



Manufacturing Site Registration Certificate

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

Certificate #	REG/MS/30389/2019	First Reg. Date	2019-02-07
Registration #	6753	Reg. Expiry Date	2024-02-06
Committee Meeting No.	32082	Meeting Date	2019-02-07
Payment Receipt No.	30389000	Payment Date	2019-01-20
Manufacturing Site Name	HEMOMONT		
Address	Ilije Plamenca St., Podgorica, Montenegro, Podgorica, MONTENEGRO		
Activities Registered For			
Manufacture of dosage forms, Packaging & Labeling, Storage & Handling, Assembly, Secondary packaging, Batch releaser (certification), Laboratory Testing			
Non Hazard Line(s) of Production Registered For			
Sterile Products-Terminally Sterilized (Dosage Forms) - Large Volume Liquids Sterile Products-Aseptically prepared (Dosage Forms) - Small Volume Liquids (Eye Drops) Sterile Products-Aseptically prepared (Dosage Forms) - Small Volume Liquids Sterile Products-Aseptically prepared (Dosage Forms) - Small Volume Liquids (Eye Wash) Non-Sterile Products-Non Sterile Products (Dosage Forms) - Liquids for external use Non-Sterile Products-Non Sterile Products (Dosage Forms) - Suspension Sterile Products-Batch certification only - Terminally Sterilized-Large Volume Liquids Sterile Products-Batch certification only - Aseptically prepared-Small Volume Liquids Non-Sterile Products-Batch certification only - Liquids for external use			
Manufacturing Site for product Class(s)	Conventional Medicines		
1. The evidence(s) of 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 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.			



هذه الشهادة صادرة من وزارة الصحة ووقاية المجتمع وتعتبر من الوثائق الحكومية الرسمية ولا تحتاج إلى توقيع، ويحظر قطعياً تقليدها أو إدخال أي تعديلات عليها سواءً بالإضافة أو الحذف أو التغيير في بياناتها أو غير ذلك من أنواع التعديل، وتعد الشهادة لاغية إذا شابهها شيء من ذلك. للتأكد من صلاحية الشهادة يرجى المسح الضوئي للرمز ثنائي الأبعاد.

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2019-03-19