Shubham Kumar

Pharmacovigilance Associate

My Contact

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Hard Skill

- Exceptional analytical research skills
- High patient data analysis
- MS-Office, MS-Excel, MS-PPT.
- Good knowledge of email writing etiquette

Soft Skill

- Observation
- Decision making
- Communication
- Multi-tasking

Education Background

- Bachelors in pharmacy
  B.Pharm
  Completed in 2018 (I DIV)
- Intermediate
  Passed in year 2013 from UP board
- HIgh School
  Passed in year 2011 from UP bpard

About Me

Dynamic Executive with 3.2 years of experience helping organizations reach their full potential. Organized and dependable candidate successful at managing multiple priorities with a positive attitude. Willingness to take on added responsibilities to meet team goals.

Professional Experience

Total Experience:5.5 Years
Relevant Experience: 3.2 Years

Wipro Ltd. | Pharmacovigilence associate 2021 – Present.

1. Pharmacovigilance Associate, Wipro Pvt. Ltd., New Delhi SEPTEMBER 2021 — PRESENT

•ICSR Case processing (Database Handled: Argus 8.2.1) of Vaccine and drug cases
•Case processing on regular basis for Solicited reports, Spontaneous report.
•Case registration (booking/ case intake), Duplicate case checking and complete case entries
•Data Entry of the adverse eventsreports (AE) form from sourcedocs including patient demographic data, suspect and concomitants, narrative, selection and detailing of adverse events, action taken, Dechallenge and Rechallenge, medical history and diagnostic details etc.)
•Adverse events coding by MedDRA browser and drug coding by WHODD browser and CDD, coding events, indications, and patient history based on MedDRA •Consistently used ARGUS features such as action item, contact log work list for communication and for the follow-up requests.
•Ensuring compliance (quality, procedures, regulations, timeliness, consistency) with local regulations and Company's global Pharmacovigilance requirements. •Processing Suspected Unexpected Serious Adverse Reaction (SUSAR) cases in clinical studies, strictly compliant to 7-Day and 15-Day timelines
•Receive, reviewand process SAE and/ Adverseevents reports from various sourceon time and quality standards.
•Triages ICSRs, evaluates ICSR data for completeness, accuracy & regulatory report ability.
Determine and perform appropriate case follow-up generating and requesting follow up information from HCP/Non-HCP reporter as per the applicable guidelines for required cases.

Hegde & Hegde Pharmaceutica LLP | Bussiness Officer.
From 2018-2021

- Expertise meetings for sale and promotion of dermatological range of medicines in Dehradun
- Communation with Doctors regarding the product promotion and development.
- Updated target competance.