



RECEIPT FOR SUBMISSION OF REGISTRATION DOSSIER
MEDICAL DEVICE PRODUCT

Local Distributor Name & City			
Contact Person			
Tel:			
Email:			
Store Licence No. & Validity			
Device Name			
Dosage Form			
Strength			
Pack Size(s)			
MAH for the U.A.E.: Name & Country			
Batch Releaser (Manufacturer) : Name & Country			
<i>The presented dossier is verified according to regulation and is</i>			
Received <input type="checkbox"/> Conditionally received <input type="checkbox"/> Rejected <input type="checkbox"/>			
<u>Reason(s) for conditional receipt or rejection</u>			
<p><i>Important:</i> <i>The dossier completion should be within 3 months from the date of submission failing which will entail dossier rejection and the contact person should withdraw the file upon notice.</i></p>			
Pre-registration Analysis:	Exempted		Not Exempted
Received by			
Signature		Date	
Receipt No.	MD / /200		