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Encapsulation Of Cell-free Bio-reactions: Delivering Dna-programmable Functions To Materials

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Encapsulation is a way to protect and co-localize molecules by packaging them into particles. Biological molecules and systems often benefit from stabilizing encapsulation treatments in examples ranging from therapeutics to foods. In this work, we seek to apply these benefits to cell-free protein synthesis reactions. Cell-free systems use crude cell extracts or purified enzymes to enact a variety of biological functions outside of living cells, with wide-ranging applications to benefit the warfighter, including enzymatic decontamination, on-demand biosynthesis of therapeutics, or eye-readable biosensors. Functions can be dictated by simply adding DNA instructions. Cell-free systems may be freeze-dried for shelf-stability and already display remarkable tolerance to integration with materials like plastic polymers or porous materials like paper, but encapsulation may push this stability even further.

Here we describe ongoing work to load cell-free reactions into particles of various types, including hydrogel resins, diblock copolymer vesicles, and microfluidic droplets. We show advantages and disadvantages of each format and propose methods to produce core-shell designs. We hypothesize these strategies will improve control over humidity exposure, re-hydration rate, and expand tolerance to material processing steps. This technology has the potential to enable integration of DNA-programmable functionality into the materials that make up warfighter equipment to introduce new capabilities like self-decontamination or reduce burden through built-in contaminant sensing.

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