

Drug Delivery and Nanotechnology: Quantum Principles for Targeted and Controlled Release

Lecture 10

Presented by:
Prof. Dr. Fouad Attia Majeed
Department of Medical Physics, Al-Mustaql University, Babil, Iraq
fouad.attia.majeed@uomus.edu.iq

Third-Year Students

Outline

- 1 Introduction to Nanotechnology in Medicine
- 2 Quantum Principles in Nanomedicine
- 3 Nanocarrier Types and Quantum-Controlled Systems
- 4 Clinical and Biomedical Applications
- 5 Multiple Choice Questions

What is Nanotechnology in Drug Delivery?

Nanotechnology involves engineering materials at the nanometer scale (1100 nm) to design intelligent drug carriers that can deliver therapeutic agents precisely to the target site. At this scale, quantum effects-like surface energy dominance and electron confinement-alter the materials optical, electrical, and chemical properties.

Key Idea: Nanotechnology merges with quantum mechanics to control how drugs are carried, released, and activated inside the human body.

Why Use Nanotechnology for Drug Delivery?

- Increases the bioavailability and solubility of poorly soluble drugs.
- Protects drugs from degradation before reaching their target.
- Allows sustained or controlled release over time.
- Enables site-specific (targeted) delivery-reducing side effects.

These nanosystems can respond to local triggers such as pH, temperature, or magnetic fields for precision therapy.

Quantum Basis of Nanotechnology

Quantum mechanics explains the unique behaviors of nanoparticles:

- **Quantum confinement:** electrons in nanocrystals have discrete energy levels, influencing reactivity and color.
- **Surface energy dominance:** atoms on nanoparticle surfaces are highly reactive, ideal for drug attachment.
- **Quantum tunneling:** allows controlled electron or ion transfer to trigger drug release.
- **Spin interactions:** used in magnetic nanoparticles for targeted movement using external fields.

These effects make quantum-scale materials powerful tools for precision medicine.

Nanodiamonds as Drug Carriers

Nanodiamonds (NDs) are carbon-based nanoparticles with a diamond-like structure and a diameter of 46 nm.

- Their large surface area enables efficient drug adsorption.
- Biocompatible and non-toxic; stable in physiological conditions.
- Functional groups on the surface (COOH, OH) allow covalent drug binding.
- The drug release rate can be controlled by modifying surface charge or applying external stimuli (light, heat, or pH).

Example: Doxorubicin-loaded nanodiamonds for targeted chemotherapy reduce toxicity and improve therapeutic efficiency.

Types of Nanocarriers

- **Liposomes:** lipid vesicles encapsulating drugs biodegradable and suitable for hydrophilic/hydrophobic drugs.
- **Polymeric nanoparticles:** provide sustained release through degradation of biodegradable polymers.
- **Quantum dots:** fluorescent nanocrystals used for imaging and therapy (theranostics).
- **Magnetic nanoparticles:** guided to target organs via external magnetic fields.

Each carrier can integrate quantum principles for controlled activation or imaging-guided drug delivery.

Quantum-Controlled Release Mechanisms

Drug release can be triggered by:

- **Photothermal activation:** nanoparticles absorb light, converting it to heat to release the drug.
- **pH-triggered release:** acidic tumor microenvironments break chemical bonds to release drugs.
- **Magnetically guided release:** magnetic nanoparticles respond to external magnetic pulses.
- **Electron tunneling control:** quantum effects enable electron transfer to cleave drugcarrier bonds.

These smart mechanisms minimize systemic exposure and improve treatment safety.

Applications of Quantum Nanomedicine

- **Cancer therapy:** nanodiamond-drug conjugates for localized chemotherapy.
- **Neurological disorders:** nanoparticles crossing the bloodbrain barrier.
- **Cardiology:** quantum magnetic particles for clot dissolution and vessel imaging.
- **Gene therapy:** nanoscale carriers deliver RNA/DNA molecules directly into cells.

Nanotechnology offers precision-level control over when and where a drug acts.

Advantages of Quantum-Based Drug Delivery

- Target-specific and efficient delivery minimizes side effects.
- Lower doses achieve higher efficacy.
- Enhanced drug stability and controlled pharmacokinetics.
- Integration with diagnostic imaging (theranostics).
- Enables real-time monitoring of drug distribution using fluorescence or magnetic resonance.

Challenges and Ethical Considerations

- Long-term safety and biodegradability of nanomaterials remain under investigation.
- Risk of nanotoxicity and accumulation in organs.
- Manufacturing reproducibility and scalability issues.
- Need for regulatory frameworks for quantum-enabled nanomedicine.

Despite challenges, research continues toward safe clinical translation.

MCQ 1-2

Q1. Nanotechnology in medicine deals with materials sized:

- A) 110 μm
- B) 1100 nm
- C) 110 mm
- D) 110 cm

Q2. Quantum confinement in nanoparticles affects:

- A) The atomic weight only
- B) Optical and electronic properties
- C) Mechanical hardness only
- D) Chemical bonds only

MCQ 3-4

Q3. Nanodiamonds are preferred drug carriers because:

- A) They are toxic
- B) They are biocompatible with large surface area
- C) They dissolve easily in blood
- D) They emit harmful radiation

Q4. Drug release from nanodiamonds can be controlled by:

- A) Surface modification and external stimuli
- B) Reducing the patients temperature
- C) Increasing gravity
- D) Sound vibration

MCQ 5-6

Q5. Liposomes are mainly composed of:

- A) Polymers
- B) Lipid bilayers
- C) Carbon nanotubes
- D) Proteins

Q6. Magnetic nanoparticles are guided to target organs using:

- A) Electric current
- B) Sound waves
- C) External magnetic fields
- D) UV light

MCQ 7-8

Q7. The surface functional groups on nanodiamonds help to:

- A) Bind drugs chemically or physically
- B) Increase toxicity
- C) Reduce surface area
- D) Dissolve in fat

Q8. Photothermal activation works by:

- A) Absorbing light and converting it to heat
- B) Reducing drug concentration
- C) Cooling the target site
- D) Changing pH

MCQ 9-10

Q9. pH-triggered drug release is suitable for:

- A) Brain tissue
- B) Acidic tumor environments
- C) Alkaline bones
- D) Muscle fibers

Q10. The main biomedical use of quantum dots is:

- A) Measuring temperature
- B) Drug imaging and theranostics
- C) Generating magnetic fields
- D) Delivering oxygen

MCQ 11-12

Q11. Quantum tunneling can control drug release by:

- A) Allowing electron transfer through molecular barriers
- B) Increasing viscosity
- C) Blocking ions
- D) Cooling nanoparticles

Q12. The primary advantage of targeted drug delivery is:

- A) Higher side effects
- B) Increased drug wastage
- C) Reduced systemic toxicity
- D) Uncontrolled release

MCQ 13-14

Q13. Nanotechnology improves bioavailability by:

- A) Increasing particle size
- B) Enhancing solubility of drugs
- C) Reducing absorption
- D) Inhibiting transport

Q14. Which property makes nanodiamonds ideal for long-term use?

- A) Chemical inertness and stability
- B) Radioactivity
- C) Fluorescence emission
- D) Thermal expansion

MCQ 15-16

Q15. Magnetic nanoparticle therapy can treat:

- A) Liver fibrosis only
- B) Cancer by heating tumors magnetically
- C) Kidney stones only
- D) Cardiac rhythm disorders

Q16. Which of the following combines diagnosis and therapy?

- A) Theranostic systems
- B) Pharmacokinetics
- C) Toxicology
- D) Immunoassays

MCQ 17-18

Q17. The major challenge in clinical nanomedicine is:

- A) Fabrication reproducibility and long-term safety
- B) Excess fluorescence
- C) Lack of funding
- D) Over-simplified chemistry

Q18. Quantum effects dominate at scales below:

- A) 1 μ m
- B) 100 nm
- C) 1 mm
- D) 10 cm

Q19. The controlled release from nanocarriers ensures:

- A) Constant therapeutic concentration
- B) Rapid overdose
- C) Irregular plasma levels
- D) Ineffective treatment

Q20. The ultimate goal of quantum nanotechnology in medicine is:

- A) Smart, personalized, minimally invasive therapy
- B) Increasing hospital stay duration
- C) Reducing accuracy of treatments
- D) Limiting research scope