GITAM INSTITUTE OF PHARMACY
VISAKHAPATNAM

Application for Certificate Program in Drug Regulatory Affairs (2019-20)

Last date to apply: 26-08-2019

1. NAME OF THE APPLICANT: (As per class X or equivalent):

____________________________________________________________________________________

2. DATE OF BIRTH: ___________________________________________________________________

3. GENDER: _________________________________________________________________________

4. NATIONALITY: ____________________________________________________________________

5. RELIGION: ______________________________________________________________________

6. CATEGORY: GENERAL OBC SC ST PH

7. MOBILE: _________________________________________________________________________

8. NAME OF THE PARENT / GUARDIAN: _______________________________________________

9. ADDRESS FOR CORRESPONDENCE:

_________________________________________________________________________________
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_________________________________________________________________________________

10. QUALIFICATION: (B.Pharm, M.Pharm, M.Sc.)

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<th>Year of Pass</th>
<th>Percentage/CGPA</th>
<th>Subject</th>
<th>University/Board</th>
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11. NAME OF THE COLLEGE LAST STUDIED/ STUDYING:

_________________________________________________________________________________

Declaration:

I hereby declare that all the particulars stated in this application are true to the best of my knowledge and belief. I shall obey the rules and regulations of GITAM (Deemed to be University).

Paste your recent passport size colour photograph

Signature of the Parent/Guardian  Signature of the Applicant
CERTIFICATE COURSE IN DRUG REGULATORY AFFAIRS

GITAM Institute of Pharmacy, GITAM Deemed to be University is offering a certificate course in Drug Regulatory Affairs. The duration of the course is 3 months. This is a value added certificate course for the students of B.Pharm, M.Pharm, M.Sc. and those professionals who are working in Pharmaceutical Industry. The course is designed to contain global regulatory requirements and practices of drugs, pharmaceuticals and medical devices. After completion of the course, the participants are expected to have in-depth knowledge and understanding of concept of global regulatory agencies and regulations on drugs and pharmaceuticals, Drug discovery and development process, Innovator and generic drug regulatory filing and approval process. The participants will be trained in GMP, GLP, GCP and quality management systems.

**Duration of Course:** 3 months

**Eligibility for admission:** A pass in B.Pharm, M.Pharm, M.Sc. and B.Pharm final year students

**Last date of Application:** 26-08-2019

**Commencement of classes:** 04-09-2019

**Timings:** 4.00 pm to 6.00 pm

**Fee:** Rs.15,000/- (Rupees fifteen thousand)

(S.Ganapaty)
Principal-GIP