

Sun Pharmaceutical Industries Limited

Walk-in Drive at Baroda



Function	Qualification & Experience	Grade	Brief JD
Regulatory Affairs -US Job Location: Baroda	M.Pharm/ 2-7 years	Executive/ Sr Executive	1) Compilation , review of dossier for US , EU , AU, Canada & IL market (ORALS/NON ORALS) 2) Review of query response prepared and compiled by regulatory associates 3) Review of all the documents received from stake holder for dossier compilation , Query response and Life cycle management
Regulatory Affairs -US (Labeling) -Baroda	B.Pharm/M. Pharm / 2-6 years	Sr Officer / Executive	1) Preparation and reviewing of labeling for ANDA and/or NDA applications and labeling query responses in accordance with the USFDA regulations. 2) Preparation of Structured Product Labeling (SPL) for drug listing in accordance with the associated application USFDA regulations
Regulatory Affairs -Plant Job Location : Halol	M.Pharm / 3-9 years	Executive / Sr Executive	1. Participates in activities related to new product registration/ new filing, re-registration, sample, queries of regulatory agency and variation filing. 2. Product life cycle management: review the Change control and provide final variation category. And arrange documents for Filing. And Activities related to variation management. 3. Approval package (Emerging Market): Co-ordination with corporate regulatory and plant team for impact and proposed change against approved package and filing of variation whenever applicable

Date of Interview : Sunday, 13th July, 2025 **Venue of Interview : Sun Pharmaceutical Inds. Ltd.**
Time : 9.30am-1.00pm **Sun Pharma Road, Tandalja , Baroda -39001**

Candidates having above experience can walk-in with their updated resume and relevant documents. Those unable to attend interview may share their cv to : HR.Tandalja@sunpharma.com

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