

Activity Report (DAY/WORKSHOP/FDP/STTP/CONFERENCE)

DEPARTMENT	HOMOEOPATHIC PHARMACY		
ACTIVITY TYPE	EDUCATION FIELD VISIT		
ACTIVITY TITLE	DR.WILLMAR SCHWABE MANUFACTURING UNIT		
DATE & TIME	11/09/25; 9:00 – 12:30PM	Duration	3 HOURS
NO. OF PARTICIPANTS	97		
EXPERT NAME WITH DESIGNATION	NA		
NAME OF EXPERT’S ORGANIZATION	PARUL INSTITUTE OF HOMOEOPATHY AND RESEARCH		
EXPERT CONTACT DETAILS	NA		
FACULTY COORDINATOR	DR.NAYANA PATEL & DR.SURAJ BHADORIYA		
FACULTY CONTACT DETAILS	8128815189		
SUSTAINABLE DEVELOPMENT GOALS (SDGs)	SDG – 3; GOOD HEALTH & WELL – BEING, SDG 4 – QUALITY EDUCATION, SDG 9 – INDUSTRY, INNOVATION, AND INFRASTRUCTURE		
COLLABORATIVE ACTIVITY UNDER MOU			
SPONSORING AUTHORITY	-	Sponsorship amount:	-

Objective:

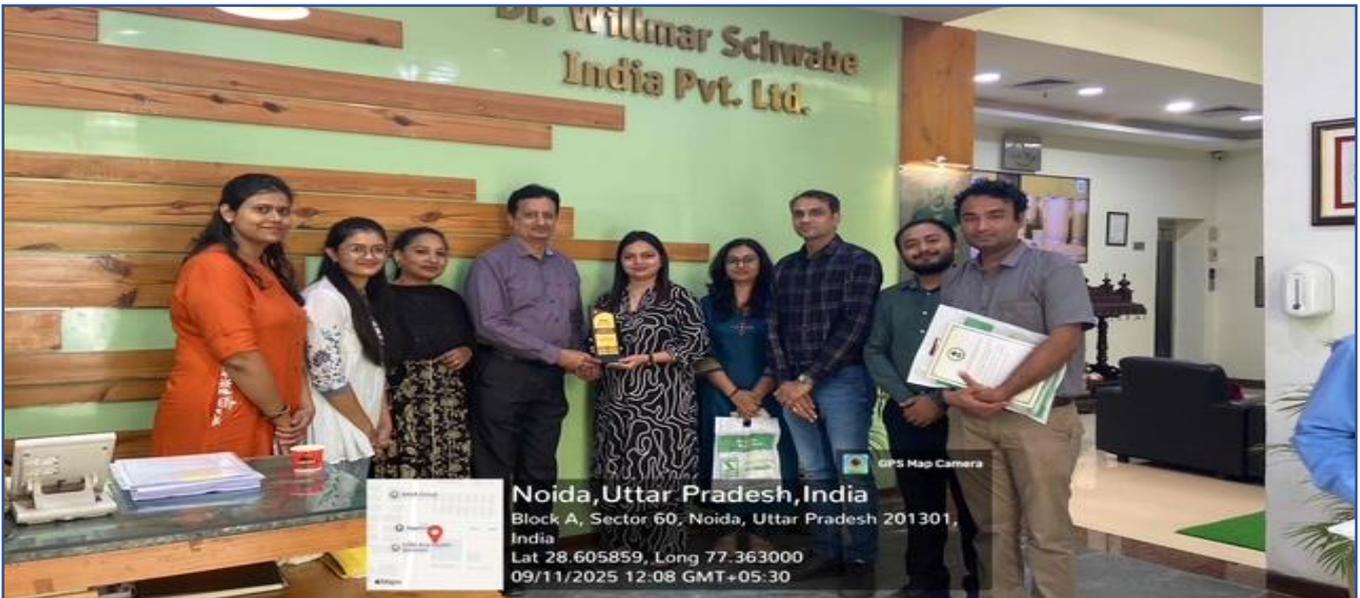
- To study the implementation of international quality standards in homoeopathic drug production.
- To learn about hygiene, safety, and regulatory compliance in pharmaceutical manufacturing.
- To observe the methods of standardization, testing, and validation of raw materials and final products.
- To understand the role of modern analytical techniques in ensuring product safety and efficacy
- To witness the use of modern equipment and automation in homoeopathic manufacturing.
- To understand the integration of traditional principles with modern technology.
- To study the documentation system for maintaining accuracy and traceability
- To understand the compliance with national and international regulatory framework
- To bridge the gap between theoretical knowledge and industrial practices.
- To develop practical insights that will be useful in professional and academic growth.

Activity Details:

- Raw Material Procurement & Storage – Understanding sourcing, identification, and storage of homoeopathic raw materials.
- Mother Tincture Preparation – Observing maceration/percolation processes and quality checks.
- Potentization Unit – Learning about serial dilution and dynamization using modern machines.
- Manufacturing of Dosage Forms – Preparation of tablets, dilutions, drops, and ointments.
- Quality Control Laboratory – Testing for purity, identity, and consistency using advanced analytical techniques.
- Packaging & Labeling Section – Observing hygienic packaging, labeling regulations, and batch numbering.
- Documentation & Record Keeping – Learning about standard operating procedures (SOPs) and regulatory compliance.
- Research & Development Division – Understanding innovation and formulation development.

Glimpses of activity:





Outcome: The visit provided students with first-hand exposure to the functioning of a hospital/pharmaceutical laboratory. Students gained practical understanding of drug dispensing, storage, compounding, and quality control procedures. The experience helped in bridging the gap between theoretical knowledge and real-world application of pharmacy practices. Students observed Good Manufacturing Practices (GMP), Good Dispensing Practices (GDP), and safety measures essential for healthcare services. Awareness about regulatory requirements, documentation, and ethical responsibilities in pharmacy was enhanced. The interaction with pharmacists and technical staff broadened the perspective on career opportunities and research avenues in the pharmaceutical sector. The visit highlighted the importance of sustainability, waste management, and responsible production, linking directly with SDG 3 (Health), SDG 4 (Education), SDG 9 (Industry), and SDG 12 (Sustainability). Overall, the visit contributed to the professional growth, confidence, and motivation of students to pursue excellence in pharmacy practice.

Name, Designation & Signature of
Coordinator with date

Name, Signature & stamp of
Head of Department / Institute