



Article

Silicon Nanoparticles in Targeted Cancer Therapy

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Abstract: Silicon nanoparticles represent a promising advancement in cancer treatment, offering precise means to target cancer cells while minimizing side effects on healthy cells. Their applications include targeted drug delivery, enhanced immunotherapy, phototherapy, and thermal therapy techniques, thereby increasing treatment efficacy and reducing tumor resistance. Despite these benefits, long-term studies are necessary to ensure the safety and effectiveness of these technologies, emphasizing the need for future innovations to overcome challenges and expand their use in medicine.

1. Introduction

Nanotechnology is a modern branch of science focusing on the study and fabrication of materials at the nanometer scale (1–100 nanometers). Emerging in the 1970s, it gained significant development in the 1990s with advancements in research and analytical tools. Nanotechnology involves the use of nanoparticles that exhibit unique properties, enabling innovative applications across various fields [1].

Nanoparticles are materials within the 1–100 nanometer range that possess unique characteristics, distinguishing them from bulk materials. These properties influence their chemical and physical behavior, making them highly applicable in fields like medicine, electronics, and energy. In drug delivery, nanoparticles enhance treatment effectiveness and reduce side effects. They are also employed to develop new materials with improved properties, such as more efficient batteries or higher-performing solar cells [2].

Silicon Nanoparticles

Silicon nanoparticles have emerged as a significant advancement in medical technology for cancer treatment. These particles have demonstrated their ability to improve drug delivery to tumors in a targeted and precise manner, reducing damage to surrounding healthy tissues. Their biodegradable and easily excretable nature makes them a promising option for medical applications, unlike other nanoparticles that may accumulate and cause side effects [3,4].

Properties of Nanoparticles

Nanoparticles exhibit unique properties that differentiate them from bulk materials. These include [5,6,7,8].

High Surface-to-Volume Ratio: Enhances chemical reactivity, making them effective in applications like catalysis and drug delivery.

Optical and Electrical Properties: Differ from bulk materials, enabling uses in electronics and photonics, such as displays and solar cells.

Magnetic Properties: Unique features like super Para magnetism are useful in magnetic resonance imaging (MRI) and data storage.

Mechanical Properties: Greater mechanical strength and durability, suitable for coatings and advanced materials.

Thermal and Chemical Stability: Some nanoparticles are more stable at specific temperatures or less prone to corrosion, increasing their longevity in harsh conditions.

Quantum Effects: At the nanoscale, quantum effects influence chemical and physical behaviors, leading to unconventional properties like color changes based on particle size.

Types of Nanoparticles

Nanoparticles vary widely in types and applications, including [9,10,11,12,13]:

Quantum Dots: Semiconductor nanoparticles with unique optical properties, used in imaging and photonics.

etallic Nanoparticles: Gold, silver, platinum, and copper particles, utilized in catalysis and medical applications.

Carbon Nanotubes: Cylindrical structures with high mechanical and electrical properties, applied in electronics and construction.

Magnetic Nanoparticles: Made of magnetic metals like iron, used in MRI and data storage.

Ceramic Nanoparticles: Resistant to heat and corrosion, used in drug delivery and water treatment.

Polymeric Nanoparticles: Biodegradable and used for controlled drug release.

Nano fibers: Employed in tissue engineering, filters, and drug delivery systems.

Nano shells: Used in drug delivery and photo thermal therapy.

Methods of Nanoparticle Synthesis

Nanoparticle synthesis methods are categorized into[14,15,16]

Physical Methods

Laser Ablation: High-energy lasers vaporize solid materials into nanoparticles.

Ball Milling: Mechanical grinding of materials to nanoscale particles.

Physical Vapor Deposition (PVD): Vaporizes and condenses materials onto surfaces.

2-Chemical Methods

Chemical Reduction: Converts metal ions into nanoparticles.

Co-precipitation: Precipitates materials into nanoparticles from solution.

Thermal Decomposition: Produces particles through high-temperature decomposition.

Sol-Gel Process: Transforms solutions into gels for nanoparticle synthesis.

Biological Methods

Biosynthesis: Utilizes microorganisms to create nanoparticles.

Enzymatic Reduction: Employs enzymes for green synthesis.

Nanoparticles in Cancer Therapy

Nanoparticles enable innovative cancer treatments, including [17,18,19,20,21]:

- Targeted Drug Delivery: Reduces side effects by delivering drugs directly to cancer cells.
- Photodynamic Therapy: Uses light-sensitive nanoparticles to kill cancer cells.
- Photo thermal Therapy: Converts light into heat to destroy tumors.
- Nanoparticle-Based Imaging: Improves diagnostic imaging.
- Immunotherapy: Enhances immune response against cancer cells.
- Gene Therapy: Delivers genetic material to modify cancer cells.
- Smart Chemotherapy: Releases drugs only in tumor environments.
- Magnetic Nanoparticles: Directs therapies to tumors using external magnetic fields.

Degradation of Silicon Nanoparticles

Silicon nanoparticles degrade safely within the body through [22,23,24,25,26]:

- 1 - Hydrolysis: Converts silicon into execrable silicic acid.
- 2 - pH Sensitivity: Accelerates degradation in acidic tumor environments.
- 3 - Enzymatic Interaction: Speeds up degradation through enzyme reactions.
- 4 -Excretion: Degraded particles are expelled via the kidneys.

Conclusions

Promising Potential: Nanoparticles provide innovative solutions in cancer therapy, enhancing drug delivery and reducing side effects.

Enhanced Efficacy: Combine therapies to increase effectiveness and lower resistance.

3-Future Studies

- Improving Precision: Developing smarter nanoparticles for cancerspecific targeting.
- Multifunctional Delivery: Designing particles to carry multiple drugs.
- Long-Term Safety: Evaluating nanoparticle safety over time.
- Enhanced Immunotherapy: Activating immune cells more effectively.
- 5-Optimized Phototherapy: Using light-sensitive nanoparticles for advanced therapies.

REFERENCES

1. Kulkarni, S. K. (2015). Nanotechnology: Principles and Practices. Springer.
2. Hornyak, G. L., Tibbals, H. F., Dutta, J., & Moore, J. J. (2008). Introduction to Nanoscience and Nanotechnology. CRC Press.
3. Cao, G., & Wang, Y. (2011). Nanostructures and Nanomaterials: Synthesis, Properties, and Applications. World Scientific Publishing.
4. Chou, L. Y. T., Ming, K., & Chan, W. C. W. (2011). Strategies for the intracellular delivery of nanoparticles. Chemical Society Reviews, 40(1), 233-245. <https://doi.org/10.1039>.
5. Zhang, X., Yin, H., Xu, Z., & Liu, Z. (2013). Recent progress in sensors based on nanomaterials. Sensors, 13(7), 9381-9403. <https://doi.org/10.3390/s130709381>.
6. Huo, S., & Jin, S. (2020). Nanoparticle interactions and toxicity in biological systems. Nature Nanotechnology, 15(3), 153-167. <https://doi.org/10.1038/s41565-020-0673-7>.

7. Lee, J. H. (2018). Applications of Nanotechnology in Medicine: Drug Delivery and Cancer Therapy (Doctoral dissertation, University of California, Berkeley).
8. Kumar, C. S. S. R. (2006). Nanostructured Materials: Processing, Properties and Applications. William Andrew Publishing.
9. Singh, P., Pandit, S., Mokkapati, V. R. S. S., Garg, A., Ravikumar, V., & Mijakovic, I. (2018). Gold nanoparticles in diagnostics and therapeutics for human cancer. *International Journal of Molecular Sciences*, 19(7), 1979. <https://doi.org/10.3390/ijms19071979>.
10. Mitragotri, S., & Lahann, J. (2009). Physical approaches to biomaterial design. *Nature Materials*, 8(1), 15-23. <https://doi.org/10.1038/nmat2344>.
11. Gupta, A. (2014). Nanoparticles for Drug Delivery: Properties and Applications (Doctoral dissertation, Stanford University).
12. Wang, Y. (2013). Synthesis and Application of Magnetic Nanoparticles in Medicine (Doctoral dissertation, University of California, Los Angeles).
13. Rai, M., Duran, N., & Ahamed, M. (2011). Metal Nanoparticles in Microbiology: Fundamentals and Applications. Springer.
14. Iravani, S. (2011). Green synthesis of metal nanoparticles using plants. *Green Chemistry*, 13(10), 2638-2650. <https://doi.org/10.1039/c1gc15386b>.
15. Mourdikoudis, S., & Liz-Marzán, L. M. (2013). Formation of stable colloidal solutions of metal nanoparticles: Critical steps and tips on nanoparticle synthesis. *Chemistry of Materials*, 25(9), 1465-1476.
16. Gupta, A. (2014). Nanoparticles for Drug Delivery: Properties and Applications (Doctoral dissertation, Stanford University).
17. Wang, Y. (2013). Synthesis and Application of Magnetic Nanoparticles in Medicine (Doctoral dissertation, University of California, Los Angeles).
18. Torchilin, V. P. (2006). Nanoparticulates as Drug Carriers. Imperial College Press.
19. Peer, D., et al. (2007). Nanocarriers as an emerging platform for cancer therapy. *Nature Nanotechnology*, 2(12), 751-760.
20. Ferrari, M. (2005). Cancer nanotechnology: Opportunities and challenges. *Nature Reviews Cancer*, 5(3), 161-171. <https://doi.org/10.1038/nrc1566>.
21. Grobmyer, S. R., & Moudgil, B. M. (2010). Cancer Nanotechnology: Methods and Protocols. Humana Press.
22. Park, J. H., Gu, L., Von Maltzahn, G., Ruoslahti, E., Bhatia, S. N., & Sailor, M. J. (2009). Biodegradable luminescent porous silicon nanoparticles for in vivo applications. *Nature Materials*, 8(4), 331-336. <https://doi.org/10.1038/nmat2398>.
23. He, Q., & Shi, J. (2011). Mesoporous silica nanoparticle-based nano drug delivery systems: Synthesis, controlled drug release and delivery, pharmacokinetics, and biodistribution. *Journal of Materials Chemistry*, 21(16), 5845-5855. <https://doi.org/10.1039/C0JM03851A>.
24. Croissant, J. G., Fatieiev, Y., & Khashab, N. M. (2017). Degradability and clearance of silicon, organosilica, and silica-based nanomaterials. *Advanced Materials*, 29(9), 1604634. <https://doi.org/10.1002/adma.201604634>.
25. Kumar, C. S. (2007). Nanoparticles for Biomedical Applications. Wiley-VCH.

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26. Slowing, I. I., Trewyn, B. G., & Lin, V. S. Y. (2007). Effect of surface functionalization of MCM-41-type mesoporous silica nanoparticles on the endocytosis by human cancer cells. *Journal of the American Chemical Society*, 129(28), 8845–8849.
<https://doi.org/10.1021/ja071978n>.