Following Documents should be submitted According to the Registration Guideline RP/01/GL (rev-5/20/2011). Reference to the Guideline is given in the parenthesis. If relevant data is submitted indicate the Annexure and page number(s).

No	Requirements in a Dossier for Pesticide Registration	Requirements fulfilled (√ Yes)	Requirements not fulfilled (√ No)	Relevant Annexure & Page No(s)	Remarks (if any)	Remarks (Office use only)
1	**Application & Dossier**					
i	Duly completed application with indication of reference to relevant annexure and page numbers in the dossier	Yes ☐	No ☐			
ii	Uniform page numbering in technical dossier (top right)	Yes ☐	No ☐			
2	**Authenticated/Original Registration Certificates**					
i	Country of origin for the technical material (s)(2.10)	Yes ☐	No ☐			
ii	Country of origin for the formulation (2.10)	Yes ☐	No ☐			
iii	Other countries where product is registered and used(2.101.1)	Yes ☐	No ☐			
3	**Declaration for supply of the technical/ formulation**					
i	Manufacturer of the technical (3.4)	Yes ☐	No ☐			
ii	Formulator of the product (3.5)	Yes ☐	No ☐			
4	**Active Ingredient**					
i	Physical Chemical properties of pure Active ingredient (s) (4.1, 4.2, 4.4)	Yes ☐	No ☐			
ii	Toxicological Impact of the pesticide & its toxicologically significant metabolites on biological	Yes ☐	No ☐			

	systems (where applicable) (4.3, 4.4)					
5	**Technical Material**					
i	Physical Chemical properties (s) (5.1)	Yes ☐	No ☐			
ii	Detailed composition including impurities-Typical recipe- Original certificate (5.2)	Yes ☐	No ☐			
6	**Formulation**					
i	Physical Chemical properties (6.1)	Yes ☐	No ☐			
ii	Detailed composition including impurities-Typical recipe- Original certificate (6.2)	Yes ☐	No ☐			
iii	Real assay result of the composition- Duly signed analytical report from accredited laboratory (6.2)	Yes ☐	No ☐			
iv	Analytical method used to determine the active ingredient in the formulation including chromatograms (6.3)	Yes ☐	No ☐			
v	Declaration on shelf life- Formulators original certificate (6.4)					
vi	Stability test report on shelf life – CIPAC elevated temperature test (54^0C/ 14 days) or ambient temperature for 2 years (6.4)	Yes ☐	No ☐			
vii	**Acute toxicity**-oral / dermal / inhalation/ skin sensitization/ skin irritation/ eye irritation Original or certified true copy of the toxicology test report summaries with Certified GLP standards and OECD or similar guideline (6.5)	Yes ☐	No ☐			
viii	Toxicological profile for the ingredients exceeding 20% of the formulation (6.5)	Yes ☐	No ☐			
ix	Analytical report for a**rsenic, cadmium, cobalt, chromium, mercury, nickel, lead, tin, thallium, cyanide** (Circular -RP/2011-07)	Yes ☐	Yes ☐			

x	Bio efficacy summary reports/test with use recommendation and/or dilution rates (6.6) (Scientifically conducted & statistically verified reports from accredited institutions)	Yes ☐	Yes ☐			
xi	CDDA/VRI/ MRI/IPSC/ Other (Specify)- Recommendation certificate	Yes ☐	No ☐			
xii	Biological Mode of Action (6.6)	Yes ☐	No ☐			
7	**Material Safety Data Sheet (MSDS)-** certified by the manufacturer and/or an accredited agency	Yes ☐	No ☐			
8	**Regulatory Actions Taken by other Governments to restrict or ban use –** Indicate if any					
9	Quality certificate issued by an internationally accepted accredited body	Yes ☐	No ☐			
10	Analytical fee (If applicable) submitted in favor of testing body					
	MRI	Yes ☐	No ☐			
	ITI	Yes ☐	No ☐			
	Other	Yes ☐	No ☐			
11	Product label (If applicable)	Yes ☐	No ☐			