College Code: 1706

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CIRCULAR

Date: 02/07/2019

This is to inform all the faculty members and students of B. Pharmacy, Pharm.D and M.Pharmacy that a SEMINAR ON RESEARCH ETHICS AND MORALS will be held on 05/07/2019 in our campus. All are requested to attend the lecture.

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Seminar on

RESEARCH, ETHICS AND MORAL

By Dr. Sudha Parimala

Date and Time
05 july.2019 11.am At
RBVRR Women's college
Seminar Hall

COORDINATOR
Dr. A.Krishana Shailaja

Patron
Prof. K. Muthyam Reddy
Hon. Secretary cum correspondent
RBVRR Women's College of pharmacy

CONVENER
Prof. M. Sumakanth
Principal
RBVRR Women's College of pharmacy

M. Sumakanth

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REPORT

Title: Session Report on Research Ethics and Morals by Dr. Sudha Parimala at RBVRR Women's College of Pharmacy on July 5, 2019

Date: July 5, 2019

Venue: RBVRR Women's College of Pharmacy

Speaker: Dr. Sudha Parimala

Introduction:

On July 5, 2019, Dr. Sudha Parimala conducted a session on Research Ethics and Morals at RBVRR Women's College of Pharmacy. The session aimed to educate students and faculty members about the importance of ethical conduct in research and the moral responsibilities associated with it.

Speaker's Profile:

Dr. Sudha Parimala is a renowned expert in the field of research ethics with extensive experience in academia and research. She holds a Ph.D. in Pharmacy and has published numerous papers on ethical considerations in research.

Overview of the Session:

The session commenced with Dr. Sudha Parimala providing an overview of the importance of research ethics and morals in the field of pharmacy. She emphasized the ethical principles that guide research conduct and highlighted the significance of upholding integrity and professionalism in all research endeavors.

Content Covered:

1. Ethical Principles in Research:
    - Dr. Sudha Parimala elucidated on key ethical principles including

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respect for persons, beneficence, and justice. She emphasized the need for researchers to prioritize the well-being and autonomy of research participants.

2. Informed Consent:
    - The speaker discussed the ethical requirement of obtaining informed consent from research participants and elaborated on the elements of valid consent. She stressed the importance of ensuring participants' understanding and voluntary participation in research studies.
3. Institutional Review Board (IRB) Approval:
    - Dr. Sudha Parimala provided insights into the role of Institutional Review Boards in evaluating research proposals to ensure compliance with ethical standards. She explained the process of obtaining IRB approval and the ethical considerations involved.
4. Data Management and Integrity:
    - The session addressed the importance of maintaining data integrity and confidentiality in research. Dr. Sudha Parimala discussed ethical guidelines for data collection, storage, and analysis, emphasizing the need for transparency and accuracy.
5. Authorship and Publication Ethics:
    - Dr. Sudha Parimala highlighted ethical issues related to authorship and publication, including criteria for authorship attribution and responsible publication practices. She emphasized the importance of avoiding plagiarism and adhering to ethical standards in disseminating research findings.

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Conclusion and Reflection:

The session concluded with a recap of the key points discussed and a call for reflection on the ethical responsibilities of researchers. Dr. Sudha Parimala encouraged participants to uphold ethical standards in their research endeavors and to prioritize integrity and professionalism.

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CIRCULAR

Date: 14/08/2019

This is to inform all the faculty members and students of B. Pharmacy, Pharm.D and M.Pharmacy that a SEMINAR ON RESEARCH PUBLICATIONS will be held on 16/08/2019 in our campus. All are requested to attend the lecture.

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Seminar on
RESEARCH PUBLICATIONS
SPEAKER- Dr. SHYAM SUNDER
(PRINCIPAL OUCT)

Date and Time
16 August 2019 2.00 pm
At RBVRR Women's college
Seminar Hall

COORDINATOR
Dr. M.VIJAY BHARGAVI

CONVENER
Prof. M. Sumakanth
Principal
RBVRR Women's College of pharmacy

Patron
Prof. K. Muthyam Reddy
Hon. Secretary cum correspondent
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M. Sumakanth

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REPORT

Title: Session Report on Research Publication by Dr. Shyam Sunder, Principal OUCT at RBVRR Women's College of Pharmacy on August 16, 2019

Date: August 16, 2019

Venue: RBVRR Women's College of Pharmacy

Speaker: Dr. Shyam Sunder, Principal OUCT

Introduction:

On August 16, 2019, Dr. Shyam Sunder, Principal of the Osmania University College of Technology (OUCT), conducted a session on Research Publication at RBVRR Women's College of Pharmacy. The session aimed to provide students and faculty members with insights into the process of publishing research in academic journals and enhancing their understanding of publication ethics.

Speaker's Profile:

Dr. Shyam Sunder is an esteemed academician with extensive experience in research and publication. As the Principal of OUCT, he has been actively involved in promoting research culture and scholarly activities among students and faculty members.

Overview of the Session:

The session commenced with an overview of the importance of research publication in academia and the significance of disseminating research findings to the scientific community. Dr. Shyam Sunder emphasized the role of publication in advancing knowledge and contributing to the academic discourse.

Content Covered:

1. Understanding the Publication Process:

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- Dr. Shyam Sunder provided an overview of the publication process, including manuscript preparation, submission, peer review, and publication. He highlighted the key stages involved in getting research published in reputable journals.

2. Selecting a Suitable Journal:
    - The speaker discussed strategies for selecting an appropriate journal for manuscript submission, considering factors such as scope, audience, impact factor, and publication policies. He emphasized the importance of targeting journals that align with the research topic and objectives.
3. Manuscript Preparation and Formatting:
    - Dr. Shyam Sunder shared insights into effective manuscript preparation and formatting, including guidelines for writing clear and concise research papers. He discussed the structure of a typical research manuscript and provided tips for improving writing quality and readability.
4. Ethical Considerations in Publication:
    - The session addressed ethical issues in research publication, such as authorship attribution, plagiarism, and conflict of interest. Dr. Shyam Sunder emphasized the importance of adhering to ethical standards and maintaining integrity in all aspects of the publication process.
5. Responding to Peer Review:
    - Dr. Shyam Sunder provided guidance on responding to peer review comments and revising manuscripts based on reviewer feedback. He discussed strategies for effectively addressing reviewer comments and

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improving the quality of research papers.

Conclusion and Reflection:

The session concluded with a reflection on the key takeaways and actionable steps for aspiring researchers and authors. Dr. Shyam Sunder encouraged participants to actively engage in the publication process and to seek mentorship and support from experienced researchers.

Overall, the session on Research Publication by Dr. Shyam Sunder at RBVRR Women's College of Pharmacy on August 16, 2019, provided valuable guidance and inspiration for students and faculty members aspiring to publish their research findings in academic journals.

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CIRCULAR

Date: 05/03/2020

This is to inform all the faculty members and students of B. Pharmacy, Pharm.D and M.Pharmacy that a SEMINAR ON CPCSEA GUIDELINES will be held on 06/03/2020 in our campus. All are requested to attend the lecture.

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Seminar on
CPCSEA GUIDELINES
Speaker- Dr. J . Archana

Date and Time
06/03/2020 at 2.00 pm
RBVRR WPC
SEMINAR HALL

COORDINATOR
Dr.M.P.KUSUMA

Patron
Prof. K. Muthyam Reddy
Hon. Secretary cum correspondent
RBVRR Women's College of pharmacy

CONVENER
Prof. M. Sumakanth
Principal
RBVRR Women's College of pharmacy

M. Sumakanth

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REPORT

Title: CPCSEA Guidelines by Dr. J Archana at RBVRR Women's College of Pharmacy on March 6, 2020

Date: March 6, 2020

Venue: RBVRR Women's College of Pharmacy

Speaker: Dr. J Archana

Introduction:

On March 6, 2020, Dr. J Archana conducted a session on the Clinical Pharmacology and Clinical Efficacy Analysis (CPCSEA) Guidelines at RBVRR Women's College of Pharmacy. The session aimed to familiarize students and faculty members with the guidelines and their implications for clinical research and practice.

Speaker's Profile:

Dr. J Archana is a respected expert in the field of clinical pharmacology with extensive experience in academia and research. She holds a Ph.D. in Pharmacology and has contributed significantly to the development and implementation of CPCSEA guidelines.

Overview of the Session:

The session commenced with an introduction to the CPCSEA guidelines and their significance in ensuring the quality and integrity of clinical research. Dr. J Archana provided an overview of the key principles and requirements outlined in the guidelines.

Content Covered:

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1. Introduction to CPCSEA Guidelines:
    - Dr. J Archana provided an overview of the CPCSEA guidelines, including their scope, objectives, and relevance to clinical research. She emphasized the importance of adhering to these guidelines to ensure the safety and efficacy of pharmaceutical products.
2. Principles of Clinical Pharmacology:
    - The speaker discussed the fundamental principles of clinical pharmacology, including pharmacokinetics, pharmacodynamics, and drug interactions. She explained how these principles are applied in clinical research to assess the safety and efficacy of drugs.
3. Study Design and Conduct:
    - Dr. J Archana elaborated on the requirements for study design and conduct outlined in the CPCSEA guidelines. She discussed key considerations such as patient selection, dosing regimens, and study endpoints, highlighting the importance of robust study protocols.
4. Data Analysis and Interpretation:
    - The session addressed the principles of data analysis and interpretation in clinical research. Dr. J Archana discussed statistical methods commonly used in clinical pharmacology studies and emphasized the importance of rigor and transparency in data analysis.
5. Regulatory Compliance:
    - Dr. J Archana provided insights into regulatory compliance requirements for clinical research, including ethical considerations, informed consent, and reporting obligations. She discussed the role of

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regulatory authorities in ensuring compliance with CPCSEA guidelines.

Conclusion and Reflection:

The session concluded with a reflection on the key takeaways and implications of the CPCSEA guidelines for students and faculty members. Dr. J Archana encouraged participants to integrate these guidelines into their research practice and to prioritize ethical conduct and scientific rigor in all aspects of clinical research.

Overall, the session on CPCSEA Guidelines by Dr. J Archana at RBVRR Women's College of Pharmacy on March 6, 2020, provided valuable insights and guidance for students and faculty members engaged in clinical research and pharmaceutical development.

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CIRCULAR

Date: 04/04/2020

This is to inform all the faculty members and students of B. Pharmacy and Pharm.D that our college will be organizing a webinar on WEBINAR ON STUDY DESIGNS IN CLINICAL RESEARCH. ON **06/04/2020**. All are requested to attend the webinar.

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Seminar on
STUDY DESIGN ON CLINICAL RESEARCH
Speaker- Dr. Neerja

Date and Time
06/04/2020 at 2.00 pm
RBVRR WPC
SEMINAR HALL

COORDINATOR
Dr.K.V.Ratnamala

Patron
Prof. K. Muthyam Reddy
Hon. Secretary cum correspondent
RBVRR Women's College of pharmacy

CONVENER
Prof. M. Sumakanth
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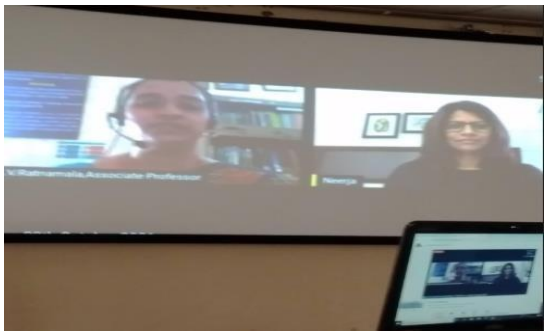

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REPORT

Title: Webinar Report on Study Design in Clinical Research by Dr. Neerja at RBVRR Women's College of Pharmacy on April 6, 2020

Date: April 6, 2020

Venue: Online (Webinar)

Speaker: Dr. Neerja

Introduction:

On April 6, 2020, RBVRR Women's College of Pharmacy hosted a webinar on Study Design in Clinical Research conducted by Dr. Neerja. The webinar aimed to educate students and faculty members about the fundamentals of study design in clinical research and its importance in generating reliable and valid scientific evidence.

Speaker's Profile:

Dr. Neerja is an accomplished researcher and academician with expertise in clinical research methodology. She holds a Ph.D. in Pharmacy and has extensive experience in designing and conducting clinical studies across various therapeutic areas.

Overview of the Webinar:

The webinar commenced with an introduction to the importance of study design in clinical research and its role in shaping the quality and validity of research findings. Dr. Neerja provided an overview of the key components of study design and their implications for research outcomes.

Content Covered:

1. Introduction to Study Design:
    - Dr. Neerja discussed the significance of study design in clinical

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research, emphasizing its role in defining research objectives, selecting appropriate methods, and ensuring the validity and reliability of study results.

2. Types of Study Designs:
    - The speaker provided an overview of common study designs used in clinical research, including observational studies (cross-sectional, case-control, cohort) and experimental studies (randomized controlled trials). She explained the characteristics and applications of each study design.
3. Key Considerations in Study Design:
    - Dr. Neerja highlighted key considerations in study design, such as research objectives, study population, sampling methods, data collection instruments, and outcome measures. She emphasized the importance of aligning study design with research objectives and addressing potential biases.
4. Ethical Considerations:
    - The webinar addressed ethical considerations in study design, including the protection of human subjects, informed consent, and adherence to ethical guidelines and regulations. Dr. Neerja emphasized the importance of ethical conduct in all phases of clinical research.
5. Practical Examples:
    - Dr. Neerja provided practical examples of study design in clinical research, illustrating how different study designs are applied to address specific research questions. She discussed real-world examples and

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their implications for research practice.

Conclusion and Reflection:

The webinar concluded with a reflection on the key takeaways and actionable insights for participants. Dr. Neerja encouraged attendees to apply the principles of study design in their research endeavors and to seek further training and mentorship in clinical research methodology.

Overall, the webinar on Study Design in Clinical Research by Dr. Neerja at RBVRR Women's College of Pharmacy on April 6, 2020, provided valuable insights and practical guidance for students and faculty members involved in clinical research.

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