BioStimulants: Product Registration*
(Manufacturers & Importers)

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BioStimulants*

"biostimulant" means a substance or microorganism or a combination of both whose primary function when applied to plants, seeds or rhizosphere is to stimulate physiological processes in plants and to enhance its nutrient uptake, growth, yield, nutrition efficiency, crop quality and tolerance to stress, regardless of its nutrient content, but does not include pesticides or plant growth regulators which are regulated under the Insecticide Act, 1968 (46 of 1968);

The biostimulants specified in Schedule VI shall be classified under any of the following categories, namely:

(a) botanical extracts, including seaweed extracts; (b) bio-chemicals;
(c) protein hydrolysates and amino acids; (d) vitamins;
(e) cell free microbial products; (f) antioxidants;
(g) anti-transpirants; (h) humic and fulvic acid and their derivatives

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BioStimulants: Product Registration*

(Manufacturers & Importers)

- Provisional Registration

&

- Permanent Registration

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BioStimulants: Provisional Registration*

(Manufacturers & Importers)

- Preparation of Product Sales Document
- Application to State Department of Agriculture for FORM G 2
- Product wise application to “The Controller of Fertilizer”, New Delhi in G 1 format.
- Product wse Provisional Registration by “The Controller of Fertilizer” in format G 3.

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BioStimulants: Provisional Registration*

(Manufacturers & Importers)

PRODUCT SALES DOCUMENT**

- One of the most Important documents
  - to get Form-G 2 from State Department of Agriculture; &
  - to get Provisional Registration (application in Form G 1) from “The Controller of Fertilizer”, GOI

To avoid any further challenges (as per our experience so far),

**to be prepared carefully

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BioStimulants: Provisional Registration

G 2 Application*: GENERAL CHECK LIST OF DOCUMENTS

(for Self Manufactured/ Imported Biostimulants)

S. No.	Details	Remarks
1	Duly filled Application for G-2 in the prescribed format (as per FCO amendment dt. 23-02-2021)	Duly Signed
2	Last 3 Years Sale report / GST Data for the Last 3 Years / Stock, Production, Trade of the product for last 3 years duly verified by Chartered Accountant.	Self Attested
3	Label of Product(s)	Self Attested
4	Self Declaration of No report ill effect or hazardous effect or any International publication or world wide acceptable test report supporting Non hazardous effect	Notarized.
5	Name of responsible person and his / her acceptance on stamp paper (as per clause 24 of FCO 1985)	Notarized.

Note: *Kindly check with the Director of Agriculture of your State for specific details needed (if any) before applying to obtain the G 2. Once we complete the documentation for G 2 application - Verification of facts, Applying to the authority, Follow up and Certificate (FORM G 2) from the Director Department of Agriculture is to be obtained directly by the client(s) from their respective States.

BioStimulants: Provisional Registration*

G 1 Application**: GENERAL CHECK LIST OF DOCUMENTS

Application in the prescribed format (FORM G 1)

1. Form G 2 from State Department of Agriculture,
2. Copy of Certificate of Incorporation / company registration (company address proof)
3. Copy of PAN card of of Authorized Signatory
4. Copy of ADHAR of Authorized Signatory
5. Technical details of product
6. Product label
7. Product sales document for last 3 years
8. Draft of FORM G 3 - Certificate of Provisional Registration

** to be applied after receiving G 2 from the State Deptt. of Agri.

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BioStimulants: Provisionally Registered Product(s)

1. Provisional registration shall be valid up to 22-02-2023.
2. Products not registered during this period not allowed after March 31st, 2022.

Ref. : Clarification from Ministry of Agri. & Farmers Welfare Dated 2nd Aug. 2021

BioStimulants: Permanent Registration*

(Manufacturers & Importers)

Provisional Registration: Valid till March 22-02-2023

Permanent Registration: Following Studies to be undertaken

- Chemistry Study: Through GLP / NABL accredited laboratory
  - Physicochemical properties of active ingredients and adjuvants, if any
  - Shelf-life study
  - Toxicity: Acute toxicity; & Eco Toxicity
  - Detection of 6 heavy metals
- Agronomic BioEfficacy: ICAR institutes and/ or State Agri. Universities
  - 1 season trial at 3 agro-ecological locations per formulation

Application in FORM G, along with study reports to be submitted to **"The Controller of Fertilizer"**

for

PERMANENT REGISTRATION.

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BioStimulants: Registration

(hFService Support)

Provisional Registration: G 1 Application

- Facilitation in documentation ,
- Customized formats and guidance w.r.t.
  - relevant information collection (for Product Sales Document),
  - preparation of separate applications (after we receive the G 2 from the client)),
  - verification and finalization (on the basis of available details; and cross checking with the client)
  - getting signed / stamped by the client,
  - submission to the authority (The Controller of Fertiliser, New Delhi),
- Regular follow up, providing additional information (if needed) in consultation with the client for provisional registration.

Note: *Once we complete the documentation for G 2 application - Verification of facts, Applying to the authority, Follow up and Certificate (FORM G 2) from the Director Department of Agriculture is to be obtained directly by the client(s) from their respective States.

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BioStimulants: Registration

(hFService Support)

Permanent Registration:

Service details for Studies from NABL / GLP Certified Lab: Chemistry, Shelf-life, Toxicity: 5 acute studies and 4 eco toxicity studies; Heavy metal detection against their prescribed limit etc.

- Suitability of lab for cost effective testing,
- finalisation of testing protocols and testing schedule,
- coordination, monitoring and review of draft reports, modifications / improvements, if needed and
- finally procuring the final report for the studies:

Service details for Agronomic BioEfficacy Study:

- Identification of SAUs / ICAR institute as per suitability of crop(s);
- Formal request and follow up to with officials for their consent
- propose trial protocols, observations (as per expectations) in consultation with the client.
- Regular follow up for timely completion of trial(s).

Note: 3rd Party Studies, Follow up, Monitoring, Report, Documentation, Application, Additional Information (if needed) and Registration from competent authority.

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BioStimulants: Product Registration*

(Manufacturers & Importers)

- Visit the Apply section to share product (BioStimulant detail)
- Kindly Apply Online

THANKS

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