Self-Assembled, Dilution-Responsive Hydrogels for Enhanced Thermal Stability of Insulin Biopharmaceuticals

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ABSTRACT: Biotherapeutics currently dominate the landscape of new drugs because of their exceptional potency and selectivity. Yet, the intricate molecular structures that give rise to these beneficial qualities also render them unstable in formulation. Hydrogels have shown potential as stabilizing excipients for biotherapeutic drugs, providing protection against harsh thermal conditions experienced during distribution and storage. In this work, we report the utilization of a cellulose-based supramolecular hydrogel formed from polymer–nanoparticle (PNP) interactions to encapsulate and stabilize insulin, an important biotherapeutic used widely to treat diabetes. Encapsulation of insulin in these hydrogels prevents insulin aggregation and maintains insulin bioactivity through stressed aging conditions of elevated temperature and continuous agitation for over 28 days. Further, insulin can be easily recovered by dilution of these hydrogels for administration at the point of care. This supramolecular hydrogel system shows promise as a stabilizing excipient to reduce the cold chain dependence of insulin and other biotherapeutics.

KEYWORDS: supramolecular, drug delivery, biotherapeutic, cold chain, biomaterials, hydrogel

INTRODUCTION

Biotherapeutics have grown to be a dominant portion of the pharmaceutical market over the past two decades. In 2018, eight of the top ten grossing drug products globally were biotherapeutics. While the complex and intricate structures of these macromolecular drugs result in pharmaceutical products with exceptional potency and selectivity, it also makes them inherently susceptible to loss of activity due to thermal denaturation and aggregation. Exposure to high, or even ambient, temperatures during transportation, storage, and use can cause partial or complete loss of bioactivity. To prevent loss of activity, most biotherapeutics require refrigerated shipping and storage throughout the supply chain, referred to as the “cold chain”, which reduces accessibility to regions of the world with limited infrastructure. Additionally, cold chain logistics are costly, with an estimated $15.7 billion spent globally by biopharmaceutical companies on cold chain logistics alone in 2019. Improved thermal stability of biotherapeutics would enable increased global access to critical drugs and reduce storage and transportation costs.

There are many approaches to improving the stability of biotherapeutic formulations, but typical strategies include modifying the drug molecule itself, altering its physical state, or formulating with excipients—often in combination—such as salts, amino acids, polymers, and surfactants. One prevalent method is to use protein engineering to design biotherapeutic analogues. While this has been successful, it carries the risk of altered drug potency or immunogenicity. Another commonly used stabilization technique is lyophilization, or freeze-drying, of the drug into a dry powder prior to shipping and long-term storage. Unfortunately, lyophilization is not ideal for many drug products because the drastic changes in temperature, pressure, and hydration state during processing can result in activity loss. Liquid formulations of biotherapeutics incorporate stabilizing excipients however, many currently used pharmaceutical excipients were developed for use with small molecule drugs and were not optimized to stabilize proteins. Therefore, there is a need to develop next-generation excipients specifically designed to enhance the stability of biotherapeutics, especially for parenteral formulations. The drug delivery field has made considerable advances in the formulation of biotherapeutics in soft materials to modulate drug pharmacokinetics; this body of knowledge and materials could be leveraged to advance materials for biostability as well.

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Recently, hydrogels have been utilized to improve the thermal stability of biotherapeutics. Nanogels, nanoparticles, microparticles, and hydrogel films are forms of hydrogel excipients used to stabilize the protein of interest and can reduce cold chain dependence.20−23 Hydrogels are promising candidates for biotherapeutic stabilization because they can maintain the proteins in their native aqueous environment while promoting stabilizing interactions between the hydrogel polymer backbone and protein. A poly(ethylene glycol) (PEG)-based covalent hydrogel network with a photolabile moiety has been reported to stabilize biotherapeutic cargo at elevated temperatures.26 Upon exposure to ultraviolet light, the hydrogel network backbone degraded to afford biotherapeutic recovery. While the system effectively stabilized the selected enzymes, the presence of degraded hydrogel material may prove to be an obstacle for clinical translation as the in vivo biocompatibility has not been assessed. Both trehalose-based hydrogels and hydrogels with trehalose side chain modifications have also shown utility as thermal stabilizing excipients for a variety of proteins, including insulin.27−29 Unfortunately, these systems have not been shown to be injectable or degradable in vivo. To use hydrogels as stabilizing excipients in parenteral biotherapeutic formulations, there is still a need to design materials that allow mild encapsulation, are injectable, are biocompatible in vivo, and easily release the drug without unintentionally altering drug pharmacokinetics.

We have developed a supramolecular hydrogel platform, based upon polymer–nanoparticle (PNP) interactions between hydrophobically modified cellulose–derived polymers and core–shell polymeric nanoparticles, as an encapsulating excipient (Figure 1a).30−33 These supramolecular hydrogels have been used for a variety of biomedical applications and are biocompatible, injectable, and readily tunable in mechanical properties as characterized by rheology.34−37 We can readily encapsulate biotherapeutic cargo into these hydrogels under mild conditions by simply mixing the drug into the bulk material. Further, the hydrogel can be diluted to disrupt the network and yield a drug-containing solution for administration, as the various hydrogel components are Generally Recognized as Safe (GRAS-listed) by the FDA and are non-toxic. These qualities indicate that our PNP hydrogel network is a promising candidate for a translatable stabilizing excipient for biotherapeutics.

In this study, we demonstrate that PNP hydrogels can stabilize insulin against thermal aggregation and denaturation. Insulin is a clinically relevant biotherapeutic that is required by over 150 million patients living with diabetes worldwide, but suffers from thermal instability and is highly prone to aggregate into insoluble, inactive and immunogenic amyloid fibrils when it is exposed to increased temperatures or agitation.30−33 Stabilized insulin capable of maintaining drug integrity in environmental conditions could increase access to this drug in regions of the world such as Africa and the Western Pacific where diabetes prevalence is increasing but reliable cold chain storage and transportation is challenging.40,41 We investigate the thermal stability of insulin encapsulated in these hydrogels with stressed aging experiments under relevant simulated environmental conditions to demonstrate the possibility for cold chain independence (Figure 1b). We show that these supramolecular hydrogels are promising for use as thermal stabilization agents in insulin formulations, and potentially other parenteral biotherapeutic formulations as well.

**MATERIALS AND METHODS**

**Materials.** Racemic lactide (3,6-dimethyl-1,4-dioxane-2,5-dione, 99%, Sigma-Aldrich) was recrystallized twice from dry ethyl acetate to remove impurities, with sodium sulfate included as a desiccant prior to the first recrystallization. Dichloromethane (DCM) was dried immediately before use by cryogenic distillation. Recombinant human insulin was purchased from Gibco as a lyophilized powder containing zinc and then reconstituted. Humulin R (100 U/ml, 3.47 mg/mL, Eli

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**Figure 1.** (a) Supramolecular polymer–nanoparticle (PNP) hydrogels composed of (hydroxypropyl)methyl cellulose (HPMC) with dodecyl side chain modifications as the network polymer that is physically cross-linked by bridging interactions with poly(ethylene glycol)-block-poly(lactic acid) (PEG-b-PLA) diblock copolymer nanoparticles. Biotherapeutics, such as insulin (shown), can be encapsulated in these hydrogels during precursor mixing. (b) Schematic indicating the desired performance of our thermally stabilizing supramolecular hydrogel excipient.
Synthesis of PEG-b-PLA block copolymers. Poly(ethylene glycol)-block-poly(actic acid) (PEG-b-PLA) block copolymers were synthesized by ring-opening polymerization of lactide onto a PEG macromer, as described previously.\textsuperscript{30} Dry PEG methyl ether (avg. $M_n = 5000$ g/mol, 0.5 g, 0.1 mmol) and 1,6-diazabicyclo[5.4.0]undec-7-ene (DBU, 98%, 15 μL, 0.1 mmol) were dissolved together in dry DCM (1.5 mL); twice recrystallized racemic lactide (2 g, 13.9 mmol) was dissolved separately in dry DCM (9 mL). The PEG/DBU solution was quickly added to the lactide solution under nitrogen atmosphere and stirred rapidly for 8 min then quenched with dilute acetic acid in acetone (100 μL). The polymer was recovered by precipitating twice from a 50:50 diethyl ether: hexanes solution and dehydrated by visual inspection. Trehalose, a common excipient, was added to the resulting NPs to achieve final concentrations of 10 mg/mL, except 2:10 hydrogels were loaded at 25 mg/mL. All NPs were subjected to 50 cycles (1 cycle = complete transfer of $20\mu L$ per well, $n = 4$), sealed, and subjected to stressed aging conditions.

In Vitro Insulin Fibrillation Assay. Insulin fibrillation was characterized in vitro by thioflavin T (ThT) assay. In the presence of insulin amyloid fibrils, ThT produces a strong fluorescence signal, yielding a binary assay result indicating the presence or absence of fibrils (Figure S5).\textsuperscript{34} Insulin-loaded hydrogels and assay controls, all preformulated with ThT, were loaded into 96-well black plates (200 μL per well, $n = 4$), sealed, and subjected to stressed aging conditions. To simulate hot ambient conditions, plated samples were subjected to continuous linear shaking (410 rpm, 5 mm) at 37 °C in a BioTek Synergy H1 Hybrid Multi-Mode Plate Reader. Data was acquired at 20 min intervals by excitation at 450 nm and collection of emitted signal at 482 nm. To simulate harsh environmental conditions, plated samples were subjected to 50 °C and continuous agitation at 150 rpm on a shaker plate inside a humidified incubator. Fluorescence readings were acquired daily (read at 10 min intervals for 1 h at 45 °C) for the first week, and every 2–3 days until the end point of the study. Aging studies were terminated when the samples appeared to be significantly dehydrated by visual inspection. Trehalose, a common excipient, was tested at 2, 6, and 12 wt % trehalose (chosen to match total polymer solid content in certain hydrogel formulations) with 10 mg/mL insulin at 37 °C. Humulin (3.47 mg/mL), a standard low-cost commercially available formulation, was tested at 37 °C. Stability data is shown as mean fluorescence signal over time ($n = 4$). Time to aggregation was determined by defining an arbitrary signal threshold (750 000 arbitrary fluorescence units, AFU).

Supramolecular PNP Hydrogel Formulation and Drug Loading. Supramolecular PNP hydrogels were formulated by combining the HPMC-C<sub>12</sub> stock solution and PEG-PLA NP stock solution diluted to the desired concentration via syringe mixing as previously described.\textsuperscript{34} PNP hydrogel formulations are designated by HPMC-C<sub>x</sub>-PEG-b-PLA, with $x$ wt % HPMC-C<sub>x</sub> and y wt % PEG-b-PLA. The total formulation mass, with the remainder comprised of PBS or drug cargo solution. HPMC-C<sub>x</sub> stock solution (400 mg) was loaded into a 3 mL luer lock syringe. NP stock solution (600 μL) was combined with PBS (200 μL) and loaded into a 3 mL luer lock syringe. The HPMC-C<sub>12</sub>-filled syringe was connected to a female luer × female luer elbow fitting, and the solution was pushed through the luer to eliminate air in the connection before attaching the NP-filled syringe. Insulin stock solution (40 mg/mL) was prepared from lyophilized powder by first dissolving in 0.12 M hydrochloric acid and then neutralizing with 0.13 M sodium hydroxide. Concentrated phosphate buffer (0.45 M), concentrated sodium chloride (135 mg/mL), and ultrapure water were added to achieve final concentrations matching PBS (10 mM phosphate, 0.9% NaCl). Neutral pH was verified using pH strips. Insulin was loaded into the hydrogels to a final concentration of 10 mg/mL except 2:10 hydrogels were loaded at 6.7 mg/mL. ThT (640 μg/mL, 2 mM stock solution in PBS) was added to a final concentration of 25 μM for all samples (15 μL per 1.2 mL sample volume).

Rheological Characterization of PNP Hydrogels. Rheological measurements were performed on a TA Instruments Discovery Hybrid Rheometer (DHR-2) with a 40 mm 2° cone geometry with a fixed gap of 51 μm. Amplitude sweeps were measured at a fixed frequency of 1 s<sup>−1</sup> (Figure S5). Flow sweeps were measured at shear rates from 100–0.01 s<sup>−1</sup>. Frequency sweeps were measured at a constant torque of 5 μN m from 0.0628–628 rad s<sup>−1</sup>. To measure the changes in hydrogel rheological properties upon dilution, the hydrogel sample was recovered from the rheometer and serially diluted with PBS using the elbow mixing method described for preparing the supramolecular PNP hydrogel formulations.

PEG-b-PLA Nanoparticle (NP) Synthesis by Nanoprecipitation. To form core–shell NPs, PEG–PLA (50 mg) was dissolved in a 75:25 acetonitrile:DMSO solution (1 mL) and added dropwise to the heated reaction vessel followed by 3 drops of acetone and purified by dialysis (MWCO 3500 kDa) for 4 days (Figure S3). Lyophilized HPMC-C<sub>12</sub> was redissolved in phosphate buffered saline (PBS) as a 6 wt % stock solution. Representative 4<sup>°</sup>H NMR analysis has been previously published.\textsuperscript{35}
of unsupplemented DMEM and starved in 100 μL of unsupplemented DMEM overnight. Insulin test samples were formulated by serial dilution in unsupplemented DMEM to the desired concentrations for the dose–response curve. The media was then removed and the cells were stimulated with 100 μL of each insulin test sample for 30 min while incubating at 37 °C. Cells were washed twice with 100 μL of cold 1X Tris-buffered saline before adding 100 μL of lysis buffer to each well and shaking for at least 10 min at room temperature to fully lyse cells. Thirty microliters of lysate was transferred to a 96-well white half-area plate for each assay. Each sample was assayed to lye cells. Thirty microliters of lysate was transferred to a 96-well white half-area plate for each assay. Each sample was assayed to determine both [pAKT] and [total AKT1]. Assays were completed by incubating at 37 °C stressed aging in vitro insulin fibrillation plate assay, n = 1 cellular assay replicates was plotted as the ratio of [pAKT]/[total AKT1] and normalized to a scale of 0 to 1. logEC50 values were determined using the log(agonist) versus response variable slope (four parameters) regression in GraphPad Prism 8.0.2.

**Statistical Analysis Methods.** Statistical significance was assessed using the Extra sum-of-squares F-test to determine if the logEC50 differed between data sets. Data sets were compared in pairs and Bonferroni posthoc tests were used to adjust for multiple comparisons (alpha = 0.01).

## RESULTS AND DISCUSSION

**PNP Hydrogel Formulation Effects on Rheology.** The ideal hydrogel drug product for insulin stabilization and delivery would be able to encapsulate the drug in a mild manner, inject easily for processability, maintain insulin stability in the vial, and readily disperse for facile subcutaneous delivery. In terms of rheological properties, the hydrogel should be shear-thinning and mechanically tunable. Polymer–nanoparticle (PNP) hydrogels are formed by bridging of hydrophobically modified (hydroxypropyl)methyl cellulose (HPMC) polymers between poly(ethylene glycol)-block-poly(lactic acid) (PEG-b-PLA) diblock copolymer NPs, whereby the HPMC polymers exhibit dynamic, multivalent interactions with the surface of the NPs (Figure 1a). These PNP hydrogels have previously been shown to exhibit both shear-thinning and self-healing properties that enable facile processing and administration by injection or spraying. In addition, the viscoelastic properties of these materials have been shown to be easily tunable by varying the PNP hydrogel formulation.

In the present work, a set of PNP hydrogels with distinct rheological properties were formulated by varying the total polymer and NP content, denoted as P:NP, where P = wt % polymer and NP = wt % nanoparticles (neat polymer solutions denoted X:0). Viscoelastic properties (storage modulus, G′, and loss modulus, G″) of the resulting hydrogels were characterized by frequency sweep within the linear viscoelastic regime and were consistent with those observed previously. Increasing NP content from 1 wt % to 10 wt % resulted in an increase in both the storage and loss modulus values. 1:5, 1:10, and 2:10 PNP hydrogels exhibited solid-like behavior (G′ > G″) across the entire frequency range; 1:1, 1:2, 2:1, and 2:2 hydrogel formulations were non-gelling (G″ > G′), though G′ and G″ approach a crossover point near 100 rad s⁻¹.

All PNP hydrogel formulations exhibited shear-thinning behavior with increasing shear rate for the range tested, with the viscosity collapsing onto that of the neat polymer solution.
as the shear rate approaches 100 s\(^{-1}\) for hydrogel formulations with NP content of 1 or 2 wt %. (Figure 2b). The neat polymer solutions behaved as viscous liquids. At low shear (0.01 s\(^{-1}\)), an approximately 3 orders of magnitude increase in viscosity was observed by increasing the NP content from 0 to 10 wt % for 1:X and 2:X hydrogel formulations, respectively. A nearly 3 order of magnitude decrease in viscosity was observed for 1:5, 1:10, and 2:10 hydrogel formulations across the measured range of shear rates, despite their comparatively high storage moduli. This shear-thinning ability is attributed to the supramolecular, non-covalent nature of the interactions between the components of the hydrogel, allowing the network to break under high shear and reform under low shear conditions. In addition, the aforementioned rheological properties have been previously shown to translate to injectability through a syringe and needle or catheter.\(^{30,34,35,44,45}\) PNP hydrogels of various formulations have been shown to be both cytocompatible, as well as biocompatible in vivo in our previous work (Figure S11).\(^{34-36,44,45}\) Taken together, these results show a rheologically tunable hydrogel platform that meets the aforementioned criteria for a suitable material for insulin stabilization.

**Figure 3.** In vitro insulin stability as determined by the thioflavin T (ThT) insulin aggregation assay. All formulations were prepared at 10 mg/mL insulin unless otherwise indicated (n = 4 per formulation). 2:10 hydrogel formulations were loaded with 6.7 mg/mL insulin. (a) ThT assay signal was verified to be unaffected by the presence of the hydrogel components. (b, c) 1:X and 2:X hydrogel formulations are compared at stressed aging conditions of 37 °C and constant agitation to simulate hot summer conditions. (d) Insulin formulated with a subset of hydrogel and polymer formulations aged at 50 °C with constant agitation to simulate extreme exposure conditions. (e) Stability comparison with the standard excipient trehalose and the commercial formulation Humulin. (f) Comparison of aggregation time for unencapsulated insulin formulation controls (n = 4, mean ± std. dev.).

Thermal Stability of Hydrogel-Encapsulated Insulin. Storage of insulin under environmental conditions can lead to significant loss of drug activity for current commercial formulations.\(^{46}\) To test the capacity of PNP hydrogels to stabilize insulin, insulin-loaded hydrogels were exposed to stressed aging conditions. Insulin was encapsulated under mild conditions by simply mixing the drug into the bulk hydrogel material at a concentration of 10 mg/mL (2:10 hydrogel at 6.7 mg/mL), which is nearly 3 times the typical commercial
clinical formulation concentration of 100 U/ml (3.47 mg/mL). The insulin-loaded samples were then subjected to stressed aging conditions (37 °C and constant agitation) and the stability of the insulin was monitored via thioflavin T (ThT) assay (Figure S7). The primary mechanism of insulin denaturation is through the formation of amyloid fibrils, and ThT interaction with these fibrils results in a sharp increase in fluorescence signal indicating insulin aggregation. Individual gel components (polymer and nanoparticles) were tested as assay controls to confirm that they did not produce background fluorescence (Figures 3a and S8).

Samples were subjected to stressed aging conditions of 37 °C and constant agitation to simulate hot summer conditions, where room temperature storage may exceed 35 °C. Under these conditions, insulin formulated in phosphate buffered saline (PBS) aggregated in 20 ± 4 h (Figure 3a). In contrast, when insulin was encapsulated in any PNP hydrogel formulation (1:X, 2:X) or formulated with neat polymer solutions (1 or 2 wt % HPMC-C12, 2 wt % HPMC), no aggregation was observed over the 28 day study (Figure 3b,c). The termination of the study at 28 days was determined by the point where substantial evaporation of the formulations had taken place. Insulin formulated with only NPs (2, 5, and 10 wt %) aggregated after less than 7 h on average (Figure S8). Together, these results suggest that the HPMC polymer contributes significantly to insulin stability in these systems.

To further test the capacity of selected top-performing formulations to withstand insulin aggregation under extreme thermal conditions, we conducted an additional stressed aging study at 50 °C with constant agitation. These conditions simulate an interruption in the cold chain where formulations are left unrefrigerated in a hot truck or shipping container. Under these conditions, insulin monitored via thioflavin T (ThT) assay detected insulin-stimulated phosphorylation of AKT and compared with the activity of fresh insulin and Humulin samples. Data shown are an average of n = 3 aged sample replicates, mean ± std. dev.

ThT assay results were corroborated by assessing both the secondary structure and in vitro bioactivity of gel-encapsulated insulin post-aging. Selected samples taken from formulations after 28 days of stressed aging (37 °C, constant agitation) were analyzed by circular dichroism to characterize the secondary protein structure and by an in vitro cellular activity assay to assess insulin function. Circular dichroism shows that the secondary structure of gel-encapsulated insulin is intact and comparable to fresh insulin, as indicated by the presence of peaks near 208 and 224 nm that are characteristic of α-helices, the dominant secondary structure in hexameric insulin (Figures 4a and S9).

To verify the activity of gel-encapsulated insulin samples, dose–response curves were determined by an AlphaLISA (Perkin-Elmer) assay to quantify the ratio of phosphorylated AKT compared to total AKT in C2C12 cells stimulated with aged insulin samples compared to fresh insulin controls. Our results indicate that insulin formulated with 2 wt % HPMC-C12 and 2:2 hydrogel exhibited activity comparable to both freshly prepared insulin, as well as Humulin, whereas aged insulin formulated only in PBS exhibited significantly reduced activity (activity, logEC50 = mean ± s.e. (n = 3): 2 wt % HPMC-C12 logEC50 = −1.3 ± 2, 2:2 hydrogel logEC50 = −4.1 ± 0.2, aged insulin logEC50 = −0.48 ± 1.3, insulin logEC50 = −2.9 ± 0.4, Humulin logEC50 = −4.1 ± 0.2) (Figure 4b). The logEC50 of insulin formulated in a 2:2 hydrogel was lower (indicating higher activity) compared to that of aged insulin in PBS (p < 0.0001). No difference was observed between insulin formulated in a 2:2 hydrogel and fresh Humulin (p = 0.49).
The ability of the PNP hydrogels to preserve insulin secondary structure and in vitro activity demonstrates their potential to reduce the cold chain dependence for insulin distribution, and moving forward, to serve as a sustained release drug delivery depot imparting enhanced drug stability at increased concentrations under physiological conditions.

Determining the precise mechanism through which these materials stabilize insulin is beyond the scope of the present work; however, it is known that the air—liquid interface can mediate insulin fibrillation by allowing for partial protein unfolding as insulin adsors to the surface, which can lead to insulin aggregate nucleation. We hypothesize that these materials improve thermal stability of insulin by limiting access to the air—water interface due to their high viscosity compared to a buffer solution. In addition, hydrogels can slow the diffusion of drug molecules, which may also contribute to their ability to promote thermal stability of biotherapeutics.

### Release of Drug Cargo by Simple Dilution

A key design component for clinical translation of hydrogel-stabilized biotherapeutics is the ability to either directly administer the hydrogel formulation, or easily separate the drug from the hydrogel encapsulant. The supramolecular PNP hydrogels can be diluted to disrupt the non-covalent polymer network and injected without any further processing steps (Video S1), reducing the potential impact on the pharmacokinetics of the encapsulated drug cargo. The biocompatible hydrogel components do not need to be separated from the biotherapeutic prior to use.

The viscosity and shear-thinning profiles of both the 2:10 hydrogel and 2:2 hydrogel were measured to demonstrate the change in rheological properties upon serial dilution with PBS (Figures 5 and S10). For the 2:10 hydrogel, which has the highest storage modulus of all of the materials evaluated, 4X dilution with PBS buffer is sufficient to cause approximately 2 orders of magnitude decrease in viscosity (Figure 5a,b). For the 2:2 hydrogel, a 2X dilution with PBS buffer decreases viscosity by an order of magnitude (Figure 5d,e and Video S1). An approximate gel-to-sol transition was determined by inverted vial test, indicating gel-to-sol transitions upon 4X and 2X dilution for the 2:10 and 2:2 hydrogel formulations, respectively (Figure 5c,f). Because these hydrogels enhance the stability of insulin, insulin could be formulated at higher concentrations and simply diluted to standard concentrations before administration. There is clinical precedent for dilution or reconstitution prior to use for lyophilized drugs and vaccines, highlighting the utility of this simple approach for treatment of diabetes in low-resource environments. Further, 1:0, 1:1, 1:2, and 2:0 formulations enhanced insulin stability and demonstrated fluid-like properties without dilution. Thus, insulin could be formulated in these materials at standard concentrations and dilution would be unnecessary. Future work could include determining a lower threshold of polymer concentration needed for effective thermal stabilization.

### CONCLUSION

We demonstrate that self-assembled physical hydrogels composed of an HPMC-C12 network non-covalently cross-linked by biodegradable core—shell nanoparticles are capable of extending the thermal stability of insulin at increased concentrations, elevated temperatures, and under constant mechanical agitation. Our physical hydrogel is mildly formulated to encapsulate biotherapeutics, effectively stabilizes insulin, and can be easily diluted for injection such that drug pharmacokinetics are unlikely to be affected. As the hydrogel components are both biocompatible and non-toxic, they would not need to be removed prior to administration. Because other biotherapeutics are also prone to denaturation and aggregation like insulin, this hydrogel platform would likely be able to provide thermal stability to other relevant research-grade and therapeutic macromolecules such as proteins and enzymes. We anticipate that the use of PNP hydrogels can help increase the global availability of insulin, and other biotherapeutics, in regions with unreliable cold chain infrastructure. Further, these hydrogels are being evaluated as injectable materials for sustained drug delivery, potentially enabling new formulation options and treatment strategies for insulin derivatives and macromolecular therapeutics more broadly as a drug class.

### ASSOCIATED CONTENT

- **Supporting Information**
  - The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsbiomaterials.0c01306.
  - Additional figures and methods (PDF)
  - Video S1: Injection and dilution of 2:2 PNP formulation (MOV)

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Notes

The authors declare no competing financial interest.

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