
A.L. 16 ta' l-2009

**ATT DWAR IL-KONTROLL TAL-PESTIĊIDI
(KAP. 430)**

**Regolamenti ta' l-2009 li jemendaw ir-Regolamenti
dwar il-Bijoċidi**

BIS-SAHHA 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 Ministru u lill-Ministru għall-Politika Soċjali, għamel dawn ir-regolamenti li ġejjin:-

1. (1) It-titolu ta' dawn ir-regolamenti hu Regolamenti ta' l-2009 li jemendaw ir-Regolamenti dwar il-Bijoċidi, u għandhom jinqraw u jiftiehm u bħala 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 jitrassponu d-Direttiva tal-Kummissjoni 2007/47/KE tal-Parlament Ewropew u tal-Kunsill tal-5 ta' Settembru 2007 li temenda d-Direttiva tal-Kunsill 90/385/KEE fuq approssimazzjoni tal-liġijiet tal-Istati Membri rigward il-mezzi mediċi attivi li jiddaħhlu f'xi parti tal-ġisem, d-Direttiva tal-Kunsill 93/42/KEE dwar mezzi mediċi u d-Direttiva 98/8/KE dwar it-tqegħid fis-suq talprodotti bijoċidali, Direttiva tal-Kummissjoni 2008/15/KE tal-15 ta' Frar, 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi l-clothianidin bħala sustanza attiva fl-Anness I għaliha, Direttiva tal-Kummissjoni 2008/16/KE tal-15 ta' Frar 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi letofenprox bħala sustanza attiva fl-Anness I tagħha, Direttiva tal-Kummissjoni 2008/75/KE ta' l-24 ta' Lulju 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi d-dijossidu tal-karbonju bħala sustanza attiva fl-Anness I tagħha, Direttiva tal-Kummissjoni 2008/77/KE tal-25 ta' Lulju 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi tthiamethoxam bħala sustanza attiva fl-Anness I għaliha, Direttiva tal-Kummissjoni 2008/78/KE tal-25 ta' Lulju 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi lpropiconazole bħala sustanza attiva fl-

Anness I tagħha, Direttiva tal-Kummissjoni 2008/79/KE tat-28 ta' Lulju 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi l-IPBC bħala sustanza attiva fl-Anness I għaliha, Direttiva tal-Kummissjoni 2008/80/KE tat-28 ta' Lulju 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi l-melħ tal-potassju tascyclohexylhydroxydiazene 1-oxide (K-HDO) bħala sustanza attiva fl-Anness I għaliha, Direttiva tal-Kummissjoni 2008/81/KE tad-29 ta' Lulju 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi d-difenacoum bħala sustanza attiva fl-Anness I tagħha, Direttiva tal-Kummissjoni 2008/85/KE tal-5 ta' Settembru 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi t-thiabendazole bħala sustanza attiva fl-Anness I għaliha u Direttiva tal-Kummissjoni 2008/86/KE tal-5 ta' Settembru 2008 li temenda d-Direttiva 98/8/KE tal-Parlament Ewropew u tal-Kunsill biex tinkludi t-tebuconazole bħala sustanza attiva fl-Anness I tagħha.

Jemenda r-regolament
2 tar-regolamenti
prinċipali.

2. Fir-regolament 2(2) tar-regolamenti prinċipali, minnufih wara l-partita (r) għandha tiżdied l-partita ġdida li ġejja:-

“(s) disposizzjonijiet dwar il-mezzi mediċi dijanjostiċi in vitro.”.

Jemenda Skeda
III li tinsab mar-
regolamenti prinċipali.

3. Fi Skeda III li tinsab mar-regolamenti prinċipali, minnufih wara l-partita 3 għandhom jiżdiedu dawn il-partiti ġodda li ġejjin:

L.N. 16 of 2009**PESTICIDES CONTROL ACT
(CAP. 430)****Biocides (Amendment) Regulations, 2009**

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 Social Policy, has made the following regulations:-

Title and scope.

L.N. 294 of 2004.

1. (1) The title of these regulations is the Biocides (Amendment) Regulations, 2009 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 2007/47/EC of the European Parliament and of the Council of 5 September 2007 amending Council Directive 90/385/EEC on the approximation of the laws of the Member States relating to active implantable medical devices, Council Directive 93/42/EEC concerning medical devices and Directive 98/8/EC concerning the placing of biocidal products on the market, Commission Directive 2008/15/EC of 15 February 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include clothianidin as an active substance in Annex IA thereto, Commission Directive 2008/16/EC of 15 February 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include etofenprox as an active substance in Annex I thereto, Commission Directive 2008/75/EC of 24 July 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include carbon dioxide as an active substance in Annex I thereto, Commission Directive 2008/77/EC of 25 July 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include thiamethoxam as an active substance in Annex I thereto, Commission Directive 2008/78/EC of 25 July 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include propiconazole as an active substance in Annex I thereto, Commission Directive 2008/79/EC of 28 July 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include IPBC as an active substance in Annex I thereto, Commission Directive 2008/80/EC of 28 July 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include cyclohexylhydroxydiazene 1-oxide, potassium salt (K-HDO), Commission Directive 2008/81/

EC of 29 July 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include difenacoum as an active substance in Annex I thereto, Commission Directive 2008/85/EC of 5 September 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include thiabendazole as an active substance in Annex I thereto, and Commission Directive 2008/86/EC of 5 September 2008 amending Directive 98/8/EC of the European Parliament and of the Council to include tebuconazole as an active substance in Annex I thereto.

2. (1) In regulation 2(2) of the principal regulations, immediately after item (r) thereof, there shall be added the following:-

Amends regulation 2 of the principal regulations.

“(s) provisions concerning in vitro diagnostic medical devices.”

3. In Schedule III to the principal regulations, immediately after item 3 thereof, there shall be added the following new items:-

Amends Schedule III to the principal regulations.

