



IIT MADRAS

Indian Institute of Technology Madras

Technology Transfer Office
TTO - IPM Cell



Industrial Consultancy & Sponsored Research (IC&SR)

Drug Loading In Nanomaterials Using Microgravity

IITM Technology Available for Licensing

PROBLEM STATEMENT

- **Nanotechnology** has made significant advancements in nanomedicine, enhancing drug bioavailability, pharmacokinetics, and targeted delivery.
- **Nanomedicine** is used for diagnosis, monitoring, control, prevention, and treatment of diseases, and regenerative medicine.
- **Nanomaterials like liposomes** can enhance drug solubility and focus on specific disease sites. However, current nanomedicines have limited drug loading efficiency, typically below 10%.
- Research has attempted to improve **drug loading efficiency** in nanomaterials, but this has been challenging, expensive, time-consuming, and requires specialized expertise.
- A universal framework is needed to optimize drug loading efficiency across all types of nanomaterials.

TECHNOLOGY CATEGORY/MARKET

Technology: Drug Loading In Nanomaterials Using Microgravity

Category: Medical & Surgical Devices, Micro & Nano Technologies

Industry: Medical & Surgical, Space Technology

Application: Nano formulation in medical field

Market: The global market size of microgravity research was estimated to be **USD 3.5 billion** in 2023 and is projected to reach approximately **USD 8.4 billion by 2032**, growing at a compound annual growth rate (CAGR) of 10.2% during the forecast period

INTELLECTUAL PROPERTY

IITM IDF Ref. - 2819

Patent No: IN – 562410 (Granted)

TRL (Technology Readiness Level)

TRL 3, Experimental proof of concept

Research Lab

Prof. Dr. Swathi Sudhakar,
Department of Applied Mechanics &
Biomedical Engineering, IIT Madras.

TECHNOLOGY

Microgravity-based Drug Loading Method

- Utilizes microgravity conditions, typically achieved using equipment like RPM, Clinostat, or Rotating Wall Vessel.
- Reduces gravitational force to 10^{-3} to 10^{-4} times standard gravity, minimizing particle agglomeration.
- Enhances mixing and distribution of drugs within nanomaterials, improving loading efficiency.
- Optimizes drug-nanomaterial interaction, potentially leading to higher drug encapsulation or adsorption onto nanocarriers.

Nanomaterials and Drug Types

- Applicable to various nanomaterials like IONP, liposomes, polymer nanoparticles, and MOFs.
- Drug types loaded include anticancer drugs(eg. Cisplatin), proteins, peptides, nucleic acids, vitamins, and other therapeutic agents.
- Supports loading multiple drug types onto the same nanomaterial platform.

Process and Efficiency

- The drug and nanomaterial mixture undergoes microgravity exposure for 6-18 hours, improving efficiency by 2-4 times compared to standard gravity or shaking.
- The nanomaterials maintain therapeutic effects and exhibit similar drug release profiles.

Characterization and Evaluation

- The technology utilizes common characterization techniques like UV-Vis spectrophotometry, SEM (Scanning Electron Microscopy), XRD (X-ray Diffraction), and FTIR (Fourier Transform Infrared Spectroscopy) to confirm the quality, size, morphology, and loading efficiency of the nanomaterials.

CONTACT US

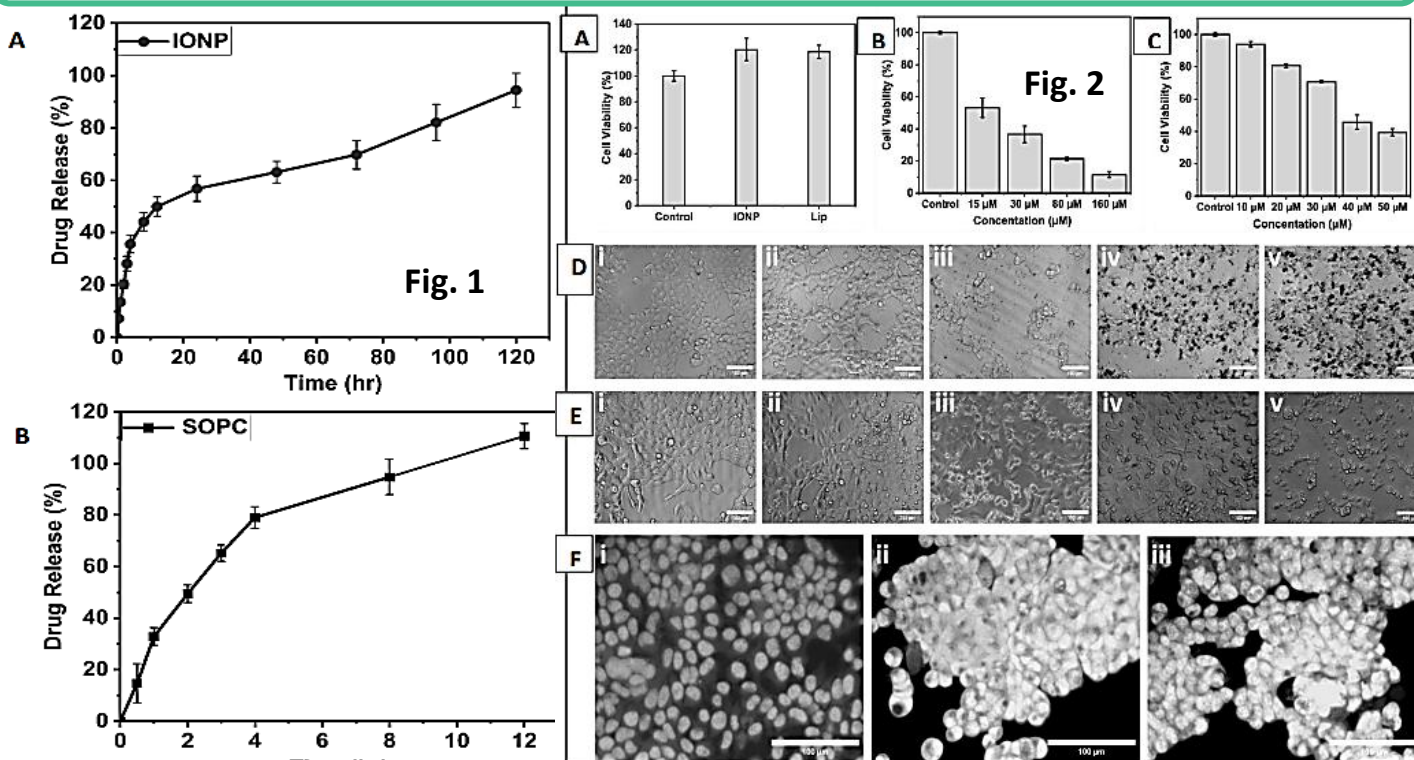
Dr. Dara Ajay, Head TTO
Technology Transfer Office,
IPM Cell- IC&SR, IIT Madras

IITM TTO Website:
<https://ipm.icsr.in/ipm/>

Email: headtto-icsr@icsrpis.iitm.ac.in
tto1@icsrpis.iitm.ac.in

Phone: +91-44-2257 9843

Fig. 1 illustrates cumulative drug release kinetics of (A) Cisplatin loaded IONP and (B) Cisplatin loaded liposomes. **Fig. 2** illustrates a) MTT assay confirming the biocompatibility of IONP and SOPC; cytotoxicity assays for b) Cis-IONP and c) Cis-Lip; microscopic images representing the cellular images at various concentrations of d) i-Control, ii-v different concentrations of Cis-IONP and e) i-v different concentrations of Cis-Lip; f) Live/dead assays fluorescence microscopic images of i) control (untreated), ii) Cis-IONP at 160 μ M, and iii) liposomes (Cis-Lip) at 50 μ M



Key Features / Value Proposition

Increased Drug Loading Efficiency

Requires lower drug doses, reducing side effects and costs.

Enhanced Drug-Carrier Interaction

Stronger drug-nanomaterial interactions for better encapsulation.

Improved Homogeneity & Distribution

Uniform drug loading with better release profiles.

Therapeutic Consistency & Safety

Maintains similar drug release and cytotoxicity to traditional methods.

Potential in Cancer Therapy

Enables targeted drug delivery to tumors, improving efficacy and reducing side effects.

Cost & Resource Efficiency

Streamlines the drug loading process, reducing labor and costs.

CONTACT US

Dr. Dara Ajay, Head TTO
Technology Transfer Office,
IPM Cell- IC&SR, IIT Madras

IITM TTO Website:
<https://ipm.icsr.in/ipm/>

Email: headtto-icsr@icsrpiis.iitm.ac.in
tto1@icsrpiis.iitm.ac.in

Phone: +91-44-2257 9843