

Estd By: Nimra Educational Society (A Muslim Minority Society)
Approved by Pharmacy Council of India (PCI), New Delhi.
Affiliated to JNTU, Kakinada, Permitted by Govt. of A.P.
E-mail: principalncp@yahoo.in Website: www.nimra.in

List of Seminars/Workshops

A.Y.2019-20

S.No	Name of the Programme	Date	No. of Participants	Details of the Resource Person
1	Adverse Drug Reactions	10-06-2019	25	Mr. P. Raja rao M.Pharm Dept. Of Pharmaceutical analysis Phone no : 7490037717 Mail Id: @gmail.com
2	2D & 3D Printing In Drug Delivery	24-06-2019	25	Mr. B. Siva Sai Kiran M.Pharm Dept. Of Pharmaceutical analysis Phone no: 9700572717 Mail Id:sivasaikiran143 @gmail.com
3	Drug Research & Development	26-08-2019	25	Mr. Sk. Abdul Saleem M.Pharm Dept. Of Pharmacology Phone no: 9959545424 Mail Id:saleem.pci @gmail.com
4	Clinical Research & Development	04-10-2019	25	Mrs. K. Yamini M.Pharm Dept. Of Pharmacognosy Phone no : 9703649067 Mail Id:yaminik.yk@gmail.com



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NIMRA COLLEGE OF PHARMACY
Nimra Nagar, JUPUDI,
Ibrahimpatnam, VIJAYAWADA-521 456

drug reactions (ADRs) Adverse effects unintended medications that can occur at therapeutic doses. They can range from mild to severe and can occur immediately or long after the medication's use. ADRs can be caused by various factors, such as drug interactions, dosage errors, or an individual's genetic makeup. Common types of ADRs include allergic reactions, drug-induced liver injury, and cardiovascular events. ADRs are a significant public health concern and can death. The detection, monitoring, and reporting of ADRs are critical to ensure the safety of medications and protect the public's health. Healthcare providers and regulatory agencies use various systems and tools, such as the MedWatch program and the Adverse Event Reporting System, to report and monitor ADRs.

Contact Us For More Info

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Organizing Committee

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Student Coordinators:

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SEMINAR ON ADVERSE DRUG REACTIONS

Date- 10.06.2019

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Courses Offered By Nimra College of Pharmacy

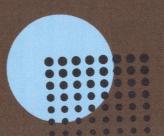
Bachelor of Pharmacy (B.Pharm)

Master of Pharmacy (M.Pharm)

Pharmaceutical Analysis

Pharmaceutics

Pharm.D



- →Principles of Toxicity
- ◆Treatment of poisoning in general
- ◆Describe idiosyncratic reactions
- ◆Pharmacovigilence
- ◆Monitoring researching
- ◆Evaluating information from health







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PROGRAM REPORT

Date: 10.06.2019

Program Name

; A Seminar on Adverse drug reactions.

Date of Activity

;- 10.06.2019

Venue

:-College Seminar Hall

Organized by

: - B. Pharmacy

Number of Participant: - 25

Objective of the program

Explain pharmacovigilence and various type of adverse drug reaction. List some common adverse effects and toxicities with examples. Discuss drug dependence and its public health importance.

Topics covered:

- Principles of Toxicity
- Treatment of poisoning in general
- Describe idiosyncratic reactions
- Pharmacovigilence
- Monitoring researching
- Evaluating information from health

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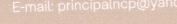
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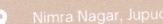
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Asst.Professor

Student Coordinators

Shaik Ayesha Shaik Yasin Shaik Rasool



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SEMINAR ON
2D & 3D PRINTING IN DRUG
DELIVERY

Date- 24.06.2019

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Courses Offered By Nimra College of Pharmacy

Bachelor of Pharmacy (B.Pharm)

Master of Pharmacy (M.Pharm)

Pharmaceutical Analysis

Pharmaceutics

Pharm.D



- ◆Pharmaceutical practices
- ◆Improve the patient care
- ◆Personalized medicine
- →Printing technologies
- ◆Bone reconstruction
- ◆Dentistry
- **♦ Personalised Medicine**
- ◆Pharmacy Application
- +Pharmacotechny
- ◆Pharmacotechnie
- +Impression 2D-3D
- **◆Bone Reconstruction**
- Dentistry





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PROGRAM REPORT

Date:24/06/2019.

Program Name

: A Seminar on 2D &3D Printing in drug delivery

Date of activity

: 24/06/2019

Venue

: College Seminar Hall

Organized by

: B Pharmacy

Number of Participants: 25

Objective of the program

2D and 3D Printing are used in the fields of health such as dentistry or for bone reconstructions. The goal of this work is to highlight relevant printing technologies for implementation in hospital pharmacies.

Topics covered

- Pharmaceutical practices
- Improve the patient care
- Personalized medicine
- Printing technologies
- Bone reconstruction
- Dentistry

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New drugs are continually required by the healthcare systems to address unmet medical needs across diverse therapeutic areas, and pharmaceutical industries primarily strive to deliver new drugs to the market through the complex activities of drug discovery and development. Discovery involves a number of processes like target identification and validation, hit identification, lead generation and optimization and finally the identification of a development. further candidate Development, on the other hand, includes optimization of chemical synthesis and its formulation, toxicological studies in animals, clinical trials, and eventually regulatory approval. These issues have had an adverse bearing on the R&D productivity in recent years, hence there is a need for innovative approaches as well as increased collaboration between industry, research governmental academia. institutions, with a common objective of constantly delivering quality medicines.

Contact Us For More Info

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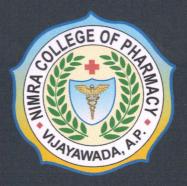
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SEMINAR ON DRUG RESEARCH & DEVELOPMENT

Date- 26.08.2019

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Bachelor of Pharmacy (B.Pharm)

Master of Pharmacy (M.Pharm)

Pharmaceutics

Pharmaceutics

Pharm.D

- **♦Introduction**
- ◆Definition of new Drug
- ◆Regulatory guideline
- ◆Steps in theDrug
- ◆Development
- **→**Timeline of Patents
- ♦ New drug discovery
- ◆Target Identification
- ◆Target validation
- **→**Drug Discovery
- ◆Drug Development
- **→**Lead Optimisation





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Program Report

DATE- 26.08.2019.

Program Name

:- A Seminar on Drug Research and Development

Date of Activity

:- 26-08-2019

Venue

:-College Seminar Hall

Organized by

:- B. Pharmacy

Number of Participant :- 25

Objective of the program

The goal of a preclinical drug discovery program is to deliver one or more clinical candidate molecules, each of which has sufficient evidence of biologic activity at a target relevant to a disease as well as sufficient safety and drug-like properties so that it can be entered into human testing

Topics Covered

- Introduction
- Definition of new Drug
- Regulatory guideline
- Steps in the Drug Development
- Timeline of Patents
- New drug discovery
- Target Identification
- Target validation

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Clinical research and development (R&D) is a critical aspect of bringing new drugs, medical devices, and therapies to market. It involves the through clinical trials. The clinical research and development process typically involves four conducted on a small group of healthy of the treatment. Phase II trials are conducted on a larger group of patients to evaluate the treatment's effectiveness and side effects. Phase III trials involve a larger group of patients to confirm the treatment's effectiveness and monitor side effects over an extended period. studies conducted after regulatory approval to monitor long-term safety and effectiveness. expensive, but it is critical to ensure that new treatments are safe and effective for patients.

Contact Us For More Info



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Nimra Nagar, Jupudi

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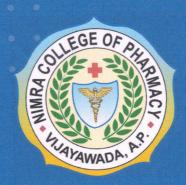
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Ms.Abdul Parveen Sultana

Asst.Professor

Student Coordinators

Shaik Rasool Shaik Ayesha Nannaka Nikitha



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SEMINAR ON
CLINICAL RESEARCH
& DEVELOPMENT

Date- 04.10.2019

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Pharmaceutical Analysis

Pharmaceutics

Pharm.D

- **♦Introduction**
- **→**Material and method
- **◆**Ethical foundation of clinical trails
- ◆Overviews of trail design
- ◆Overview of drug designs
- ◆Preclinical, phase 1,phase 2
- **♦Phase III trails**
- **♦Phase IV trails**
- +Conclusion





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PROGRAM REPORT

Date:04/10/2019

Program Name

:A Seminar on Clinical Research and Development

Date of activity

: 04/10/2019

Venue

: College Seminar Hall

Organized by

: B Pharmacy

Number of Participants: 25

Objective of the program

The clinician investigator integrates all the Can Med competencies including the roles of a scholar, communicator, health advocate, manager, collaborator and professional in their roles as a medical expert to conduct ethical research, establish and maintain skills and attitudes appropriate to their research practices, using ethical and effective research methodologies to address questions.

Topic Covered

- Introduction
- Material and method
- Ethical foundation of clinical trails
- Overviews of trail design
- Overview of drug designs
- Preclinical, phase 1 , phase 2 trails
- Phase III trails
- Phase IV trails
- Conclusion

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