FORM 1 MEDICINE IDENTIFICATION SHEET

Use one form per DIN

Please Spec 1 NAME(S) AND USE(S) OF TH		
Brand Name:		
Generic Name:		
Therapeutic use(s) of the medicine		
Approved by Health Canada		
Human	(if the medicine is for human use and is a controlled substance as defined in the Controlled Drugs Prescription And Substances Act or contains a substance listed or described in Schedules C or D to the Food and Drugs Act or Schedule F to the Food and Drug Regulations)	
<u>OR</u>	OR (if the medicine is for human use and is not a controlled substance as defined in the Controlled Over the counter Over the counter	r D
Veterinary		
2 NOTICE OF COMPLIANCE	(N.O.C.)	
First N.O.C. Y M I	Check if Special Access Program or applicable Or Clinical Trial Application or Investigational New Drug	
3 DRUG IDENTIFICATION NU	JMBER (DIN)	
Drug Identification Number	Dosage Form Strength/Unit	
4 DATE OF FIRST SALE	5 PRODUCT MONOGRAPH	
Date of 1st Sale Y M I	Product Monograph OR Draft Product Monograph (Copy Included) OR (Copy Included) Other (Copy Included)	
6 PATENT NUMBER OF PATE	ENTEE'S OR FORMER PATENTEE'S INVENTIONS PERTAINING TO THE MEDICINE	
Patent Number	Date Granted Expiration Date	
1 1 1 1 1 1	Y M D Y M D	
	Y M D Y M D	
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	Y M D Y M D	
	Y M D Y M D	

Priviledged s87 Patent Act

FORM 1 MEDICINE IDENTIFICATION SHEET

7 PATENT APPLICATION NUMBER OF PATENTEE'S INVENTIONS PERTAINING TO THE MEDICINE

Patent Application Number					Date of Filing of Application									
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									Y		N	Л	I)
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8 REPORTING PATENT	TEE or FORMER PATENTEE
Patentee Name	
Patentee Address	
Identity if the reporting	
patentee is:	the patent holder person holding a licence other (specify)
9 CERTIFIED BY: (in ac	cordance with Section 7 of the Patented Medicines Regulations)
	I hereby certify that the information presented is true and correct.
Signature of reporting patentee,	
former patentee or its corporate	officer:
Title:	
Date:	
Tel. Number: ()	- Fax Number: () -
E-mail:	