Reprint

as at 1 November 2011

Agricultural Compounds and Veterinary Medicines Amendment Regulations 2005

(SR 2005/90)

Agricultural Compounds and Veterinary Medicines Amendment Regulations 2005: revoked, on 1 November 2011, pursuant to regulation 16 of the Agricultural Compounds and Veterinary Medicines (Exemptions and Prohibited Substances) Regulations 2011 (SR 2011/327).

Pursuant to section 75 of the Agricultural Compounds and Veterinary Medicines Act 1997, Her Excellency the Governor-General, acting on the advice and with the consent of the Executive Council, makes the following regulations.

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Note

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Changes authorised by section 17C of the Acts and Regulations Publication Act 1989 have been made in this eprint.

A general outline of these changes is set out in the notes at the end of this eprint, together with other explanatory material about this eprint.

These regulations are administered by the Ministry of Agriculture and Forestry.

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Schedule 1 New Schedules 1 to 4 substituted in principal regulations

Schedule 2 New Schedule 7 substituted in principal regulations

1 Title

- (1) These regulations are the Agricultural Compounds and Veterinary Medicines Amendment Regulations 2005.
- (2) In these regulations, the Agricultural Compounds and Veterinary Medicines Regulations 2001¹ are called "the principal regulations".

2 Commencement

These regulations come into force on the 28th day after the date of their notification in the *Gazette*.

3 Agricultural compounds exempt from registration if conditions complied with

- (1) Regulation 5 of the principal regulations is amended by omitting the words "Schedules 2 and 3", and substituting the expression "Schedule 2".
- (2) Regulation 5(b) of the principal regulations is amended by omitting the word "schedules", and substituting the word "schedule".

4 New regulation 5A inserted

The principal regulations are amended by inserting, after regulation 5, the following regulation:

"5A Combined agricultural compounds exempt from registration

"(1) An agricultural compound is exempt from registration under section 21 or section 27 of the Act if the agricultural compound is a combination of 2 or more agricultural compounds that are exempt from registration under these regulations.

¹ SR 2001/101

- "(2) The combined agricultural compound is subject to any—
 - "(a) conditions set out in column 2 of Schedule 2; and
 - "(b) other provision of these regulations that applies to each agricultural compound that comprises the combined agricultural compound."

5 Information requirements

- (1) Regulation 6 of the principal regulations is amended by omitting the words "Schedules 2 and 3", and substituting the expression "Schedule 2".
- (2) Regulation 6(f) of the principal regulations is amended by omitting the words "or Schedule 3".

6 Regulation 7 revoked

Regulation 7 of the principal regulations is revoked.

7 New Schedules 1 to 4 substituted

The principal regulations are amended by revoking Schedules 1, 2, 3, and 4 and substituting the Schedules 1, 2, 3, and 4 set out in Schedule 1 of these regulations.

8 New Schedule 7 substituted

The principal regulations are amended by revoking Schedule 7, and substituting the Schedule 7 set out in Schedule 2 of these regulations.

Schedule 1

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New Schedules 1 to 4 substituted in principal regulations

Schedule 1

Agricultural compounds exempt from registration under sections 21 and 27

of Act if applicable codes of practice complied with

1 Compounds (including homeopathic, herbal, chemical, and oral nutritional compounds) prepared by a person for use by the person as an agricultural compound on animals or plants owned by the person, or in any land, place, or water owned or occupied by the person.

> The following compounds are excluded unless there is an applicable code of practice in force under section 28 of the Act:

- (a) active ingredients that are prescription medicines or restricted medicines (as those terms are defined in the Medicines Act 1981):
- (b) antibiotic active ingredients:
- (c) hormones:
- (d) substances that are prohibited by countries importing New Zealand primary produce:
- (e) vertebrate toxic agents.
- 2 Homeopathic plant compounds used commercially.
- 3 Non-medicated topical hoof preparations used solely to maintain or improve hoof condition.
- 4 Non-medicated topical skin preparations used solely to maintain or improve skin condition.
- 5 Non-absorbent masking agents used to disguise odours.
- 6 Topical non-absorbent and non-solvent cleaning products, including non-medicated shampoos, soaps, tear-stain removers, and toothpaste.
- 7 In vitro diagnostics used to confirm the presence or absence of disease or as an aid in the diagnosis of disease or abnormal conditions.

- 8 Compounds used in the production of plant tissue cultures.
- 9 Compounds (not containing biologically active ingredients) used to protect plant grafts.
- 10 Vertebrate and invertebrate attractants and repellants that are not applied directly to animals or plants.
- 11 Invertebrate mating disrupters that are not applied directly to animals or plants.
- 12 Antisapstains.
- 13 A preparation of 2 or more ingredients if each ingredient is an agricultural compound described in this schedule and the combination of ingredients does not increase or change any of the risks described in section 19 of the Act.

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Agricultural compounds exempt from registration under sections 21 and 27 of Act if conditions in column 2 and requirements in regulation 6 complied with

Column 1	Column 2
Agricultural compound	Conditions
Oral and topical preparations—	If used as a veterinary medicine, the label information must—
(a) prepared by a process of solution, extraction, or titration of an active ingredient followed by strictly regimented serial dilution; and	(a) identify the compound as a homeopathic preparation; and

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(b) that do not claim to prevent, control, or cure a specific disease characterised by pain or distress in animals	(b) include a statement that, if the preparation fails to alleviate the condition being treated, the user should seek veterinary advice
Oral and topical preparations—	If used as a veterinary medicine, the label information must—
(a) prepared from either any part of a plant or an unrefined extract from a plant, except a plant listed in Schedule 6; and	(a) identify the compound as a herbal preparation; and
(b) that do not claim to prevent, control, or cure a specific disease characterised by pain or distress in animals	(b) include a statement that, if the preparation fails to alleviate the condition being treated, the user should seek veterinary advice
	Must not be used on the teats of lactating animals if the milk of the animals is intended for human consumption
Markers, paints, and dyes used as pigments or colourants for topical application to identify animals temporarily	of lactating animals if the milk of the animals is intended for
as pigments or colourants for topical application to identify	of lactating animals if the milk of the animals is intended for

	The veterinarian must act in accordance with any applicable code of practice in force under section 28 of the Act
Preparations compounded and used by veterinarians	The veterinarian must act in accordance with any applicable code of practice in force under section 28 of the Act
	Preparations must not be used on animals except under the direct care, authority, or prescription of a veterinarian
Substance or biological compound or mix of substances or biological compounds (to which this schedule does not otherwise apply) to be used as a veterinary medicine	May be imported only if the Director- General is satisfied that—
	(a) there is no equivalent veterinary medicine registered under the Act; and
	(\mathbf{h}) it is as series of the event of the

(b) it is required to ensure the immediate welfare of animals

Must not contain any substance or biological compound that is prohibited for use as an agricultural compound

Must not be used on animals except under the direct care, authority, or prescription of a veterinarian

Schedule 1

	The veterinarian must act in accordance with any applicable code of practice in force under section 28 of the Act
Topical preparations—	Must not be used on the teats of lactating animals if the milk of the animals is and intended for human consumption
(a) containing ingredients not able to be absorbed through the skin;	
(b) used solely to treat minor injuries or to prevent dermatological abnormalities; and	Must be manufactured in accordance with good manufacturing practice
(c) that do not include any ingredient listed in Schedule 3	
Non-medicated anti-diarrhoea preparations	Must be manufactured in accordance with good manufacturing practice
	The label information must include a statement that if the preparation fails to alleviate the condition being treated the user should seek veterinary advice
Non-medicated oral laxatives and lubricants	Must be manufactured in accordance with good manufacturing practice
	The label information must include a statement that if the preparation fails to alleviate the condition being treated the user should seek veterinary advice

Cauterising preparations used or applied superficially	Must be manufactured in accordance with good manufacturing practice
	The label must include a statement that if the preparation fails to stop bleeding the user should seek veterinary advice
Urinary tract modifiers (acidifiers and alkalisers) that are oral preparations used solely for modification of urinary pH	Must not be used on animals from which animal material is intended to be used for the production of human food or human pharmaceutical products
	Must be manufactured in accordance with good manufacturing practice
	Must be packaged for sale in dosage-size packages (not in bulk or concentrated form) appropriate for the animals for which the agricultural compound is recommended
Respiratory tract modifiers (expectorants and cough suppressants) that—	Must not be used on animals from which animal material is intended to be used or the production of human food or human pharmaceutical products
(a) have only a locally acting, superficial effect on the respiratory tract; and	
(b) are given orally, applied topically to the nose, or inhaled; and	Must be manufactured in accordance with good manufacturing practice

(c) are used solely in companion Must be packaged for sale in animals to promote mucolysis, dosage-size packages (not in bulk or concentrated form) cough suppression (by alleviating only irritation) appropriate for the animals and relieve compromised for which the agricultural airways and upper respiratory compound is recommended tract congestion Compounds used to extend An applicable code of practice animal semen or to be used as must be must in force under media for animal ova section 28 of the Act and be complied with Spray markers that are coloured Must not be used on food crops unless they do not indicators to show where liquid agri-chemicals have been produce residues in primary applied to help prevent overlaps produce that fail to comply with applicable food residue standards set in or under any enactment Plant compound adjuvants, Must not be used on food crops including unless wetting and they do not produce residues sticking agents, pH buffers, in primary produce that fail to drift retardants, and water comply with applicable food conditioners residue standards set in or under any enactment Repellants applied directly to Must not be used on food plants and used solely to repel crops unless they do not vertebrates or invertebrates produce residues in primary produce that fail to comply with applicable food residue standards set in or under any enactment

Attractants applied directly to plants and used solely to attract vertebrates or invertebrates	Must not be used on food crops unless they do not produce residues in primary produce that fail to comply with applicable food residue standards set in or under any enactment
Mating disrupters applied directly to plants and used solely to interfere with the reproduction of invertebrates	Must not be used on food crops unless they do not produce residues in primary produce that fail to comply with applicable food residue standards set in or under any enactment
Anti-transpirants used solely to prevent drying of plants	Must not be used on food crops unless they do not produce residues in primary produce that fail to comply with applicable food residue standards set in or under any enactment
Frost protectants of a chemical nature used solely to prevent frost damage	Must not be used on food crops unless they do not produce residues in primary produce that fail to comply with applicable food residue standards set in or under any enactment
Sunblocks used solely to prevent or reduce sunburn in plants	Must not be used on food crops unless they do not produce residues in primary produce that fail to comply with applicable food residue standards set in or under any enactment

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Schedule 3

Schedule 2

Ingredients excluded from topical preparations containing ingredients not able to be absorbed through skin and used solely to treat minor injuries or to prevent dermatological abnormalities

Ingredient

Antibiotics

Schedule 4

Oral nutritional compounds exempt from registration under sections 21 and 27 of Act if following conditions complied with

- 1 They must be supplied with a label containing the following information:
 - (a) trade name:
 - (b) the name and address of the producer, if applicable:
 - (c) the name and address of the manufacturer, if applicable:
 - (d) ingredients:
 - (e) directions for use, including the species, type, and class of animal intended to be used for:
 - (f) details of any precautions to be taken to prevent or manage risks described in section 19 of the Act when being used, particularly potential hazards to animals fed with or exposed to them:
 - (g) batch number, if applicable:
 - (h) manufacturing date, if applicable:
 - (i) use by date or expiry date, if applicable.
- 2 They must be fit for the purpose of feeding to the species, type, and class of animal specified under clause 1(e).
- 3 They are fit for their purpose only if they are used as recommended and do not do any of the following:
 - (a) produce residues in primary produce that fail to comply with applicable food residue standards set in or under any enactment:

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- (b) result in toxic reactions causing pain or distress in the animal:
- (c) result in malnutrition causing pain or distress in the animal:
- (d) result in physical harm causing pain or distress in the animal:
- (e) contain pathogenic micro-organisms at levels that could cause disease resulting in pain and distress in the animal.
- 4 Agricultural compounds that are therapeutic or pharmacological substances or preparations may be incorporated into oral nutritional compounds only if—
 - (a) the agricultural compounds are registered under the Act; and
 - (b) the incorporation of the agricultural compounds is consistent with any conditions of their registration.
- 5 Feed additives may be used in oral nutritional compounds only if the feed additives are described in Schedule 7 and used in accordance with any relevant limitations specified in that schedule.
- 6 Oral nutritional compounds that are feed commodities are subject only to clauses 2 and 3. For the purposes of this clause, feed commodities means plants (or any part or parts of those plants) that are raised in an agricultural context and used as feed or for feed production for animals.

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New Schedule 7 substituted in principal regulations

Schedule 7

Schedule 4

Substances generally recognised as safe if used in accordance with any applicable conditions in Schedule 4

Substances generally regarded as safe feed additives in oral nutritional compounds Interpretation

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A reference to a substance is to all forms of the substance unless a chemical abstract (CAS) number is specified or otherwise stated. Where the first column refers to an organism (including plants), the reference means the whole or any part or any extract of the organism.

General limitations

Each substance in this schedule is subject to the following limitations:

- •• that it is included in a trade name product formulated in accordance with good manufacturing practice; and
- •• that it is of an appropriate grade, and the amount added to the product must not exceed the amount reasonably required to accomplish the intended technical effect.

Substance	Identification	Limitations
	CAS number (if appropriate) unless otherwise stated	
Acacia (Gum arabic)	5/01/9000	
Acetic acid	64-19-7	
Acetophenone	98-86-2	
Adipic acid	124-04-9	
Aldehyde C-18	104-61-0	
Allium sativum		
Almond shell meal		
Aloe vera		

	nes Amendment Regulation	Belledule 2
alpha-galactosidase		From the following sources: Aspergillus niger, var. Morteirella vinaceae, var. raffinoutiliser. Sacchcaromyces sp.
alpha-pinene	80-56-8	When used at no more than 2% by weight of pesticide formulations
Aluminium hydroxide	20768-67-6	
Ammonium chloride	12125-02-9	
Ammonium formate	540-69-2	
Ammonium hydroxide	1336-21-6	
Ammonium phosphate (mono or dibasic)	7722-76-1	
Ammonium propionate		
Ammonium sulphate	7783-20-2	
Amyl butyrate	540-18-1	

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Amylase		From the following sources: Animal pancreatic tissue, Aspergillus oryzae, var. Aspergillus niger, var. Bacillus amyloliquefaciens, B. lentus, B. licheniformis, B. licheniformis containing a B. stearothermophilus gene for alpha- amylase, B. stearothermophilus. B. subtilis containing a B. megatrium gene for alpha-amylase, B. subtilis containing B. stearothermophilus gene for alpha- amylase, B. subtilis, var. Barley malt, Rhizopus oryzae, var.
Anethole	4180-23-8	
p-Anisaldehyde	123-11-5	
A · 1 ·1	0007 70 2	

Allethole	4160-23-6
p-Anisaldehyde	123-11-5
Aniseed oil	8007-70-3
Anisole	100-66-3
Apple flavour	
Ascorbic acid	50-81-7

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Ascorbyl palmitate	137-66-6		
Aspartame	22839-47-0		
Aspergillus niger			
Aspergillus oryzae			
Astaxanthin			
Bacillus licheniformis			
Bacillus subtilis		Non-patho strains	ogenic
Beeswax			
Beetroot			
Bentonite	1302-78-9		
Benzaldehyde	100-52-7		
Benzoic acid	65-85-0	Not more of final fe	than 0.1% ed
Benzyl acetate	140-11-4		
Benzyl alcohol	100-51-6		
beta-apo-8 carotenoic acid ethyl ester	1109-11-1		
beta-carotene			

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beta-glucanase		From the following sources: Aspergillus niger, var. Bacillus lentius, B. subtilis, var. Humicola insolens, Trichoderma longibrachiatum
Betaine hydrochloride	590-46-5	
Bifidobacterium spp.		
Biospernum montanum		
Birch oil		
Boerhavia diffusa		
Brilliant Black BN	2519-30-4	
Brilliant Blue FCF	3844-45-9	
Bromolain	9001-00-7	
Bronopol		
Butterscotch flavour		
Butylated hydroxy-anisole	25013-16-5	Total content of antioxidants must be not more than 0.02% fat content of feed
Butylated hydroxy-toluene	64742-46-7	Total content of antioxidants must be not more than 0.02% fat content of feed

	ltural Compounds and Vete ines Amendment Regulation	
Butyric acid	107-92-6	
Calcium carbonate	471-34-1	
Calcium caseinate	9005-46-3	
Calcium chloride	10035-04-8	
Calcium disodium EDTA	662-33-9	
Calcium formate	544-17-2	
Calcium hydroxide	1305-62-0	
Calcium lactate	814-80-2	
Calcium lignosulfonate	8061-52-7	
Calcium oxide	1305-78-8	
Calcium propionate	4057-81-4	
Calcium silicate	1344-95-2	
Calcium sulphate	7778-18-9	
Camphor		Maximum of 5% in premixes used in production of animal feeds

Candida pintolepesii

Canthaxanthin	514-78-3
Capric [decanoic] acid	334-48-5
Caproic acid	142-62-1
Caprylic acid	124-07-2
Capsanthin	465-42-9

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Caramel		
Caraway		
Carbon black	1333-86-4	
Carminic acid	1260-17-9	
Carmosine [azorubine]	3567-69-9	
Carnauba wax	8015-86-9	
Carob	9000-40-2	
Carophyll pink	514-78-3	
Carrageenan	1/07/9000	
i-carrageenan	1/07/9062	
k-carrageenan	11114-20-8	
1-carrageenan	9064-57-7	
Cassia gum	5373-11-5 / 8013-11-4	
Cayenne peppe	r	
Cedrus deodura	L	
Cellulase		From the following sources: Aspergillus niger, var. Bacillus lentus, Humicola insolens, Trichoderma longibrachiatum
Cellulose	9004-34-6	
Charcoal, activa	ated	
Chlorophyll	1406-65-1	

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Chocolate brown	4553-89-3	
Choline chloride	68-48-1	
Chromium proprionates		
Chymotrypsin		
Cinnamic aldehyde	104-55-2	
Cinnamon		
Citranaxanthin		
Citric acid	77-92-9	
Clove oil	8000-34-8	
Cobalt carbonate	513-79-1	
Colour Amaranth	915-67-3	
Colour Brown HT	4556-89-3	
Colour Green S	3087-16-9	
Colour Indigo Carmine Blue	860-22-0	
Copper carbonate	1184-64-1	
Corn sugar		
Cryptoxanthin	465-42-9	
Curcuma domestics		
Curcuma longa		
Cyperus scarriosus		Specify source
Dandelion		
Diacetyl	431-03-8	

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Diatomaceous earth	7631-86-9	
Dicalcium phosphate	7789-77-7	
2,4-Dichlorobenzyl	1777-82-8	To be used as a preservative alcohol only
Didecyl dimethyl ammonium bromide	2390-68-3	
Dimethyl polysiloxane	8050-81-5	
Disodium EDTA	139-33-3	No more than 240 mg/kg in finished feed
Disodium guanylate	9/12/5550	
Disodium inosinate	4691-65-0	
Dolomite	16389-88-1	
Echinacea		
Elephantopous scaber		
Embelia ribes		Maximum of 5% in premixes used in production of animal feeds
Enterococcus faecium		
Erythorbic acid	7378-23-6	
Erythrosine	16423-68-0	

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Ethoxyquin	91-153-2	Maximum quantity used and to remain in feed must be not more than 0.015%
Ethyl acetate	141-78-6	
Ethyl alcohol	64-17-5	Not more than 10% of the formulation
Ethyl butyrate	105-54-4	
Ethyl formate	109-94-4	
Ethyl-o- aminobenzoate		
Ethyl phenylacetate	101-97-3	
Ethyl propionate	105-37-3	
Ethyl sorbate	2396-84-1	
Ethyl vanillin	121-32-4	
Ethylene diamine tetra-acetic acid	60-00-4	
Fennel	8006-84-6	
Fenugreek		
Ferric chloride	7705-08-0	
Ferrous oxide	1345-25-1	
Ferrous sulphate	7720-78-7	
Food starch and food starch (modified)		Use at a level not in excess of the amoun reasonably required to accomplish the intended effect

Schedule 2	Agricultural Compounds and Medicines Amendment Regula	Veterinary ations 2005Reprinted as at 1 November 2011
Formaldehyde	50-00-0	Not more than 0.25% of final feed
Formic acid	64-18-6	
Fumaric acid	110-17-8	
Garlic	8000-78-0	
Ginger	7/08/8007	
Glucose		Includes dextrose and its hydrated and anhydrous forms
Glutamic acid	617-65-2	
Glycerides (mot and di)	no	
Glycerine	56-81-5	
Glycerol	56-81-5	
Glycerol mono- oleate	25496-72-4	
Glycerol monostearate	31566-31-1	
Guar gum	9000-30-0	
Gypsum	10101-41-4 / 3397-24-5	
Haematococcus algae		

Hemicellulase		From the following sources: Aspergillus niger, var. A. aculeatus, Bacillus lentus, B. subtilis. var. Humicola insolens. Trichoderma longibrachiatum
Holarrhena antidys-tenterica		
3-Hydroxy-2- methyl-4-pyrone (Palatone/Maltol)	118-71-8	Use at a level not in excess of the amount reasonably required to accomplish the intended effect
Hydroxypropyl cellulose	9004-64-2	
Inulin	9005-80-5	
Iron oxides (black)	1317-61-9	
Iron oxides (red)	1309-37-1	
Iron oxides (yellow)	5 1 274-00-1	
Iso-eugenol	97-54-1	
Isopropyl alcohol	67-63-0	
Kaolin	1332-58-7	
Kon jac gum	9000-36-6	
Lactic acid	50-21-5	
Lactobacillus acidophilus		

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Lactobacillus bifidus		
Lactobacillus bulgaricus		
Lactobacillus c	casei 68333-14-2	
Lactobacillus fermentum		
Lactobacillus plantarum		
Lactobacillus rhannosus		
Lactose	63-42-3	
Lauric acid	143-07-7	
Lecithin	8002-43-5	
Lemon grass		
Lemon oil	8008-56-8	
Licorice (Glycyrrhiza)		Includes all licorice derivatives. Not more than 0.1% in final feed
Lignosulphona	tes	
Lime oil	8008-26-2	
Limonene	138-86-3	
Linalool	78-70-6	

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Lipase		From the following sources: Animal pancreatic tissue. aspergillus niger, var. A. oryzae, var. Candida rugosa, edible forestomach of calves, kids. and lambs
Locust bean gum	9000-40-2	
Lutein	57-83-0	
Lycopene	502-65-8	
Macrogol esters (poly-ethylene esters)	9000-99-3	
Magnesium acetate	142-72-3	Includes hydrated forms. Only added to the levels needed
Magnesium aluminium silicate	1327-43-1	Includes hydrated forms. Only added to the levels needed
Magnesium aspartate	18962-61-3	Includes hydrated forms. Only added to the levels needed
Magnesium carbonate	546-93-0	Includes hydrated forms. Only added to the levels needed
Magnesium chloride	7791-18-6	Includes hydrated forms. Only added to the levels needed

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Magnesium citrate	3344-18-1	Includes hydrated forms. Only added to the levels needed
Magnesium gluconate	3632-91-5	Includes hydrated forms. Only added to the levels needed
Magnesium glutamate	64407-99-4	Includes hydrated forms. Only added to the levels needed
Magnesium glycerophosphate	927-20-8	Includes hydrated forms. Only added to the levels needed
Magnesium hydroxide	12141-11-6	Includes hydrated forms. Only added to the levels needed
Magnesium hypophosphite		Includes hydrated forms. Only added to the levels needed
Magnesium orotate	34717-03-8	Includes hydrated forms. Only added to the levels needed
Magnesium oxide	1309-48-4	Includes hydrated forms. Only added to the levels needed
Magnesium phosphate	10043-83-1	Includes hydrated forms. Only added to the levels needed
Magnesium silicate	1343 88-0	Includes hydrated forms. Only added to the levels needed

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Magnesium stearate	557-04-0	Includes hydrated forms. Only added to the levels needed	
Magnesium sulphate	7487-88-9	Includes hydrated forms. Only added to the levels needed	
Magnesium trisilicate	14987-04-3	Includes hydrated forms. Only added to the levels needed	
Malic acid	6915-15-7		
Maltodextrin	9050-36-6		
Maltol	118-71-8		
Mannitol	87-78-5		
Marigold (Aztec)			
Menthol		Not for use in cats	
Methyl alcohol	67-56-1		
6-Methyl-5-hepten- 2-one	110-93-0		
3-Methyl-3-phenyl glycidic acid, ethyl ester	77-83-8		
Methyl salicylate	119-36-8		
Mineral oil		High viscosity	
Monoisopropyl citrate	1321-57-9		
Monopotassium phosphate	7778-77-0		

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Monosodium glutamate	32221-81-1	
Myrica nagi		
Neohesperidine dihydrochalcone	20702-77-6	When used at no more than 30 ppm in finished feed
Neotame	165450-17-9	
Nonyl phenol ethoxylate		
Oatflour		
Octyl gallate	1034-01-1	
Onion oil	2179-59-1	
Operculina turpethum		
Orange oil	8008-57-9	
Oregano		
Pancreatin		
Papain	9001-73-4	
Paprika		
Parabens, includ the following:	ling	
Benzyl paraben	94-26-8	
Calcium methyl paraben	83542-69-2	
Calcium propyl paraben	94-18-8	
Ethyl paraben	120-47-8	

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4-Hydroxy benzoic acid	99-96-7	
Methyl paraben	99-76-3	
Propyl paraben	94-13-3	
Sodium butyl paraben	40167-95-1	
Sodium methyl paraben	5026-62-0	
Sodium propyl paraben	35285-69-9	
Para-formaldehyde	30525-89-4	Not more than 0.25% of final feed
Patent Blue V	129-17-9	
Pectinase	9032-75-1	
Pediococcus acidilactici		
Pediococcus		
pentosaceus		
Peppermint oil		Not for use in cats
Pericol black	2519-30-4	
Phosphoric acid	7664-38-2	
Phyllanthus emblica		
Phytase		From the following sources: Aspergillus niger, var. A. oryzae, var.
Picorhiza kurroa		

Piper longum		
Piper nigrum		
Piper officinarum		
Pistacia integerima		
Plumbago zeylanica		
Polyoxyl 35 castor oil (Cremphor EL)		
Polyoxyl 40 castor oil (Cremphor RH 40)		
Polyoxyl 60 castor oil (Cremphor RH 60)		
Polyoxyethylene nonyl phenol ether	9016-45-9	Only to be used as a wetting agent. Not more than 0.5% of formulated product
Polyethylene oxide, polyproylene glycol block copolymer	6/11/9003	
Polyoxyethylene (20) sorbitan monooleate	9005-65-6	
Polyvinylpyrroli- done	9003-39-8	
Ponceau 4R	2611-82-7	When used at no more than 50 mg/kg in finished feed
Potassium carbonate	584-08-7	

ıltural Compounds and ines Amendment Regula	Veterinary ations 2005 Schedule 2
7447-40-7	
1310-58-3	
996-31-6	
590-00-1	
79-09-4	
109-60-4	
71-23-8	Not more than 55 g/head/day
2315-68-6	
121-79-9	Total content of antioxidants must be not more than 0.02% fat content of feed
57-55-6	
	From the following sources: Aspergillus niger, var. A. oryzae. var. Bacillus amyloliquefaciens, B. licheniformis. B. subtilis, var. B. subtilis, containing a B. amyloliquefaciens gene for protease
	ines Amendment Regula 7447-40-7 1310-58-3 996-31-6 590-00-1 79-09-4 109-60-4 71-23-8 2315-68-6 121-79-9

Pumice

Schedule 2	Agricultural Compounds and Veterinary Medicines Amendment Regulations 2005	Reprinted as at 1 November 2011
Raffinase		
Rapeseed oil	8002-13-9	
Raspberry flavo	ur	
Rennet		
Rosemary	8000-25-7	
Rum ether	8030-89-5	
Rutin	153-18-4	
Saccharin sodiu	m 128-44-9	
Saccharomyces cerevisiae		
Sage oil		
Silica (silicon dioxide)	7631-86-9	
Silicone antifoa	m 63148-62-9	
Skatole	83-34-1	
Smokey bacon flavour		
Sodium acid pyrophosphate		
Sodium alginate	9005-38-3	
Sodium alkyl benzene sulphor	25155-30-0 nate	
Sodium aluminosilicate	73987-94-7	
Sodium ascorba	te 134-03-2	

	lltural Compounds and Vet ines Amendment Regulatio		Schedule 2
Sodium benzoate	532-32-1	Not more t of final fee	
Sodium bicarbonate	144-55-8		
Sodium carbonate	497-19-8		
Sodium carboxy methylcellulose	9004-32-4		
Sodium chloride	7647-14-5		
Sodium citrate	68-04-2		
Sodium cyclamate	139-05-9		
Sodium erythorbate	6381-77-7		
Sodium formate	141-53-7		
Sodium hex- ametaphosphate	10124-56-8		
Sodium hydroxide	1310-73-2		
Sodium lignosulphonate	8061-51-6		
Sodium metabisulphite 7681-57-4			
Sodium nitrite	7632-00-0	Not more t of final fee	
Sodium propionate	137-40-6		
Sodium silico aluminate	1344-00-9	Not more t of final fee	
Sodium tri- polyphosphate	7758-29-4		
Sorbic acid	110-44-1		

Schedule 2	Agricultural Compounds and Veterinary Medicines Amendment Regulations 2005	Reprinted as at 1 November 2011
Sorbitan monostearate	1338-41-6	
Sorbitol	50-70-4	
Strawberry flavo	our	
Streptococcus (Enterococcus) salivarius subspecies thermophilus		
Sulphamic acid	5329-14-6	
Sulphuric acid	7664-93-9	
Sunflower oil	8001-21-6	
Sunset yellow	2783-94-0	
Tagetes oil	8016-84-0	
Tangerine oil	8008-31-9	
Tartaric acid	87-69-4	
Tartrazine	1934-21-0	
Terminalia baler	rica	
Terminalia cheb	ula	
Tertiary butylhydroqui n (TBHQ)	1984-33-0 one	
Tetra potassium pyro-phosphate	7758-87-4	
Tetra sodium pyrophosphate	7722-88-5	
Thaumatin	53850-34-3	

November 2011 Medici	nes Amendment Regula		Schedule
Thyme oil	8007-46-3		
Thymol	89-83-8	When ad levels con with good practice	nsistent
Titanium dioxide	13463-67-7		
Tocopherols (extracts of natural origin)	1406-66-2		
Tricalcium phosphate	7758-87-4		
Trimethylamine	75-50-3		
Trypsin			
Turmeric	8024-37-1		
Undecylenic alcohol	112-43-6		
Valerian			
Valeric acid	109-52-4		
Vanillin	121-33-5		
Vermiculite	1318-00-9		
Vitamin B1			
Vitamin B12			
Xanthan gum	11138-66-2		
Xanthophyll	127-40-2		

	Agricultural Compounds and Veterinary	Reprinted as at
Schedule 2	Medicines Amendment Regulations 2005	1 November 2011

Xylanase	From Aspergillus oryzae carrying a gene from Thermomyces lanuginosus coding for xylanase		
Yucca schidigera			
Zeaxanthin			
Zinc oxide	1314-13-2		
Zinc proprionates			
Zingiber officinale			
Diane Morcom, Clerk of the Executive Council.			

Explanatory note

This note is not part of the regulations, but is intended to indicate their general effect.

These regulations, which come into force on the 28th day after their notification in the *Gazette*, amend the Agricultural Compounds and Veterinary Medicines Regulations 2001 ("the principal regulations").

The amendment—

- inserts new regulation 5A into the principal regulations, which provides that a combined agricultural compound—
 - is exempt from registration under section 21 or 27 of the Agricultural Compounds and Veterinary Medicines Act 1997 (the Act) if the agricultural compound is a combination of 2 or more agricultural compounds that are exempt from registration under these regulations:

is subject to any—

• prescribed conditions; and

- other provision of these regulations that applies to each agricultural compound that comprises the combined agricultural compound:
- revokes regulation 7 of the principal regulations (which relates to reports on agricultural compounds):
- substitutes the following schedules:
 - Schedule 1 (which lists agricultural compounds that are exempt from registration if applicable codes of practice are complied with) has been amended to insert 3 new compounds into, and to exclude 5 compounds from, item 1 of the former Schedule 1:
 - *Schedule 2* (which lists agricultural compounds that are exempt from registration if certain conditions are complied with) has been amended to include the compounds listed in the former Schedule 3 of the principal regulations, and to insert—
 - a new condition in relation to oral and topical preparations:
 - a new compound into the schedule, namely, any compound that is used to extend animal semen or to be used as media for animal ova:
 - New *Schedule 3* (which lists excluded ingredients for topical preparations):
 - Schedule 4 (which lists conditions that apply to oral nutritional compounds exempt from registration) has been amended to include a new criteria for determining whether an oral nutritional compound is fit for its purpose, namely, that its use does not result in physical harm causing pain or distress in an animal:
 - Schedule 7 (which lists substances that are generally recognised as safe feed additives in oral nutritional compounds) has been amended by removing Part B and inserting new items into the schedule.

Issued under the authority of the Acts and Regulations Publication Act 1989. Date of notification in *Gazette*: 14 April 2005.

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Notes

1 General

This is an eprint of the Agricultural Compounds and Veterinary Medicines Amendment Regulations 2005. The eprint incorporates all the amendments to the regulations as at 1 November 2011. The list of amendments at the end of these notes specifies all the amendments incorporated into this eprint since 3 September 2007.

Relevant provisions of any amending enactments that contain transitional, savings, or application provisions that cannot be compiled in the eprint are also included, after the principal enactment, in chronological order.

2 About this eprint

This eprint has not been officialised. For more information about eprints and officialisation, please *see* http://www.pco.parliament.govt.nz/eprints/.

3 List of amendments incorporated in this eprint (most recent first)

Agricultural Compounds and Veterinary Medicines (Exemptions and Prohibited Substances) Regulations 2011 (SR 2011/327): regulation 16