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Government of India /Ministry of Agriculture & Farmers Welfare

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(Department of Agriculture, Co-operation & Farmers Welfare)

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DIRECTORATE OF PLANT PROTECTION, QUARANTINE & STORAGE

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Central Insecticides Board and Registration Committee

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N.H. IV, FARIDABAD (HARYANA)-121001

CERTIFICATE OF REGISTRATION UNDER SECTION 9(4) OF THE INSECTICIDES ACT.,1968.

Certified that the Insecticide Neem Based Granular Formulation Containing Azadirachtin 0.15% (1500 ppm) w/w min indigenous manufacture indigenous manufacture has been registered under section 9(4) of the act in the name of the Person/Undertaking whose particulars are specified below:

1.	**Name of the person/Undertaking :**	**M/s M/s Comis Biotech Pvt. Ltd. Plot No-3B ,Part 3/1/2,Basement Floor ,Rahul Enterprises Building ,D-Block,MIDC-Akurdi.Pimpri Chinchwad Pune, Pune, Maharashtra 411019**
2.	**Address of the manufacturing premises :**	**Plot No-3B ,Part 3/1/2,Basement Floor ,Rahul Enterprises Building ,D-Block,MIDC-Akurdi.Pimpri Chinchwad Pune MH-411019 IN.**
3.	**Registration No.**	**CIR-174233/2020-Neem Based Granular Formulation (Containing Azadirachtin) (423)-1727 F.No. 101450-F/9(4)/2020**
4.	**Name of the Insecticide :**	**Neem Based Granular Formulation Containing Azadirachtin 0.15% (1500 ppm) w/w min indigenous manufacture**

5. Conditions :

i) The insecticide shall be manufactured indegenously.

ii) The registration is subject to the strict compliance of various provisions of the Insecticides Act, 1968 as amended from time to time and Rules, bye-laws framed and notifications issued there under and as amended from time to time.

iii) The registration certificate is further subject to such conditions which may be varied and specified from time to time by the Registration Committee under section 9(3c).

iv) Non-compliance of the conditions set out herein before and hereinafter will entail action under section 17 of the Act.

v) The insecticide shall have the composition (kind, name and percentage of the ingredients) as given below :-

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Neem Extract Concentrate Containing Azadirachtin (a.i.)	**0.15% % w/w min.**
Color (Crystal Violet)	**0.02 % % w/w**
Binder (Paraffin Wax)	**1.50% % w/w**
Inert Material (Sand)	**Q.S. %**
	Total : 100.00 % W/W

vi) The insecticides shall contain the maximum impurities as quantified/identified and submitted to the Registration Committee.

vii) The Product shall confirm to the specification submitted by you and also to the IS vide No. and amendment thereof as and when the same are formulated and published.

viii) A sample quantity of the insecticide being registered alongwith a small quantity of reference analytical standard should be sent to the Director, Central Insecticides Laboratory, Directorate of Plant Protection, Quarantine & Storage, N.H. IV, Faridabad, as and when required, for verifcation.

ix) The shelf-life of the insecticide shall be one year(s).

x) A copy, each of the approved label and leaflet is enclosed. No change, addition, alteration, modification or deletion with respect to the inscriptions on the lables/leaflets shall be done without the prior approval of the Registration Committee.

xi) The labels and leaflets shall be printed by using letters that are BOLD enough for a man of ordinary/normal vision to read them without any external help.

xii) The Registration Committee does not itself responsible for the use of trade name by you. The use of the trade name shall be regulated as per the existing laws on the subject.

xiii) The licence should be granted subject to the conditions that the licensee shall comply with the provisions of the Act and the Rules made thereunder and the conditions of registration for the time being in force.

xiv) No licence to manufacture an insecticide shall be granted unless the licensing officer is satisfied that necessary plant and machinery, safety devices, first-aid facilities, quality control measures, the requirement laid down under Chapter VIII of the Rules, etc. exist in the premises where the insecticide is proposed to be manufactured.

xv) Inspection of the manufacturing unit should be undertaken to collect in-process samples of the insecticides. The samples should be analysed to verify the claims made by the licensee relating to chemical parameters, and the report thereof should be submitted to the Registration Secretariat within a period of six months.

xvi) No export should take place in contravention to the provisions of the Rotterdam Convention on prior informed consent procedure for certain hazardous chemicals and pesticides in international trade.

xvii) If a pesticide is banned or severely restricted in India, before exporting such pesticide, permission from Designated National Authority for Pesticide of the Country under Rotterdam Convention may be obtained.

xviii) Health records of Industrial workers may be maintained in Appendix E (conditions) of Form-III as prescribed in the Insecticides Rules 1971. In case any untoward/adverse effect is noticed, then the same may be reported to Registration Secretariat by the Licensing Officer.

xix) The registrant shall have to commence actual production of the pesticide within three years from the date of issue of registration certificate and is required to produce a certificate from the concerned State/UT Governments as a proof of production, failing which the certificate of registration shall automatically lapse.

xx) In case of export, the packaging shall be as per the requirement of the importing country and conforming to IMDG guidelines.

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xxi) The registrant has to submit detail of Import/Export or indigenous manufacturing (as the case may be) of this pesticide month wise mandatory to the Secretariat of CIB&RC as the case may be. In case non compliance of this condition is observed this CR shall be cancelled immediately without any Notice.

xxii) The product is registered for domestic use as well as for Export.

xxv) The product is registered for domestic as well as for export and in case of export primary packaging shall be as per the requirement of importing country.

xxvi) The registrant is required to use the label and leaflets as per GSR 355(E) dated 04.06.2020 and applicable Insecticides Rules. The Issuance of the Certificate of Registration is strictly subject to the outcome of the W.P(C) No. 4136/2020 and W.P(C) No. 4137/2020.

Specific Conditions :

The product Neem based Granular Formulation Containing Azadirachtin 0.15% (1500 ppm) w/w min is approved of Rice leaf folder and rice stem borer in Paddy crops only. The finished formulation shall be packed in milky white LDPE bag/pouches of 100 gms, 250gm, 500gm, 1 kg and 5 kg capacity with film thickness not less than 0.062 mm, as a primary pack as per IS: 2508-1984. This pouches shall be heat sealed at both the ends to make it leak-proof and shall be inserted individually in cardboard carton confirming to IS:-6604- 1991 up to 500gm, 1 kg and 5 kg shall be individually packed in E-fluted, 3 ply cartons. CFB boxes of double flute (A+B) of capacity 10 kg maximum, conforming to IS: 2771 (Pt.-I): 1990 shall be use for transportation purpose. No other packaging system shall be used without approval of Registration Committee.

Faridabad
Dated : 16/12/2020

(Dr. J P Singh)
Secretary
Central Insecticides Board
and Registration Committee

Copy To: The Director of Agriculture Maharashtra

AUTHENTICATED

(CIR I)
Section Officer

(Dr. J P Singh)
Secretary
Central Insecticides Board
and Registration Committee

"Note: The Labels and leaflets have been generated through newly created databank. Notwithstanding deligence exercised in creation of the databank, the possibility of errors creeping into labels and leaflets cannot be ruled out. The same is subjected to necessary correction."

Signature Not Verified
Digitally Signed By DR J P SINGH
Secretary, CIB&RC
Thu, 17 Dec 2020 19:16:26 IST
Location: CIB&RC, NH4, CGO Complex, Faridabad