RESUME

Mr. Vishal Sharma

Phone:+91-9953758831

E-mail :
vishal.sharma995375@gmail.com

Personal Data:

Father Name Krishan Sharma

Gender : Male

Nationality : Indian

DOB: : 11th Jan 1998

Marital Status : Unmarried

Language: English and Hindi.

Permanent address:
K-190/4 Saurabh Vihar Jaitpur Ext. Hari Nagar Badarpur New Delhi-44

Hobbies:

Meeting new person's,
Traveling ,MUSIC,
PIANO and GUITAR

Strengths:

- Hard working honesty
- Decision Making
- Leadership
- Manage with people very effectively.

Resume Headline:

Individuals by providing guidance and establishing myself as an distinct with strong leadership skills to put my efforts and skills together for the growth of an organization.

Professional Qualification:

- Advanced Diploma in Drug Regulatory Affairs from Jamia Humdard university Year of passing 2023
- Bachelor of pharmacy Sharda university, Greater Noida Year of passing July-2022 Taken 8.0 CGPA
- Diploma of pharmacy Sharda university, Greater Noida Year of passing July-2019 Taken 7.8 CGPA
- Higher secondary Examination Completed from Central Board of Secondary Education at August-2017
- Senior secondary Examination Completed from Central Board of Secondary Education at May-2015

Bachelor of pharmacy Project Work :

- Review on formulation and evaluation of liposomal drug delivery system.

Knowledge About:

- Microsoft Excel
- Power Points Presentation
- MS-Words

Key Skills:

- Knowledge about regulatory guidelines. (USFDA, MHRA, EMA and DCGI etc.)
- Knowledge about data review and report preparations

Work Experience

- Presently associated with - Raptim Research Pvt. Ltd-Mumbai as an officer in Technical documentation Department.
  Period: 14th Nov 2023 to till date
- Have done 1-month Industrial training at Synokem pharmaceuticals limited haridwar from 20 June 19 July -2021.
- Have done 3-month holy family hospital at New Delhi From 27 June05 November -2019.

Key Responsibilities

- Responsible for preparation and review of DCGI application for BENOC, Test License and Working Standard import license application
- Responsible for query reply to all DCGI related matters like BENOC, Test License and Working Standard
- Responsible for DCGI requirement (feasibility of molecules)
- Record keeping of all the submission and approval
- Responsible for preparation and review of CBN application
- Responsible for NDPS-2, and NDPS-6 License from state FDA
- Responsible for dossier submission (Module 5) in e-CTD and Ne-CTD format to Regulatory Agencies of US, Europe, WHO, TGA, MCC, MHRA, Canada & ASEAN countries as per the latest regulatory guidelines of the respective countries.
- Preparation and review of module 2.7 for USFDA, Module 2.7.1 for EU and CSBE summary for Canada
- Preparation and review of DBE table for USFDA
- Preparation and review of module-5 and module 2.7.
- Preparation and review of subject information sheet and informed consent form.
- Preparation and review of clinical study report.
- Preparation and review of case report form.
- Preparation and review of CDISC datasets as per regulatory requirements.

Area Of Interest :

- Business development
- Regulatory Affairs
- Research and Development

Extra Curriculum Activity:

- Diploma in Music
- Singing course from T-Series
- Stage work academy
- Awarded E-certificates for many competition

Declaration

- I hereby declare that the above written particulars are true to the best of my knowledge and belief.

DATE……

PLACE……

Yours sincerely

(Vishal Sharma)