

FORMAT No: ARI/06/FOR/171/E

CERTIFICATE OF MANUFACTURER

Name of the Product and Strength/Potency : PREGABALINA CAPSULAS 75 MG 30 BLS RFF	
Lot/Batch Number: NK1337	Dosage form: Solid-Oral
Date of fabrication/manufacture: SEP 2017	Expiry date: AUG 2019
Importing country: CHILE	Package size: 30 BLS
SFG Batch No: NK1339	Marketing Authorization/ANDA/ NDC/ DIN/ARTG Ref. Number : F-22.319/15
Finished Product Specification No. : PRB-CA-PFP-CAP-75MG-APO-INT	
Manufacturing Formula Reference Number: MFNPREGBAC, MFNPREGCDC	
Master Manufacturing record Number: PRB-MFMX-625-00-I, PRB-MFCA-75-00-I	
Master Packaging record Number: PRB-PO-75CK-I	
Name and address of fabricator(s)/manufacture(s)-manufacturing site(s): APOTEX RESEARCH PVT. LTD. Plot No.1& 2, Bommasandra Industrial Area, 4 th Phase, Jigani Link Road, Bangalore- 560 099.	
Number of Manufacturing Authorization/License or Certificate of GMP Compliance of a manufacturer /fabricator: KTK/25/521/2006 and KTK/28/374/2008	
Results of analysis: • This batch product complies as per Ph. Eur / USP / In-house Specifications.	
Comments/remarks: Not Applicable.	
Certification Statement: <i>I hereby certify that the above information is authentic and accurate. This batch of product has been fabricated/manufactured, including packaging and quality control at the above mentioned site(s) in full compliance with the GMP requirements of the local Regulatory Authority and with the specifications in the Marketing Authorization of the importing country. The batch processing, packaging and analysis records were reviewed and found to be in compliance with GMP.</i>	
Name and Position/title of person authorizing the batch release: Sreesha G Senior Reviewer-I - Quality Assurance (Product Release & Logistics)	
Signature of person authorizing the batch release: 	
Date of signature: 25-08-17	