

*Suppliment tal-Gazzetta tal-Gvern ta' Malta Nru. 18,623, 23 ta' Lulju, 2010*

*Taqsimi B*

## A.L. 371 ta' l-2010

### ATT DWAR IL-KONTROLL TAL-PESTIĆIDI (KAP. 430)

#### Regolamenti ta' l-2010 li jemendaw ir-Regolamenti dwar il-Bijoċidi (Emenda Nru. 3)

BIS-SAĦHA tas-setgħat mogħtija bl-artikoli 4 u 5 ta' l-Att dwar il-Kontroll tal-Pestiċidi, il-Ministru għar-Riżorsi u Affarijiet Rurali, wara li kkonsulta lill-Prim Ministro u lill-Ministro għas-Saħħha, l-Anzjani u Kura fil-Komunità, għamel dawn ir-regolamenti li ġejjin:-

**1.** (1) It-titolu ta' dawn ir-regolamenti hu Regolamenti ta' l-2010 li jemendaw ir-Regolamenti dwar il-Bijoċidi (Emenda Nru. 3), u għandhom jinqraw u jiftieħmu haġa waħda mar-Regolamenti ta' l-2004 dwar il-Bijoċidi, hawn iżjed 'il quddiem imsejħin "ir-regolamenti prinċipali".

Titolu u skop.

A.L. 294 ta' l-2004.

(2) L-iskop ta' dawn ir-regolamenti hu li jittrassponu d-Direttiva tal-Kummissjoni 2010/5/UE tat-8 ta' Frar 2010 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi l-*acrolein* bħala sustanza attiva fl-Anness I għaliha, Direttiva tal-Kummissjoni 2010/7/UE tad-9 ta' Frar 2010 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi l-*magnesium phosphide releasing phosphine* bħala sustanza attiva fl-Anness I għaliha, Direttiva tal-Kummissjoni 2010/8/UE tad-9 ta' Frar 2010 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi l-*warfarin sodium* bħala sustanza attiva fl-Anness I għaliha, Direttiva tal-Kummissjoni 2010/9/UE tad-9 ta' Frar 2010 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex testendi l-inklużjoni fl-Anness I tagħha tas-sustanza attiva *aluminium phosphide* li jirrilaxxa l-*phosphine* għat-tip ta' prodott 18 kif definit fl-Anness V tagħha, Direttiva tal-Kummissjoni 2010/10/UE tad-9 ta' Frar 2010 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi l-*brodifacoum* bħala sustanza attiva fl-Anness I għaliha, Direttiva tal-Kummissjoni 2010/11/UE tad-9 ta' Frar 2010 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi l-*warfarin* bħala sustanza attiva fl-Anness I għaliha.

**2.** Minflok il-Parti I ta' l-Iskeda III li tinsab mar-regolamenti prinċipali, għandu jidħol dan li ġej:

Jemenda Skeda  
III li tinsab mar-regolamenti prinċipali.

**"PARTI I**

Tabella għall-Implimentazzjoni ta' Miżuri fi Stati Membri għal Sustanzi Attivi awtorizzati biex jintużaw fil-Bijoċidi

No	Isem Komuni	Direttiva ta' inklužjoni	Data ta' implementazzjoni (*)
1	Sulfuryl Flouride	Direttiva tal-Kummissjoni 2006/140/KE	1 ta' Jannar 2009
		Direttiva tal-Kummissjoni 2009/84/KE	1 ta' Lulju 2011
2	Dichlofluanid	Direttiva tal-Kummissjoni 2007/20/KE	1 ta' Marzu 2009
		2007/20/KE	
3	Difethialone	Direttiva tal-Kummissjoni 2007/69/KE	1 ta' Novembru 2009
4	Clothianidin	Direttiva tal-Kummissjoni 2008/15/KE	1 ta' Frar 2010
5	Etofenprox	Direttiva tal-Kummissjoni 2008/16/KE	1 ta' Frar 2010
6	Tebuconazole	Direttiva tal-Kummissjoni 2008/86/KE	1 t'April 2010
7	Carbon Dioxide	Direttiva tal-Kummissjoni 2008/75/KE	1 ta' Novembru 2009
8	Propiconazole	Direttiva tal-Kummissjoni 2008/78/KE	1 t'April 2010
9	Difenacoum	Direttiva tal-Kummissjoni 2008/81/KE	1 t'April 2010
10	K-HDO	Direttiva tal-Kummissjoni 2008/80/KE	1 ta' Lulju 2010
11	IPBC	Direttiva tal-Kummissjoni 2008/79/KE	1 ta' Lulju 2010
12	Chlorophacinone	Direttiva tal-Kummissjoni 2009/99/KE	1 ta' Lulju 2011
13	Thiabendazole	Direttiva tal-Kummissjoni 2008/85/KE	1 ta' Lulju 2010
14	Thiamethoxam	Direttiva tal-Kummissjoni 2008/77/KE	1 ta' Lulju 2010
15	Alphachloralose	Direttiva tal-Kummissjoni 2009/93/KE	1 ta' Lulju 2011
16	Brodifacoum	Direttiva tal-Kummissjoni 2010/10/UE	1 ta' Frar 2012
17	Bromadiolone	Direttiva tal-Kummissjoni 2009/92/KE	1 ta' Lulju 2011
18	Thiacloprid	Direttiva tal-Kummissjoni 2009/88/KE	1 ta' Jannar 2010

19	Indoxacarb	Direttiva tal-Kummissjoni 2009/87/KE	1 ta' Jannar 2010
20	Aluminium phosphide	Direttiva tal-Kummissjoni 2009/95/KE	1 ta' Settembru 2011
		Direttiva tal-Kummissjoni 2010/9/UE	1 ta' Frar 2012
21	Fenpropimorph	Direttiva tal-Kummissjoni 2009/86/KE	1 ta' Lulju 2011
22	Boric acid	Direttiva tal-Kummissjoni 2009/94/KE	1 ta' Settembru 2011
23	Boric Oxide	Direttiva tal-Kummissjoni 2009/98/KE	1 ta' Settembru 2011
24	Disodium tetraborate	Direttiva tal-Kummissjoni 2009/91/KE	1 ta' Settembru 2011
25	Disodium octaborate tetrahydrate	Direttiva tal-Kummissjoni 2009/96/KE	1 ta' Settembru 2011
26	Magnesium phosphide releasing phosphine	Direttiva tal-Kummissjoni 2010/7/UE	1 ta' Frar 2012
27	Nitrogen	Direttiva tal-Kummissjoni 2009/89/KE	1 ta' Settembru 2011
28	Coumatetralyl	Direttiva tal-Kummissjoni 2009/85/KE	1 ta' Lulju 2011
29	Tolylfluanid	Direttiva tal-Kummissjoni 2009/151/KE	1 t'Ottubru 2011
30	Acrolein	Direttive tal-Kummissjoni 2010/5/UE	1 ta' Settembru 2010
31	Flocoumafen	Direttiva tal-Kummissjoni 2009/150/KE	1 t'Ottubru 2011
32	Warfarin	Direttiva tal-Kummissjoni 2010/11/UE	1 ta' Frar 2012
33	Warfarin sodium	Direttive tal-Kummissjoni 2010/8/UE	1 ta' Frar 2012".

**L.N. 371 of 2010****PESTICIDES CONTROL ACT  
(CAP. 430)****Biocides (Amendment) (No. 3) Regulations, 2010**

IN exercise of the powers conferred by articles 4 and 5 of the Pesticides Control Act, the Minister for Resources and Rural Affairs, in consultation with the Prime Minister and with the Minister for Health, the Elderly and Community Care, has made the following regulations:-

Title and scope.

L.N. 294 of 2004.

**1.** (1) The title of these regulations is the Biocides (Amendment) (No. 3) Regulations, 2010 and they shall be read and construed as one with Biocides Regulations, 2004, hereinafter referred to as “the principal regulations”.

(2) The scope of these regulations is to transpose Commission Directive 2010/5/EU of 8 February 2010 amending Directive 98/8/EC of the European Parliament and of the Council to include acrolein as an active substance in Annex I thereto, Commission Directive 2010/7/EU of 9 February 2010 amending Directive 98/8/EC of the European Parliament and of the Council to include magnesium phosphide releasing phosphine as an active substance in Annex I thereto, Commission Directive 2010/8/EU of 9 February 2010 amending Directive 98/8/EC of the European Parliament and of the Council to include warfarin sodium as an active substance in Annex I thereto, Commission Directive 2010/9/EU of 9 February 2010 amending Directive 98/8/EC of the European Parliament and of the Council to extend the inclusion in Annex I thereto of the active substance aluminium phosphide releasing phosphine to product type 18 as defined in Annex V thereto, Commission Directive 2010/10/EU of 9 February 2010 amending Directive 98/8/EC of the European Parliament and of the Council to include brodifacoum as an active substance in Annex I thereto, Commission Directive 2010/11/EU of 9 February 2010 amending Directive 98/8/EC of the European Parliament and of the Council to include warfarin as an active substance in Annex I thereto.

Amends Schedule III to the principal regulations.

**2.** For Part I of Schedule III to the principal regulations, there shall be substituted the following new part:-

## “PART I

Time Table for Implementation Measures in Member States for Active  
Substances authorized for use in Biocides

No	Common Name	Inclusion Directive	Date of implementation (*)
1	Sulfuryl Flouride	Commission Directive 2006/140/EC	1 January 2009
		Commission Directive 2009/84/EC	1 July 2011
2	Dichlofluanid	Commission Directive 2007/20/EC	1 March 2009
3	Difethialone	Commission Directive 2007/69/EC	1 November 2009
4	Clothianidin	Commission Directive 2008/15/EC	1 February 2010
5	Etofenprox	Commission Directive 2008/16/EC	1 February 2010
6	Tebuconazole	Commission Directive 2008/86/EC	1 April 2010
7	Carbon Dioxide	Commission Directive 2008/75/EC	1 November 2009
8	Propiconazole	Commission Directive 2008/78/EC	1 April 2010
9	Difenacoum	Commission Directive 2008/81/EC	1 April 2010
10	K-HDO	Commission Directive 2008/80/EC	1 July 2010
11	IPBC	Commission Directive 2008/79/EC	1 July 2010
12	Chlorophacinone	Commission Directive 2009/99/EC	1 July 2011
13	Thiabendazole	Commission Directive 2008/85/EC	1 July 2010
14	Thiamethoxam	Commission Directive 2008/77/EC	1 July 2010
15	Alphachloralose	Commission Directive 2009/93/EC	1 July 2011
16	Brodifacoum	Commission Directive 2010/10/EU	1 February 2012
17	Bromadiolone	Commission Directive 2009/92/EC	1 July 2011
18	Thiacloprid	Commission Directive 2009/88/EC	1 January 2010

19	Indoxacarb	Commission Directive 2009/87/EC	1 January 2010
20	Aluminium phosphide	Commission Directive 2009/95/EC	1 September 2011
		Commission Directive 2010/9/EU	1 February 2012
21	Fenpropimorph	Commission Directive 2009/86/EC	1 July 2011
22	Boric acid	Commission Directive 2009/94/EC	1 September 2011
23	Boric Oxide	Commission Directive 2009/98/EC	1 September 2011
24	Disodium tetraborate	Commission Directive 2009/91/EC	1 September 2011
25	Disodium octaborate tetrahydrate	Commission Directive 2009/96/EC	1 September 2011
26	Magnesium phosphide releasing phosphine	Commission Directive 2010/7/EU	1 February 2012
27	Nitrogen	Commission Directive 2009/89/EC	1 September 2011
28	Coumatetralyl	Commission Directive 2009/85/EC	1 July 2011
29	Tolylfluanid	Commission Directive 2009/151/EC	1 October 2011
30	Acrolein	Commission Directive 2010/5/EU	1 September 2010
31	Flocoumafen	Commission Directive 2009/150/EC	1 October 2011
32	Warfarin	Commission Directive 2010/11/EU	1 February 2012
33	Warfarin sodium	Commission Directive 2010/8/EU	1 February 2012”.

