GMP DESKTOP ASSESSMENT APPLICATION FORM



National Pharmaceutical Regulatory Agency Ministry of Health Malaysia

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Tel. No. : 03-78835400 Fax No. : 03-79571200 Vebsite : http://npra.gov.my

For Official Use Only	
Application No. (GDA)	
Application No. (FI)	
Date Received:	
Date Completed:	

	Website: http://npra.gov.my			Date Completed:			
PART A APPLICANT / PRODUCT REGISTRATION HOLDER INFORMATION							
Name of App							
Name of Prod	duct Registration H	lolder:			-		
Address:		w					
Addic55.							
Company/Bu	siness Registration	Number:					
Contact Tele	ohone:	Contact Fax:	Email Ad	Email Address:			
PART B	FOREIGN MA	NUFACTURER INFO	ORMATION				
Name:							
Address:							
					<u> </u>		
DADTC		PORTING DOCUME			Tick (√) if	For Official	
PART C	non-exhaustive list	cuments MUST be submit; other documents may be	ted together with this requested as evaluati	application. However, this is on progress.)	provided	Use Only	
1.	Current Certificate of Outsourced Laboratory (If applicable)						
2.	Current Manufacturing Licence						
3.	Most recent GMP Inspection Report issued by local authority agency						
4.	Corrective Action and	Prevention Action (CAPA) repo	ort for inspection stated in	(3) above			
5.		Report(s) for on-site inspection	(s) performed by PIC/S P	articipating Authority (related			
6.	report) Quality Manual (or equivalent documentation)						
7.	One sample investigation report for product complaint and recall (related complaints and recall)						
8.	Latest Product Quality Review report (related products)						
9.	Process Validation protocol and report (related products)						
10.	Batch Manufacturing/Packaging Record (BMR/BPR) for batches produced within the last 6-12 months (related products)						
					<u> </u>		
PART D	APPLICANT D	ECLARATION					
1. I am he	reby authorised by	the company to make th	nis application.				
2. I understand that Foreign GMP Inspection by NPRA will be conducted if the evaluation is found to be unsatisfactory.							
3. I hereby declare that details furnished on this form are true, accurate and complete; the supporting documents are authentic or true							
copies.							
	(Signature)		(Date)	(0	Company Stam	p)	
(1)		m)					
(N	ame & Designation	II <i>)</i>					