BHARGAVI REDDY NANDYALA
nandyalabhargavi18@gmail.com
Ph: +91 7675 8989 55

CAREER OBJECTIVE:

To prove myself dedicated, worthy, full and energetic in a progressive organization that gives me scope to apply my creative thoughts and skills and be a member of a team that dynamically works towards organizational and personal worth.

JOB SUMMARY: Pharmacovigilance , Medical transcription and virtual medical scribe

- Pharmacovigilance Professional (Drug safety associate) with 2 years of hands-on experience in capturing and assessing single adverse event reports associated with client products on a worldwide basis on the clients drug safety database, in accordance with international and local regulatory requirements.
- Medical and drug terminology in-depth knowledge in using MedDRA and WHO-DD.
- Successfully faced multiple client audits.
- Thorough understanding of adverse event reporting and safety databases.
- Good understanding of the quality and confidentiality of patient data and process involved in processing the data to make it presentable as per regulatory requirements.
- Managed Individual Case Safety Reports of all case types (serious and non-serious cases from spontaneous and solicited programs, literature and E2B cases, which includes the Triage of confidential to determine whether the qualify for expedited reporting or not and the timelines within which they need to be submitted to the regulatory authorities.
- Excellent interpersonal communication, organizational and leadership skills.
- Good at handling modules independently and as a team.
- Excellent verbal and written communication skills.

Medical transcription

- ❖ Listened to voice recordings by physicians and other doctors, and converted them into written reports
- ❖ Provided assistance to review and edit previous medical documents using speech recognition technology
- ❖ Performed checking of the written documents by using medical reference sites and other educational material to ensure maximum accuracy in medical terminologies

- ❖ Transcribed and edited dictation by various medical personnel and healthcare providers to document patient care, and assisted in maintaining updated records.
- ❖ Prepare medical documentation of audio files regarding treatment of chronic ailments received from specialists on mail
- ❖ Attended medical seminars as directed by the specialists to record guest lectures on treatment of diseases under research, and prepared reports
- ❖ Completed daily operations associated with transcription workload ensuring a high-quality verbatim qualitative work
- ❖ Coordinated with the team to perform additional administrative duties such as answering phones, and scheduling patient appointments as required.

Virtual medical scribe(Aquity solutions)

November-2023 present

Maintaining 100% accuracy of medical records by entering patients information into EMR with typing speed 55 WPM with 97% accuracy with uptodate medical terminologies

SKILLS: Good Knowledge on Pharmacovigilance, Clinical Research, Quality Assurance, medical transcription and scribe,Argus safety database ver 8.0, Trained in SAS Clinical program, MS Office Suite.

- Self paced training on clinical sas base
- Certified A00-231 SAS 9.4 Base programming-performance based exam
- Training taken on medical transcription,drug safety associate and clinical research fields.

PROFESSIONAL EXPERIENCE

Aurobindo Pharma|Hyderabad
Pharmacovigilance Specialist(Drug safety associate-I)

Responsibilities:

- PV mailbox monitoring for case intake.
- Perform initial evaluation of reported adverse events (serious and non-serious) from all sources Maintenance of triage tracker as per SOP.
- Prioritization of business partner sharing cases Based on timelines. Handling and monitoring of EVDMS case and MLM cases.
- Downloading of Evdms and mlm cases from Eudravigilence website on daily basis. Monitoring of ICSR search on each working day on the Eudravigilence website.

- Screening of number of Evdms cases and mlm cases received from Eudravigilence on daily basis.
- Ensure preliminary identification of business partner sharing cases and book in of the cases and prioritization of cases based on timelines.
- Ensure e2b reports (e.g evdms cases, mlm cases and mhra) acceptance and rejection in argus database and performing duplicate check.
- Communicating with ema service desk for ICSR related queries and suspected duplicate cases received from Eudravigilence.
- Handling and monitoring of Canada cases from Canada vigilance adverse reaction online database.
- Ensure Canada cases download from Canada vigilance adverse reaction online database and screening of downloaded cases from Canada vigilance adverse reaction online database.
- Ensure book in the valid cases on regular basis to prioritization of cases based on regulatory timelines.
- Book in of case reports including spontaneous cases, literature reports, MHRA E2B reports and cases downloaded from EVDMS database and in safety database by checking validness of case.
- ICSR case processing
- Case intake, duplicate check and registration. Maintain log of source documents and other communication.
- Data entry of individual case safety reports (ICSRs) into database (including serious and non-serious cases from spontaneous and clinical trials origin, literature and E2B cases as applicable).
- Code all medical history, events, drugs/procedures/indications and laboratory tests according to the appropriate dictionary (e.g., MedDRA, company product dictionary, WHO-DD).
- Write medically relevant safety narrative of cases and checking the completeness and accuracy of the data entered in the various fields.
- Perform peer review of all book in cases and resolve the queries related to book in (IRD wrongly captured, wrong source document attachment) and evaluate seriousness criteria as per IME list and CTC.

ACADEMIC PROFILE:

- Pharm.D (Doctor of pharmacy) | 2014-2020 | Nirmala college of pharmacy, Vijayawada, A.P
- Intermediate (BiPC) | 2012-2014 | Sri Chaitanya College, Vijayawada, A.P

Training Undertaken:

- Completed 10 months Internship program at Manipal Super Specialty Hospital,Tadepalli,Vijayawada in various Departments along with Physician as "CLINICAL ASSISTANT ".

PROJECTS:

- Title: A PROSPECTIVE COMPARATIVE STUDY ON PROBIOTIC TO ENHANCE THE BIOAVAILABILITY OF IRON IN YOUNG FEMALE ADULTS

- Duration: 3 months
Role: Author
- Description:Our project was based upon studies that related to young females are more suffering with iron deficiency due to low iron diet and blood loss during menstrual periods. Iron absorption concept based upon LP299v in combination with iron and folic acidis clinically proven to increase iron uptake.

PUBLICATIONS:

- A Prospective comparative study on probiotic:To enhance bioavailability of iron in young female adults ,Nandyala. Bhargavi, J. Global Trends Pharm Sci, 2020; 11 (2):7881 - 7894
- Role of immunosuppressants in organ transplantation, EUROPEAN JOURNAL OF PHARMACEUTICAL AND MEDICAL RESEARCH.
- Congenital Myasthenia Syndrome: A Case Report, IJPPR,Human Journals Case Report March 2020 Vol.:17, Issue:4 by Nandyala Bhargavi
- Assesment and drug utilization pattern on anti platelets agents in cardiovascular patients.