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THE PEST CONTROL PRODUCTS (REGISTRATION) REGULATIONS

NO. 46 OF 1984

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Pest Control Products Act

The Pest Control Products (Registration) Regulations Legal Notice 46 of 1984

Legislation as at 31 December 2022

Note: This Legal Notice was **repealed** on 1984-04-06 by [The Pest Control Products \(Registration\) Regulations](#) (Legal Notice 46 of 1984).

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The Pest Control Products (Registration) Regulations (Legal Notice 46 of 1984)
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PEST CONTROL PRODUCTS ACT
THE PEST CONTROL PRODUCTS
(REGISTRATION) REGULATIONS
LEGAL NOTICE 46 OF 1984

Published in Kenya Gazette Vol. LXXXVI—No. 15 on 6 April 1984

Commenced on 6 April 1984

[Amended by [Pest Control Products \(Registration\) \(Amendment\) Regulations, 1984 \(Legal Notice 109 of 1984\)](#) on 6 April 1984]

[Amended by [Pest Control Products \(Registration\) \(Amendment\) Regulations, 2006 \(Legal Notice 123 of 2006\)](#) on 13 October 2006]

[Amended by [Pest Control Products \(Registration\) \(Amendment\) Regulations, 2014 \(Legal Notice 122 of 2014\)](#) on 12 September 2014]

[Amended by [Pest Control Products \(Registration\) \(Amendment\) Regulations, 2015 \(Legal Notice 124 of 2015\)](#) on 17 July 2015]

[Revised by [24th Annual Supplement \(Legal Notice 221 of 2023\)](#) on 31 December 2022]

[Revoked by [The Pest Control Products \(Registration\) Regulations \(Legal Notice 46 of 1984\)](#) on 6 April 1984]

1. Citation

These Regulations may be cited as the Pest Control Products (Registration) Regulations.

2. Interpretation

In these Regulations, unless the context otherwise requires—

“accredited scientists/institution” means a person or an institution that has been officially recognized by the Board as having the capacity and competence to undertake biological efficacy trials;

“active ingredient” means an ingredient of a pest control product to which the effects of the pest control product are attributed and includes a synergist; but does not include a component that by itself is not primarily responsible for the control effect of the pest control product;

“biochemical pesticide” means a pest control product whose active ingredient constitutes a chemical derived from naturally occurring plant or animal intended to control invertebrate pests;

“certificate of registration” means a certificate issued by the Board under regulation 7;

“device” means any article, instrument, apparatus, contrivance or gadget that by itself or in conjunction with a pest control product is used as a means to control pests directly or indirectly;

“experimental permit” means a permit issued by the Board for small quantity of a pest control product imported or produced locally for purposes of research and efficacy trials prior to consideration for registration;

“microbial and macrobial biopesticide” means a pest control product of naturally occurring micro-organisms (microbiological agents viruses and rickettsia; bacteria, protozoa, fungi,) and macro-organisms (macro biological agents such as predators, parasitoids and entomopathogenic nematodes), respectively intended for the control of invertebrate pests weeds, pathogens of crops, and pests of livestock and public health and to which effects of the pest control products or active agent are attributed but does not include a component that by itself is not primarily responsible for the control effect of the pest control product or genetically modified living microorganism and macro-organism;

“national collection number” means the unique code given to a culture or an isolate by the National Museums of Kenya;

“parallel/daughter registration” means a registration of a trade name based on the strength of an existing fully registered product from the same manufacturer and source and with authorization from the registrant;

“residue” means the ingredient of a pest control product that remains after the pest control product has been used and includes substances resulting from degradation or metabolism.

[L.N. 123/2006, r. 2, L.N. 124/2015, r. 2.]

3. Exemption from registration

A pest control product shall be exempt from registration if—

- (a) it is for use by a person for research purposes if that use has been approved by the Board;
- (b) it is a type or kind set out in the First Schedule and meets the conditions relevant to that substance as set out in that Schedule.

3A. Efficacy testing

- (1) Every person desiring to introduce a pest control product for efficacy testing shall—
 - (a) make application to the Board for an experimental permit in Form C set out in the Second Schedule;
 - (b) provide all the details required in the form;
 - (c) on request supply any further information which may be required by the Board; and
 - (d) pay the prescribed application fees determined by the Board from time to time therefor.
- (2)
 - (a) The Board shall evaluate the data provided and if satisfied that the application merits approval, shall issue an experimental permit in the prescribed Form D in the Second Schedule
 - (b) The Board shall, in addition, give the applicant information relating to the existing accredited scientists or institutions in the field of trial whom the applicant will work with;
- (3) When the efficacy trials are complete the accredited scientist or institution shall submit efficacy reports to the Board.

[L.N. 123/2006, r. 3.]

4. Application for registration of pest control products

- (1) Every person desiring to register a pest control product shall make application to the Board in either Forms A, A1, A2, A3, A3a, A3b, A4 or A5 set out in the Second Schedule and shall, on request, supply any further information which may be required by the Board.
- (2) An applicant who is not resident in Kenya shall appoint an agent permanently resident in Kenya to whom any notice or correspondence may be sent.
- (2A) An application for registration of an agent shall be submitted in Form A6 set out in the Second Schedule.
- (2B) An application for registration for change of an agency shall be submitted in Form A7 set out in the Second Schedule.
- (3) An application for the registration of a pest control product shall be accompanied by five copies of the proposed label for the pest control product or reasonable facsimiles thereof.

- (4) An applicant who is not resident in Kenya shall be required to deposit with the Board a binding agreement entered with the agent permanently resident in Kenya.

[L.N. 123/2006, rr. 4, 5 & 6, L.N. 124/2015, r. 3.]

4A. Application for registration of a synthetic or conventional pest control product

- (1) The application for registration of a synthetic or conventional pest control product (non-generic) under regulation 4 (1) shall be in the prescribed Form A set out in the Second Schedule completed by the applicant or duly authorized person and submitted in triplicate.
- (2) The application for registration of generic conventional pest control products shall be submitted in Form A4.
- (3) The Board shall supply the applicant with checklists and an index to ensure that the applicant has supplied the relevant data required in Form A set out in the Second Schedule.

[L.N. 124/2015, r. 4.]

4B. Application for the registration of microbial biopesticide

- (1) An application for the registration of microbial biopesticide shall be in Form A1 set out in the Second Schedule.
- (2) Information in support of a request for registration, both published and unpublished (fully cited), shall be supplied in the form of a summary data sheet laid out in the format given in Form A1 set out in the Second Schedule.
- (3) Pre-registration consultations between the applicant and the Board shall be undertaken after the application is made.
- (4) All applicants intending to import or export live organisms into or out of the country shall comply with all other existing laws governing such organisms.
- (5) The use of genetically modified organisms as microbial biopesticides shall comply with any other existing laws governing such organisms before an application is made to the Board.
- (6) The Board shall supply checklists and an index to ensure that the applicant has provided all relevant data and cited material.

[L.N. 124/2015, r. 5.]

4C. Application for the registration of macrobial biopesticide

- (1) An application for the registration of macrobial biopesticide shall be as set out in Form A2 in the Second Schedule.
- (2) Information in support of a request for registration, both published and unpublished (fully cited), shall be supplied in the form of a summary data sheet laid out in the format given in Form A2 set out in the Second Schedule.
- (3) Pre-registration consultations between the applicant and the Board shall be undertaken after the application is made.
- (4) The applicant shall be required to-
 - (a) submit a sample of the pest control product to the National Museums of Kenya or the national collection number obtained if the culture is already in collection;
 - (b) provide a sample of the technical grade of its active agent;

- (c) send an additional sample to the National Agricultural Research Laboratories, Biological Control Unit, Kenya Agricultural Research Institute and Kenya Plant Health Inspectorate Service;
 - (d) supply any other sample as may be requested by the Board.
- (5) All applicants intending to import or export live organisms into or out of the country shall comply with any other existing laws governing such organisms.
- (6) The use of genetically modified organisms and living modified organisms as microbial biopesticides shall comply with any other existing laws governing such organisms before an application is made to the Board.
- (7) The Board shall supply checklists and an index to ensure that the applicant has provided all relevant data and cited material.

[L.N. 124/2015, r. 6.]

4D. Application form for the registration of a biochemical pesticide other than semiochemicals

- (1) The application form for the registration of a biochemical pesticide other than semiochemicals, shall be in Form A3 in the Second Schedule.
- (2) Information in support of a request for registration both published and unpublished shall be supplied in form of a summary data sheet laid out according to the format given in Form A3.
- (3) Pre-registration consultation between the applicant and the registration authority shall be undertaken.
- (4) The application for registration of semiochemicals pest control products for monitoring and control of pests in crops, livestock and public health shall be in Form A3a.
- (5) The application for introduction of straight chained lepidopteran pheromone for monitoring purposes shall be in Form A3b.

[L.N. 124/2015, r. 7.]

4E. Application form for registration of spray adjuvants

- (1) The application form for registration of spray adjuvants shall be in Form A5 as set out in the Second Schedule.
- (2) Information in support of a request for registration, both published and unpublished (fully cited) shall be supplied in the form of a summary data sheet laid out in Form A5 in the Second Schedule.
- (3) The Board shall supply checklists and an index to ensure that the applicant has provided all relevant data and cited material.

[L.N. 124/2015, r. 8.]

4F. Registration of a parallel/daughter pest control product

- (1) The Board may upon such terms and conditions as it may specify, on payment of the introductory and registration fees, register a parallel/daughter pest control product where the applicant-
 - (a) completes Form C set out in the Second Schedule;
 - (b) provides a letter of access from registrant and a letter of no objection from the local agent;
 - (c) submits a letter of no objection from the trade name owner where the trade name owner is not the registrant;

Repealed

- (d) borrows approved label of the original registered product and only changes the trade name.
- (2) Each parallel/daughter registration shall have its own registration number which shall be linked to the original registered product by indicating the original registered product number on the parallel registration certificates.
 - (3) Voluntary cancellation of a product shall apply to the registered product and the parallel/daughter products.
 - (4) A company may leave an area without canceling the parallel/daughter registration and may transfer access of the original dossier thereto.
 - (5) The parallel/daughter registration shall be automatically revoked when the registrant withdraws the letter of access.
 - (6) The parallel/daughter registration shall not be used to register other different products.
 - (7) A new dossier shall not be required for the registration of the parallel/daughter products;
 - (8) The original registered product and the parallel/daughter registration shall be required to originate from the same source.
 - (9) Parallel/daughters registrations shall be exempt from local efficacy trials if the intended use is identical to that of the original registered product.
 - (10) Efficacy trials shall be undertaken where new uses which are different from those of the original registered products.
 - (11) The Board shall exercise discretion in determining the number of parallel/daughter products to be registered on a case-by-case basis but not more than five such products shall be registered in respect of one original product.

[L.N. 124/2015, r. 8.]

4G. Extension of the use of a registered pest control product

The Board may, upon such terms and conditions as it may specify, extend the use of a registered pest control product through a label extension where the applicant submits-

- (a) successful two-season efficacy trial data on the respective area of use such as crop or pest combination;
- (b) residue data based on the Good Agriculture Practice in the efficacy trial, if the new use is on edible crops or animals;
- (c) a copy of the previously approved label;
- (d) revised commercial label with the proposed new uses, rates, pre-harvest interval and re-entry interval and the final commercial version label;
- (e) proposed maximum residue limits and pre-harvest intervals for edible commodities, withdrawal period for livestock and re-entry interval for greenhouse use.

[L.N. 124/2015, r. 8.]

4H. Summary dossier

The application for registration of pest control products shall be accompanied by a copy of a summary dossier as prescribed in form-

- (a) B for conventional pest control products;
- (b) B1 for microbial pest control products;

- (c) B2 for macrobial pest control products;
- (d) B3 for biochemical pest control products other than semiochemicals; and
- (e) B4 for semiochemicals.

[L.N. 124/2015, r. 8.]

5. Applicant to provide samples

- (1) An applicant shall, when requested to do so by the Board, provide—
 - (a) a sample of the pest control product;
 - (b) a sample of the technical grade of its active ingredient;
 - (c) a sample of the laboratory standard of its active ingredient; and
 - (d) any other sample as may be required by the Board.
- (2) The samples shall be submitted to the Board in Form A8 set out in the Second Schedule.

[L.N. 124/2015, r. 9.]

6. Registration fees

The fees payable by an applicant for the registration of a pest control product shall be the prescribed fees determined by the Board from time to time where a pest control product is a device or it contains an active ingredient that has been previously assessed or evaluated for the purpose of the Act and these Regulations.

[L.N. 123/2006, r. 7.]

7. Issue of certificate of registration

- (1) The Board shall consider the application under regulation 4, and, if it is satisfied of the safety, efficacy, quality and economic value of the pest control product, shall register the pest control product, and issue a certificate of registration which shall be in Form B set out in the Second Schedule.
- (1A) The applicant shall submit the proposed trade name for consideration by the Board.
- (1B) The trade name in sub-regulation (1A) may be changed upon request to the Board in Form A9 set out in the Second Schedule.
- (1C) The fees payable by an applicant for the change of a trade name of a pest control product shall be ten thousand shillings.
- (2) If the Board is not satisfied as to the safety, efficacy, quality and economic value of the pest control product it may, after providing an opportunity for the applicant to be heard, reject the application for the registration of the pest control product and inform the applicant the reasons for the rejection in writing.
- (3) No person to whom a certificate of registration has been issued under this Regulation shall lend, hire, sell, transfer or otherwise dispose of the certificate to any other person without the approval of the Board, which approval shall be endorsed on the certificate of registration.

[L.N. 124/2015, r. 10.]

8. Duration and renewal of certificate of registration

- (1) A certificate of registration issued under these Regulations shall, unless earlier suspended or revoked, be valid for a period of three years from the date of issue and may thereafter be renewed for periods not exceeding two years at any one time.
- (2) The fee for the renewal of a certificate of registration shall be the prescribed fees determined by the Board from time to time, and an application for renewal shall be accompanied by five copies of the current label for the pest control product.
- (3) A holder of a certificate of registration issued under these Regulations shall give a notice to the Board in writing at least 3 months before the expiry of the registration of any intentions to keep the product registration in abeyance for a period not exceeding 5 years and the notice shall—
 - (a) give reasons for temporary withdrawal; and
 - (b) show the records of all quantities of the pest control product in stock, manufactured or sold by him.
- (4) The Board shall consider the notification under subregulation (3) and if it is satisfied with the reasons for temporary withdrawal shall suspend the registration of the pest control product for a period not exceeding 5 years.
- (5) The information on suspended registration under subregulation (4) shall be made known to the holder in writing and the general public by Gazette notice.
- (6) A person whose certificate of registration has been suspended under subregulation (4) shall withdraw the product from the market within a period of 3 months from the date of expiry of registration.
- (7) A person whose certificate of registration has been suspended under these Regulations shall give a notice to the Board in writing of any intentions to reintroduce the product registration and the notice shall—
 - (a) give reasons for reintroduction,
 - (b) be accompanied by a fee for the renewal of a certificate of registration for the preceding two years and the current year,
 - (c) be accompanied by five copies of the current label for the pest control product.
- (8) A holder of a certificate of registration issued under these Regulations whose product registration has been suspended for a period exceeding five years shall apply for registration afresh and shall, on request supply any further information, which may be required by the Board.

[L.N. 123/2006, rr. 8 & 9.]

9. Temporary registration

- (1) The Board may upon such terms and conditions as it may specify, on payment of a prescribed fee determined by the Board from time to time, register a pest control product for a period not exceeding one year where—
- (2) Any terms and conditions specified by the Board under paragraph (1) shall be contained in the temporary certificate of registration.

[L.N. 123/2006, r. 10.]

10. Refusal to register pest control

The Board may refuse to register a pest control product if in its opinion—

- (a) the applicant for registration or the label for the pest control product does not comply with, the provisions of the Act and these Regulation;
- (b) the information provided to the Board by the applicant is insufficient to enable the pest control product to be assessed or evaluated;
- (c) the applicant fails to establish that the pest control product has merit or value for the purpose claimed when the pest control product is used in accordance with its label directions; or
- (d) the use of the pest control product would lead to an unacceptable risk or harm to—
 - (i) things on or in relation to which the pest control product is intended to be used; or
 - (ii) public health, plants, animals or the environment.

11. Suspension and revocation of certificates of registration

- (1) The Board may suspend, amend or revoke a certificate of registration issued under these Regulations for such time as the Board may determine.
- (2) The powers conferred by paragraph (1) shall not be exercised by the Board except on one or more of the following grounds—
 - (a) that the matters stated in the application on which the certificate of registration was granted were false or incomplete in a material particular;
 - (b) that new information has become available to the Board which renders the pest control product unsafe or dangerous;
 - (c) that the premises on which, or on part of which, the pest control product is manufactured, assembled or stored by or on behalf of the holder of the certificate of registration are unsuitable for the manufacturing, assembly or storage of pest control products;
 - (d) that the holder of a certificate of registration has given a notice to the Board in writing of any intentions to suspend product registration for a period not exceeding 5 years;
 - (e) that new information has become available to the Board indicating that the pest control product is sourced from a manufacturer, formulator, repacker or any facility other than that specified in the application forms and dossier for registration for the respective pest control product or sources authorized by the Board;
 - (f) that the principal or registrant withdraws the technical support to the local agent or distributor on the basis of which a pest control product was registered, in writing;
 - (g) that the new information has become available to the Board which renders the pest control product ineffective, not efficacious or of no economic value for the purposes for which it is intended;
 - (h) that a holder of a registered pest control product has in connection with the registration concerned, contravened or failed to comply with a provision of this Act;
 - (i) that a holder of such registration has contravened or failed to comply with a condition to which the registration concerned is subject to;
 - (j) that it is contrary to the public interest that such pest control product should remain registered;

- (k) that any incorrect or misleading advertisement is used in connection with any registered pest control product.

[L.N. 123/2006, r. 11, L.N.122/2014, r. 2.]

12. Notice to holder of certificate of registration, etc.

Where the Board—

- (a) refuses to register a pest control products; or
- (b) suspends or revokes the certificate of registration,

it shall send to the applicant or the holder of a certificate of registration, as the case may be, a notice by registered post notifying him of the refusal, suspension or revocation.

13. Appeals

An applicant or holder of a certificate of registration who has received a notice under regulation 12 may within thirty days from the date which the notice is received by him appeal to the Minister, who may amend or vary the decision as he thinks fit and whose decision shall be final.

14. Records

A holder of a certificate of registration issued under these Regulations shall keep a record of all the quantities of pest control products stored, manufactured or sold by him and the record shall—

- (a) be maintained for five years from the time it is made; and
- (b) be made available to the Board at such times and in such manner as the Board may require.

15. Declared source of product

- (1) Where a certificate of registration is issued under this regulation, the product shall be obtained from the declared source at the time of registration.
- (2) An application for change of source for the product shall be submitted to the Board as prescribed in Form E set out in the Second Schedule.

[L.N. 124/2015, r. 11.]

FIRST SCHEDULE [r. 3]

ITEMS EXEMPTED FROM REGISTRATION

1. Garment bags, cabinets or chests that are manufactured, represented or sold as a means to protect clothing or fabrics from pests.
2. Electronic apparatus that is manufactured, represented or sold as a means to attract or destroy flying insects.
3. Devices of products that are manufactured, represented or sold to repel birds and other pests by causing physical discomfort by means of sound or touch.
4. Devices of attachment to garden watering hoses that are manufactured, represented or sold as pest control product.
5. Devices that are manufactured, represented or sold as a means of providing the automatic or unattended application of a pest control product.
6. Devices that are sold for use with chemical products containing cyanide as a means to control animal pests.

SECOND SCHEDULE [r. 4]

FORMS

[L.N. 123/2006.]

FORM A

(r. 4A(3))

APPLICATION FOR REGISTRATION OF A PEST CONTROL PRODUCT (CONVENTIONAL)

TRADE NAME OF THE PRODUCT

PURPOSE OF APPLICATION (tick as appropriate)

(a) Pest control product containing a new active ingredient <input type="checkbox"/>	
(b) Pest control product where source of active and/or formulation is not identical to that of a registered product	<input type="checkbox"/>
(c) Registration transfer	<input type="checkbox"/>
(d) Amendments to existing registration	<input type="checkbox"/>
(e) Other (Explain)	

Repealed

Will the product be marketed under own label?	Yes <input type="checkbox"/> No <input type="checkbox"/>
If no, Specify	
Proposed date of marketing	

1. APPLICANT	
1.1 Identification	
Name of applicant/Corporate name of company	
Business Reg No.	
Name of registration holder	
Name of local agent in country: (If different from registration holder)	
1.2 Status (Importer/formulator/distributor)	
Business Registration No.	
1.3 Physical Address	
1.4 Postal Address	
1.5 Telephone: (and area code)	
1.6 Fax: (and area code)	
1.7 e-mail	
2. PRODUCT	
2.1 Designation (Description of product)	Trade name:
	Trade mark:

Repealed

		Trade mark holder:	
2.2 Function of product: (eg. Insecticide, herbicide etc.)			
2.3 Intended use: (Veterinary, public health, industrial, agriculture, forestry, etc.)			
2.4 Target pest(s) and host(s)			
2.5 Method, dosage rates and frequency of application:			
2.6 Types of formulation (e.g. EC, WP, etc.)		Crop life International (CLI*) Code (if available)	
2.7 (a) Is the product registered in country of manufacture?		Yes <input type="checkbox"/> No <input type="checkbox"/>	
(b) Is the product registered in the country of formulation?		If no, give reasons	
		Yes <input type="checkbox"/> No <input type="checkbox"/>	
		If no, give reasons	
2.8 Registration in SEARCH* country (ies) (names)			
2.9 Existing registration No(s) and country (s)			
2.10. Customs Tariff Code: (Brussels Tariff Nomenclature)			
3. COMPOSITION OF ACTIVE INGREDIENT(S) (Technical grade) (information on a.i. may be attached in sealed envelope)			
Active ingredients(s)(Common name(s))	Manufacture(Name and address)	Minimum a.i.%	purity a.i. Range %
4. FORMULATION			
4.1 Formulator: (Name)			
Postal Address			
Physical address			
4.2 Internal code			

Repealed

4.3 Composition (Information on composition may be attached in sealed envelope)			
Ingredients and Function	g/l	g/kg	Range

*Formerly GCPF

*SEARCH- Southern and Eastern African Regulation Committee on Harmonization of Pesticide Registration.

5.TOXICOLOGY (formulated product)					
5.1 Rat	Acute Oral (LD50mg/kg)	Acute Denmal (LD50 mg/kg)			Inhalation LC50 (mg/1/hour)
	Experimental	Experimental			Experimental
	Calculated	Calculated			Calculated
5.2 Rabbit	Skin Irritation	Eye irritation			
None					
Mild					
Moderate					
Severe					
5.3 Skin Sensitization in guinea pig: (tick)	None <input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe <input type="checkbox"/>				
5.4 WHO classification	Ia	Ib	II	III	Others
5.5. Summary of other mammalian toxicological studies : eg. livestock, wildlife, poultry, pets					
5.6. Summary of environmental effects					
5.6.1 Toxicity to bees					
5.6.2 Toxicity to fish and other aquatic organisms					

Repealed

5.6.3 Toxicity to birds	
5.6.4 Toxicity to earthworms and soil micro-organisms	
5.6.5 Toxicity to other non-target organisms	
5.6.6 Persistence in environment	
5.6.7 Other effects: Specify:	
6. PACKAGING	
6.1 Packaging material/ container	
6.2 Pack size(s)	
6.3 Disposal of empty container(s)	
7. OTHER SPECIFIC REQUIREMENTS	
7.1 Human exposure	
(a) Dermal absorption. (b) Likely human exposure under field conditions. (c) Available toxicological data relating to other ingredients in formulation (non-active additives in formulation).	
8. DECLARATION	
For and on behalf ofI hereby certify that the above mentioned information and data provided in support of this application are to the best of my knowledge true, correct and complete.	
..... Name in full (printed) Signature
..... Official Title Date

Repealed

Official Stamp of Applicant/Company	FOR OFFICIAL USE Remarks				
	<table style="width: 100%; border: none;"> <tr> <td style="width: 50%; border: none;">.....</td> <td style="width: 50%; border: none;">.....</td> </tr> <tr> <td style="width: 50%; border: none; text-align: center;">Signed</td> <td style="width: 50%; border: none; text-align: center;">Date</td> </tr> </table>	Signed	Date
.....				
Signed	Date				
NOTE: The format of this application is recognized by all SEARCH countries.					

FORM A1

(r. 4B(2))

**APPLICATION FOR REGISTRATION OF A MICROBIAL
BIOPESTICIDE (PEST CONTROL PRODUCTS-MICROBIAL AGENT)**

TRADE NAME OF THE PRODUCT

PURPOSE OF APPLICATION (tick as appropriate)

(a) Biopesticides containing a new active agent
(b) Biopesticides where source of active and/or formulation is not identical to that of a registered product
(c) Registration transfer
(d) Amendments to existing registration
(e) Other (Explain)
Will the product be marketed under own label? Yes <input type="checkbox"/> No <input type="checkbox"/> If No, specify.....
1. APPLICANT
Name of applicant
Corporate name of company

Repealed

Reg No.	
Name of registration holder:	
Name of local agent in country: (If different from registration holder)	
Status: (Importer/formulator/distributor etc.)	
Physical Address	
Postal Address	
Telephone (and area code)	
Fax (and area code)	
E-mail	
2. PRODUCT	
2.1 Identity and stage (s) of active agent and culture collection code.	
2.2 Concentration of active agent in technical material.	
2.3 Designation (Description of product).	Trade name
	Trade mark
	Trade mark holder
	Internal code
2.4 Function of product (e.g. Insecticide, herbicide etc).	
2.5 Intended use: (Veterinary, horticultural, public health, industrial, agriculture, forestry, etc).	
2.6 Target pests (s) and host(s).	

Repealed

2.7 Method, dosage rates and frequency of application.			
2.8 Types of formulation (e.g. Suspension, WP, etc.).			
2.9 Is the product registered in country of-		Yes <input type="checkbox"/> No <input type="checkbox"/>	
(a) origin		If No, specify	
(b) manufacture		Yes <input type="checkbox"/> No <input type="checkbox"/>	
(c) formation		If No, specify	
2.10 Registration in SEARCH country/ies (country name, product name and registration number).			
2.11 Registration in other country/ies, particularly OECD countries: (country name, product name and registration number).			
2.12 Custom Tariff Code (Brussels Tariff Nomenclature).			
3. IDENTIFICATION			
3.1 Identification of Micro-organism		Life stage (spore, hyphae etc.)	
3.2 Identification		Genus	Species
Scientific name			Sub species
Common name(s)			
3.3 Contents (number per unit)			
4. COMPOSITION OF MICROBIAL PEST CONTROL AGENT(S) (Technical grade) (Information on active agent may be attached in sealed envelope)			
Active agent(s) (Common name/s)	Manufacturer: (Name and address)	Minimum a.1.% purity	a.i. Range %
5. FORMULATION			
5.1 Formulator: (Name)		Postal Address	
Internal code		Physical address	

Repealed

(8) Composition (Information on composition may be attached in sealed envelope)			
Ingredients and Function	Units (w/w, w/v etc.)	Units (e.g. cfu or IUP)	Range
6. BIOLOGICAL PROPERTIES OF ACTIVE AGENT			
6.1 History and geographical distribution of active agent			
6.2 Mode of action and host range			
6.3 Life cycle			
6.4 Infectivity, dispersal and colonizing ability			
6.5 Relationships to know plant, animal or human pathogens			
6.6 Genetic stability			
6.7 Information on the production of metabolites, especially antibiotics and toxins			
7. TOXICOLOGY (active agent)			
7.1 Rat	Acute Oral (LD 50 mg/kg)	Inhalation LC ₅₀ (mg/4/hour)	Intra-peritoneal injection for infectivity (LD 50 g/kg)
	Experimental	Experimental	Experimental
	Calculated	Calculated	Calculated
Hypersensitivity/ allergies in humans			
8. TOXICOLOGY (formulated product)			

Repealed

	Acute Oral (LD50 mg/kg)	Acute Dermal (LD50 g/kg)	Inhalation LC50 (mg/4/hour)		
	Experimental	Experimental	Experimental		
	Calculated	Calculated	Calculated		
8.2 Rabbit	Skin Irritation	Eye Irritation			
None					
Mild					
Moderate					
Severe					
8.3 Skin Sensitization in guinea pig: (tick)	None	Mild	Moderate	Severe	
8.4 WHO classification (tick)	la	lb	II	III	Others
8.5 Summary of other mammalian toxicological studies: e.g. livestock, wildlife, poultry, pets					
9.ECOTOXICOLOGY					
9.1 Toxicity to bees:					
9.2 Toxicity to fish and other aquatic organisms					
9.3 Toxicity to birds					
9.4 Toxicity to earthworms or other soil invertebrates and soil micro-organisms					
9.5 Toxicity to other non-target organisms					
9.6 Persistence in environment					
9.7 Available toxicological data relating to other ingredients in formulation (non-active additives in formulation)					

Repealed

9.8 Other effects: Specify	
10. PACKAGING	
10.1 Packaging material/container	
10.2 Pack size(s)	
10.3 Disposal of empty container (s)	
11. OTHER SPECIFIC REQUIREMENTS	
11.1 Operator exposure	
11.2 Sanitary and phytosanitary measures	
11.3 Has the product been cleared by the phytosanitary authorities? (tick) a. In the country of origin b. the recipient country	Yes <input type="checkbox"/> (provide evidence) No <input type="checkbox"/> (give reasons)
12. DECLARATION	
For and on behalf of	
I hereby certify that the above mentioned information and data provided in support of this application are to the best of my knowledge true, correct and complete.	
..... <i>Name in full (printed)</i> <i>Official Title</i> <i>Signature</i> <i>Date</i>

FORM A2

(r. 4C(1))

**APPLICATION FOR REGISTRATION OF A MICROBIAL BIOPESTICIDE
(PEST CONTROL PRODUCTS-MACROBIOLOGICAL AGENT)**

PRODUCT TRADE NAME

PURPOSE OF APPLICATION (tick as appropriate)

(a) Biopesticides containing a new active agent		
(b) Biopesticides where source of active and/or formulation is not identical to that of a registered product		
(c) Registration transfer		
(d) Amendments to existing registration		
(e) Other (Explain)		
Will the product be marketed under own label Yes <input type="checkbox"/> No <input type="checkbox"/>		
If No, specify.....		
Proposed date of marketing.....		
.....		
1. APPLICANT		
1.1 Name of applicant		
1.2 Corporate name of company		
1.3 Reg. No. of the company		
1.4 Name of registration holder		
1.5 Name of local agent in country (If different from registration holder)		
1.6 Status : (Importer/formulator/distributor etc.)		
1.7 Physical Address	1	2
1.8 Postal Address	1	2
1.9 Telephone (and area code):	1	2

Repealed

Fax (and area code):	1	2
E-mail	1	2
2. PRODUCT		
2.1 Identity and stage(s) of active agent and culture collection code		
2.2 Concentration of active agent in technical material.		
2.3 Description of product.	Trade name	
	Trade mark	
	Trade mark holder	
	Internal code	
2.4 Function of the product: (e.g. predator, parasitoid, entomopathogenic nematode).		
2.5 Intended use: (veterinary, horticultural, public health, industrial, agriculture, forestry, etc.).		
2.6 Target pest(s) and host (s).		
2.7 Method, dosage rates and frequency of application.		
2.8 Type of formulation: (if any).		
2.9 Is the product registered in country of:		
(a) origin	Yes <input type="checkbox"/> If no, specify	No <input type="checkbox"/>
(b) manufacture	Yes <input type="checkbox"/> If no, specify	No <input type="checkbox"/>

Repealed

(c) formulation	Yes <input type="checkbox"/>	No <input type="checkbox"/>
	If no, specify	
2.10 Registration in SEARCH country(ies): (country names, product name and registration number).		
2.11 Registration in other country (ies), particularly OECD countries: (country names, product name and registration number)		
2.12 Customs Tariff Code: (Brussels Tariff Nomenclature).		
3. IDENTIFICATION		
Identification of Microbiological agent	Life stage (egg/adult/larva etc)	
3.1 Identification	Genus	Species
Scientific name		
Common name(s)		
3.2 Contents (number per Unit)		
4. SOURCE		
Source (original isolation)		
5. FORMULATION		
5.1 Formulator: (Name)	Postal Address:	
5.2 Internal code	Physical address:	
5.3 Composition (information on composition may be attached in sealed envelop)		
Ingredients and Function	Units	Range
6. SUMMARY OF ENVIRONMENTAL EFFECTS (BIOSAFETY)		

Repealed

6.1 Risk assessment for replacement of indigenous or endangered species in same niche (exotic microbials only)	
6.2 Risk to bees	
6.3 Risk to fish and other aquatic organisms	
6.4 Risk to birds	
6.5 Risk to earthworms and soil micro-organisms	
6.6 Risk to other non-target organisms	
6.7 Other effects : specify (human health problems)	
7. PACKAGING	
7.1 Packaging material/container	
7.2 Pack size(s)	
8. OTHER SPECIFIC REQUIREMENTS	
8.1 Operator exposure	
8.2 Likely operator exposure under field conditions	
8.3 Sanitary and phytosanitary measures	
8.4 Has the product been cleared by the phytosanitary authorities?	Yes <input type="checkbox"/> No <input type="checkbox"/>
9. DECLARATION	
For and on behalf of I hereby certify that the above mentioned information and data provided in support of this application are to the best of my knowledge true, correct and complete	
..... Name in full (printed) Signature

Repealed

..... Official Title Date
-------------------------	---------------

NOTE: The format of this application form is recognized by all SEARCH countries.

FORM A3

(r. 4D)

**APPLICATION FOR REGISTRATION OF A BIOCHEMICAL
PESTICIDE (PEST CONTROL PRODUCT-BIOCHEMICAL PRODUCTS)**

PRODUCT TRADE NAME

PURPOSE OF APPLICATION (tick as appropriate)

a. Biochemical pesticides containing a new active ingredient <input type="checkbox"/>	
b. Biochemical pesticides where source of active and/or formulation is not identical to that of a registered product <input type="checkbox"/>	
c. Registration transfer <input type="checkbox"/>	
d. Amendments to existing registration <input type="checkbox"/>	
e. Other (Explain)	
.....	
Will the product be marketed under own label Yes <input type="checkbox"/> No <input type="checkbox"/>	
If no, specify	
1. APPLICANT	
1.1 Identification	
1.2 Name of applicant/Corporate name of company	

Repealed

1.3 Reg No.		
1.4 Name of registration holder		
1.5 Name of local agent in country (if different from registration holder)		
1.6 Status (Importer/formulator/distributor etc.)		
1.7 Physical Address		
1.8 Postal Address		
1.9 Telephone (and area code)		
1.10 Fax (and area code)		
1.11 E-mail		
2. PEST CONTROL PRODUCTS		
2.1 Identify		
2.2 Concentration of a.i.		
2.3 Designation (Description of product)	Trade name	
	Trade mark	
	Trade mark holder	
	Internal code	
2.4 Function of product (e.g. Insecticide, herbicide etc.)		
2.5 Intended use (Veterinary, public health, industrial, agricultural, forestry etc.)		
2.6 Target pest (s) and host (s)		

Repealed

2.7 Method, dosage rates and frequency of application			
2.8 Types of formulation (e.g. EC, WP, etc.)			
2.9 Is the product registered in country of			
(a) origin	Yes <input type="checkbox"/>	No <input type="checkbox"/>	
	If no, specify.....		
(b) manufacture	Yes <input type="checkbox"/>	No <input type="checkbox"/>	
	If no, specify.....		
(c) formulation	Yes <input type="checkbox"/>	No <input type="checkbox"/>	
	If no, specify.....		
2.10 Registration in SEARCH** country/ies (names)			
2.11 Registration in other country/ies, especially OECD countries (names)			
2.12 Custom Tariff Code (Brussels Tariff Nomenclature)			

*Formerly GCPF.

**SEARCH-Southern and Eastern African Regulatory Committee on Harmonization of Pesticide Registration

3. COMPOSITION OF ACTIVE INGREDIENT (S) (Technical grade) (Information on a.i may be attached in sealed envelope)			
Active ingredient (s): (Common name/s)	Manufacturer: (Name and address)	Minimum a.i.%	Purity a.i. Range %
4. TOXICOLOGY OF ACTIVE INGREDIENTS (Technical grade)	Acute Oral (LD50 mg/kg)	Acute dermal (LD50 mg/kg)	Inhalation LC50 (mg/1/hour)

Repealed

	Experimental	Experimental	Experimental		
	Calculated	Calculated	Calculated		
5. FORMULATION					
5.1 Formulator: (Name)		Postal Address			
5.2 Internal code:		Physical Address			
5.3 Composition (Information on compensation may be attached in sealed envelope)					
Ingredients and Function	Units	Units	Range		
6. TOXICOLOGY (formulated product)					
6.1 Rat	Acute Oral (LD50 mg/kg)	Acute Demal (LD50 g/kg)	Inhalation LC50 (mg/l/kg)		
	Experimental	Experimental	Experimental		
	Calculated	Calculated	Calculated		
6.2 Rabbit	Skin irritation	Eye irritation			
None					
Mild					
Moderate					
Severe					
6.3 Skin Sensitization in guinea pig (tick)	None <input type="checkbox"/>	Mild <input type="checkbox"/>	Moderate <input type="checkbox"/>	Severe <input type="checkbox"/>	
	la	lb	II	III	Others

Repealed

6.5 Summary of other mammalian toxicological information may be required	
6.6 Summary of environmental effects	
6.6.1 Toxicity to bees	
6.6.2 Toxicity to fish and other aquatic organisms	
6.6.3 Toxicity to birds	
6.6.4 Toxicity to earthworms and soil micro-organisms	
6.6.5 Toxicity to other non-target organisms may be required	
6.6.6 Persistence in environment	
6.6.7 Other effects: Specify	
7. PACKAGING	
7.1 Packaging material/container	
7.2 Pack size (s)	
7.3 Disposal of empty container(s)	
8. OTHER SPECIFIC REQUIREMENTS	
8.1 Operator exposure	
8.2 Dermal absorption	
8.3 Likely operator exposure under field conditions	
8.4 Available toxicological data relating to other ingredients in formulation (non-active additives in formulation)	
9. DECLARATION	
For and on behalf of I hereby certify that the above mentioned information and data provided in support of this application are to the best of my knowledge true, correct and complete.	

Repealed

..... Name in full (printed) Signature
..... Official Title Date
Official Stamp of Applicant/ Company	FOR OFFICIAL USE Remarks Signed Date

NOTE: The format of this application is recognized by all SEARCH countries.



FORM A3a

(r. 4D(4), (5))

APPLICATION FOR THE REGISTRATION OF A SEMIOCHEMICAL PEST CONTROL PRODUCT

Introduction

1. These guidelines are for any proposed use of the semiochemicals (e.g. Pheromone, Allomone, Kairomone, Sysndrome, etc) of naturally occurring organisms for the monitoring and control of pests, livestock and public health.
2. Information in support of a request for registration, both published and unpublished should be supplied in the form of a summary data sheet laid out according to the format given in Form A3a.
3. List II of this form applies to a semiochemical not physically combined with a pesticide. Where a semiochemical is physically combined with a synthetic insecticide then the requirements under list II of Form A will apply.

Where a semiochemical and an insecticide are used separately in a device then the requirements under List I and II in Form A will apply for the insecticide.

4. A pre-registration consultation between the applicant and the registration authority is strongly recommended.

Information for Applicants

1. The application form must be completed by a person duly authorized by the applicant/company

2. The application form must be submitted in triplicate to:

The Secretary, Pest Control Products Board (PCPB) P.O. BOX. 13794, 00800 Nairobi.

Email address:pcpboard@todays.co.ke/md@pcpb.or.ke Tel: 254-020- 4446115/4450242

Fax: 254-020-4449072

3. Every application must be accompanied by-

- (a) registered fee as prescribed.
- (b) three copies of the draft label as per PCPB requirements.

4. The applicant shall be required to submit:-

- (a) a sample of the pest control product;
- (b) a sample of the technical grade of its active ingredient.
- (c) a sample of the reference standard of its active ingredient of known purity.
- (d) any other sample as may be required by PCPB.

5. List I and II are supplied as check list and an index to ensure that the applicant has provided all relevant data.

6. The application must be accompanied by a technical dossier as per PCPB data requirements i.e. Lists I and II.

7. An applicant who is not a resident in Kenya must appoint an agent permanently resident in Kenya.

PURPOSE OF APPLICATION (tick as appropriate)

a. Semiochemical pest control product containing a new active ingredient <input type="checkbox"/>
b. Semiochemical pest control product where source of active and/or formulation is not identical to that of a registered product <input type="checkbox"/>
c. Registration transfer <input type="checkbox"/>
d. Amendments to existing registration <input type="checkbox"/>

Repealed

e. Other (Explain)
.....

Will the product be marketed under own label Yes <input type="checkbox"/> No <input type="checkbox"/>
If no, specify.....

1. APPLICANT		
1.1 Identification		
Name of applicant/ Corporate name of the company		
Registration Number.		
Name of registration holder.		
Name of local agent i country: (if different from registration holder)		
1.2 Status: (Importer/formulator/distributor) etc.		
1.3 Physical Address		
1.4 Postal Address		
1.5 Telephone: (and area code)		
1.6 Fax: (and area code)		

2. PRODUCT	
2.1 Identity	
2.2 Concentration of a.i.	

Repealed

2.3 Designation (Description of product)	Trade name:		
	Trade mark:		
	Trade mark holder:		
	Internal code:		
2.4 Function of product: (e.g. attractant, repellent, Mating disruptor, etc.)			
2.5 Intended use: (Veterinary, public health, industrial, agriculture, forestry, etc.)			
2.6 Target pest(s) and host(s)			
2.7 Method, dosage, rates and frequency of application:			
2.8 Types of formulation: (e.g. EC,WP, etc.)			
FOR INFORMATION			
2.9 Is the product registered in country of (a) origin: (b) manufacture: (c) formulation:	Yes <input type="checkbox"/> No <input type="checkbox"/> If no, specify..... Yes <input type="checkbox"/> No <input type="checkbox"/> If no, specify..... Yes <input type="checkbox"/> No <input type="checkbox"/> If no, specify.....		
2.10 Registration in SEARCH* countries/ies: (names)			
2.11 Registration in other country/ies, especially OECD** countries: (names)			

Repealed

2.12 Customs Tariff Code: (Brussels Tariff Nomenclature)	
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*SEARCH- Southern and Eastern African Regulatory Committee on Harmonisation of Pesticide Registration

** OECD- Organisation for Economic Cooperation Development

3. COMPOSITION OF ACTIVE INGREDIENT(S) (Technical grade) (Information on a.i may be attached in a sealed envelope)			
Active ingredient (s): (Common name(s))	Manufacturer: (Name and address)	Minimum a.i% purity	a.i% Range%

4. TOXICOLOGY OF ACTIVE INGREDIENTS (Technical grade) (Exempt for SCLPS2)	Acute Oral (LD50 mg/kg)	Acute dermal (LD50 mg/kg)	Inhalation LC50 mg/1/ hour
	Experimental	Experimental	Experimental
5. FORMULATION	Calculated	Calculated	Calculated
5.1 Formulator: (Name)		Postal Address:	
5.2 Internal code:		Physical address:	
5.3 Composition (Information on composition may be attached in sealed envelope)			
Ingredients and Function:	Units	Units	Range

6. TOXICOLOGY (formulated product) (Exempt for SCLPs if inert are of known toxicity)			
6.1 Rat:	Acute Oral (LD50 mg/ kg)	Acute Dermal LD50 g/kg)	Inhalation LC50 (mg/1/ hour)
	Experimental	Experimental	Experimental
	Calculated	Calculated	Calculated

Repealed

6.2 Rabbit:	Skin irritation		Eye irritation		
None					
Mild					
Moderate					
Severe					
6.3 Skin Sensitisation in guinea pig (<i>tick</i>)			None Mild Moderate Severe <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>		
6.4 WHO classification:	Ia	Ib	II	III	Others
6.5 Summary of other mammalian toxicological information may be required					
6.6 Summary of environmental effects					
6.6.1 Toxicity to bees:					
6.6.2 Toxicity to fish:					
6.6.3 Toxicity to birds:					
6.6.4 Toxicity to earthworms:					
6.6.5 Toxicity to other non-target organisms may be required:					
6.6.6 Other effects: Specify					
PACKAGING					
Type of packaging (packaging material/ container, compatability with content)					
7.1 Pack size (s)					
7.2 Manner of packaging					

Repealed

7.3 Specification for primary packaging	
7.4 Disposal of empty container (s):	

Please note that the product must be sold only in the package size and type notified to the Pest Control Products Board and for which the label is approved.

8. OTHER SPECIFIC REQUIREMENTS

8.1 Operator exposure	
8.2 Likely operator exposure under field conditions	
8.3 Available toxicological data relating to other ingredients in formulation (<i>non-active additives in formulation</i>).	
9. DECLARATION	
For and on behalf of..... I hereby certify that the above mentioned information and data provided in support of this application are to the best of my knowledge true, correct and complete.	
..... Name in full (<i>printed</i>) Signature
..... Official Title Date
Official Stamp of Applicant/Company	FOR OFFICIAL USE Remarks..... Signed: Date

NOTE: The format of this application form is recognized by all SEARCH countries.

FORM A3b

(r. 4(1), 4D(5))

**APPLICATION FOR INTRODUCTION OF A STRAIGHT CHAIN
LEPIDOPTERAN PHEROMONE FOR PEST MONITORING PURPOSES**

Introduction

1. These guidelines are for any proposed use of the straight chain lepidopteran pheromones (SCLP) of naturally occurring organisms for the monitoring and control of pests of crops, livestock and public health.
2. Information in support of a request for introduction, both published and unpublished should be supplied in the form of a summary data sheet laid out according to the format given in Form A3b.
3. This form applies to a SCLP not physically combined with a pesticide. Where a SCLP is physically combined with an insecticide then the requirements under List II of Form A will apply.
4. Where a SCLP and an insecticide are used separately in a device then the requirements under Lists I and II in Form A will apply for the insecticide.
5. A pre-introduction consultation between the applicant and the regulatory authority is strongly recommended.

Information for Applicants

1. The application form must be completed by a person duly authorized by the applicant/company
2. The application form must be submitted in triplicate to:
The Secretary, Pest Control Products Board (PCPB) P.O Box 13794, 00800 Nairobi.
E-mail address: pcpboard@todays.co.ke/md@pcpb.or.ke Tel: 254-020-4446115/4450242
Fax: 254-020-4449072
3. Every application must be accompanied by:-
 - (a) introduction fee as prescribed.
 - (b) three copies of the draft label as per PCPB requirements.
4. The applicant shall be required to submit-
 - (a) a sample of the pest control product;
 - (b) a sample of the reference standard of its ingredient of known purity.
 - (c) any other sample as may be required by PCPB.
5. An applicant who is not a resident in Kenya must appoint an agent permanently resident in Kenya.

PURPOSE OF APPLICATION (*tick as appropriate*)

a. SCLP pest control product containing a new active ingredient <input type="checkbox"/>
b. SCLP pest control product where source of active and/or formulation is identical to that of an existing product <input type="checkbox"/>
c. Amendments to existing product <input type="checkbox"/>

Repealed

d. Other (Explain)

1. APPLICANT		
1.2 Identity		
Name of applicant/Corporate name of company		
Registration Number of company:		
Name of registration holder of product.		
Name of local agent in country: (if different from registration holder)		
1.3 Status: (Importer/formulator/distributor)etc.		
1.4 Physical Address		
1.5 Postal Address:		
1.6 Telephone: (and area code)		
1.7 Fax (and area code):		

2. PRODUCT	
2.15 Identity (name(s) of a.i	
2.16 Concentration of a.i.	
2.17 Designation (Description of product)	Trade name:
	Trade mark:

2. PRODUCT	
	Trade mark holder:
	Internal code:
2.18 Function of product:	Is it an attractant? Yes <input type="checkbox"/> No <input type="checkbox"/>
2.19 Intended use: (Veterinary, public health, industrial agriculture, forestry, etc.)	
2.20 Target pest (s) and host (s)	
2.21 Method of use, frequency of replacements of the SCLP and No. of devices per unit area.	
2.22 Degree of species specificity	
2.23 Time of application	

FOR INFORMATION

2.24 Is the product used in country of (a) origin: (b) manufacture: (c) formulation:	Yes <input type="checkbox"/> No <input type="checkbox"/> If yes, attach evidence..... If no, explain..... Yes <input type="checkbox"/> No <input type="checkbox"/> If yes, attach evidence..... If no, explain..... Yes <input type="checkbox"/> No <input type="checkbox"/> If yes, attach evidence If no, explain
2.25 Use in SEARCH* country/ies:(names)	If yes, attach evidence
2.26 Use in other country/ies, especially OECD** COUNTRIES: (names)	If yes, attach evidence

Repealed

2.27 Customs Tariff Code (Brussels Tariff Nomenclature):	
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*SEARCH – Southern and Eastern African Regulatory Committee on Harmonisation of Pesticide Introduction

**OECD- Organisation for Economic Cooperation and Development

3. COMPOSITION OF ACTIVE INGREDIENT(S) (<i>Technical grade</i>) (Information on a.i may be attached in a sealed envelope)			
Active ingredient(s): (Common name/s)	Manufacture: (Name and address)	Minimum a.i.% purity	a.i. Range%
3.1 Molecular structures (to be attached)			
3.2 Molecular formula			
3.3 Mode of action			

5. FORMULATION			
5.1 Formulator: (<i>Name</i>)	Postal Address:		
5.2 Internal code:	Physical address:		
5.3 Composition (<i>Information on composition may be attached in sealed envelope</i>)			
Ingredients and Function:	units	units	Range
5.4 Information on storage stability			
5.5 Information on shelf life			
5.6 Information on use and efficacy			

6. Method and certificate of Analysis (<i>to be attached</i>)

Repealed

PACKAGING	
Type of packaging (<i>packaging material/ container, compatability with content</i>)	
Pack size (s)	
Manner of packaging	
Specification for primary packaging	
7.4 Disposal of empty container (s):	

Please note that the product must be sold only in the package size and type notified to the Pest Control Products Board.

8. INFORMATION ON DEVICE (e.g. TRAP)	
8.1 Type	
8.2 Weight	
8.3 Thickness/height	
8.4 Diameter/size	
8.5 Colour	

9. OTHER SPECIFIC REQUIREMENTS

9.1 Operator exposure	
9.2 Available toxicological data relating to other ingredients in formulation (non-active additives in formulation).	
10. DECLARATION	
For and on behalf of..... I hereby certify that the above mentioned information and data provided in support of this application are to the best of my knowledge true, correct and complete.	

Repealed

..... Name in full (printed) Signature
..... Official Title Date
Official Stamp of Applicant/ Company	FOR OFFICIAL USE Remarks Signed: Date:

NOTE: The format of this application form is recognized by all SEARCH countries.



FORM A4

(r. 4A(2))

APPLICATION FOR THE REGISTRATION OF A PEST CONTROL PRODUCT (GENERIC)

Introduction

These guidelines are for registration of identical products that are manufactured after the expiry of the patent of an original/proprietary registered product. These identical products are generally referred to as generics and will include conventional and biochemical pesticides. A pre-registration consultation between the applicant and the registration authority is strongly recommended.

Information for Applicants

1. The application form must be completed by a duly authorized person.
2. The application must be submitted in triplicate to:
 The Managing Director/Secretary
 Pest Control Products Board (PCPB)
 P.O. Box 13794-00800 Nairobi.
 E-mail address: pcpboard@todays.co.ke/md@pcpb.or.ke
 Tel: 254-020-8021846/7/8 Fax: 254-020-8021865
 Website: www.pcpb.or.ke
3. Every application must be accompanied by:-
 - a) application fee as prescribed (Registration fee is payable upon approval by the Board).
 - b) 3 copies of the draft label as per PCPB requirements.
4. The applicant may be required to submit to:-
 - (a) a sample of the pest control product;
 - (b) a sample of the technical grade of its active ingredient;
 - (c) a sample of the laboratory standard of its active ingredient;
 - (d) any other sample as may be required by the Board.
5. List I and II are supplied as a check list and an index to ensure that the applicant has provided the relevant data.
6. The application must be accompanied by a technical dossier as per PCPB data requirements (Lists I and II)
7. An applicant who is not a resident in Kenya must appoint an agent permanently resident in Kenya and duly recognized by the Pest Control Products Board.

TRADE NAME

PURPOSE OF APPLICATION (tick/fill as appropriate)

a. Pest control product containing a generic active ingredient <input type="checkbox"/> i) Date of expiry of patent..... ii) Name of former patent holder.....
b. Pest control product where source of active and/or formulation is not identical to that of a registered product <input type="checkbox"/>
c. Registration transfer <input type="checkbox"/>
d. Amendments to existing registration (e.g inerts, source of technical material e.t.c) <input type="checkbox"/>

e. Other (Explain).....
f. Will the product be marketed under own label? Yes <input type="checkbox"/> No <input type="checkbox"/> If no specify.....
Proposed date of marketing.....

1. APPLICANT	
1.1 Identification	
Name of applicant/ Corporate name of company	
Business Registration No:	
Name of registration holder	
1.12 Status: (Manufacturer formulator/other)	
1.13 Physical Address	
1.14 Postal Address:	
1.15 Telephone: (and area code)	
1.16 Fax: (and area code)	
1.17 e-Mail:	
2. Name of local agent in country: (if different from registration holder)	
Business Registration No:	

Repealed

2.1 Status: (Importer/formulator/ distributor)			
2.2 Physical Address			
2.3 Postal Address			
2.4 Telephone: (and area code)			
2.5 Fax: (and area code)			
2.6 e-mail:			
3 PRODUCT			
Designation (Description of product)		Trade name:	
		Trade mark:	
		Trade mark holder:	
Function of product: (eg. Insecticide, herbicide etc.)			
3.1 Intended use: (Veterinary, public health, industrial)			
3.2 Target pest(s) and host (s)			
3.3 Method, dosage rates and frequency of application:			
3.4 Type of formulation: (eg. EC, WP, etc.)		Crop Life International(CLI) Code (if available)	

Repealed

3.5 a) Is the technical grade (active ingredient) registered in country of manufacture?	Yes <input type="checkbox"/> No <input type="checkbox"/> If no, give reasons		
b) Is the product registered in the country of formulation	Yes <input type="checkbox"/> No <input type="checkbox"/> If no, give reasons		
3.6 Proof of registration in SEARCH** country/ies (names)			
3.7 Proof of registration in other countries.			
3.8 Customs Tariff Code: (Brussels Tariff Nomenclature)			
4 COMPOSITION OF ACTIVE INGREDIENT (S) (Technical grade) (Information on a.i may be attached in sealed envelope			
Active ingredient (s): (Common name/s)	Manufacturer: (Name and address)	Minimum a.i.%purity	a.i. Range%

FORMULATION			
5.1 Formulator: (Name) Postal Address: Physical address:			
5.2 Internal code:			
5.3 Composition (Information on composition may be attached in sealed envelope)			
Ingredients and Function:	g/L	g/Kg	Range
6. TOXICOLOGY (formulated product)			

Repealed

6.1 Rat:	Acute Oral (LD50 mg/Kg)	Acute Dermal (LD50 mg/Kg)	Inhalation LC50 (mg/L/4 hour)		
	Experimental	Experimental	Experimental		
6.2 Rabbit:	Skin Irritation	Eye irritation			
None					
Mild					
Moderate					
Severe					
6.3 Skin Sensitization in guinea pig: (tick)		None <input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe <input type="checkbox"/>			
6.4 WHO		Ia	Ib	II	III
Table V					
7 Summary of environmental effects					
7.1 Toxicity to bees:					
7.2 Toxicity to fish and other aquatic organisms:					
7.3 Toxicity to birds:					
7.4 Toxicity to earthworms and soil micro-organisms:					
7.5 Toxicity to other non-target organisms:					
7.6 Persistence in environment:					
7.7 Other effects: Specify					
8. PACKAGING					
8.1 Packaging material/container:					
8.2 Pack size (s):					

Repealed

8.3 Disposal of empty container (s):	
9. OTHER SPECIFIC REQUIREMENTS	
9.1 Operator exposure	
9.1.1 Dermal absorption.	
9.1.2 Likely operation exposure under field conditions	
9.2 Available toxicological data relating to other ingredients in formulation (non-active additives in formulation)	
10. DECLARATION	
For and on behalf of I hereby certify that the above mentioned information and data provided in support of this application are to the best of my knowledge true, correct and complete.	
..... Name in full (printed)	
..... Signature	
..... Official Title	
..... Date	
..... Official Stamp of Applicant/ Company	
FOR OFFICIAL USE	
Remarks	
.....	
.....	

Repealed

<p>.....</p> <p>Signed</p> <p>.....</p> <p>Date</p>

NOTE: The format of this application is recognized by all SEARCH countries.

FORM A5

(r. 4E(1), (2))

APPLICATION FOR THE REGISTRATION OF A PEST CONTROL PRODUCT (SPRAY ADJUVANT)

A spray adjuvant: Is a compound or substance than enhances or modifies or is intended to enhance or modify the physical or chemical characteristics of a pest control product to which it is added.

Information for Applicants

1. The application form must be completed by a duly authorized person.
2. The application must be submitted in triplicate to:

The Managing Director/Secretary
Pest Control Products Board (PCPB)
P.O. Box 13794-00800 Nairobi.
E-mail address: pcpboard@todays.co.ke/md@pcpb.or.ke
Tel: 254-020-8021846/7/8 Fax: 254-020-8021865
Website: www.pcpb.or.ke
3. Every application must be accompanied by:-
 - (a) application fee as prescribed (Registration fee is payable upon approval by the Board).
 - (b) 3 copies of the draft label as per PCPB requirements.
4. The applicant may be required to submit;-
 - (a) a sample of the pest control product;
 - (b) a sample of the laboratory standard of its active ingredients;
 - (c) any other sample as may be required by the Board.
5. List I is supplied as a check list and an index to ensure that the applicant has provided the relevant data.
6. The application must be accompanied by a technical dossier as per PCPB data requirements(dossier index.)
7. An applicant who is not a resident in Kenya must appoint an agent permanently resident in Kenya and duly recognized by the Pest Control Products Board.

TRADE NAME

PURPOSE OF APPLICATION (tick as appropriate)

a. A pest control product which is an adjuvant <input type="checkbox"/>
b. Pest control product where source of active and/or formulation is not identical to that of a registered product <input type="checkbox"/>
c. Registration transfer <input type="checkbox"/>
d. Amendments to existing registration
e. Other (Explain)
f. Will the product be marked under own label? Yes <input type="checkbox"/> No <input type="checkbox"/> If no specify
Proposed date of marketing

1. APPLICANT

Repealed

1.1 Identification	
Name of applicant/ Corporate name of company	
Business Registration No:	
Name of registration holder	
1.18 Status: (manufacturer/ formulator/other)	
1.19 Physical Address	
1.20 Postal Address:	
1.21 Telephone: (and area code)	
1.22 Fax: (and area code)	
1.23 E-mail:	
2. Name of local agent in country: (if different from registration)	
Business Registration No:	
3.1 Status: (Importer/formulator/distributor)	
3.2 Physical address	
3.3 Postal Address:	
3.4 Telephone: (and area code)	
3.5 Fax: (and area code)	
3.6 e-mail:	
3. PRODUCT	51
3.1 Designation	Trade name:

Repealed

	Trade mark:		
	Trade mark holder:		
3.2 Spray adjuvant function: (wetter, surfactant, etc)			
3.3 Intended use: (Veterinary, public health, industrial, agriculture, forestry, etc.			
3.4 Target use e.g product and crop/animal			
3.5 Method, dosage rates and frequency of application:			
3.6 Type of formulation: (e.g. EC,WP, etc.)		Crop Life International (CLI) Code (if available)	
3.7 (a) Is the technical grade (active ingredient) registered in country of manufacture? (b) Is the product registered in the country of formulation?	Yes <input type="checkbox"/> No <input type="checkbox"/> If no, give reasons Yes <input type="checkbox"/> No <input type="checkbox"/> If no, give reasons		
3.8 Registration in SEARCH** country/ies: (names)			
3.9 Proof of existing registration in other country (ies)			
3.10 Customs Tariff Code: (Brussels Tarrif Nomenclature)			
4. SPRAY ADJUVANT FORMULA (attach confidential formula)			
Active ingredient(s): (Common name/s)	Manufacturer: (Name and address)	Spray adjuvant function	Percentage

5. FORMULATION			
5.1 Formulator: (Name)			
Postal Address:			
Physical address:			
5.2 Internal code:			
5.3 Composition (Information on composition formula may be attached in sealed			
Ingredients	g/L	g/KG	Range
6. TOXICOLOGY (formulated product)			
6.1 Rat:	Acute Oral (LD50 mg/Kg)	Acute Dermal (LD50 mg/Kg)	Inhalation LC 50 (mg/L/4/hour)
	Experimental/Calculated	Experimental/Calculated	Experimental/Calculated
6.2 Rabbit:	Skin irritation	Eye irritation	
None			
Mild			
Moderate			
Severe			
6.3 Skin Sensitization in guinea pig: (tick)		None <input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe <input type="checkbox"/>	
6.4 WHO classification:	Ia	Ib	II III Table V
6.5 Summary of other mammalian toxicological studies: eg. Livestock, wildlife, poultry, pets			
7. Summary of environmental effects (where applicable e.g. sensitive areas)			
7.1 Toxicity to bees:			

Repealed

7.2 Toxicity to fish and other aquatic organisms:	
7.3 Toxicity to birds:	
7.4 Toxicity to earthworms and soil microorganisms:	
7.5 Toxicity to other non-target organisms:	
7.6 Persistence in environment:	
7.7 Other effects: Specify	
8 PACKAGING	
8.1 Packaging material/container:	
8.2 Pack size(s):	
8.3 Disposal of empty container(s):	
9. OTHER SPECIFIC REQUIRMENTS	
9.1 Operator exposure	
9.1.1 Dermal absorption.	
9.1.2 Likely operator exposure under field conditions	
9.2 Available toxicological data relating to other ingredients informulation (non-active additives in formulation).	
10. DECLARATION	
For and on behalf of I hereby certify that the above mentioned information and data provided in support of this application are to the best of my knowledge true, correct and complete.	
..... Name in full (<i>printed</i>) Signature
..... Official Title Date

Repealed

Official Stamp of Applicant/Company	FOR OFFICIAL USE Remarks Signed Date
--	---

FORM A6

(r. 2)

APPLICATION FOR REGISTRATION AS A LOCAL AGENT

information for Applicants

1. "Agent" means a person or company who has/that has been appointed to act on behalf of a registrant in accordance with regulation 4(2) of the Pest Control Products (Registration) Regulations.
2. The application form must be completed by a duly authorized person.
3. The application must be submitted to:
 The Managing Director/Secretary
 Pest Control Products Board (PCPB)
 P.O.Box 13794-00800 Nairobi
 E-mail address:md@pcpb.or.ke
 Tel:254-020-8021846/7/8 Fax: 254-020-8021865
 Website: www.pcpb.or.ke
4. Every application must be accompanied by:-
 - (a) An original letter from the registrant,
 - (b) A binding agreement entered between the registrant and the agent.
5. The applicant may be required to submit:-
 - (a) A sample of the pest control product;
 - (b) A sample of the technical grade of its active ingredient;
 - (c) A sample of the laboratory standard of its active ingredient;
 - (d) Any other information as may be required by the Board.
6. For each product, there can only be one local agent.
7. The local agent shall appoint in the distributor(s)
 Product Details
 Trade Name.....

Name of Manufacturer.....
 Name of Registrant.....
 Name of agent.....
 Name of distributor.....
 Signature of applicant..... Date

Official Stamp of Applicant/Company

FORM A7

(r. 4(2)(b))

APPLICATION FOR CHANGE OF AGENCY.

Information for Applicants

1. "Agent" means a person or company who has/that has been appointed to act on behalf of a registrant in accordance with regulation 4(2) of the Pest Control Products (Registration) Regulations.
2. The application form must be completed by a duly authorized person.
3. The application must be submitted to:
 The Managing Director/Secretary
 Pest Control Products Board (PCPB)
 P.O.Box 13794-00800 Nairobi.
 E-mail address:md@pcpb.or.ke
 Tel:254-020-8021846/7/8 Fax: 254-020-8021865
 Website: www.pcpb.or.ke
4. Every application must be accompanied by:-
 - (a) An original letter from the registrant,
 - (b) A binding agreement entered between the registrant and the agent.
 - (c) An original letter of no objection from the current agent,
 - (d) Application fee of Ksh 20,000 per product (change of agency fee is payable upon approval by PCPB after meeting the other requirements),
 - (e) A copy of the draft label as per PCPB requirements,
 - (f) Proof of licensing of the new agent by PCPB.
5. The applicant maybe required to submit:-
 - (a) A sample of the pest control product;
 - (b) A sample of the technical grade of its active ingredient;
 - (c) A sample of the laboratory standard of its active ingredient;

(d) Any other information as may be required by the Board.

Trade name

Name of Manufacturer

Name of Registrant

Registration Number (If registered) Status of registration

Name of former agent

Name of new agent

Signature of applicant Date

Official Stamp of Applicant/Company

For official use only

Please check whether the following documents have been provided:

Registration Department:

1. Has an original letter from the registrant been provided Yes No
2. Has an original letter of no objection from the former agent been provided?
Yes No
3. Has the applicant attached a copy of the draft label?
Yes No

Inspection Department

4. Is the applicant licensed as a pesticide dealer/agent with PCPB in the current year?
 Yes No if yes indicate licence No.

Accounts:

5. Has the applicant paid the change of agency fee? Yes No.....
Indicate Receipt Number Date
6. Has the applicationt paid the dealers/agency license fee? Yes..... No.....
Indicate Receipt Number..... Date.....

Recommended <input type="checkbox"/>	Not Recommended <input type="checkbox"/>	Recommended <input type="checkbox"/>	Not Recommended <input type="checkbox"/>	Recommended <input type="checkbox"/>	Not Recommended <input type="checkbox"/>
Date		Date		Date	
Registration Officer		Inspector		Accountant	

.....

Approved by Managing Director Date

Pest Control Products Board

Changes effected by (IT Officer)..... Date

FORM A8

(r. 5(2))

THE PEST CONTROL PRODUCTS (MISCELLANEOUS FORMS) SUBMISSION OF SAMPLE (S) FOR EFFICACY TESTING

This form should be filled in duplicate: Part I and II to be filled by the Applicant. The completed form MUST be accompanied with a copy of:

- i. the trial permit
- ii. the Material Safety Data Sheet (MSDS)

The trial sample MUST be labeled in accordance with the Pest Control Products Board guidelines for experimental labeling.

I) Product Details

Trade name

Formulation type

Active ingredient (s)

Concentration of active ingredient(s)

Quantity of sample (Liters or grams)

Expiry date

Number of packages

REF: (Permit No. and date)

Recommended storage conditions of temperature

Name of Applicant (Local agent)

II) Submission details

Submitted to PCPB by:

Name Signature.....Date.....

Delivery details

Received on behalf of PCPB by:

Name Signature Date

(Tick Appropriately)

- 1. Is the application form attached with a copy of the trial permit? YES NO
- 2. Is the form attached with a copy of the Material Safety Data Sheet? YES NO
- 3. Does the label conform to the PCPB guidelines for experimental labeling? YES NO

Institution(s) of destination

Means of delivery

A. PCPB personnel

Name of Person delivering

Date of delivery

Signature

B. Courier Service

Name of company

Contact person

Charges (Attach receipt)

Date of delivery

Official stamp.....

C. Receiving Institution

Name of Receiving Institution

Date of receipt

Person receiving

Signature

Official Stamp

FORM A9

(r. 7(1)(b))

APPLICATION FOR CHANGE OF TRADE NAME FOR A PEST CONTROL PRODUCT.

Information for Application

1. The application form must be completed by a duly authorized person/agent
2. The application must be submitted to:
 The Managing Director/Secretary
 Pest Control Products Board (PCPB)
 P.O Box 13794 - 00800 Nairobi.
 E-mail address: md@pcpb.or.ke
 Tel: 254- 020 - 8021846/7/8 Fax: 254- 020- 8021865
 Website: www.pcpb.or.ke
3. Every application must be accompanied by:-
 - (a) An original consent letter from the owner of the trade name (with evidence from a trade mark registering body);
 - (b) A copy of the draft label as per PCPB requirements;
 - (c) Proof of licencing as a dealer with Pest Contro Products by PCPB
5. The applicant may be required to submit:
 - (a) A sample of the technical grade of its active ingredient:
 - (b) A sample of the laboratory standard of its active ingridients;
 - (c) Any other information as may be required by the Board.
6. Evidence from Kenya Intellectual Property Institute (KIPI) that the new trade name is available for use.
7. Payment of a change of trade name fee of Kshs. 10,000

Product Details

Current Trade Name.....

Proposed Trade Name*

Reason for change

.....

.....

Stage of Registration

i) Registered (Indicate registration No.)

(Trade name to be in use 6 months after approval to allow exhaustion of old stock)

ii) Undergoing trials (State institution carrying out trials and permit No.)

iii) Other (indicate)

Name of Manufacturer

Name of Registrant (Proprietary owner of Technical information)

.....

Name of agent

Signature of applicant Date

Official Stamp of Applicant/Company

For official use only

Please check whether the following documents have been provided:

Registration Department:

- 7. Has an original letter from the owner of the trade name been provided? Yes No
- 8. Has the applicant attached a copy of the draft label?
 Yes No
- 9. Has KIPI confirmed availability of new trade name? Yes No
- 10. Has the applicant submitted a sample of the pest control product? (not mandatory) Yes No.....

Inspection Department:

- 11. Is the applicant licensed as a pesticide dealer/agent with PCPB in the current year?
 Yes No

Indicate Receipt Number..... Date.....

- 13. Has the applicant paid the change of trade name fee? Yes... no...

Indicate Receipt Number Date.....

- 14. Departmental recommendations

Recommended <input type="checkbox"/>	Not Recommended <input type="checkbox"/>	Recommended <input type="checkbox"/>	Not Recommended <input type="checkbox"/>	Recommended <input type="checkbox"/>	Not Recommended <input type="checkbox"/>
Date.....		Date.....		Date.....	
Registration Officer		Inspector		Accountant	

* superlatives are not allowed and Trade Names should be simple, append additional information

- 15. Recommendation of the Technical and Registration Committess of the Board

Recommended Not Recommended

Reason for rejection.....

Date.....

16. Decision by the Board of Management

Approved Not Approved Date.....

Reasons for rejection.....

17. Changes affected on Lists of Products and data base by (IT Officer)

Signature..... Date.....

FORM B

(r. 4H)

PEST CONTROL PRODUCTS BOARD

The managing Director/Secretary

Pest Control Products Board (PCPB)

P.O Box 13794 - 00800 Nairobi.

E-mail address: pcpboard@todays.co.ke/md@pcpb.or.ke

Tel: 254- 020 - 8021847/7/8 Fax: 254- 020- 8021865

Website: www.pcpb.or.ke

SUMMARY OF THE DATA SUBMITTED TO THE PCPB FOR REGISTRATION OF A PEST CONTROL PRODUCT.

PART I

Trade Name

The Name and Address of Formulator

Common Name of the active ingredient (s).....

Concentration of active ingredients(s).....

Source of active ingredients(s).....

Chemical Name.....

Formulation type.....

Proposed Uses.....

Packaging/Containers (Material size).....

Registrant (Name, Address, Status).....

Agents/Distributors in Kenya.....

Premises (Reg. No. Date of issue).....

PART II

CHEMISTRY DATA

- (a) Physical/Chemical Properties of the a.i
- (b) Physical/Chemical properties of the technical grade material
- (c) Composition of the technical product (purity%, nature and content of impurities, isomers, by-products - other details should be provided in the dossier)
- (d) Physical/Chemical Properties of the Formulated Products
- (e) Composition of the Formulated Product (Concentration of a.i in the formulation. other details should be provided in the dossier).....
- (f) Method of analysis for determination of the a.i in the technical and formulated products

PART III

Biological(efficacy) Data

- (a) Target Pest(s), Diseases(s), Hosts(s).....
- (b) Method, Rate, Frequency of application.....
- (c) Recommendation for use in Kenya.....
- (d) Recommendations for use by authorized bodies outside Kenya.....

PART IV

Toxicological data

- (a) Acute Toxicological Data of the active ingredient(s)
.....
.....
- (b) Acute toxicity data of the formulated product:..
.....
.....
- (c) Short term toxicity studies
- (d) Other toxicological studies:

- (1) Reproduction studies
- (2) Teratological studies
- (3) Neurotoxicity studies
- (4) Mutagenicity studies
- (5) Long term toxicity/carcinogenicity studies
- (6) Accumulation of compound in tissues
- (7) Metabolic studies
- (8) Effects on the livestock, poultry
- (9) Toxicity Data on impurities
- (10) Toxicity Data on metabolites
- (11) Human toxicology and medical aspects:
 - (1) Hazard to humans
 - (2) Symptoms of poisoning
 - (3) Antidote
 - (4) Treatment.....
 - (5) First Aid Measures
 - (6) Safety Precautions/Restrictions

TV - RESIDUE DATA

- (a) Principal Residues
- (b) Disappearance and fate of residues
- (c) Method(s) of analysis (crops, soil, water, feedsuffs etc.)

PART VI

Environment and wildlife hazards

- (a) Degradation and mobility studies (soil, water, air)
- (b) Toxicity to birds
- (c) Toxicity to fish
- (d) Toxicity to honeybees/beneficial insects
- (e) Toxicity to earthworms, other soil invertebrates
- (f) Changes in soil ecology

PART VII

Information on Approvals/Registrations in other countries

.....

.....

PART VIII

Draft of local label (as per Legal Notice No. 89/1984).

.....
.....

PART IX

Brief prepared by

Signature

Official stamp.....

Date

PART X

Decision of the PCPB registration Sub-Committee

Recommended/Not Recommended for registration

Reasons:-

.....

Date

FORM B1

(r. 4H)

PEST CONTROL PRODUCTS BOARD

P.O. BOX 13794-00800, NAIROBI, WAIYAKI WAY

Tel: 254-020 4446115/4450242 Fax: 254-020 4449072

E-MAIL: pcpboard@todays.co.ke

WEBSITE ADDRESS: www.pcpb.or.ke

SUMMARY OF THE DATA SUBMITTED TO THE PCPB FOR REGISTRATION OF A MICROBIAL PEST CONTROL PRODUCT.

PART I

- Trade Name
- Name of the manufacturer and address
- The Name and Address of Formulator
- Common Name of the active Agent(s)
- Concentration of the active ingredient(s)
- Source of active ingredient(s)
- Scientific name of the microbial Agent
- Formulation type
- Proposed Uses
- Packaging/Containers(Material size)
- Agents/Distributors in Kenya
- Premises (Reg. No. Date of issue)

PART II

PHYSICAL/CHEMICAL PROPERTY OF THE ACTIVE AGENT

- (a) Physical/Chemical Properties of the active agent
-
- (b) Physical/Chemical properties of the technical grade material add (incase different from point a) above)
-
- (c) Composition of the technical product (purity%, nature and content of impurities/contaminants, by-products - other details should be provided in the dossier)
-
-
- (d) Physical/Chemical Properties of the Formulated products
-
-
- (e) Concentration of active agents in the formulation. (Other details should be provided in the dossier)
-
-
- (f) Method if Identification, Enumeration and Bioassay
-

.....

PART III

BIOLOGICAL PROPERTIES OF THE MICROOGANISM

- (a) Origin of microorganisms and its uses.....
- (b) Effect in the non-target organisms.....
- (c) Life cycle of the microorganisms.....
- (d) Infectivity (plant and animals).....
- (e) Dispersal and colonization.....
- (f) Effect of environmental parameters (UV, temperature, soil pH, humidity, nutrition requirements, etc.) on stability and survival
- (g) Relationships to known plant, animal or human pathogens.....
- (h) Genetic stability and factors affecting it (potential mutant).....
- (i) Information on the production of metabolites (especially toxins).....
- (j) Show antibiotics and other anti-microbial properties

PART IV

Biological (efficacy) Data

- (a) Target Pest(s), Diseases(s), Host(s).....
- (b) Mode of action of the microorganism.....
- (c) Method, Rate, Frequency of application.....
- (d) Recommendations for use in Kenya.....
- (e) Recommendation for use by authorized bodiesoutside Kenya.....

PART V

Toxicology data

- (a) Acute Toxicological/Infectivity Data of the active agent(s)
.....
.....
- (b) Acute toxicity data of the formulated product:
.....
.....

- (c) Short term toxicity studies (if there is concern under Tier 1 studies)
- (d) Other toxicological studies(if concerns on Tier 1 and 2)
- (1) Reproduction studies
- (2) Tertological studies
- (3) Neurotoxicity studies
- (4) Long term toxicity/carcinogenicity studies
- (6) Metabolic studies (if microorganism organism is known to produce metabolites)
- (7) Effects on livestock, poultry (if exposure is expected)
- (8) Toxicity information on impurities/contaminants if pathogenic significant
- (9) Toxicity Data on metabolites (if applicable)
-
-
- (10) Human toxicology and medical aspects:
 - (1) Hazards to humans
 - (2) Sysmptoms of poisoning or allergic reactions
 - (3) Antidote
 - (4) Treatment
 - (5) First Aid Measures
 - (6) Safety Precautions/Restrictions

PART VI - RESIDUE DATA

(Data is required if the microorganism produces metabolites)

- (a) Principal Residues
-
- (b) Disappearance and fate of residues
-
- (c) Method(s) of analysis (crops, soil, water, feedstuffs etc.).....
-
-

PART VII

Environment and wildlife hazards

- (a) Degradation and mobility studies (soil, water, air)
-
- (b) Toxicity to birds
- (c) Toxicity to fish
- (d) Effects on aquatic-invertebrates.....
- (e) Toxicity to honeybees/beneficial insects.....
- (g) Effects on other soil microorganisms

PART VIII

- (a) Information on Approvals by local phytosanitary authorities
- (b) registrations in other countries
- (c) Information on approval by national biosafety authority if GMO's
-
-

PART IX

Draft of local label (as per Legal Notice No. 89/1984).

.....

.....

.

PART X

Brief prepared by.....

Signature.....

Official stamp.....

Date.....

PART XI

Decision of the PCPB Technical and registration Committee

Recommended/Not Recommended for registration

Reasons:-

.....

.....

Date

FORM B2

(r. 4H)

PEST CONTROL PRODUCTS BOARD

P.O BOX 13794-00800, NAIROBI, WAIYAKI WAY

Tel: 254-020 4449115/4450242 Fax 254-020 4449072

E-MAIL: pcpboard@todays.co.ke

WEBSITE ADDRESS: www.pcpb.or.ke

**SUMMARY OF THE DATA SUBMITTED TO THE PCPB FOR
REGISTRATION OF MICROBIAL PEST CONTROL PRODUCT.**

PART I

Trade Name

Collection Number (National museum of Kenya)

The name and address of Formulator.....

Common Name of the active agent(s)

Description of unit.....

Source of active agent.....

scientific name of the agent.....

Form of presentation (Stage of development, carrier material

.....

Proposed Uses

Packaging/Containers (Material, size)

Registrant (Name, Address, Status)

- Agents/Distributors in Kenya
- Premises (Reg. No. Date of issue)

PART II

BIOLOGICAL

- (a) Description of the agent as presented (stage, colour,)
- (b) Taxonomy
- (c) Descriptive identification of the agent
- (e) Host specificity range and effects on non-target species...(including invasiveness, dispersal, colonization ability)
- (f) Development stages/life cycle
- (g) Stability in proposed packaging
- (i) Method of qualification

PART III

Efficacy Data

- (a) Target Pest(s), Disease(s), Host(s).....
- (b) Mode of action
- (b) Method, Rate, Frequency of application.....
- (c) Recommendation for use in Kenya.....
- (d) Recommendations for use by authorized bodies outside Kenya.....

PART IV

Biosafety data

- (a) Bio-Surveillance data available.....
- (b) Relationships to known plant, animal or human parasites.....
- (c) Hazards to humans.....
- (d) Safety precautions/Restrictions.....
- (e) Recommended methods and precautions concerning handling, storage, or storage.....
- (f) Procedures for destructions.....
- (g) measures in case of an accident.....

PART VI

Environmental Data

- (a) Effects of environmental parameters on stability and survival (IV, temperature, soil, pH, Humidity, etc).....

PART VII

- (a) Clearance by phytosanitary authority.....
- (b) Information on approvals/Registrations in other countries

PART VIII

Draft of local label (as per Legal Notice No. 89/1984).....

PART IX

Brief prepared by.....

Signature.....

Official Stamp.....

Date.....

PART X

Decision of the PCPB registration Sub-Committee

Recommended/Not Recommended for registration

Reasons:-

.....

.....

Date

PEST CONTROL PRODUCTS BOARD
P.O BOX 13794-00800, NAIROBI, WAIYAKI WAY
Tel: 254-020 4446115/1150242 Fax: 254-020 4449072
E-MAIL: pcpboard@todays.co.ke
WEBSITE ADDRESS: www.pcpb.or.ke

**SUMMARY OF THE DATA SUBMITTED TO THE PCPB FOR
REGISTRATION OF A BIOCHEMICAL PEST CONTROL PRODUCT**

PART I

1. Trade Name.....
2. The Name and Address for Formulator.....
3. Common Name of the active ingredient(s).....
4. Concentration of active ingredients(s).....
5. Biological source of a.i. (for botanical specify the plant part, stage of growth etc.)
6. Name & Location of producer of a.i.....
7. Chemical Name.....
8. Formulation type
9. Proposed Uses.....
10. packaging/Containers (Material, size).....
11. Registrant (Name, Address, Status).....
12. Agents/Distributors in Kenya.....
13. Premises (Reg.No. Date of issue).....

PART II

CHEMISTRY DATA

14. Physical/Chemical Properties of the a.i
- (b) Physical/Chemical properties of the technical grade material.....
- (c) Composition of the technical product (purity %, natures & identity of impurities other details should be provided in the dossier).....
- (d) Physical/Chemical Properties of the Formulated Product.....
- (e) Composition of the Formulated Product (Concentration of a.i in the formulation. other details should be provided in the dossier).....
- (f) Method of analysis for determination of the a.i in technical and formulated products (State all the methods for different components)

.....
.....

PART III

Biological(efficacy) Data

- (a) Target Pest(s), Diseases(s), Host(s).....
- (d) Mode of action
- (c) Method, Rate, Frequency of application.....
- (d) Recommendation for use in Kenya.....
- (e) Recommendations for use by authorized bodies outside Kenya.....

PART IV

Toxicological data

- (a) Acute Toxicological Data of the active ingredient(s).....
- (b) Acute toxicity data of the formulated product:-.....
- (c) Short term toxicity studies.....
- (d) Other toxicological studies:
 - (1) Reproduction studies.....
 - (2) Teratological studies.....
 - (3) Neurotoxicity studies.....
 - (4) Mutagenicity studies.....
 - (5) Long term toxicity/carcinogenicity studies.....
 - (6) Accumulation of compound in tissues.....
 - (7) Metabolic studies.....
 - (8) Effects on livestock, poultry.....
 - (9) Toxicity Data on Impurities.....
 - (10) Toxicity Data on metabolites.....
 - (11) Human toxicology and medical aspects:
 - (1) Hazards to humans.....
 - (2) Symptomps of poisoning.....
 - (3) Antidote.....
 - (4) Treatment.....
 - (5) First Aid Measures.....

(6) Safety Precautions/Restrictions.....

PART V

- (a) Principal Residues.....
- (b) Disappearance and fate of residues.....
- (c) Method(s) of analysis (crops, soil, water, feedstuffs etc.).....

PART VI

- (a) Degradation and mobility studies (Soil, water, air).....
- (b) Toxicity to birds.....
- (c) Toxicity to fish.....
- (d) Toxicity to honeybees/beneficial insects.....
- (e) Toxicity to earthworms, other soil invertebrates.....
- (f) Changes in soil ecology.....

PART VII

Information on Approvals/Registrations in other countries.....

PART VIII

Draft of local label (as per Legal Notice No. 89/1984).
.....

PART IX

Brief prepared by.....
 Signature.....
 Official stamp.....
 Date.....

PART X

Decision of the PCPB registration Sub-Committee
Recommended/Not Recommended for registration..

Reasons:-

.....
.....

Date.....

FORM B4

(r. 4H)

P.O BOX 13794-00800, NAIROBI, WAIYAKI WAY

Tel: 254-020 444611/4450242 Fax: 254-020 4449072

E-MAIL: pcpboard@todays.co.ke

WEBSITE ADDRESS: www.pcpb.or.ke

**SUMMARY OF THE DATA SUBMITTED TO THE PCPB FOR
REGISTRATION OF A SEMIOCHEMICAL PEST CONTROL PRODUCT**

PART I

1. Trade Name.....
2. The Name and Address of Formulator.....
3. Common Name of the active ingredient(s).....
4. Concentration of active ingredient(s).....
5. Source of a.i (natural or synthetic).....
6. Name & Location of producer of a.i.....
7. Chemical Name.....
- 8a. Formulation type.....
- 8b. Associated Device.....
9. Proposed Uses.....
10. Packaging/Containers (Material, size).....
11. Registrant (Name, Address,).....

- 12. Agent/Distributors in Kenya.....
- 13. Premises (Registration No. Date of issue).....

PART II

CHEMISTRY DATA

- 14(a) Physical/Chemical propoerties of the technical grade material.....
- (b) Composition of the tecnical product (purity %, nature & identity of impurities other details should be provided in the dossier).....
- (c) Physical/Chemical Properties of the Formulated Product.....
- (d) Composition of the Formulated Product (Concentration of a.i in the formulation, Other details should be provided in the dossier).....
- (e) Method of analysis for determination of the a.i in technical and formulated products (State all the methods for different components).....

PART IV

TOXICOLOGICAL DATA

(A) Technical grade active ingridient(s)

TIER I Requirements

- (a) Acute Toxicological Data of the Technical grade active ingridient(s) Straight-Chain Lepidopteran Pheromones (SCLPs) are exempt from all toxicological data requirements. The following studies are required for non-SCLPs

Acute oral LD50

Acute dermal LD50.....

Inhalation LC50.....

- (b) Short term toxicity studies.....

- (c) Mutagenecity studies.....

TIER II Requirements (Information is required are triggered by TIER I studies

- (1) Reproduction studies.....
- (2) Teratological studies.....
- (3) Neurotoxicity studies.....
- (4) Additional mutagenecity studies.....
- (5) Carcinogenicty studies.....
- (6) Chronic toxicity.....
- (7) Hypersensitivity/allergies in human or any other human exposure data
- (8) Metabolic studies.....

Repealed

(B) Acute toxicity data of the formulated product:

SCLPs are exempt provided the co-formulants are not of toxicological concern (MSDS must be provided.)
The Acute toxicity studies will be provided for non-SCLPs if any of the co-formulants are of toxicological concern.

.....
.....

PART V

EMERGENCY PROCEDURE IN CASE OF A ACCIDENTAL EXPLOSURE OR POISONING

- (a) Hazards to humans.....
- (b) Symptomps of poisoning
- (c) Antidote.....
- (d) Treatment.....
- (e) First Aid Measures.....
- (f) safety Precautions/Restrictions.....

PART VI

- (a) Toxicity to birds (Required if the product could be ingested by birds, e.g. a granular formulation)
- (b) Toxicity to fish (Required if product is applied by air, or directly to water or at a rate exceeding natural background levels)
- (c) Freshwater invertebrates (Required if product is applied by aircraft, or directly to water or at a rate exceeding natural background levels)
- (d) Algae (Waived for products in affixed dispensers and if exposure is unlikely to exceed natural background levels)
- (e) Toxicity to bees (Information/discussion, to address whether behaviour or reproduct would be affected, is required if exposure is likely to exceed natural background levels)
- (f) Toxicity to earthworms (Required if product is applied to soil and can accumulate in soil. Required if exposure exceeds natural background levels)

PART VII

Information on Approvals/Registrations in other countries.....

PART VIII

Draft of local label (as per Legal Notice No. 89/1984).

.....

PART IX

Brief prepared by.....

Signature.....

Official stamp.....

Date.....

PART X (For official use only)

Decision of the PCPB registration Sub-Committee

Recommended/Not Recommended for registration

Reasons:-

Date.....

FORM C

(r. 4F(1)(a))

PEST CONTROL PRODUCTS ACT

[Cap. 346]

PEST CONTROL PRODUCTS (REGISTRATION) REGULATIONS, 2006

APPLICATION FOR THE INTRODUCTION OF NEW PEST CONTROL PRODUCT

(To be completed and submitted in triplicate)

To: The Managing Director
Pest Control Products Board
P.O. Box 13794-00800
Westlands, Nairobi

APPLICANT'S NAME AND ADDRESS.....

Tel No. Fax No.

STATUS OF APPLICANT (Manufacturer, agent etc.).....

1. Approved Common Name(s).....
2. Chemical Name.....
3. Chemical formula.....
4. Chemical structure.....
5. Trade Name(s).....
- 6: Proposed Kenyan Name(s).....
- 7: Formulation Type (W.P.E.C., Dust etc.).....
8. Concentration of Each Active Ingredients.....
9. Quantity required for testing.....
10. Proposed Uses (Agricultural, Health, Veterinary, Forestry etc.).....
11. Location and Area of Test Plots.....
12. Target pest(s) Host(s) or Area of Application.....
13. Mode of action.....
14. Toxicity of the product to test animals (Acute Oral and Dermal LD50 Inhalation LC50 etc.).....
15. The effects of the product on the environment-
 - (a) Toxicity to bees.....
 - (b) Toxicity to fish.....
 - (c) Toxicity to birds.....
 - (d) Toxicity to soil micro-organism.....
16. Proposed precautions to users.....
17. Antidote, Treatment of poisoning.....
18. Shelf life of the product.....
19. Country of Origin of the product.....
20. Name and Address of manufacturer.....
21. Name and address of formulator.....
22. Countries where tested and registered

Repealed

- 23. Ownership of data (name, address).....
- 24. Patent sit (lilt ion/patent holder)..... Confirm that the information contained herein is true to the best of my knowledge and belief.

.....

Date of application

Signature of applicant

Name..... Designation/Position held

Note

Every applicant must be accompanied by

- 1. Supporting data and information which should include -
 - (a) Chemistry specifications, composition of the product, and the technical a.i method of analysis of the a.i determination
 - (b) information on biological activity on the product, directions for use
 - (c) Metablisim, Residues, methods of analysis for residues
 - (d) Toxicological data on the technical and formulated product(s)
 - (e) Environmental toxicity
- 2. Experimental labels (typed)
- 3. Analytical standards (approximate 100% a.i) 1.0 gram.

FORM D

(r. 3A(2)(a))

PEST CONTROL PRODUCTS ACT

[Cap. 346]

PEST CONTROL PRODUCTS (REGISTRATION) REGULATION, 2006

Date

REF: PERMIT NO.

PERMIT FOR EXPERIMENTAL AND EFFICACY TRIALS OF NEW PEST CONTROL PRODUCTS

This is to grant permission as requested for your Centre/Organization to carry out efficacy trials of the new pest control product(s) as indicated below -

Pest Control Product(s)

Crop(s)/Commodity(ies)/Use(s)/Target Pest(s)

You are requested to inform the Pest Control Products Board of the commencement of the experimental/efficacy trials and also periodically submit to the Board progress reports. The trial should be carried out using a Pest Control Products Board approved trial protocol. At the conclusion of the experimental/efficacy trial, a detailed confidential report on the performance of the candidate pesticide and recommendations for its use shall be submitted to the Board quoting the above reference and date.

It would be highly appreciated if trials are completed as quickly as possible to avoid delays in introducing suitable products in the market. The company will provide you with the required trial samples/materials but the Board shall not meet expenses for the trials.

It is the responsibility of the applicant to ensure that the efficacy trials are carried out to the satisfaction of the Board.

Managing Director

Pest Control Products Board

FORM E

(r. 14(2))

CHANGE OF SOURCE

PRODUCT INFORMATION

1. Trade Name:.....
2. Common name of active ingredient(s) and Concentration:.....
3. PCPB registration No:.....
4. Technical Grade Active ingredient
 - (a) Name of basis manufacture:.....
 - (b) Physical Location of basic manufacture:.....
 - (c) Address:
 - (i) Postal Address:
 - (ii) Telephone No:
 - (iii) E-Mail:
 - (iv) Fax No:
 - (v) Street/Road:
 - (d) Specifications (with analytical proof)
 - (e) Relationship with Registrant, if different
5. Formulated Product
 - (a) Name of Formulator:
 - (b) Physical Location:

(c) Address

(i) Postal Address:.....

(ii) Telephone No:

(iii) Fax No:

(iv) E-Mail:

(v) Street/Road:

(d) *Composition (with analytical proof)*

6. Name of new source

7. Relationship with old source/manufacturer/formulator, if more than one.