

Design and Development of a Targeted Molecular Probe for Ultrasensitive Early Detection of Breast Cancer



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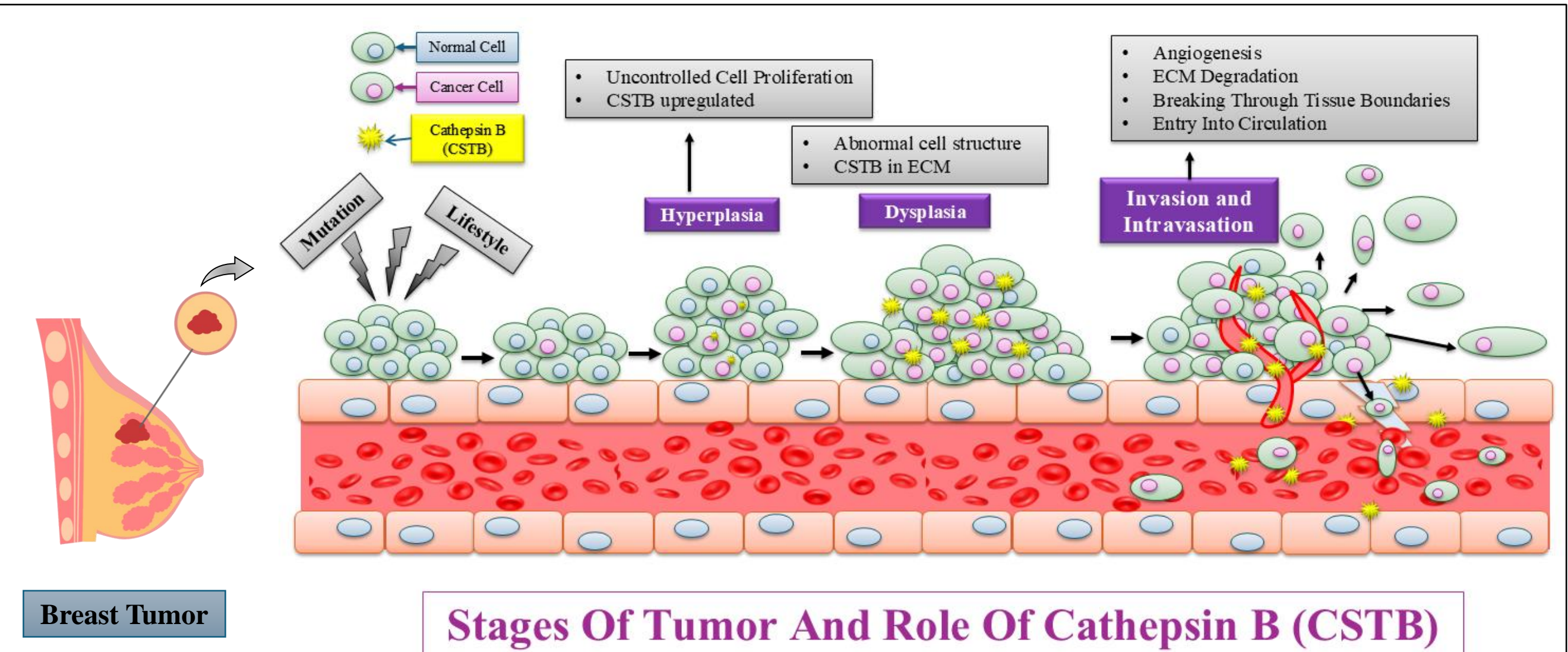
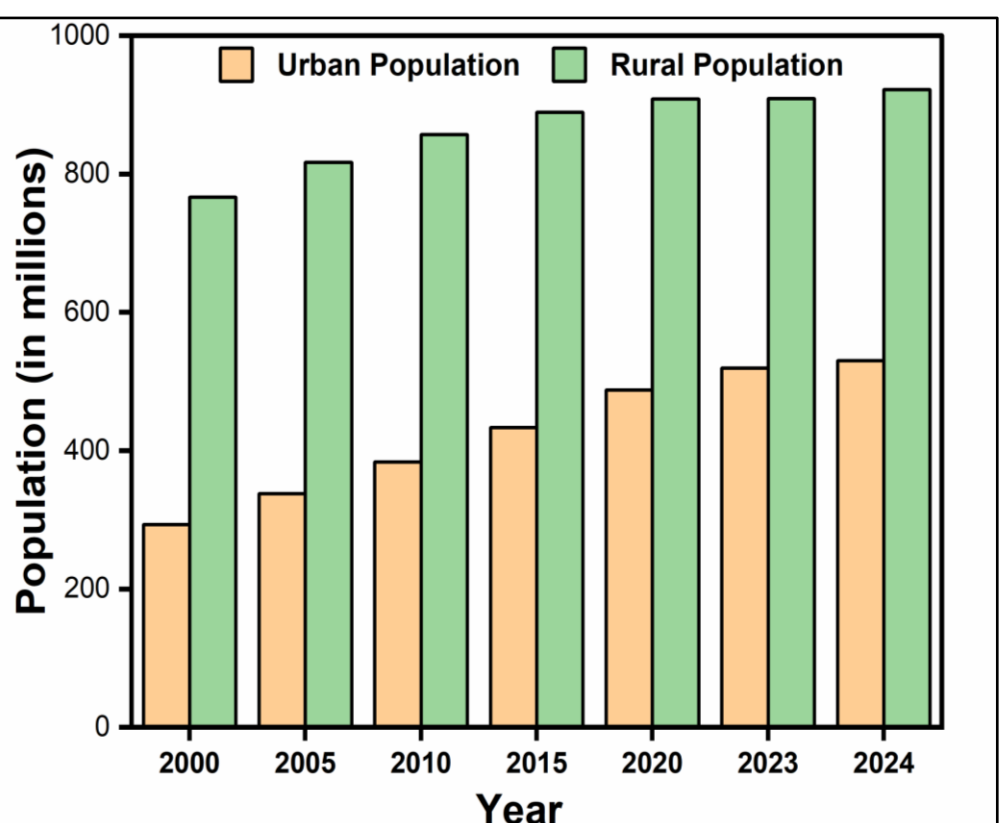
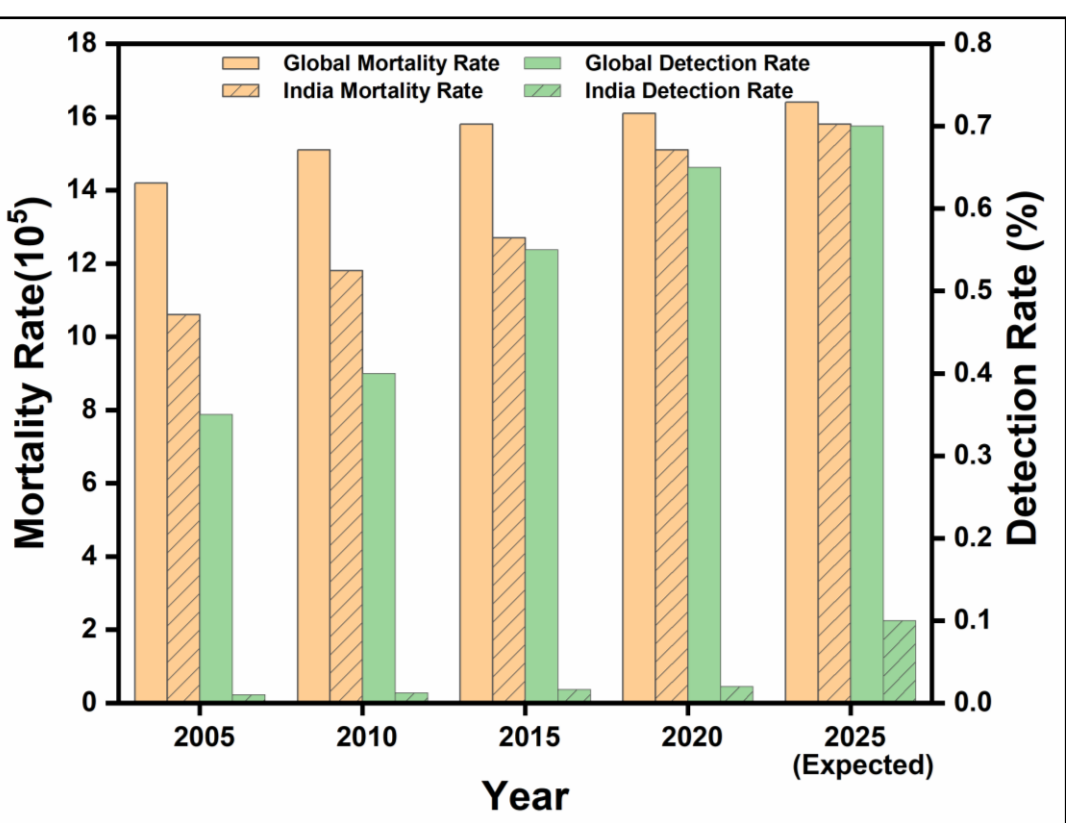
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Background

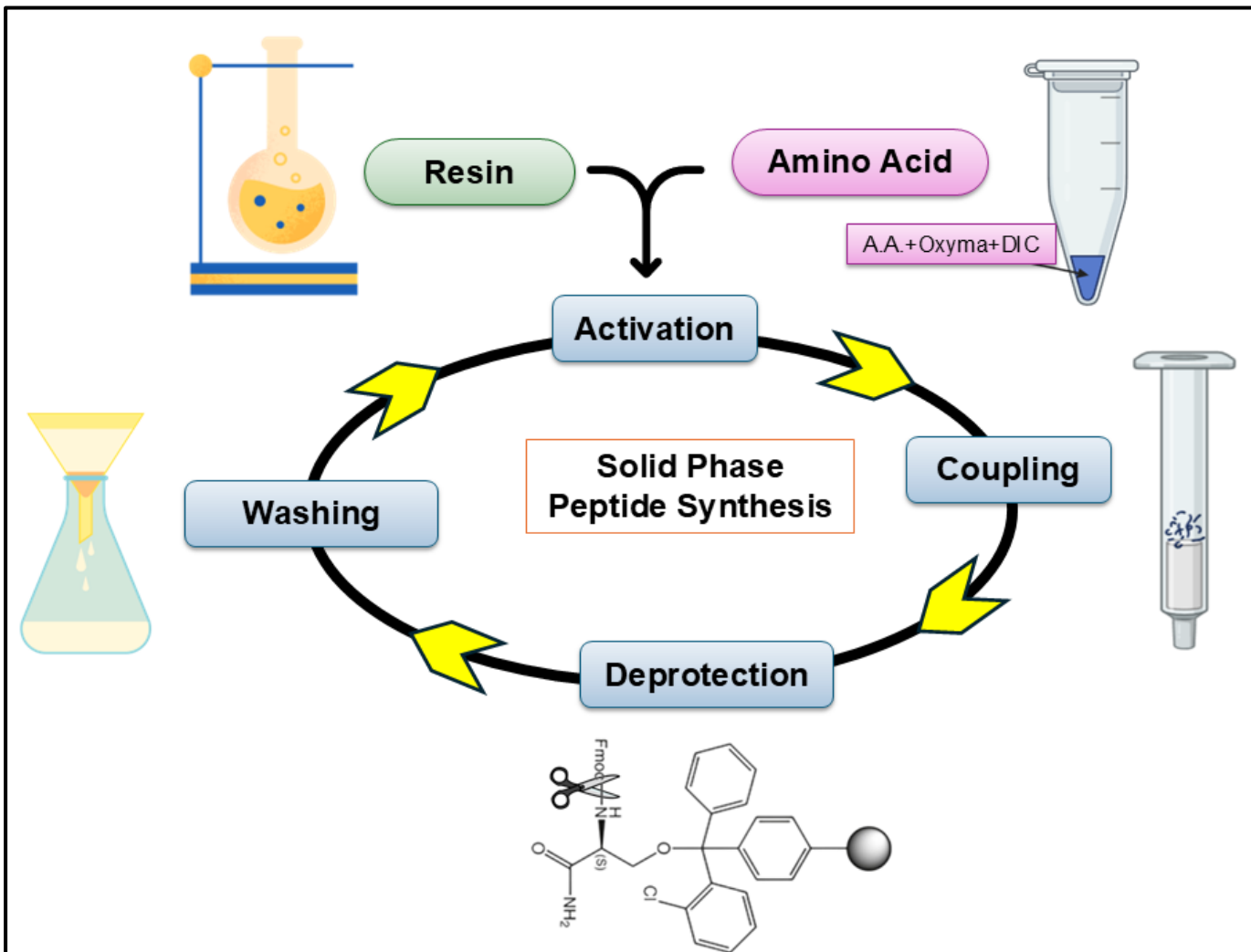
“While completely preventing breast cancer remains a challenge, but the casualties due to breast cancer can be reduced, through early detection.”

According to the Global Cancer Observatory (ICAR-WHO, 2022), breast cancer is the most prevalent cancer worldwide, with an estimated 660,000 deaths among females in 2022. India recorded the highest number of estimated breast cancer fatalities, with ~100,000 deaths in the same year.

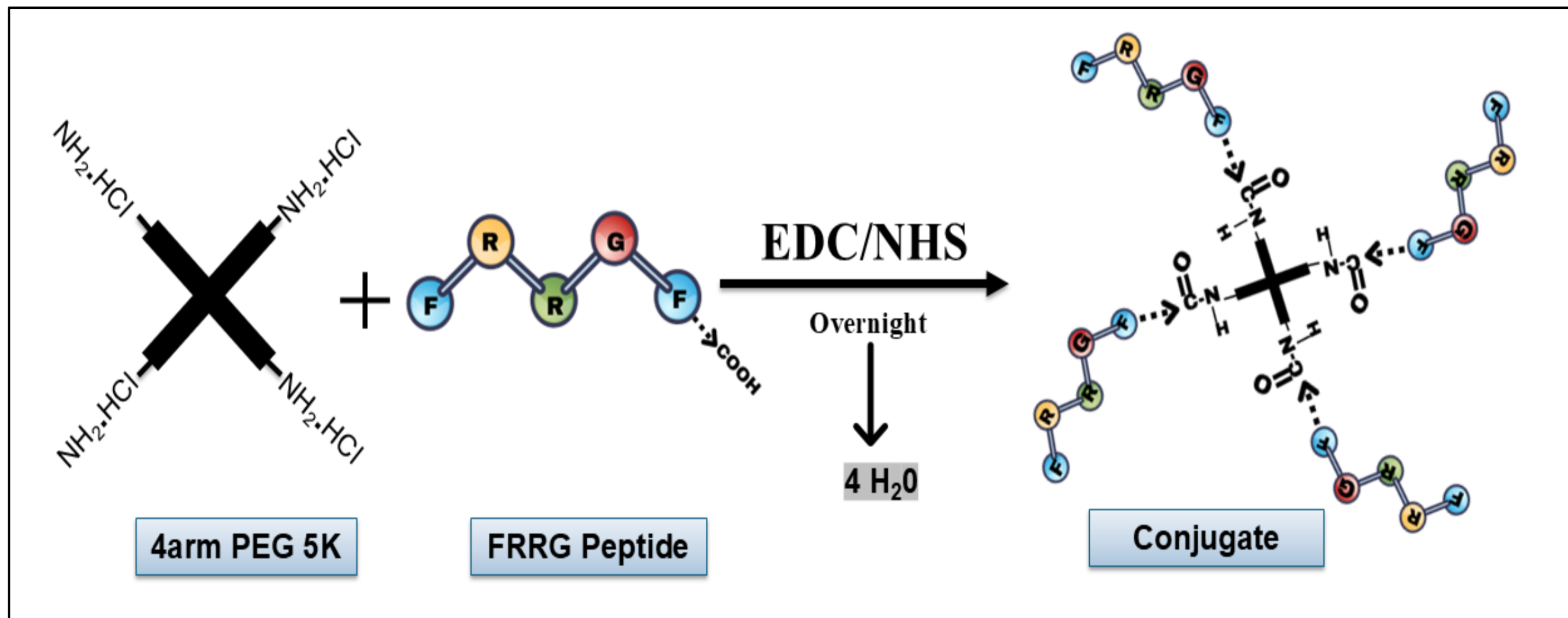


Objectives and Methodologies

Objective 1: Probe Designing



Objective 2: Probe Conjugation with Polymer and Nanoparticle Synthesis



Objective 3: Encapsulating the MRI Contrast agent with the Probe and Polymer conjugate - Future Work

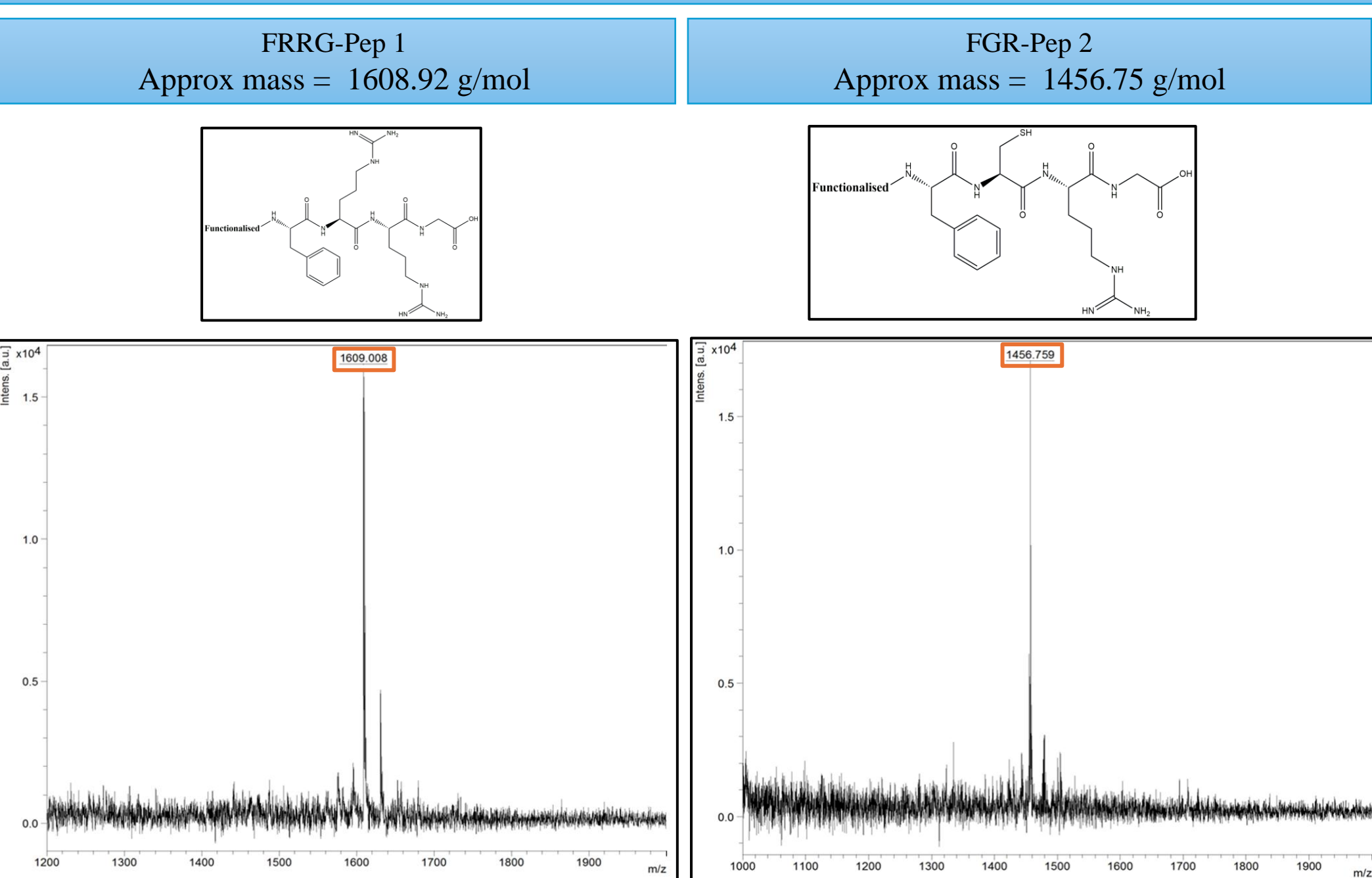
Studies

- Synthesis of probe for CSTB using Solid Phase Peptide Synthesis and characterized with LC-MS and MALDI.
- Conjugation of Biomarker and 4arm-PEG5K Using EDC/NHS Coupling: Analysis via MALDI, FTIR, UV-Vis Spectrophotometry, and Zeta Potential Measurement.
- Probe's Nanoparticles synthesis and its characterization with SEM, EDS and DLS.

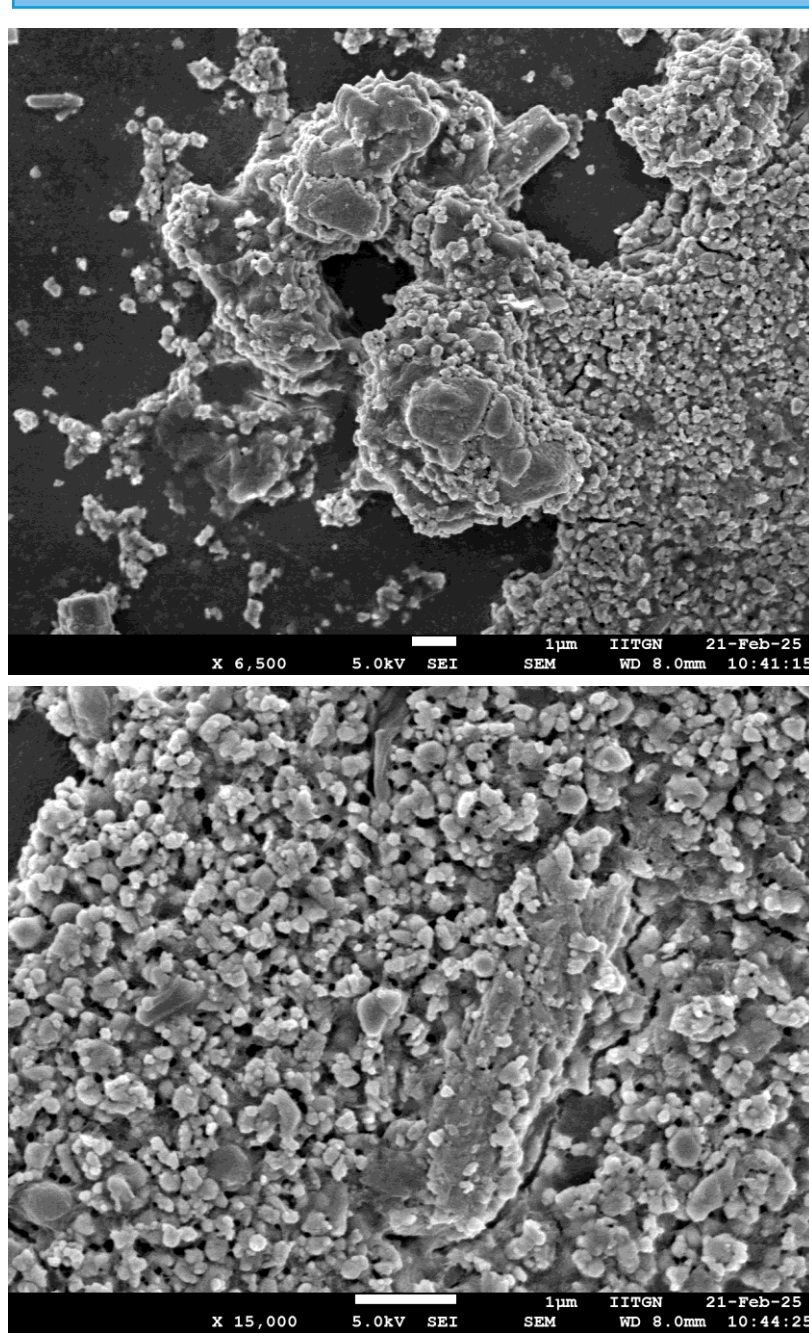


Results

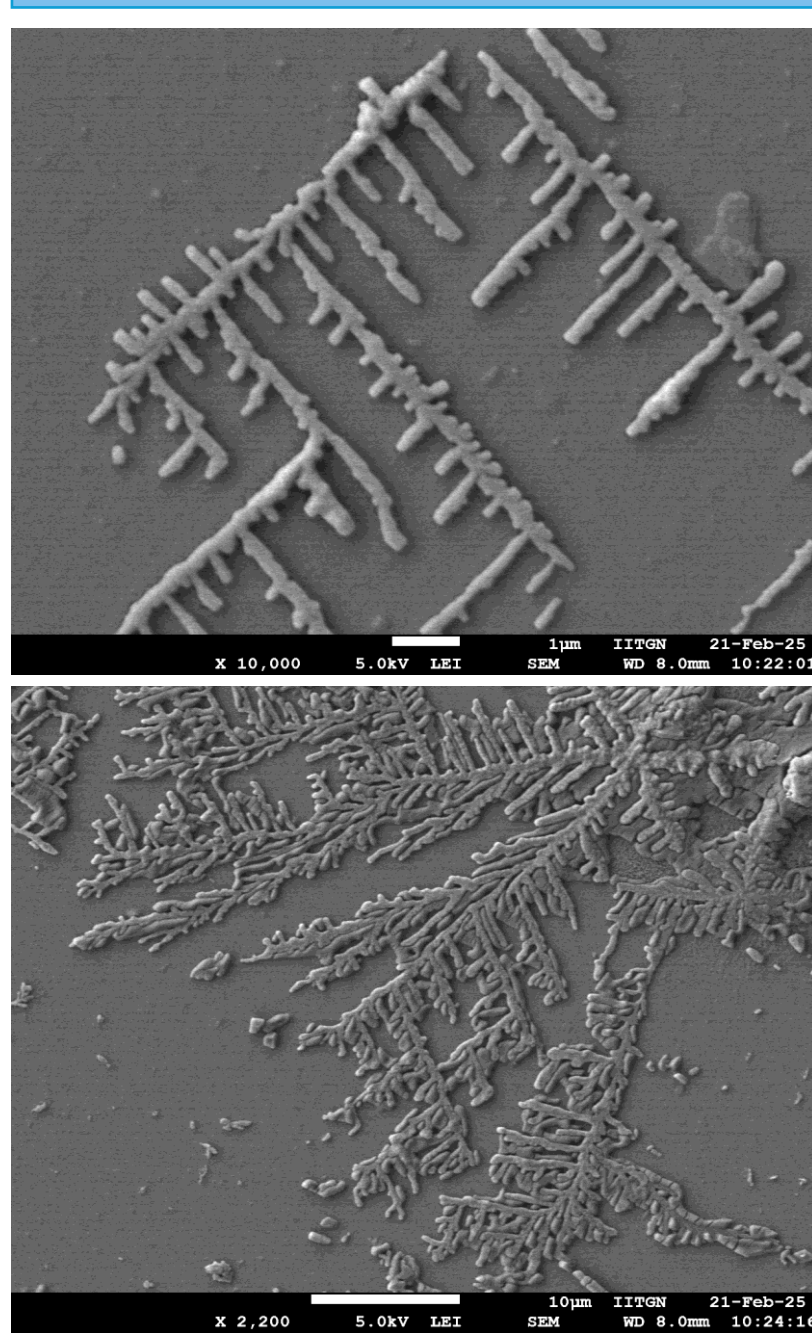
MALDI



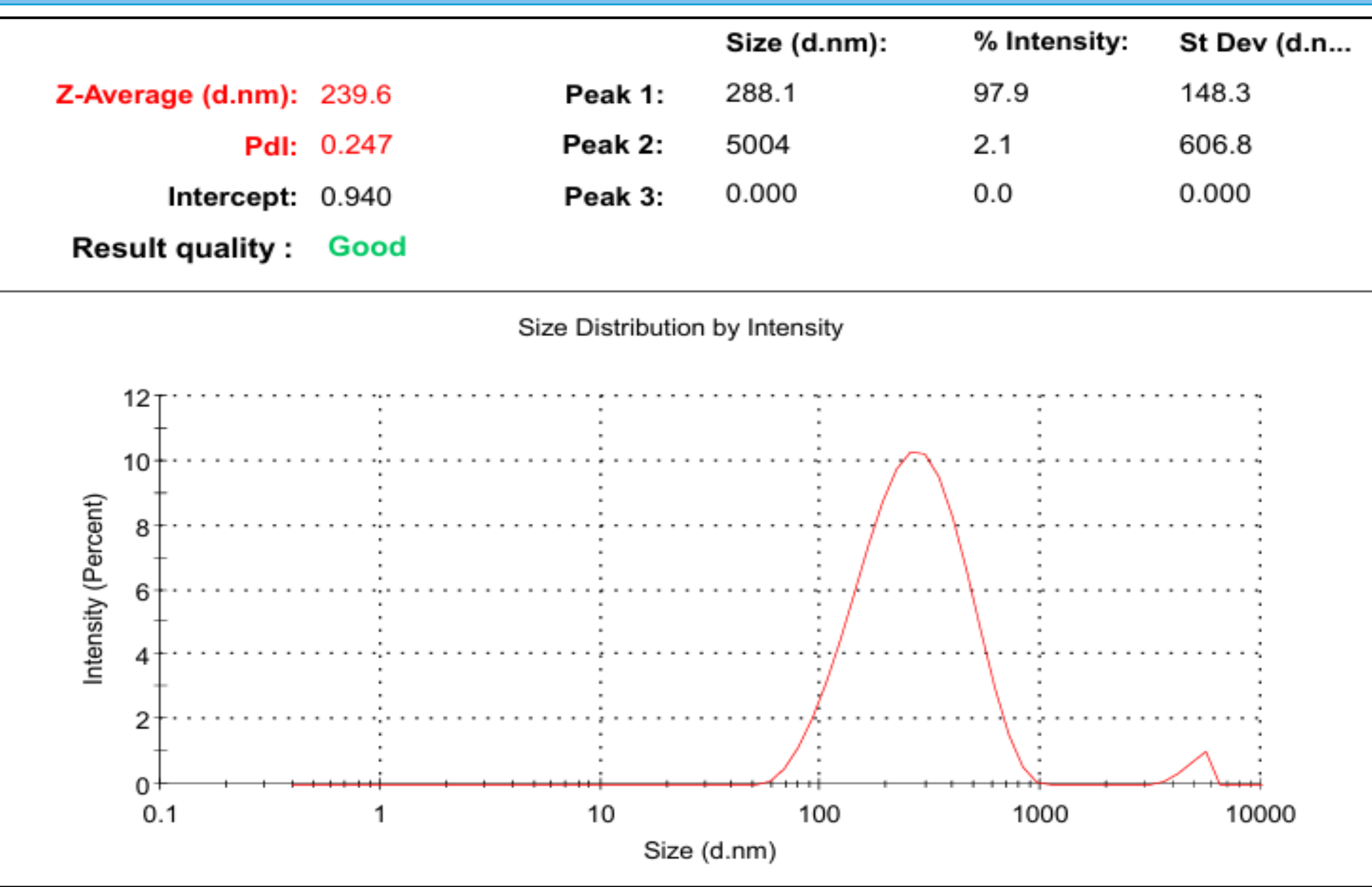
SEM of PEG5K



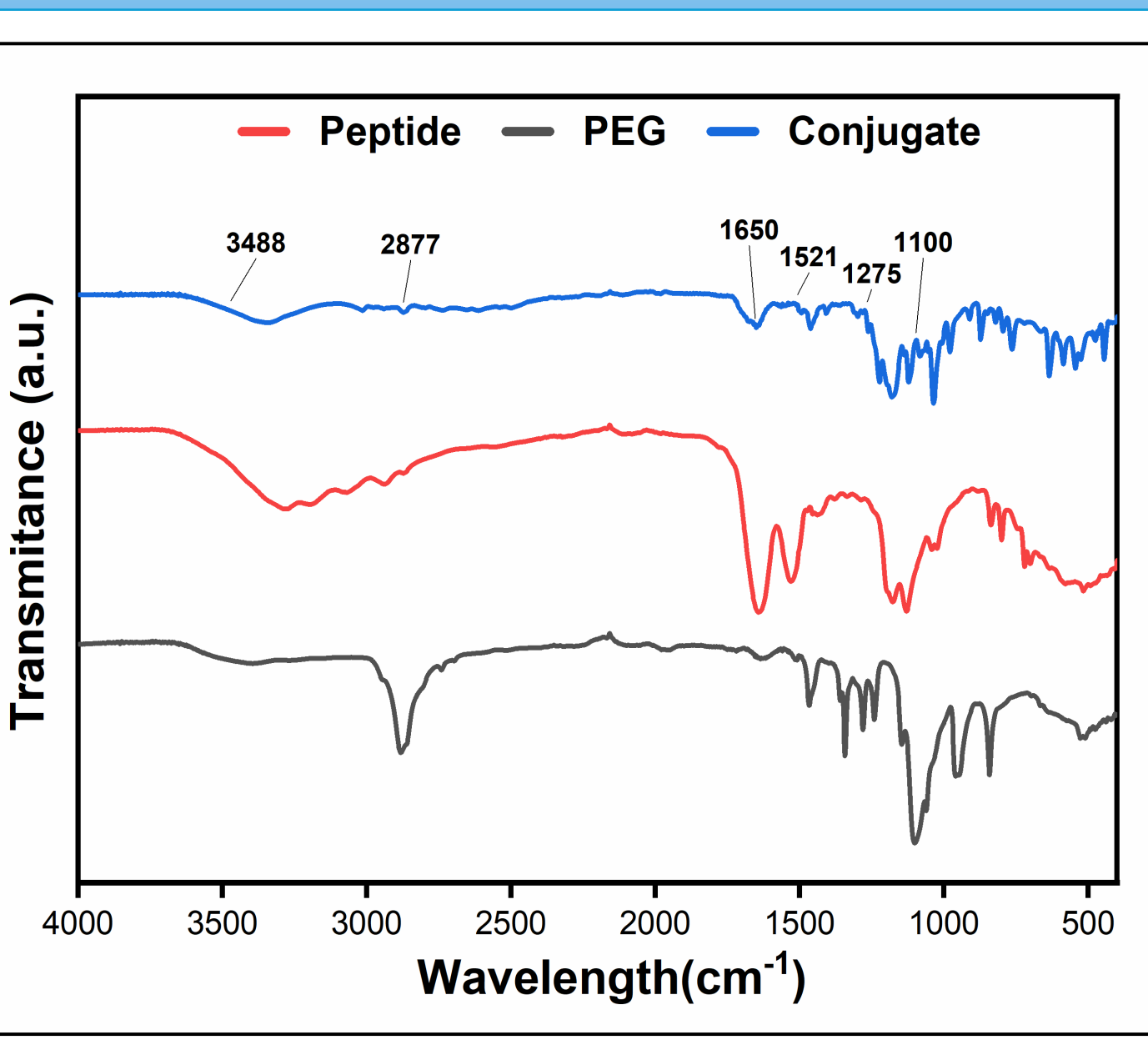
SEM of conjugate



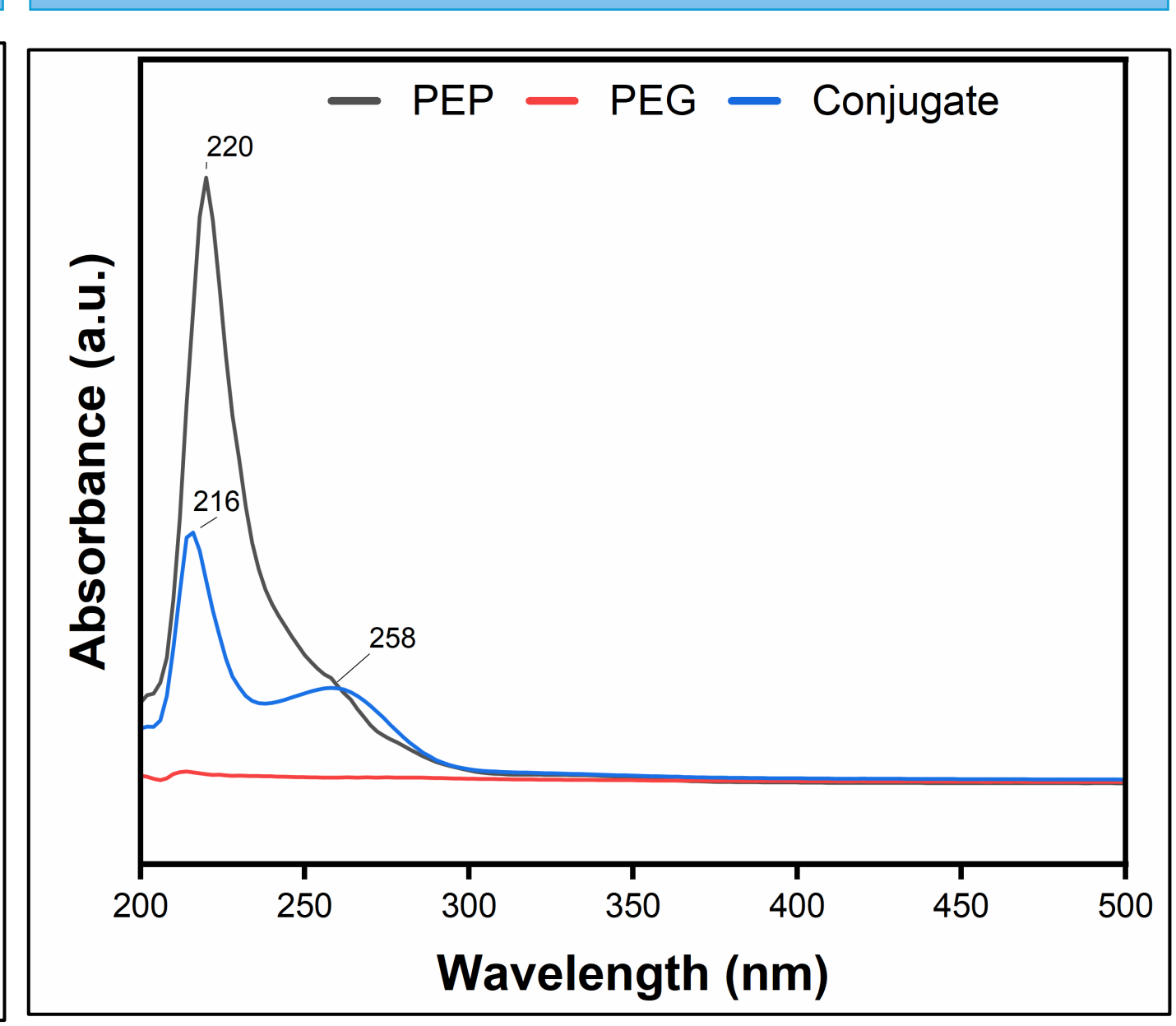
MRI Contrast Agent (SEM, EDS and DLS)



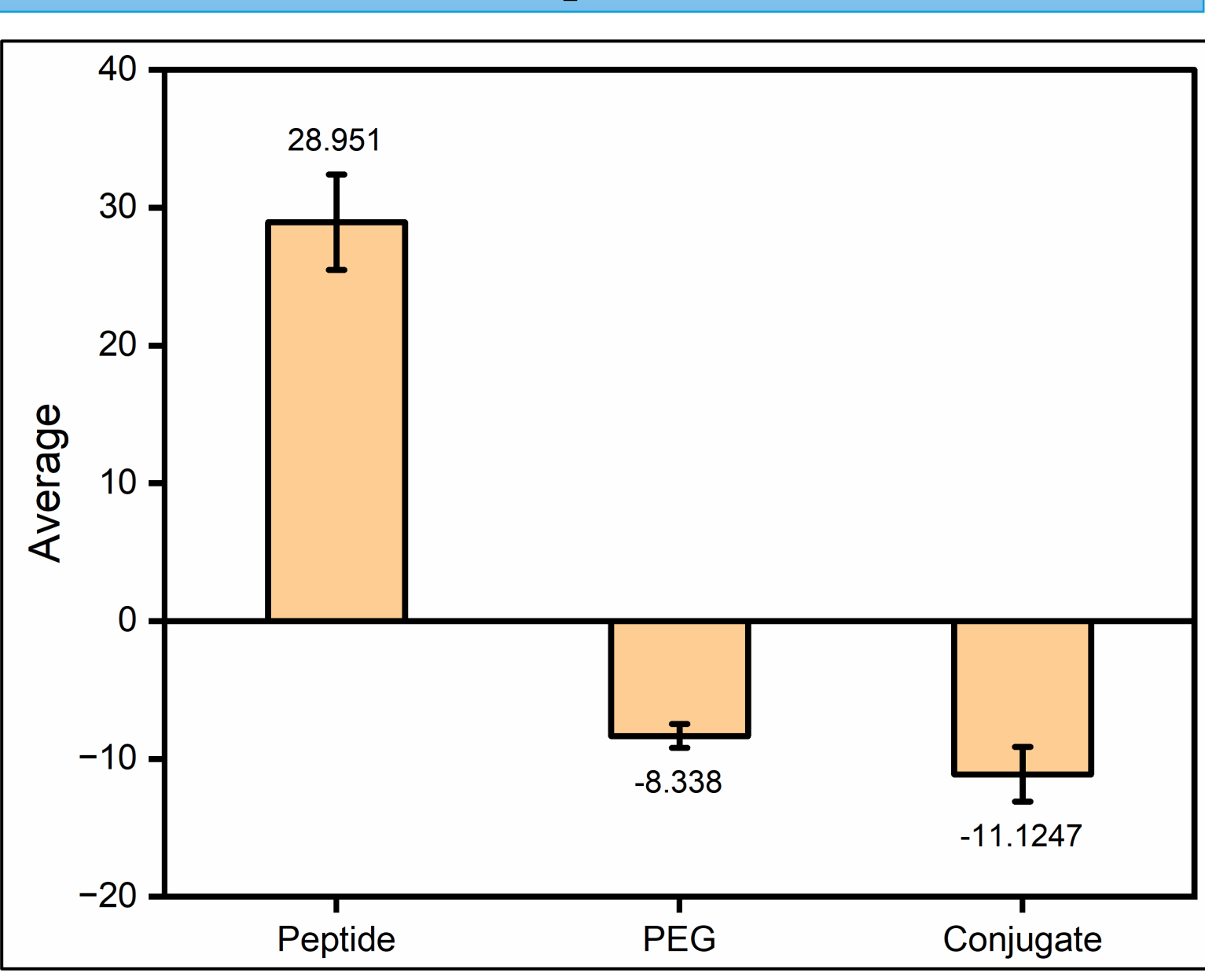
FT-IR



UV-Vis Spectrophotometer



Zeta-potential

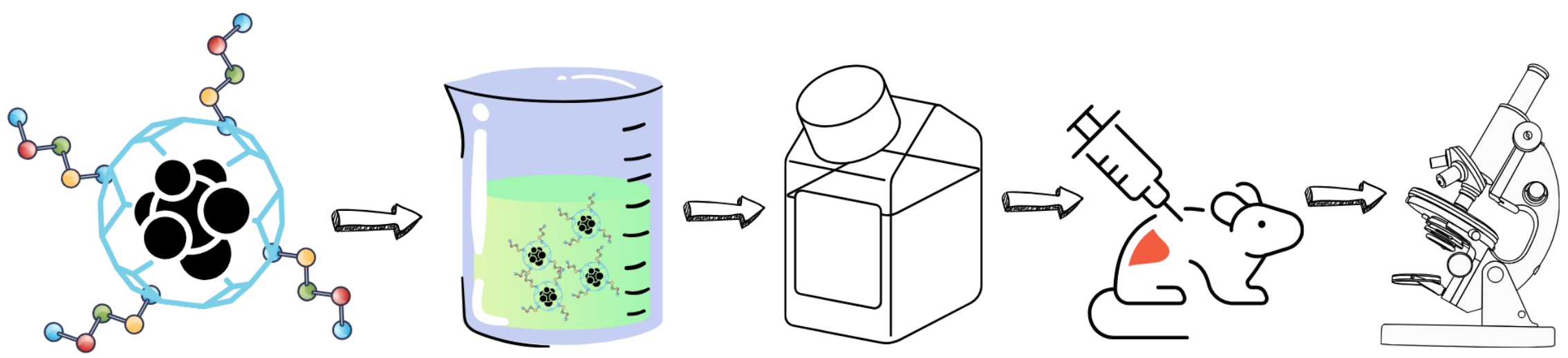


Future Studies

Synthesis of the probe complex for early breast cancer detection

Structural characterization

In-Vitro and In-Vivo study



Conclusion

- A 12-amino acid peptide probe targeting Cathepsin B and a control peptide were synthesized using solid-phase peptide synthesis and characterized via LC-MS and MALDI.
- The peptides were conjugated with functionalized 4arm-PEG5K using EDC/NHS coupling, confirmed through FTIR, UV-Vis spectrophotometry, and zeta potential measurements.
- MRI contrast agent nanoparticles were synthesized and characterized using SEM, EDS and DLS meeting physicochemical criteria, with the next step involving *In-vitro* and *In-vivo* assays for biomarker targeting efficiency.

Acknowledgment

- I would like to express my sincere gratitude to Dr. Sharad Gupta, Dr. Ramasamy Mayilmurugan and Dr. Dhiraj Bhatia, for their invaluable guidance and encouragement throughout this research.
- I also extend my thanks to my colleagues Shritilekha Dash, Sebika Panja, Abhijit Biswas, Ashish Kumar Samal, and SG Lab members for their guidance and support.
- Finally, I am grateful to the organizers of the Indian Peptide Symposium for providing this platform to present my research.

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