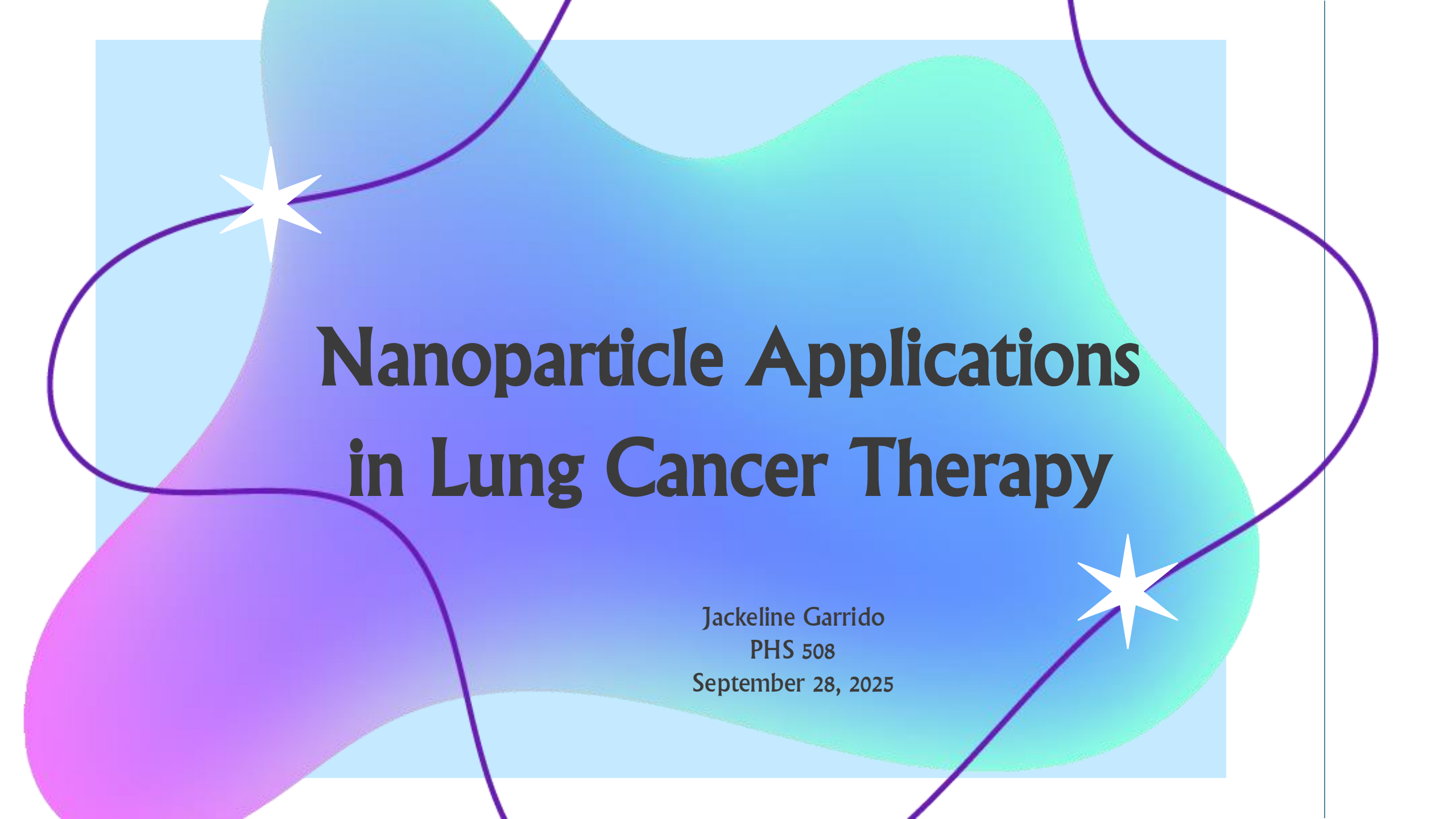




Nanoparticle Applications in Lung Cancer Therapy

Jackeline Garrido
PHS 508
September 28, 2025

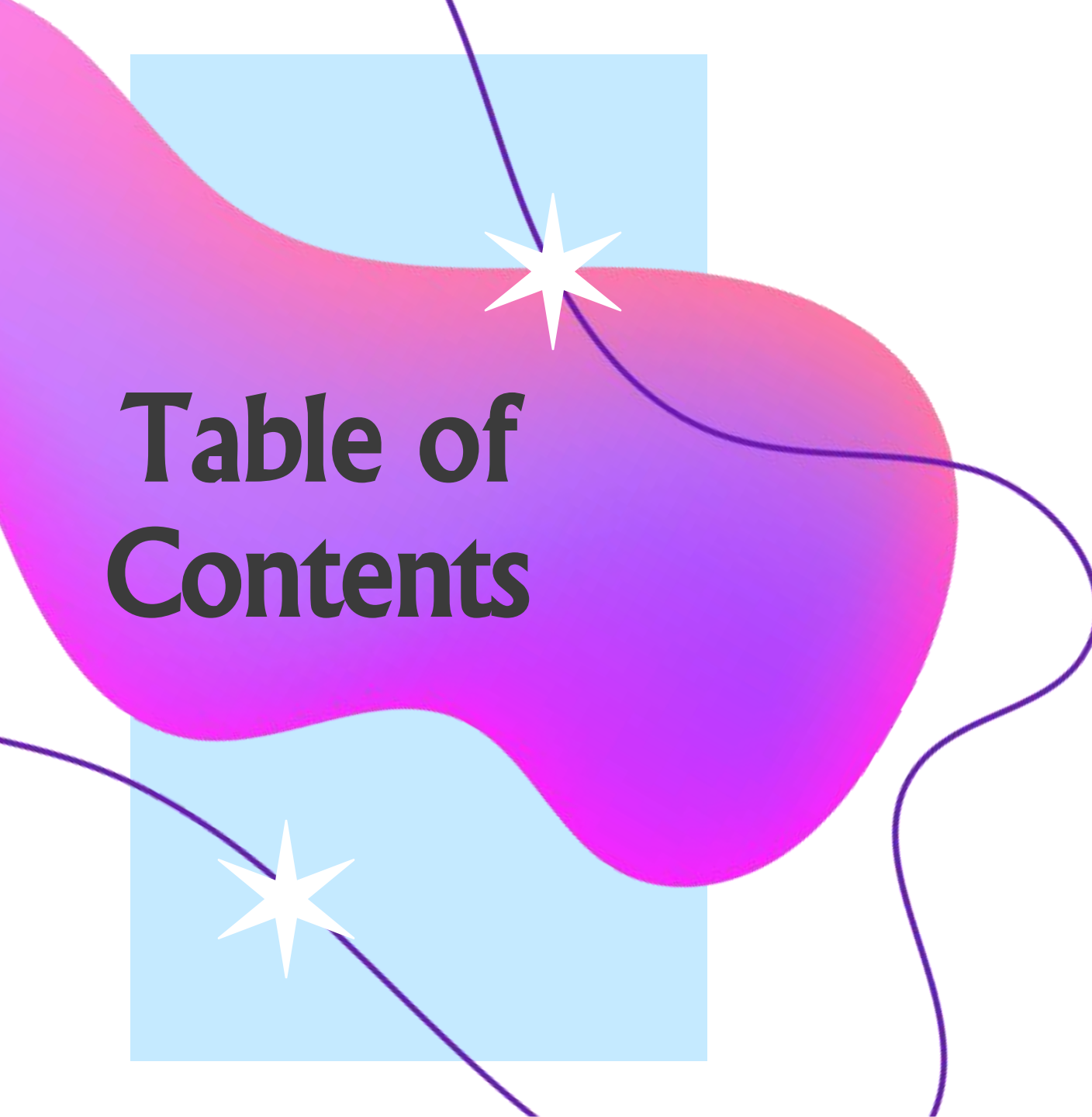
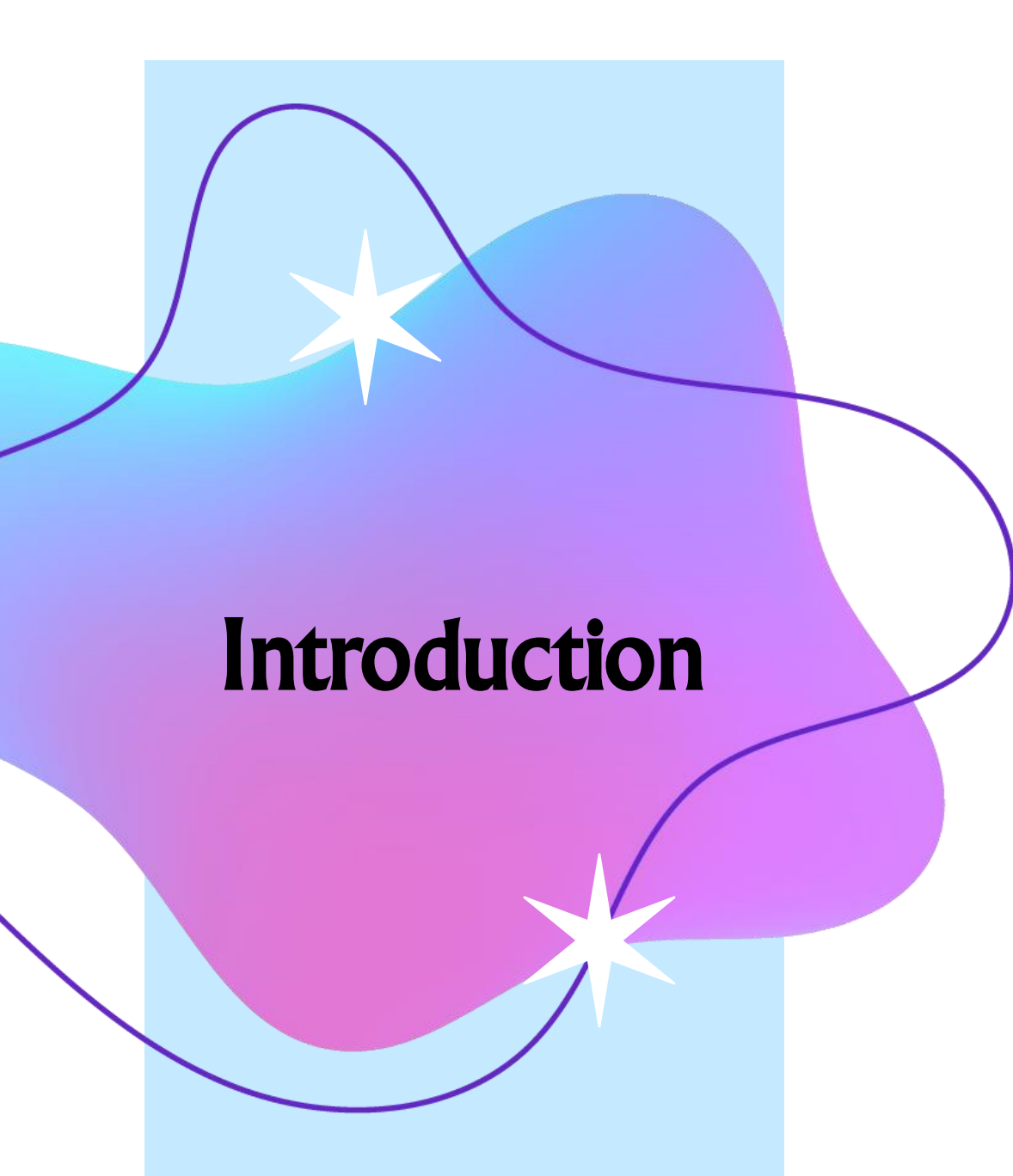


Table of Contents

- **Introduction**
- **Overview of Nanotechnology in Medicine**
 - Types of Nanoparticles
 - Liposomes & Lipid Nanoparticles
 - Dendrimers
 - Gold Nanoparticles
 - Polymeric Nanoparticles
- **Advantages of Nanomedicine**
- **Extracellular Vesicles (EVs) in Cancer**
- **FDA-Approved Nanomedicines**
- **Challenges in Nanomedicine Development**
- **Clinical Trials in Nanotechnology**
- **Future Applications**
 - Smart Nanocarriers
 - AI & Machine Learning
 - CRISPR-based Therapies
 - Theranostics
- **Ethical & Regulatory Considerations**
- **Conclusion**
- **References**



Introduction

- Nanotechnology: frontier in modern medicine
- Focus: cancer treatment, esp. lung cancer
- Modalities: liposomes, polymeric NPs, gold NPs, extracellular vesicles (EVs)
- Potential: personalized & targeted therapy
- Challenges: toxicity, immune reactions, regulatory hurdles

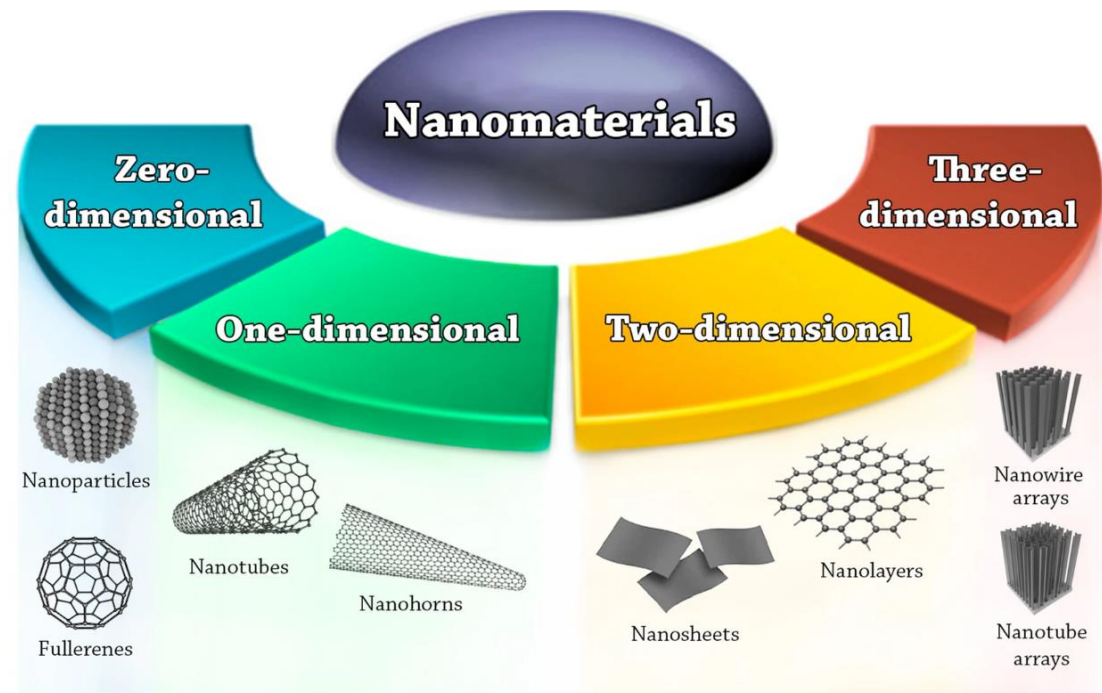
- Nanoparticles = 1–100 nm, unique properties
- Used in drug delivery, imaging, regenerative medicine
- Three classes: Organic, Carbon-based, Inorganic
- Enable new approaches not possible with traditional drugs



Overview of Nanotechnology in Medicine

Types of Nanoparticles

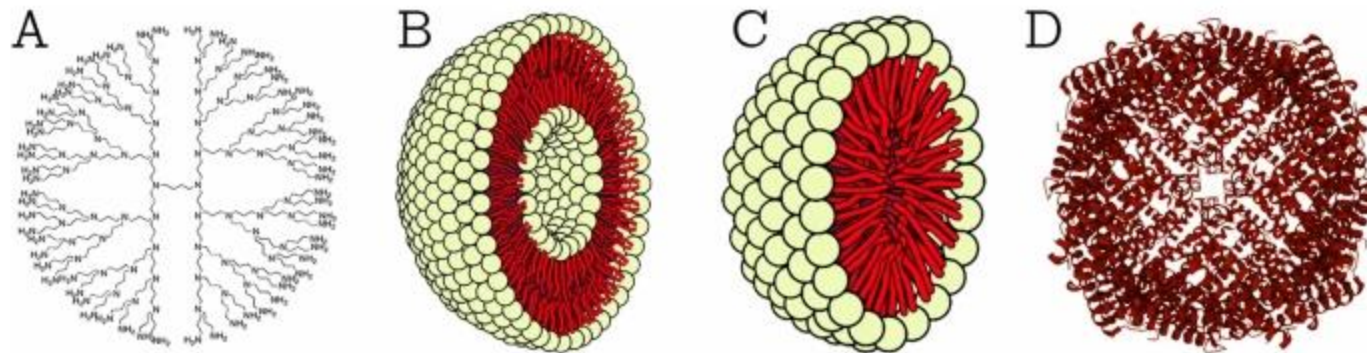
- Organic: liposomes, dendrimers, lipid carriers
- Inorganic: gold nanoparticles
- Polymeric: versatile, controlled-release systems
- Each type has unique advantages and limitations



From: [Nanoparticle classification, physicochemical properties, characterization, and applications: a comprehensive review for biologists](#)

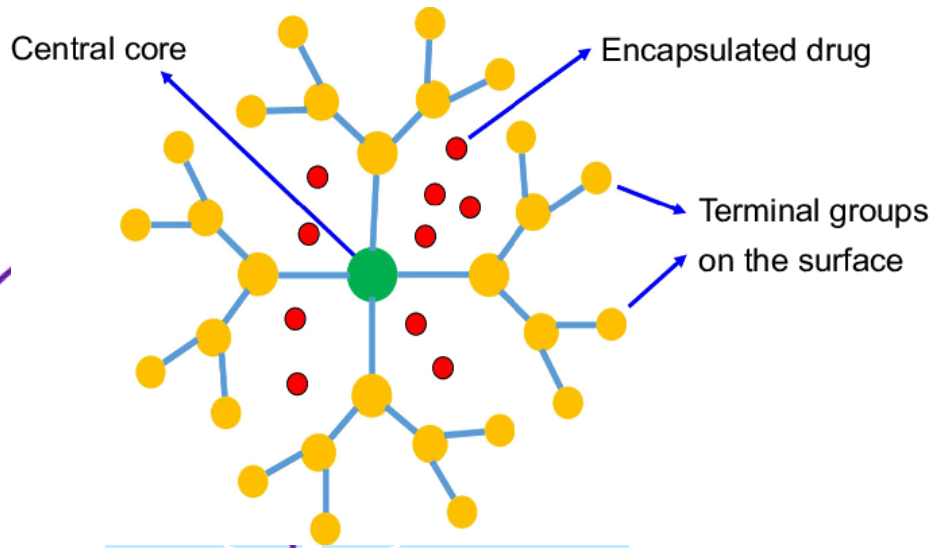
Liposomes & Lipid Nanoparticles

- Lipid bilayer spheres encapsulating drugs
- Enhance drug stability & targeting
- Categories: liposomes, niosomes, transfersomes, SLN, NLC
- FDA-approved example: Doxil (liposomal doxorubicin)
- Limitation: storage stability, dermal delivery barriers



Types of organic NPs. **A** Dendrimers; **B** liposomes; **C** micelles; and **D** ferritin

From: [Nanoparticle classification, physicochemical properties, characterization, and applications: a comprehensive review for biologists](#)



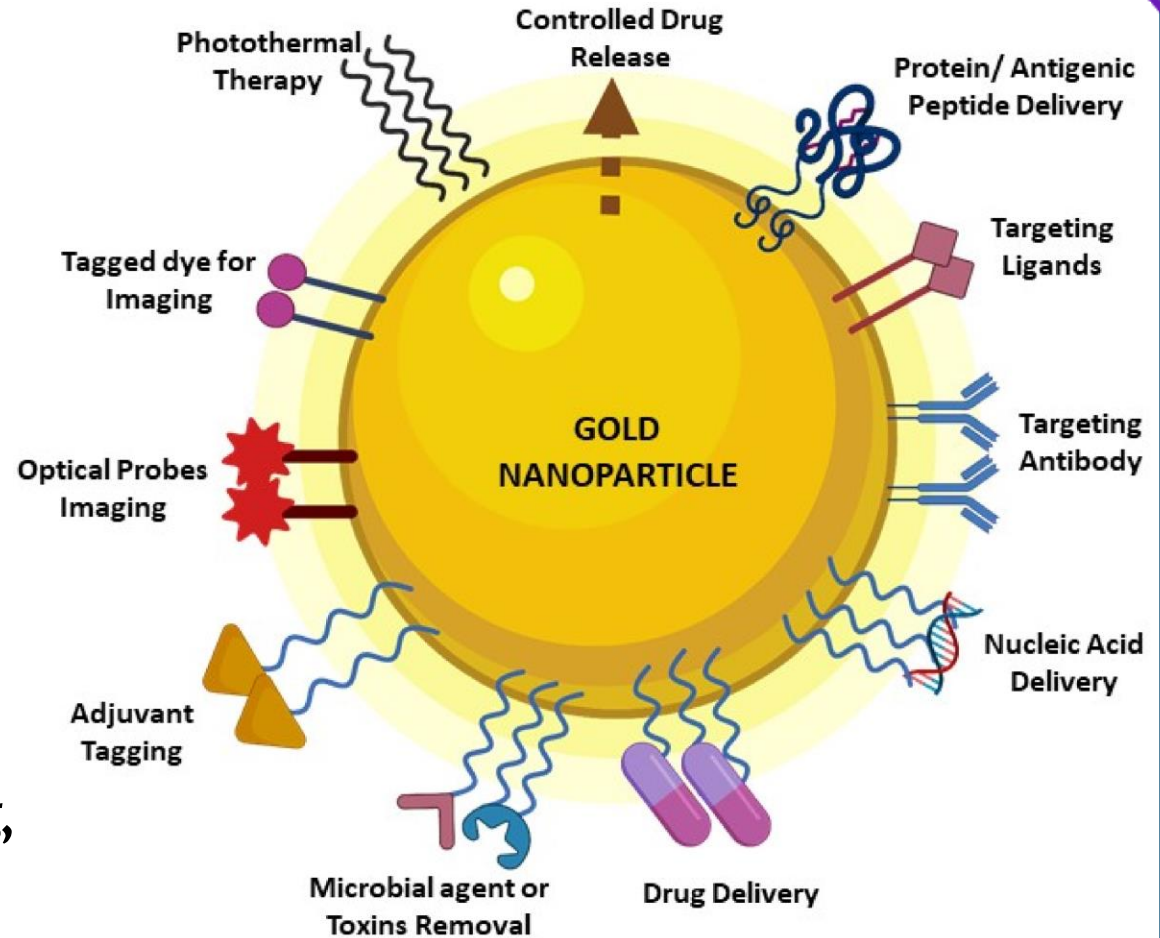
Zeb, Alam & Khan, Fakhar & Aman, Waqar & Qureshi, Omer. (2017). Effective use of nano-carriers as drug delivery systems for the treatment of selected tumors. International Journal of Nanomedicine. 12. 7291–7309.

Dendrimers

- Tree-like, branched structures
- Functional groups allow high drug loading
- Delivery modes:
 - Physical encapsulation
 - Electrostatic interactions
 - Covalent conjugation
- Example: PAMAM dendrimers for anticancer drugs

Gold Nanoparticles

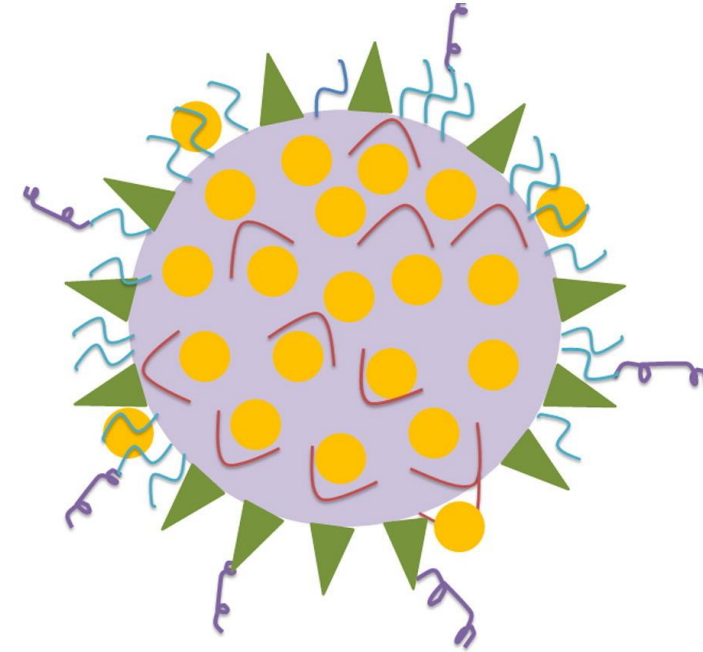
- Small, stable, non-toxic, biocompatible
- Tunable shapes (spheres, rods, shells)
- Surface modifications: citrate, thiols, PEG
- Unique property: surface plasmon resonance (SPR)
- Applications: photothermal therapy, imaging, drug delivery



From: **Efficacy and Immune Response Elicited by Gold Nanoparticle- Based Nanovaccines against Infectious Diseases**
<https://www.mdpi.com/2076-393X/10/4/505>

Polymeric Nanoparticles

- Controlled drug release systems
- Adaptable for inhalation delivery
- Improve solubility, stability, tumor-specific release
- Barriers: toxicity, scalability, regulatory approval



Key:

 Polymeric matrix (Polyhydroxyalkanoates, poly-lactide-co-glycolide, cyclodextrin etc.)

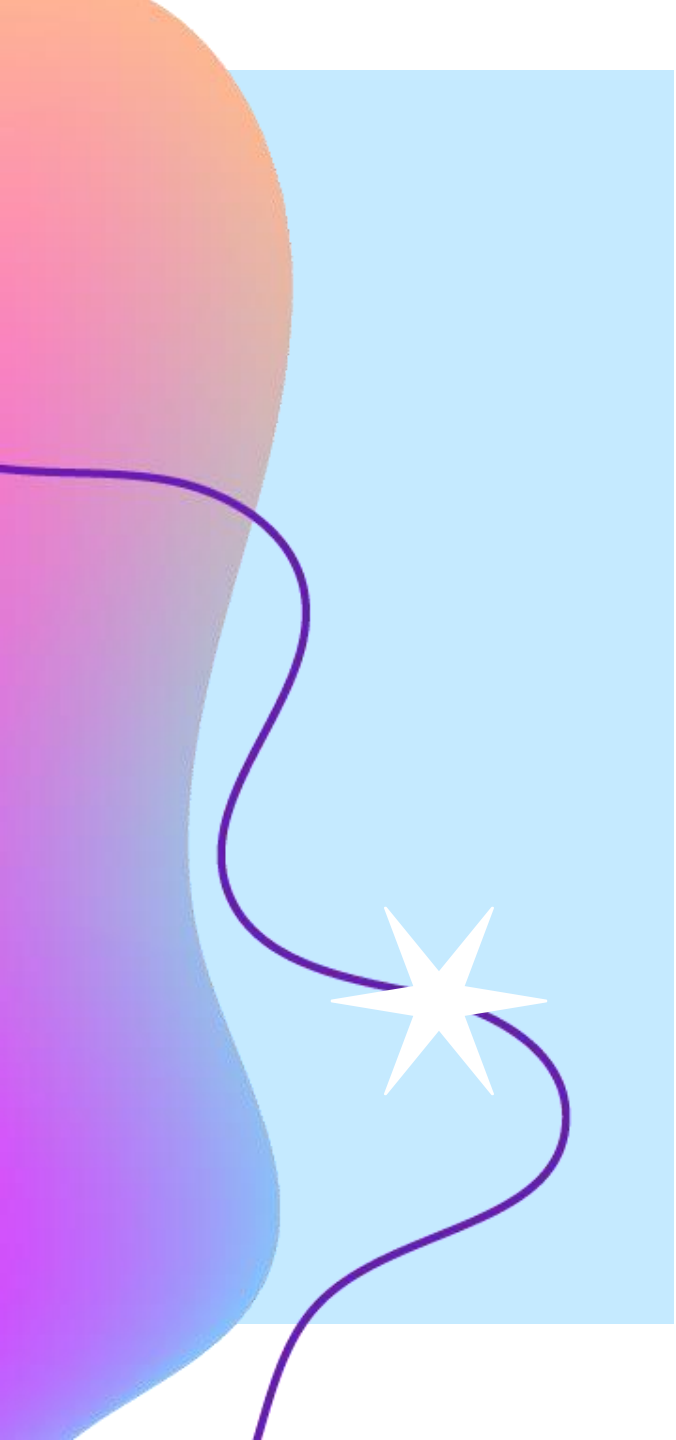
 Drug (Paclitaxel, Cisplatin, Doxorubicin etc.)

 Short interfering RNA

 Ligand (small molecules, peptides, antibodies, aptamers etc.)

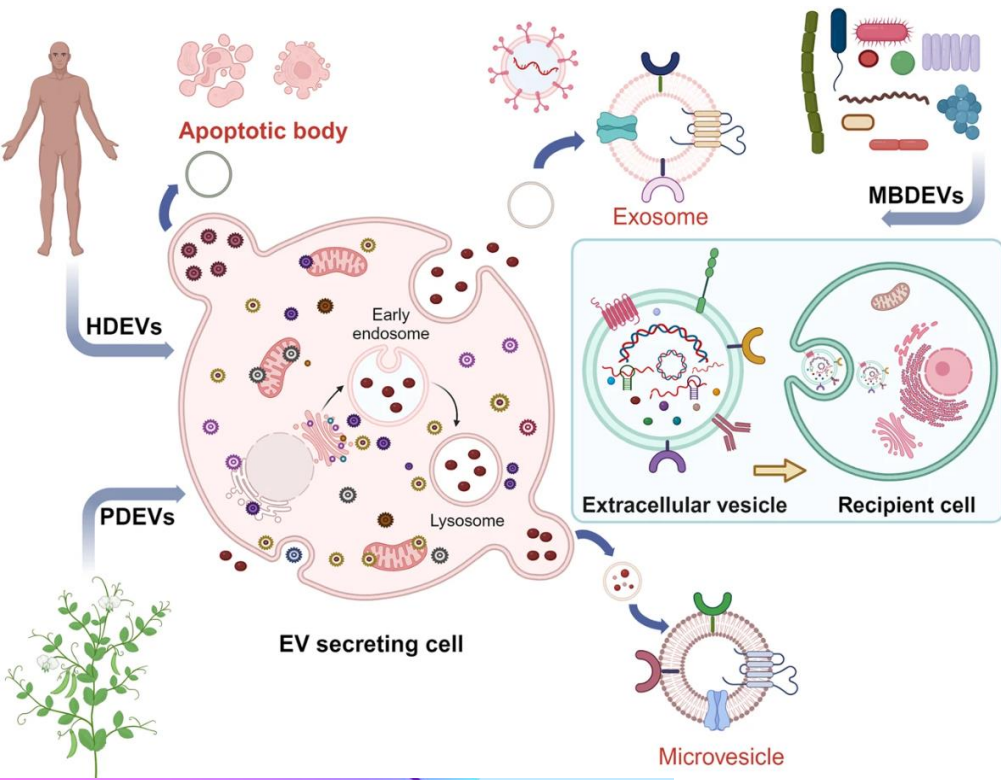
 Hydrophilic outer shell (Polyethylene glycol, polyvinyl alcohol etc.)

 Peptides (RGD, ATN-161 etc)



Advantages of Nanomedicine

- Improved drug bioavailability
- Reduced systemic toxicity
- Targeted delivery via EPR effect & modifications
- Potential for personalized medicine
- Example: EVs as autologous nanomedicine



Extracellular Vesicles (EVs) in Cancer

- Nano-sized, lipid-bound particles released by cells
- Subtypes: exosomes, microvesicles, apoptotic bodies
- Functions: intercellular communication, immune modulation
- Clinical uses: liquid biopsies, drug delivery, cancer vaccines
- Challenges: large-scale production & standardization

<https://www.nature.com/articles/s41392-024-01735-1>



FDA-Approved Nanomedicines

- **Abraxane:** albumin-bound paclitaxel, better tolerated than solvent-based forms
- **Genexol-PM:** polymeric micelle paclitaxel, improved solubility & dosing
- **Doxil:** liposomal doxorubicin, reduced cardiotoxicity
- Limited approvals compared to systems in research

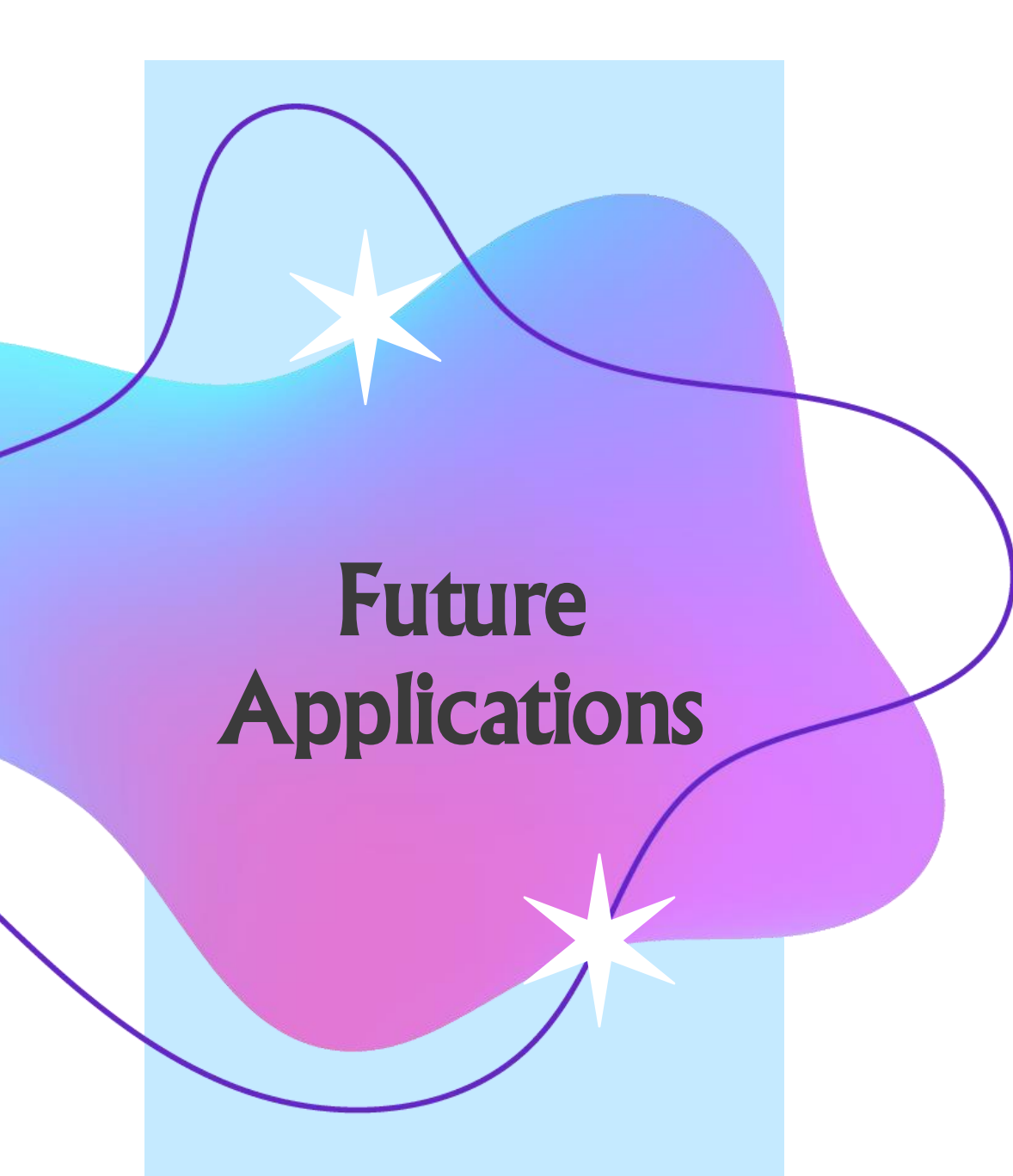
Challenges in Nanomedicine Development

- Low drug encapsulation efficiency
- Stability issues during circulation/storage
- Immune rejection & clearance
- Long-term safety: organ accumulation, chronic toxicity
- Tumor variability → need for personalized approaches



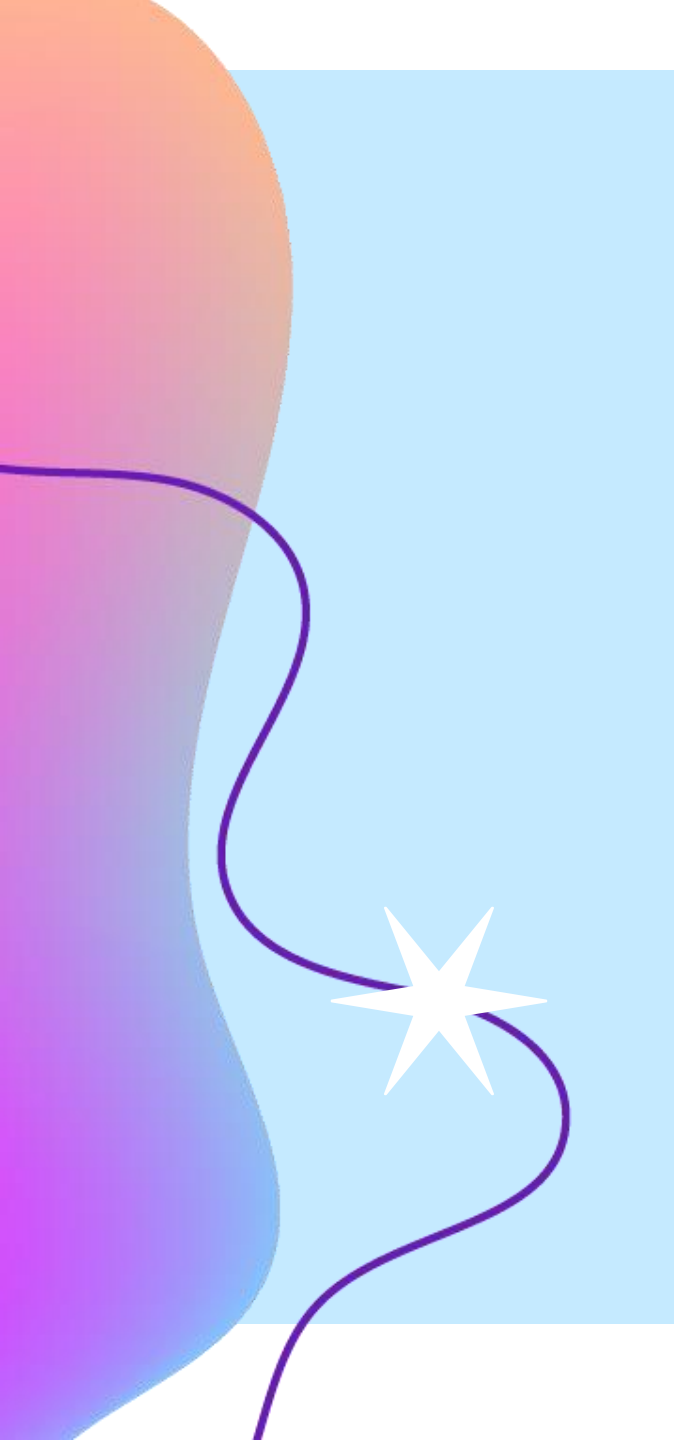
Clinical Trials in Nanotechnology

- Nanoparticle immunotherapies with ICIs
- Nanobiotix (NBTXR3): radioenhancer in trials
- EV-based liquid biopsies for real-time monitoring
- CRISPR-loaded lipid nanoparticles under evaluation
- Ongoing trials show promise but need long-term safety data

A graphic on the left side of the slide featuring a light blue square background. Overlaid on this is a large, irregular, organic shape with a purple-to-pink gradient. A dark purple line meanders through the shape, forming a loop. Two white, multi-pointed starburst icons are placed on the purple line: one in the upper left and one in the lower right. The text 'Future Applications' is centered within the pinkish-purple area of the shape.

Future Applications

- **Smart nanocarriers:** stimuli-responsive (pH, enzymes, temperature)
- **AI/ML models:** optimize design, size, release profiles
- **CRISPR delivery:** gene editing for cancer mutations
- **Theranostics:** dual therapy + diagnostic systems



Ethical & Regulatory Considerations

- Long-term patient safety: organ accumulation, biodegradation
- Equitable access: high costs & limited availability
- Genetic manipulation risks (CRISPR misuse)
- Regulatory gaps: EVs & multifunctional systems need harmonization

Conclusion

- Nanotechnology shifts lung cancer therapy from broad toxicity → precision targeting
- Modalities: liposomes, dendrimers, gold NPs, polymeric NPs, EVs
- Goal: maximize tumor drug delivery, minimize side effects
- Future: AI-guided design, CRISPR therapies, theranostic tools
- Needs: safety frameworks, global EV standards, equitable access



Thank You



References

(Condensed, 10 key sources)

- Abdullah, A. (2025). *Cancer nanomedicine: From bench to bedside*. *Frontiers in Oncology*, 15, 110238. <https://doi.org/10.3389/fonc.2025.110238>
- Asleh, R., et al. (2023). Extracellular vesicle isolation and analysis for liquid biopsies in cancer. *Biomarker Research*, 11(1), 67. <https://doi.org/10.1186/s40364-023-00540-2>
- Crintea, A., Melinte, C., & Turcu-Stolicea, A. (2021). Nanomedicine in cancer therapy: Advances and challenges. *Journal of Clinical Medicine*, 10(15), 3350. <https://doi.org/10.3390/jcm10153350>
- Inner, R., Schubert, A., Kretschmer, V., & Pfaffl, M. W. (2023). Extracellular vesicle isolation methods: The need for standardization in clinical applications. *International Journal of Molecular Sciences*, 24(3), 2102. <https://doi.org/10.3390/ijms24032102>
- Kato, T., Yoshioka, Y., & Ochiya, T. (2021). Exosomes: Emerging diagnostic and therapeutic targets in human cancer. *Cancers*, 13(2), 307. <https://doi.org/10.3390/cancers13020307>
- Masarwy, R., et al. (2025). CRISPR/Cas9 lipid nanoparticle delivery for gene editing in head and neck cancer. *Advanced Science*, 12(15), 1032. <https://doi.org/10.1002/advs.202411032>
- Nanobiotix. (2025). *NBTXR3 clinical development pipeline*. Retrieved from <https://nanobiotix.com>
- Shen, C., Frakes, J. M., et al. (2023). Efficacy from ongoing Phase I trial with NBTXR3 + anti-PD-1 in advanced cancers. *Journal of Clinical Oncology*, 41(16_suppl), 6038. https://doi.org/10.1200/JCO.2023.41.16_suppl.6038
- Svenson, S. (2018). Dendrimers as versatile nanocarriers for drug delivery. *Journal of Controlled Release*, 262, 200–215. <https://doi.org/10.1016/j.jconrel.2017.07.031>
- Jain, S., Hirst, D. G., & O'Sullivan, J. M. (2012). Gold nanoparticles as novel agents for cancer therapy. *British Journal of Radiology*, 85(1010), 101–113. <https://doi.org/10.1259/bjr/59448833>