Of interest, phthalimide is a mammalian metabolite of folpet, and therefore considered to be included in the parent compound of folpet for the Acceptable Daily Intake (ADI).

Based on the toxicity profile, the Scheduling Delegate has included folpet in Schedule 6 of the Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP), with no concentration cut-off to a lower schedule.

The APVMA has considered the ecotoxicological aspects of folpet. Folpet is not persistent and the metabolites do not bioaccumulate to toxicologically relevant concentrations in terrestrial, aquatic or soil food chains. Folpet has high toxicity to

fish and aquatic invertebrates, moderate toxicity to algae and aquatic plants, and low toxicity to birds, mammals bees and soil macro-organisms. Environmental risks will be mitigated by label instructions the product.

On the basis of the data provided, and the toxicological and ecotoxicological assessments, it is proposed that the following active constituent standard be established for folpet:

**Table 10: Proposed APVMA Active Constituent Standard for folpet**

Constituent	Specification	Level
Folpet	Folpet	940 g/kg minimum

Impurities of toxicological significance are not expected to occur in folpet as a result of the raw materials and the synthetic route used.

### Further information

A Public Release Summary (PRS) of the evaluation of this product is available from the [APVMA website](#) or from the contact listed below.

### Making a submission

In accordance with section 12 of the Agvet Code, the APVMA invites any person to submit a relevant written submission as to whether folpet should be approved. Submissions should relate only to matters that are considered in determining whether the safety criteria set out in section 5A of the Agvet Code have been met. Submissions should state the grounds on which they are based.

Submissions must be received by the APVMA within 28 days of the date of this notice and be directed to the contact listed below. All submissions to the APVMA will be acknowledged in writing via email or by post.

**Please note:** Submissions will be published on the APVMA's website, unless you have asked for the submission to remain confidential (see [public submission coversheet](#)).

Please lodge your submission with a [public submission coversheet](#), which provides options for how your submission will be published.

Note that all APVMA documents are subject to the access provisions of the *Freedom of Information Act 1982* and may be required to be released under that Act should a request for access be made.

Please send your written submission and coversheet by email or post to:

Email: [casemanagement@apvma.gov.au](mailto:casemanagement@apvma.gov.au)

Post:

Director Chemistry and Manufacture  
Australian Pesticides and Veterinary Medicines Authority  
GPO Box 574  
Canberra ACT 2601

### Privacy

For information on how the APVMA manages personal information when you make a submission, see our [Privacy Policy](#).

## Application for registration of a new product – Folpan 800 WG Fungicide containing folpet

The APVMA has before it an application for registration of a new product, Folpan 800 WG Fungicide, containing a new active constituent, folpet.

**Table 11: Particulars of the application**

<b>Proposed product name</b>	Folpan 800 WG Fungicide
<b>Applicant company</b>	Adama Australia Pty Limited
<b>Name of active constituent</b>	Folpet
<b>Signal heading</b>	Poison
<b>Summary of proposed use</b>	A preventative treatment for control of downy mildew, powdery mildew and grey mould in grapes.
<b>Pack sizes</b>	1 – 1000L
<b>Withholding period</b>	Harvest (H): DO NOT HARVEST FOR 7 DAYS AFTER APPLICATION. Grazing (G): TREATED CROPS ARE NOT TO BE GRAZED OR FED TO ANIMALS. DO NOT GRAZE VINEYARDS AFTER APPLICATION.

A summary of the APVMA's evaluation of Folpan 800 WG Fungicide in accordance with the requirements of section 14(1)(C) of the Agricultural and Veterinary Chemicals Code (the 'Agvet Code'), scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994*:

- 1) The APVMA has evaluated the application and in its assessment in relation to whether the safety criteria have been met in accordance with the definition set out in section 5A of the Agvet Code, proposes to determine that:
  - i. The APVMA is satisfied that the proposed use of Folpan 800 WG Fungicide would not be an undue hazard to the safety of people exposed to it during its handling and use.
  - ii. The APVMA is satisfied that the proposed use of Folpan 800 WG Fungicide will not be an undue hazard to the safety of people using anything containing its residues.
  - iii. The APVMA is satisfied that the proposed use of Folpan 800 WG Fungicide is not likely to have an unintended effect that is harmful to animals, plants or the environment if used according to the product label directions.
- 2) The APVMA has evaluated the application and in its assessment in relation to whether the efficacy criteria have been met in accordance with the definition set out in section 5B of the Agvet Code, and proposes to determine that:
  - i. The APVMA is satisfied that when used in accordance with the label approved by APVMA the proposed use of Folpan 800 WG Fungicide would be effective for its proposed uses.
- 3) The APVMA has evaluated the application and in its assessment in relation to whether the trade criteria have been met in accordance with the definition set out in section 5C of the Agvet Code, proposes to determine that:
  - i. The APVMA is satisfied that when used in accordance with the label approved by APVMA the proposed use of Folpan 800 WG Fungicide would not adversely affect trade.

### Further information

A Public Release Summary (PRS) of the evaluation of this product is available from the [APVMA website](#) or by contacting the APVMA as listed below.

## Making a submission

In accordance with section 13 of the Agvet Code, the APVMA invites any person to submit a relevant written submission as to whether Folpan 800 WG Fungicide should be registered. Submissions should relate only to matters that are required by the APVMA to be taken into consideration in determining whether the safety, efficacy or trade criteria have been met. Submissions should state the grounds on which they are based.

Submissions must be received by the APVMA within 28 days of the date of this notice and be directed to the contact listed below. All submissions to the APVMA will be acknowledged in writing via email or by post.

Relevant comments will be taken into account by the APVMA in deciding whether the product should be registered and in determining appropriate conditions of registration and product labelling.

**Please note:** Submissions will be published on the APVMA's website, unless you have asked for the submission to remain confidential (see [public submission coversheet](#)).

Please lodge your submission with a [public submission coversheet](#), which provides options for how your submission will be published.

Note that all APVMA documents are subject to the access provisions of the *Freedom of Information Act 1982* and may be required to be released under that Act should a request for access be made.

Please send your written submission and coversheet by email or post to:

Email: [casemanagement@apvma.gov.au](mailto:casemanagement@apvma.gov.au)

Post:  
Executive Director Agricultural Chemicals Branch  
Australian Pesticides and Veterinary Medicines Authority  
GPO Box 574  
Canberra ACT 2601, Australia

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## Application for registration of a new products – Basta Ultra Herbicide and Liberty Ultra Herbicide, containing 211 g/L Glufosinate-P-ammonium

The APVMA has before it an application for registration of a new products, Basta Ultra Herbicide and Liberty Ultra Herbicide, containing a new active constituent, Glufosinate-P-ammonium.

**Table 12: Particulars of the application**

<b>Proposed product name</b>	Basta Ultra Herbicide and Liberty Ultra Herbicide
<b>Applicant company</b>	BASF AUSTRALIA LTD
<b>Name of active constituent</b>	Glufosinate-P-ammonium
<b>Signal heading</b>	Schedule 5
<b>Summary of proposed use</b>	<p>Basta Ultra Herbicide: For non-residual control of broadleaf and grass weeds in various situations.</p> <p>Liberty Ultra Herbicide: For non-residual control of certain broadleaf and grass weeds in LibertyLink® Canola varieties (with tolerance to Liberty® Ultra Herbicide), fallow and optical spot spray technologies (OSST).</p>
<b>Pack sizes</b>	1 – 1000 L
<b>Withholding period</b>	<p><b>Basta Ultra Herbicide:</b></p> <p>Harvest:</p> <p>Blackberry, blackcurrants, blueberries, boysenberry, citrus fruit, grapes, loganberry, olives, raspberry, strawberries, tomatoes, tree nuts: NOT REQUIRED WHEN USED AS DIRECTED.</p> <p>Tropical and sub-tropical fruits – inedible peel, (avocado, banana, feijoa, guava, kiwifruit, litchi, mango, pawpaw, passionfruit, pineapple, pitaya (dragon fruit) and rambutan): NOT REQUIRED WHEN USED AS DIRECTED.</p> <p>Pome and stone fruit: DO NOT HARVEST FOR 21 DAYS AFTER APPLICATION.</p> <p>Sugarcane: DO NOT HARVEST FOR 16 WEEKS AFTER APPLICATION.</p> <p>Green Bean / French Bean: DO NOT HARVEST FOR 4 WEEKS AFTER APPLICATION.</p> <p>Date palms, green tea, native foods: DO NOT HARVEST FOR 1 DAY AFTER APPLICATION. DO NOT HARVEST LEAVES FROM NATIVE PEPPER OR WATTLES THAT ARE CLOSE TO THE GROUND FOR FOOD USES.</p> <p>Coffee: NOT REQUIRED WHEN USED AS DIRECTED.</p> <p>Grazing:</p> <p>Sugarcane: DO NOT GRAZE OR CUT FOR STOCK FOOD FOR 16 WEEKS AFTER APPLICATION.</p> <p>Green Bean / French Bean: DO NOT GRAZE OR CUT TREATED AREAS FOR STOCKFOOD FOR 4 WEEKS AFTER APPLICATION.</p> <p>All other crops: DO NOT GRAZE OR CUT TREATED AREAS FOR STOCK FOOD FOR 8 WEEKS AFTER APPLICATION.</p> <p><b>Liberty Ultra Herbicide:</b></p> <p>HARVEST (H):</p> <p>Not required when used as directed.</p> <p>GRAZING (G):</p> <p>CANOLA: DO NOT GRAZE OR CUT FOR STOCK FOOD FOR 6 WEEKS AFTER APPLICATION.</p> <p>CANOLA GROWN FOR SEED PRODUCTION: DO NOT GRAZE OR CUT FOR STOCK FEED.</p> <p>SUMMER FALLOW: (including optical spot spraying technology): DO NOT GRAZE OR CUT FOR STOCK FOOD A CROP SOWN FOLLOWING A FALLOW SPRAY FOR 6 WEEKS AFTER SOWING.</p> <p>All other situations: DO NOT GRAZE OR CUT TREATED AREAS FOR STOCK FOOD FOR 8 WEEKS AFTER APPLICATION</p>

A summary of the APVMA's evaluation of Basta Ultra Herbicide and Liberty Ultra Herbicide in accordance with the requirements of section 14(1)(C) of the Agricultural and Veterinary Chemicals Code (the 'Agvet Code'), scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994*:

- 1) The APVMA has evaluated the application and in its assessment in relation to whether the safety criteria have been met in accordance with the definition set out in section 5A of the Agvet Code, proposes to determine that:
  - i. The APVMA is satisfied that the proposed use of Basta Ultra Herbicide and Liberty Ultra Herbicide would not be an undue hazard to the safety of people exposed to it during their handling and use.
  - ii. The APVMA is satisfied that the proposed use of Basta Ultra Herbicide and Liberty Ultra Herbicide will not be an undue hazard to the safety of people using anything containing their residues.
  - iii. The APVMA is satisfied that the proposed use of Basta Ultra Herbicide and Liberty Ultra Herbicide is not likely to have an unintended effect that is harmful to animals, plants or the environment if used according to the product label directions.
- 2) The APVMA has evaluated the application and in its assessment in relation to whether the efficacy criteria have been met in accordance with the definition set out in section 5B of the Agvet Code, and proposes to determine that:
  - i. The APVMA is satisfied that when used in accordance with the label approved by APVMA the proposed use of Basta Ultra Herbicide and Liberty Ultra Herbicide would be effective for their proposed uses.
- 3) The APVMA has evaluated the application and in its assessment in relation to whether the trade criteria have been met in accordance with the definition set out in section 5C of the Agvet Code, proposes to determine that:
  - i. The APVMA is satisfied that when used in accordance with the label approved by APVMA the proposed use of Basta Ultra Herbicide and Liberty Ultra Herbicide would not adversely affect trade.

## Further information

A Public Release Summary (PRS) of the evaluation of this product is available from the [APVMA website](#) or by contacting the APVMA as listed below.

## Making a submission

In accordance with section 13 of the Agvet Code, the APVMA invites any person to submit a relevant written submission as to whether Basta Ultra Herbicide and Liberty Ultra Herbicide should be registered. Submissions should relate only to matters that are required by the APVMA to be taken into consideration in determining whether the safety, efficacy or trade criteria have been met. Submissions should state the grounds on which they are based.

Submissions must be received by the APVMA within 28 days of the date of this notice and be directed to the contact listed below. All submissions to the APVMA will be acknowledged in writing via email or by post.

Relevant comments will be taken into account by the APVMA in deciding whether the product should be registered and in determining appropriate conditions of registration and product labelling.

**Please note:** Submissions will be published on the APVMA's website, unless you have asked for the submission to remain confidential (see [public submission coversheet](#)).

Please lodge your submission with a [public submission coversheet](#), which provides options for how your submission will be published.

Note that all APVMA documents are subject to the access provisions of the *Freedom of Information Act 1982* and may be required to be released under that Act should a request for access be made.

Please send your written submission and coversheet by email or post to:

Email: [casemanagement@apvma.gov.au](mailto:casemanagement@apvma.gov.au)

Post:

Executive Director Agricultural Chemicals Branch  
Australian Pesticides and Veterinary Medicines Authority  
GPO Box 574  
Canberra ACT 2601, Australia

## **Privacy**

For information on how the APVMA manages personal information when you make a submission, see our [Privacy Policy](#).

## Licensing of veterinary chemical manufacturers

Pursuant to Part 8 of the Agricultural and Veterinary Chemicals Code (Agvet Code), scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994*, the APVMA hereby gives notice that it has taken action with respect to the licensing of the following veterinary chemical manufacturers with effect from the dates shown.

For a comprehensive listing of all licensed manufacturers please see the [APVMA website](#).

### New licenses

The APVMA has issued the following licenses under subsection 123(1) of the Agvet Code:

**Table 13: New licenses issued by the APVMA under subsection 123(1) of the Agvet Code**

Company name	Licence number	Company ACN	Address	Product types	Steps of manufacture	Date issued
International Animal Health Products Pty Ltd	2015	003 185 699	18 Healey Circuit Huntingwood NSW 2148	Category 2: Creams / lotions, ointments, pastes, powders, pellets, sprays, bolus, and liquids Category 3: Liquids, sprays, and powders Category 4: Premixes, supplements, and probiotics	Quality assurance (QA) of raw materials, formulation including blending, dry milling, granulation, filling, packaging, labelling, pellet extrusion, analysis and testing (physical), secondary packaging, secondary labelling, repackaging, relabelling, storage and release for supply.	19 December 2025
Mars Australia Pty Ltd	2106	008 454 313	1 Petcare Place Wodonga VIC 3690	Category 2: Palatable ingestible chews	Quality assurance (QA) of raw materials, formulation including blending, dry milling, filling, packaging, labelling, strip, blister or sachet packaging, microbiological reduction treatment (heat) analysis and testing (physical), storage, and release for supply.	20 January 2026

### Licence cancellations

The APVMA has cancelled the following licenses under subsection 127(1) of the Agvet Code:

**Table 14: Licenses cancelled by the APVMA under subsection 127(1) of the Agvet Code**

Company name	Licence number	Company ACN	Address	Date cancelled
Advanced Feeds Pty Ltd	2117	009 403 623	13 Meliador Way Midvale WA 6056	10 February 2026

### APVMA contact

Manufacturing Quality and Licensing  
Australian Pesticides and Veterinary Medicines Authority  
GPO Box 574  
Canberra ACT 2601

**Phone:** +61 2 6770 2301  
**Email:** [mls@apvma.gov.au](mailto:mls@apvma.gov.au)

## **Notice of proposed chemical products standard for chemical products containing fenitrothion**

The Australian Pesticides and Veterinary Medicines Authority (APVMA) is proposing a new standard for chemical products containing fenitrothion as an active constituent in the Agricultural and Veterinary Chemicals Code (Chemical Products) Standard 2026 (Chemical Products Standard).

The APVMA may make standards for chemical products by legislative instrument under section 6E of the Agricultural and Veterinary Chemicals Code (Agvet Code) scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994* (Agvet Code Act).

Before making standards for chemical products, the APVMA must undertake the consultation and notice requirements set out in regulation 8AF of the Agricultural and Veterinary Chemicals Code Regulations 1995 (Regulations).

### **Statement as to why the APVMA determines that the standard for chemical products containing fenitrothion is necessary**

In undertaking its recent reconsideration of fenitrothion, the APVMA identified a risk to safety of users, animals, the environment and trade from products containing fenitrothion as an active constituent. To manage this risk, the APVMA imposed several new controls including deciding to impose a standard to limit impurities in chemical products containing fenitrothion as an active constituent. Further information on the APVMA's final regulatory decision on fenitrothion can be found in the [APVMA Gazette No 17, Tuesday 19 August 2025](#).

The proposed standard for chemical products containing fenitrothion as an active constituent specifies the proposed acceptable levels (minimum, maximum or range as appropriate) for concentrations of fenitrothion active constituents and toxicologically significant impurities consistent with the scientific assessments conducted as part of the chemical review of fenitrothion. If the toxic impurities in chemical products containing fenitrothion as an active constituent exceed prescribed limits, they will pose an unacceptable risk to safety, particularly the safety of people using these products.

Due to the effect of the existing legislation, if the standard for chemical products containing fenitrothion as an active constituent is exceeded, a person may commit an offence or be liable to pay a civil penalty if the person then supplies the product, or causes or permits such a product to be supplied (see paragraph 87(2)(a) of the Agvet Code and subregulation 42(3) of the Regulations).

Accordingly, the standard for chemical products containing fenitrothion as an active constituent will help ensure that continued use of these products are safe for people, plants, animals and the environment by setting limits for toxic impurities.

In considering whether it is necessary to make a standard for chemical products containing fenitrothion as an active constituent, the APVMA has had regard to the FAO and WHO specification for fenitrothion as it related to relevant fenitrothion product formulation types, as required by regulation 8AF(a) of the Regulations. The proposed standard for chemical products containing fenitrothion as an active constituent is consistent with the FAO and WHO specification for fenitrothion. There are no differences between the FAO specifications and the proposed standard in the impurities specified and their limits. This is supported by the toxicological and stability data assessments conducted for the reconsideration of fenitrothion, as described in the [Fenitrothion Final Review Technical Report](#).

## **How to obtain more information**

More information, including a copy of the [draft Agricultural and Veterinary Chemicals Code \(Chemical Products\) Amendment \(Fenitrothion\) Standard 2026](#) and [how to make a written submission](#) on the draft, is available on [apvma.gov.au](http://apvma.gov.au)

## Agvet chemical voluntary recall: Frequency Herbicide

**Product name:** Frequency Herbicide

**APVMA registration number:** 86267

**APVMA approved label number:** 131170, 145930

**Batch numbers:** 224V110001, 224V120002, 224V120006, 224V120008, 244V020001, 244V020003, 244V020005, 244V020006, 244V020007, 244V020009, 244V020010, 244V020011, 244V020012, 244V020013, 244V020014, 244V020015, 254V020001, 254V020002, 254V020003, 254V020004, 254V020005, 254V020006, 25E9040001, 25E9040002, 25E9040003, 25E9040004, 25E9050001, 25E9050002

**Sold by:** agricultural retailers nationally between 1 June 2023 to 16 February 2026

On 16 February 2026, BASF Australia Ltd (ACN 008 437 867) initiated a voluntary recall under section 106 of the Agricultural and Veterinary Chemicals Code scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994* (Cth) in relation to the chemical product described above.

### Reason for voluntary recall

A number of Frequency Herbicide batches do not meet BASF's quality requirements.

### Hazard

As the affected product does not meet BASF's quality requirements, it may not perform as per label directions.

### What to do if in possession of this chemical product

DO NOT use the product if from an affected batch number (listed above). Contact your retailer to arrange return and credit for the product. Retailers should immediately quarantine all stock of the recalled product and contact your BASF Key Account Manager or BASF Australia Ltd Customer Service (1800 558 399) to organise a stock return.

### More information

Visit the APVMA website to [view the notice](#) of voluntary recall for the chemical product described above.

The APVMA publishes a list of [agvet chemical recall notices](#) on its website and provides a [subscription option](#) to be notified by email when a new recall notice is published.

### Contact

Questions about this voluntary recall should be directed to:

BASF Australia Ltd Customer Service

**Phone:** 1800 558 399

## Recall Notice: Augustine Approved ACN 130 045 368

Recall Notice under section 101 of the Agricultural and Veterinary Chemicals Code scheduled to the *Agricultural and Veterinary Chemicals Code Act 1994 (Cth) (Agvet Code) No 1 of 2026*.

On 17 February 2026, the Australian Pesticides and Veterinary Medicines Authority (APVMA) issued a recall notice to Augustine Approved Pty Ltd, ACN 130 045 368 ('the Company'), pursuant to section 101 of the Agvet Code in relation to the unregistered chemical products labelled or known as:

- Augustine Approved Faith's Cleanse & Detox
- Augustine Approved Raw Coconut Oil
- Augustine Approved Turmeric Powder 5-6%
- Augustine Approved Lecithin Powder
- Augustine Approved Wakame Powder
- Augustine Approved Augustine's SuperBalm
- Augustine Approved SuperHeal
- Augustine Approved SuperBath

This notice requires the Company, within 30 days of being issued the notice, to stop supply of the above identified products. It also requires the Company to notify any person who has possession or custody of those products – whether they received them directly from the Company or through someone else – that they must stop supply of the above identified products and return them to the Company. The Company is also required to take all necessary steps to recover all remaining stock from any person that the Company has either directly or indirectly supplied the above identified products to.

NOTE (included on 29 April 2026):

- On 25 February 2026, the company applied to the Administrative Review Tribunal (**Tribunal**) for review of the decision to issue the recall notice. The proceedings are ongoing.
- On 13 March 2026, at the request of the company, the Tribunal ordered pursuant to s 32(2) of the *Administrative Review Tribunal Act 2024 (Cth)* that the operation of the recall notice is stayed until the decision of the Tribunal on the application for review comes into operation or until further order of the Tribunal.

### Supplier details

Augustine Approved Pty Ltd ACN 130 045 368  
435 Robinsons Road  
Langwarrin Vic 3910, Australia

### APVMA contact

For any enquiries or further information about this matter, please contact:

Director, Compliance Operations  
Australian Pesticides and Veterinary Medicines Authority  
GPO Box 574  
Canberra ACT 2601

**Phone:** +61 2 6770 2300

**Email:** [compliance@apvma.gov.au](mailto:compliance@apvma.gov.au)

### **More information**

The APVMA publishes a list of [recall notices](#) on its website, and provides a [subscription option](#) to be notified by email when a new recall notice is published.

## Notice of cancellation at the request of the holder

At the request of the holder, in accordance with section 42(1) of the *Agricultural and Veterinary Chemicals Code Act 1994* (Agvet Code), the APVMA has cancelled the approvals and/or registrations set out in Table 15:

**Table 15: Active constituent approval/product registration/label approval cancelled at the request of the holder**

Approval or registration number	Name	Type of approval or registration	Holder	Reason for cancellation (if relevant pursuant to s45A(3))	Date of cancellation
69683/61437 69683/107601 69683/112417 69683/126178	Timbertreat A Azole Insecticide/Fungicide Wood Preservative	Label	Kop-Coat Australia Pty. Ltd.	May not meet labelling criteria	18 February 2026

In accordance with section 45A(1)(b) of the Agvet Code, the APVMA publishes this notice of the cancellation, including the following instructions which set out how a person can deal with the cancelled active constituent, cancelled product or product bearing a cancelled label referred to in Table 15.

### Instructions

Instructions for persons who possess, have custody of or use the cancelled active constituent, cancelled product, or the product bearing a cancelled label under section 45B(3) of the Agvet Code.

A person who possesses, has custody of or uses the cancelled active constituent, cancelled product or product bearing a cancelled label referred to in Table 15 in accordance with the instructions contained in this notice, is taken to have been issued with a permit under section 45B(3) of the Agvet Code to possess, have custody of or use the cancelled active constituent, cancelled product or product bearing a cancelled label, in accordance with those instructions.

### Possession or custody

A person may possess the cancelled active constituent, cancelled product or product bearing a cancelled label referred to in Table 15 in accordance with its label instructions for 12 months from the date of cancellation.

### Use, supply or otherwise deal with

A person may use the cancelled active constituent, cancelled product or products bearing a cancelled label referred to in Table 15 according to its label instructions, including any conditions relating to shelf life or expiry date, for 12 months after the date of cancellation.

A person may supply or cause to be supplied at wholesale or retail level the cancelled active constituent, cancelled product, or product bearing a cancelled label referred to in Table 15, for 12 months after the date of cancellation.

### Contraventions

After the day that is 12 months from the date of cancellation it will be an offence against the Agvet Code to have possession or custody of the cancelled active constituents, cancelled products or products bearing a cancelled label with the intention to supply, or to supply the cancelled active constituent, cancelled product, or product bearing a cancelled label.

It is an offence to possess, have custody of, use, or otherwise deal with the cancelled active constituents, cancelled products or products bearing the cancelled label listed in Table 15 in a manner that contravenes the above instructions.

### **APVMA contact**

For any enquiries or further information about this matter, please contact:

Chemical Review  
Australian Pesticides and Veterinary Medicines Authority  
GPO Box 574  
Canberra ACT 2601

**Phone:** +61 2 6770 2400

**Email:** [chemicalreview@apvma.gov.au](mailto:chemicalreview@apvma.gov.au)

### **More information**

The APVMA publishes a list of [voluntary cancellations at the request of the holder](#) on its website, and provides a [subscription option](#) to be notified by email when the list is updated.