



Stimuli-responsive polydopamine nanoparticle disassembly for drug delivery applications

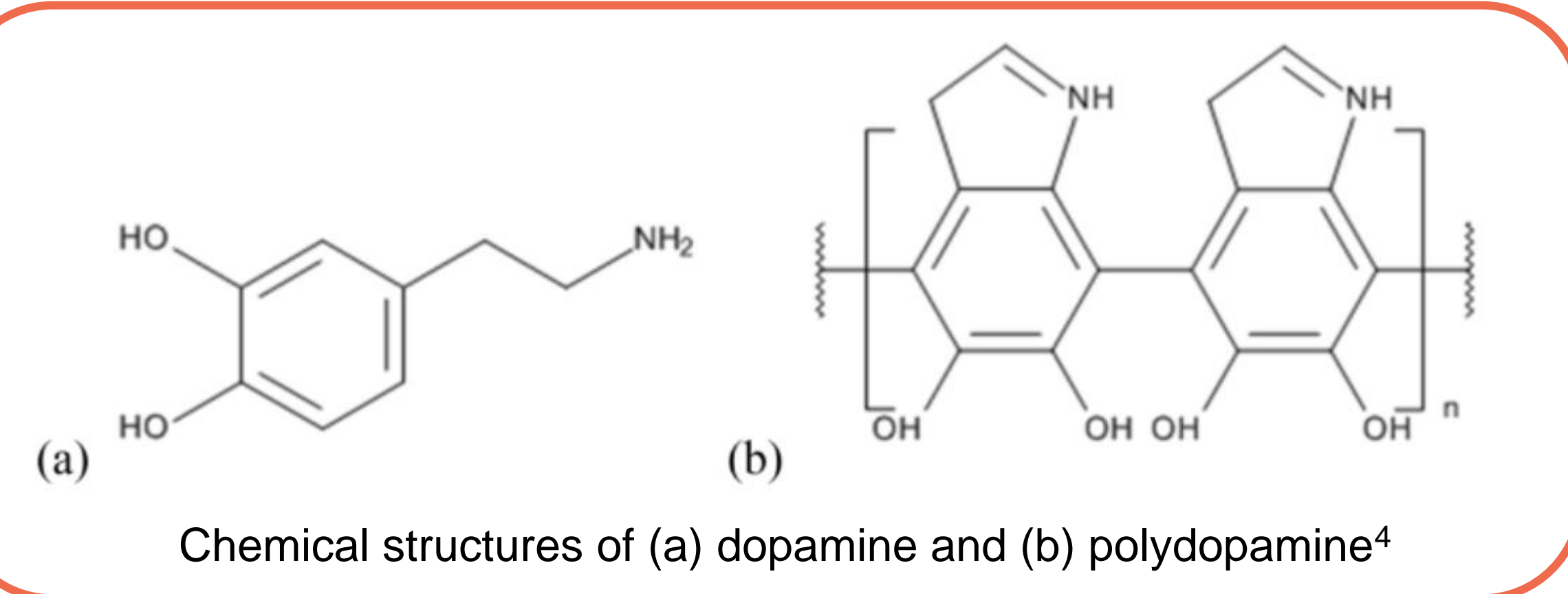
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Introduction

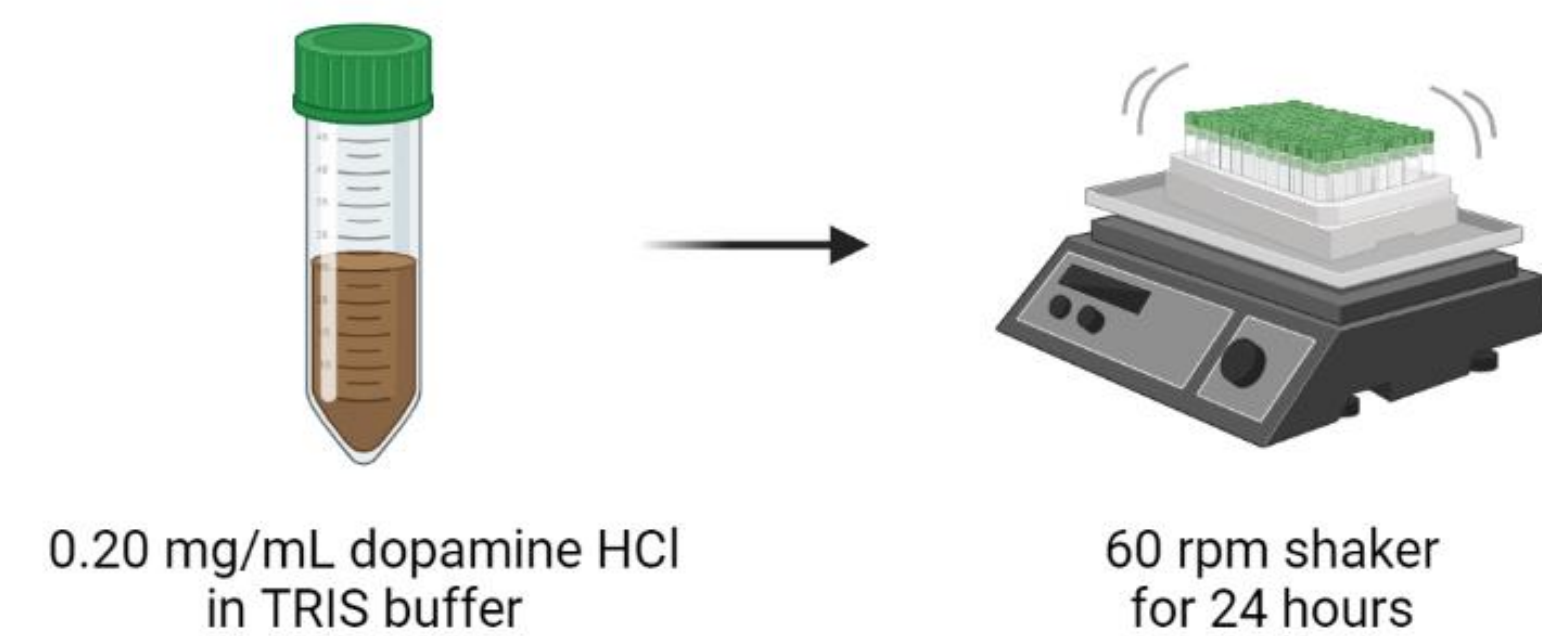
- Polydopamine (PDA) has attracted significant research attention in biomedical fields due to its simple synthesis and multifunctional properties.¹
- Its chemical structure allows for strong adhesion,² enabling drug loading.³
- Polydopamine's inherent biocompatibility¹ combined with its drug retention abilities³ make it ideal for applications in drug delivery systems.



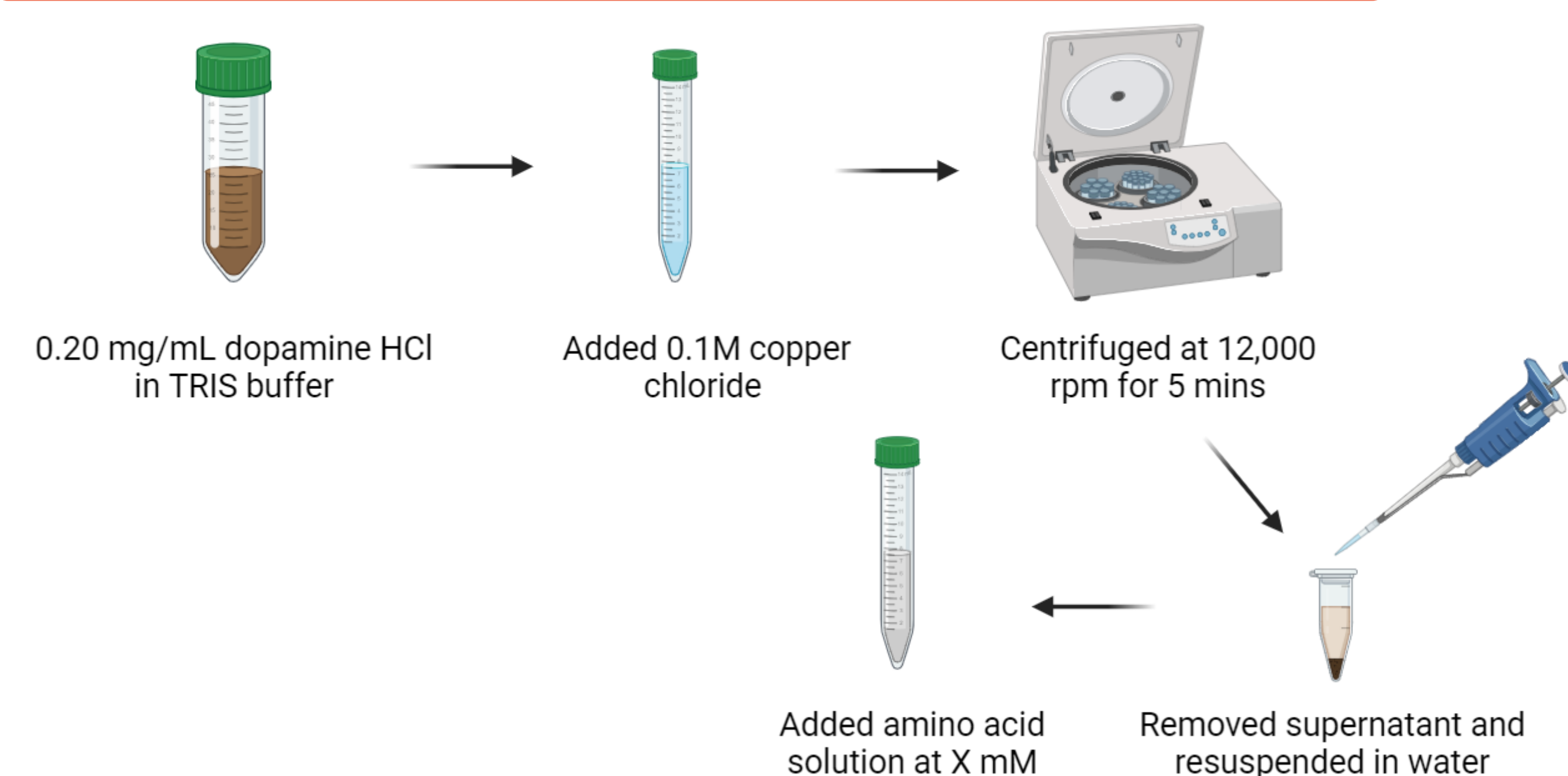
This project explores the disassembly of polydopamine nanoparticles using amino acids that can facilitate the on-demand release of drugs

Methods

Preparation of Polydopamine Oligomers



Assembly and Disassembly of Polydopamine Nanoparticles



Instruments

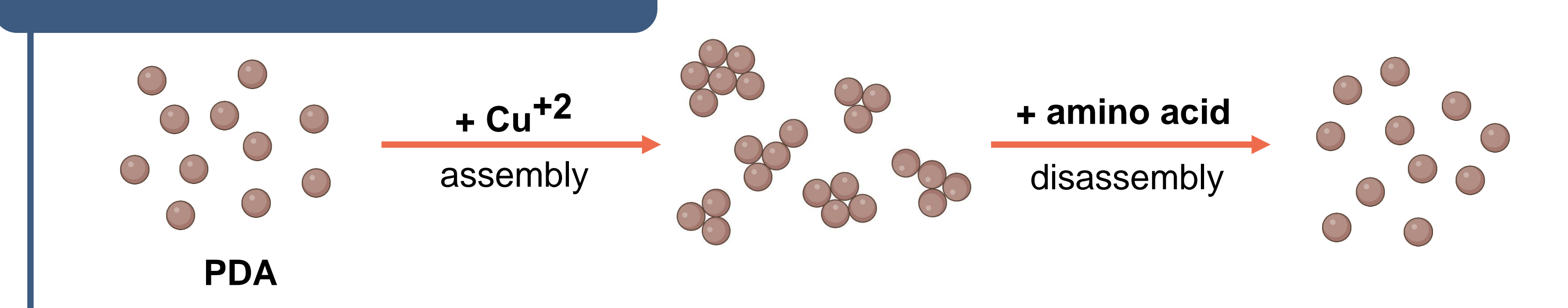


DLS measures the random particle movements to determine its size⁵

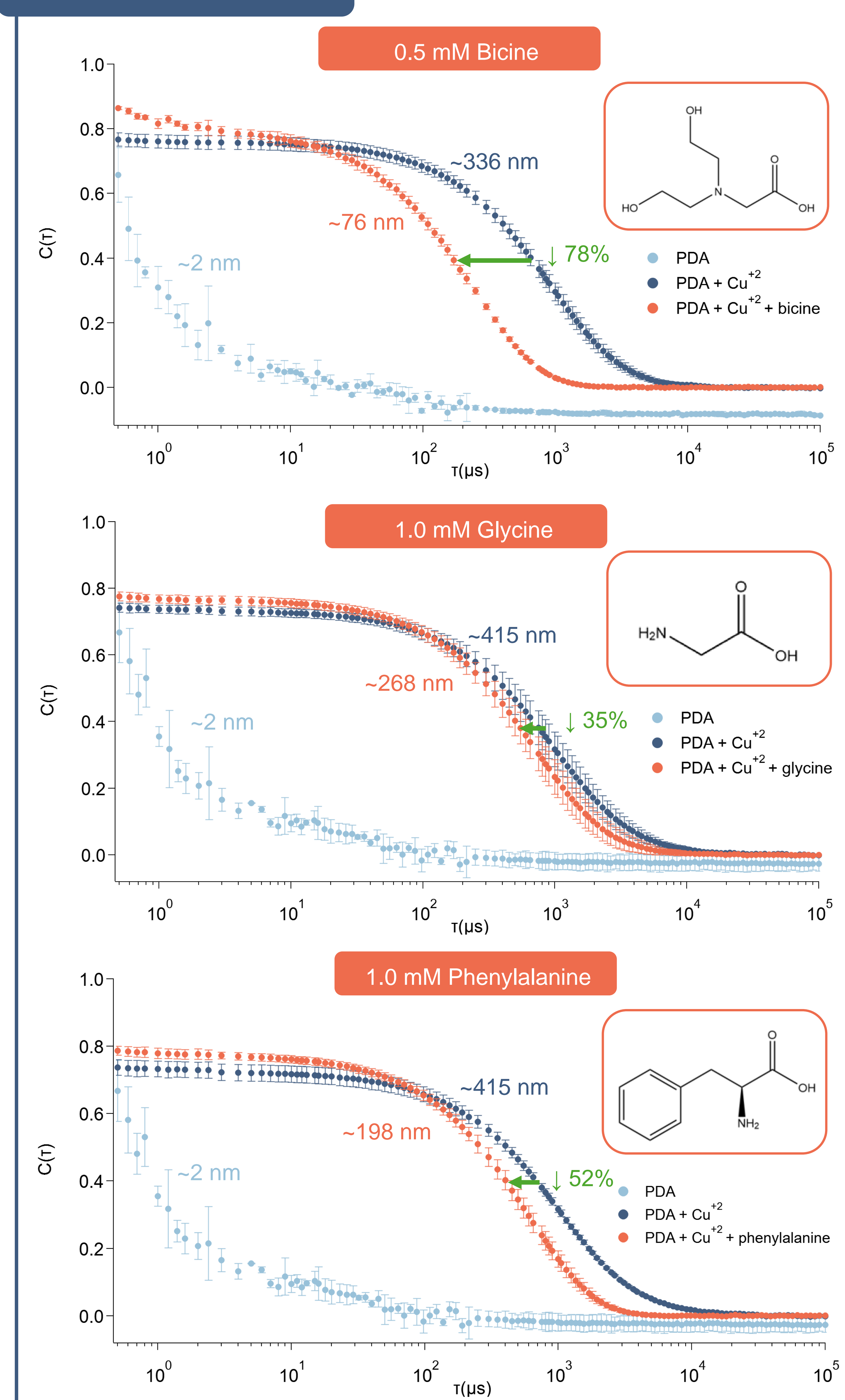
UV-vis measures the light absorbed at a wavelength range⁶

TEM uses transmitted electrons to image and size particles⁷

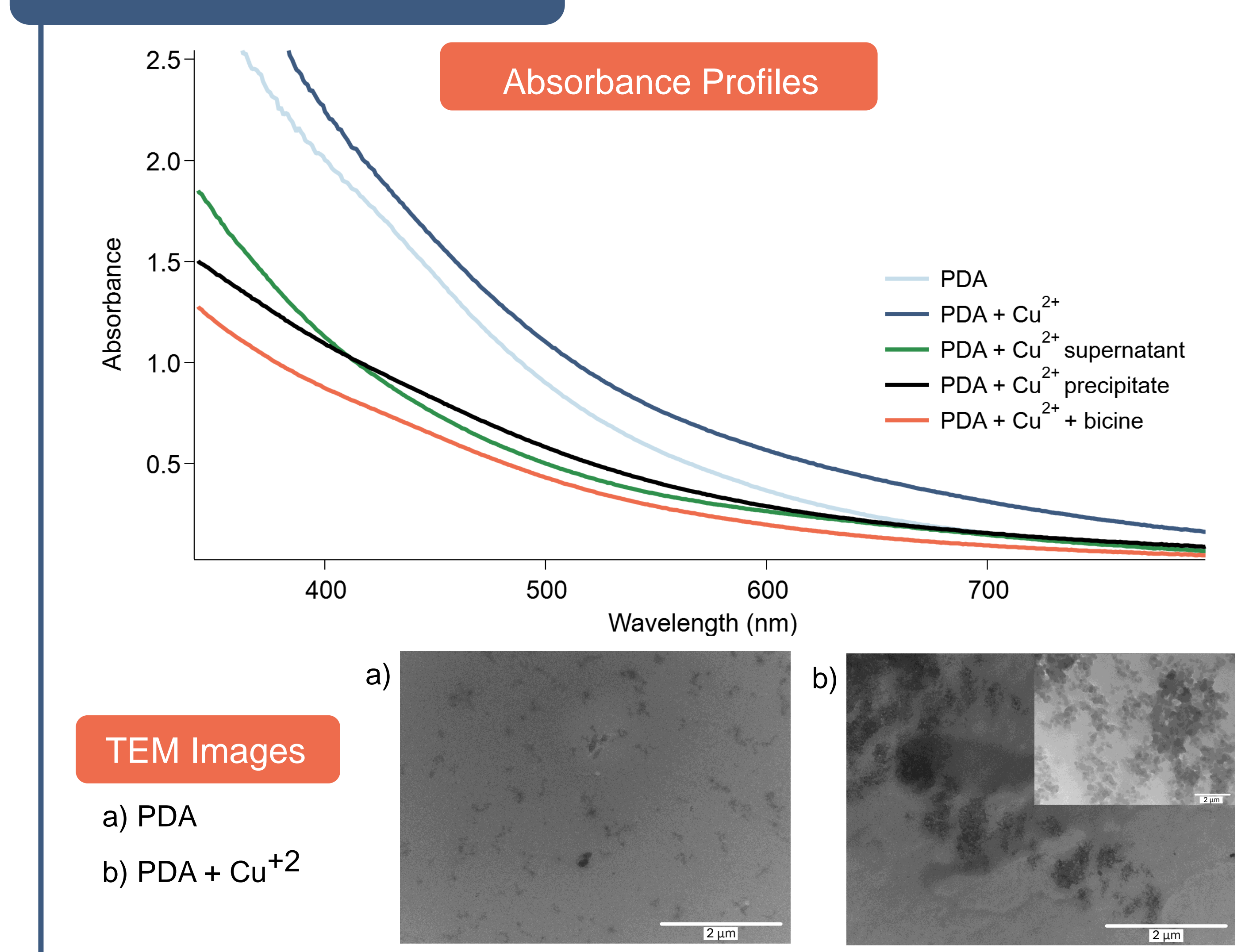
Schematic



Results



Results



Conclusion

Relative polydopamine disassembly strength of amino acids

Glycine < **Phenylalanine** < **Bicine**

- Hydrophilic
- Zwitterionic*

Phenylalanine

- Hydrophobic
- Zwitterionic*

Bicine

- Hydrophilic
- Zwitterionic*
- Glycine derivative

*Has a net 0 charge at neutral pH

Future Directions

- This project investigates using amino acids to disassemble nanoparticles formed from polydopamine and the divalent metal cation, Cu²⁺ – facilitating on-demand drug release.
- Further investigation will examine the disassembly process using amino acids with varying physical characteristics, such as those with acidic and basic side chains.

References

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