Report No.

Summary Sheet for Data Submission for the Pesticides Subcommittees (DPH/IP)

01. Product name		
02. Common name & Strength		
03. Registrant (Local)		
04. Request /Use classification		
05. Product Category		
06. Chemical Class of active ingredient (AI)		
07. Manufacturer/Country		
08. Reg. Status in Manufacturing country		
09. Other Countries Registered		
10. Formulation Type		
11. Types of containers to be used		
12. Intended Use	**Pests**	
	Disease	
13. Mode of Action (Summary)		
14. Toxicity Data (Acute) / Formulation		Oral Toxicity in rat- …….. ……mg/kg body weight Dermal Toxicity in rat -……. mg/kg body weight Inhalation 4h LC50 in rat – …….mg/l
15. Sub Chronic & Chronic (three months-two years) feeding studies in rat, dog and mouse; Carcinogenicity /Mutagenicity /Reproductive Toxicity/ Teratogenicity; if any		
16. Chemical & Physical Properties		Solubility in water-
		Octanol/water Partition Coefficient, pH, Flash point, etc.
17. Ecological Information		Birds, acute LD_{50} or dietary LC_{50} toxicity
		Bees, Acute oral & contact toxicity
		Fish, acute LC_{50} (96h)
		Aquatic invertebrates, daphniabEC50 (48h)
18. Fate in the Environment		Degradation in soil
		Degradation in the aquatic environment
19. Hazard Class (WHO/GHS)		
20. Estimated amount to be imported per year		
21. Proposed Disposal Method (Container & expired products)		
22. WHO Specifications		
23. Rate of Application		
24. Method of Application		
25. Period of effectiveness		
26. Registration Status		**** US EPA:** ***** NMRA:** ****** VDCA:**

For Office Use:

ROP concerns	

Note: Detail compositions of the technical materials and the <u>product</u> and <u>MSDS</u> of the product also need to be submitted with this summary sheet.

*** Use separate sheet, if allocated space is not enough. **** Veterinary Drugs Control Authority.**

**** Environmental Protection Agency, *** National Medicines Regulatory Authority (NMRA)**