

## CURRICULUM VITAE

**Shreya Parashar**

**6 part Bareja**

**Near water tank**

**Email:shreyaparashar2609@gmail.com**

**Mob no: 9824013642**

### PERSONAL DETAILS :

<b>Date of birth</b>	26 Sep1996
<b>Marital Status</b>	UnMarried
<b>Languages Known</b>	English, Gujarati, Hindi
<b>Hobbies</b>	Music,Dancing,singing
<b>Nationality</b>	Indian

### EDUCATIONAL QUALIFICATION :

<b>Examination</b>	<b>University/Institute/Board</b>	<b>Year of passing</b>	<b>Grade</b>
M. Sc (Clinical Research)	GUJARAT UNIVERSITY	2019	1 <sup>st</sup> Class
B.Sc	GUJARAT UNIVERSITY	2017	1st class
H. Sc	G.H.S.B	2014	68%

### AREA OF INTEREST :

- Quality Assurance /Quality Control
- CRA
- Crc
- Clinical Data Management
- Pv

### **STRENGTH :**

- Ability to coordinate multiple clinical research projects
- Work efficiently with diverse group of people
- Excellent assessment, problem solving, planning and evaluation skills
- Proficiency in computer software programs(e.g. Word, Excel, Power point)
- Team work

### **WORKING EXPERIENCE :**



#### **Job Responsibilities:**

- To work according applicable guidelines.
- Request controlled document from QA.
- Screening related activities.
- Site preparation for study and ensuring the enrollment of eligible volunteer.
- Communication with Hospital and Cro study team
- Responsible for the check-in and check-out process of study
- Make study related documentation
- Reporting of AEs and SAEs.
- Reviewing of Protocols, CRF, ICF and SOPs.
- Preparation and Compilation of Pre-Study, During Study and Post-Study Trial Master File (TMF) as per ICH GCP.
- Counseling and taking informed consent of the subject for screening and study.
- Drug administration to the study volunteer.
- Dosing supervision of the project.
- Ensuring compliance with the Protocol, SOPs, ICH-GCP and other applicable regulatory requirements..
- Working as a clinical research coordinator at Jivrajmehta hospital, Ahmedabad, August 2018
- Now working as a clinical research coordinator at Rudraksha Multispeciality

**hospital Bareja.**

**Job Responsibilities:**

- Handling of Investigational product at clinical site
- Documentation and eCrf (BizNet)
- Communications with CRO and PI
- Ensure that all the study documents are accurate, complete, timely and legible and data entered on to the CRFs in consistent with source data and submitted to QC/QA at stipulated time line
- Ec coordinator at Rudraksha Multispeciality hospital
- Routine and non-routine Maintenance of equipment's at clinical site
- To work according to the application guidelines and applicable Sop

**PROJECTS :**

- Works on government programs (GEMI) the topic of **Biomedical waste** on clinic or hospitals and pathology lab for Ahmadabad as well Baroda

**TRAINING :**

- I have (Nida) "Good clinical practice" training certificate

**SKILLS :**

- Good Communication skill
- Good Representation skill
- Adaptable type of person
- Always eager to learn new things

**DECLARATION :**

Thereby state that the Information compiled above is precise and accurate in its entirety to my knowledge.

**DATE :**

**PLACE :**

***Yours sincerely  
Shreya Parashar***