



MANUFACTURING SITE REGISTRATION CERTIFICATE

It is certified that the following Manufacturing Site has been registered in the UAE Ministry of Health, in accordance with the Article 65 of the Federal Law No. 3 of 1984.

Certificate #	04155C01	First Reg. Date	31-August-2015
Registration #	4155	Reg. Expiry Date	31-August-2020
Committee Meeting No.	11868	Meeting Date	31-August-2015
Payment Receipt No.	020955300715203	Payment Date	30-July-2015
Manufacturing Site Name	ASPIRO PHARMA LIMITED		
Address	Sy.No.321, Biotech Park, Phase- III, Karkapatla, Mulugu Mandal, Medak District, Telangana[INDIA;Hyderabad]		
Activities Registered for			
Manufacture of dosage forms , Packaging & Labeling , Storage & Handling , Laboratory Testing , Batch releaser (certification)			
Non Hazard Line(s) of Production Registered for			
(1) Sterile Products-Aseptically prepared (Dosage Forms) Lyophilisates , Small Volume Liquids , Powder (1) Sterile Products-Terminally Sterilized (Dosage Forms) Small Volume Liquids			
Manufacturing Site for product class(s)	Conventional Medicines		
<ol style="list-style-type: none">1. The evidence(s) for GMP for the above mentioned activity(s) is/are acceptable.2. Failure to provide current acceptable GMP or any equivalent evidence prior to the expiry / as & when requested could result in refusal to receive product registration dossier or removal of the affected company and its product(s) from the register, according to the case.3. Registration as Manufacturing Site makes it eligible to involve in the registered activities in respect to the products to be registered in the U.A.E.4. The Manufacturing Site should apply for minor variation as and when an amendment is needed in registration status.5. This registration applies only to the above site name and address.			


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www.moh.gov.ae

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