DEPARTMENT OF PHARMACEUTICAL SERVICES MINISTRY OF HEALTH

BRUNEI DARUSSALAM

Form No. DPS/DRS/05

Effective Date:

APPLICATION FOR AN IMPORT PERMIT OF A MEDICINAL PRODUCT ON **CONSIGNMENT BASIS**

Application No. (for official use only):				
A) PARTICULARS OF APPLICANT				
1.	Name of Firm/Company: (in BLOCK letters)			
2.	Business Address:			
3.	Telephone No: Fax No:			
	Official E-mail:			
4.	*Company/Business Registration No.:			
5.	Authorised Person making the Application on behalf of Firm/Company:			
	Name:			
	Residential Address:			
	Designation:			
	Designation.			

^{*}Delete where applicable

B) PARTICULARS OF PRODUCT TO BE IMPORTED

1.	Brunei Product Registration No. of the locally registered product:								
	BRU	L			F				
2.	Name of Product (state strength & dosage form):								
			•••••			• • • • • • •			••
3.									
3.	Name & Strengt	h of Active Ingred	dient(s):						
4.	Name of Manufacturer:								
	T								
5.	Country of Manufacturer:								
,	T								
6.	Quantity to be In	nported:							
	Unit No.	Pack Size	Bato	h / Lot No	0.		Expi	ry	
							•		
7.	Product to be Imported from (country):								
8.	Purpose of Impo	rt (please tick wh	ere applicable)					
	For Supply to:								
	☐ Quotation ☐ Private Hospital ☐ Private Clinic ☐ Others, please state:								

C) PARTICULARS OF POISONS LICENCE HOLDER (For Poisons Licence Holder Only)

1.		lame of Licence Holder:				
	1 433	port 1.0	1.0.110.			
2.	Poisons Licence No.:					
	1					
3.	a) Ir	nport Permit No. for Psychotropic (where applicable)	Drugs:	b) Import Permit No. for Narcotic Drugs: (where applicable)		
	Р					
D) I	DECL	ARATION				
I, or that:		alf of the company named in Sect	tion A, h	ereby declare that/ undertake to/ confirm		
i)		All particulars given in this application are true and that the documents enclosed are authentic or true copies.				
ii)		Notify Drug Administration Section within one week of any change in the particulars submitted in this application and new safety information or any adverse drug reactions related to the product.				
iii)		Comply with the conditions imposed in the import permit and guideline issued by the Department of Pharmaceutical Services, Ministry of Health, that are related to the marketing of the medicinal product in Brunei Darussalam.				
iv)		Responsible for the quality, safety and efficacy of the product to be imported and that it is in all respects identical to the medicinal product registered in Brunei Darussalam.				
I understand that a willfully false statement is an offence under the Medicines Order.						
Signature of Authorised Person:						
Name of Authorised Person:						
Company		Stamp:	Date:			

FOR OFFICIAL USE ONLY

Application No:	Date Received:				
Signature of Verifying Officer: Name of Verifying Officer:					
Remarks:					

No. of attachments:

LIST OF ATTACHMENT S

Attachment No. (Corresponds to the section & no. on the application form)	Nature of Attachments	For Official Use Only