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Molecular Dynamics-Assisted Design of High Temperature-Resistant Polyacrylamide/Poloxamer Interpenetrating Network Hydrogels

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Abstract: Polyacrylamide has promising applications in a wide variety of fields. However, conventional polyacrylamide is prone to hydrolysis and thermal degradation under high temperature conditions, resulting in a decrease in solution viscosity with increasing temperature, which limits its practical effect. Herein, combining molecular dynamics and practical experiments, we explored a facile and fast mixing strategy to enhance the thermal stability of polyacrylamide by adding common poloxamers to form the interpenetrating network hydrogel. The blending model of three synthetic polyacrylamides (cationic, anionic, and nonionic) and poloxamers was first established, and then the interaction process between them was simulated by all-atom molecular dynamics. In the results, it was found that the hydrogen bonding between the amide groups on all polymers and the oxygen-containing groups (ether and hydroxyl groups) on poloxamers is very strong, which may be the key to improve the high temperature resistance of the hydrogel. Subsequent rheological tests also showed that poloxamers can indeed significantly improve the stability and viscosity of nonionic polyacrylamide containing only amide groups at high temperatures and can maintain a high viscosity of 3550 mPa·s at 80 °C. Transmission electron microscopy further showed that the nonionic polyacrylamide/poloxamer mixture further formed an interpenetrating network structure. In addition, the Fourier transform infrared test also proved the existence of strong hydrogen bonding between the two polymers. This work provides a useful idea for improving the properties of polyacrylamide, especially for the design of high temperature materials for physical blending.

Keywords: acrylamide polymers; high temperature-resistant; hydrogel; hydrogen bonding; molecular dynamics; poloxamers

1. Introduction

Polyacrylamide (PAM), a water-soluble material polymerized from acrylamide monomers [1,2], is widely used in many fields, such as oil extraction, metallurgy, papermaking, and medicine [3,4]. Due to its excellent properties such as flocculation, thickening, resistance reduction, and dispersion, PAM with high molecular weight can increase the viscosity of oil field injection fluid and improve the heterogeneity of reservoirs, so it is widely used as an oil displacement agent in the process of oil recovery [5–7]. However, due to the curl shrinkage of the PAM molecular chain and the partial hydrolysis of the amide group on PAM at higher temperatures, the viscosity of the polymer decreases significantly with an increase in temperature, which greatly limits its application range. The viscosity drops rapidly, which seriously restricts its application [8–10]. Therefore, the design and preparation of high temperature-resistant PAM have attracted more and more attention.
In recent decades, scientists have carried out numerous innovative works on improving the temperature resistance of polymers. The research directions mainly include two aspects: 1. Synthesize copolymers with temperature-resistant structural units [11–13]. The special properties of polymers obtained by copolymerizing one or more temperature-resistant monomers with acrylamide monomers, such as hydrophobically associative, molecular complex, and zwitterionic [14,15], enable the hydrolysis process under restricted high temperature conditions, thereby improving the stability of the material. 2. Blend with some special substances [16–18]. Through chemical interaction (Schiff base, grafting, and substitution reaction) or physical interaction (hydrophobic association, hydrogen bonding, and electrostatic interaction) [18,19], it is easier to form a cross-linked structure between the PAM molecules. This can make the polymer stronger and more difficult to transform conformation, thereby improving its stability at high temperature. Compared with the complicated synthesis process, the addition of foreign substances to form the composite system of cross-linked polymers is simpler, and it is also beneficial for large-scale preparation and application. Although the optional additives are also various, such as organic chromium, phenolic, and polyethyleneimine, etc., these materials require caution due to their cytotoxicity and environmental concerns [19–21]. Furthermore, developing a class of efficient and safe additives undoubtedly requires more effort and time, which is very challenging for researchers.

Herein, with the assistance of molecular dynamics simulations, we report a strategy to improve the thermal stability of polyacrylamide by using poloxamer as a crosslinker to form an interpenetrating network hydrogel. The so-called poloxamer is a new type of polymer nonionic polyol with low toxicity, low irritation, and good biocompatibility, which can be used in biological fields such as pharmaceutical excipients, gene therapy, and the inhibition of postoperative intestinal adhesions [22–24]. First, multiple models were established to simulate the interaction between poloxamers and three common polyacrylamides, which were anionic polyacrylamide (APAM) [25], cationic polyacrylamide (CPAM) [26], and nonionic polyacrylamide (NPAM) [27]. The results predicted strong hydrogen bonds between the amide groups on the polymer chain and the poloxamers. Subsequently, the different viscosity characteristics and morphological structures of the three composite systems at high temperature were verified experimentally. Finally, further rheological tests also indicated that the NPAM/poloxamer composite system is a high-viscosity hydrogel with ideal thermal stability and high temperature resistance. This facile mixing method does not require further complicated processing steps, which facilitates large-scale production and the rapid preparation of materials and has broad application prospects in the fields of oil extraction, papermaking, and environmental protection. Furthermore, this work utilizes molecular dynamics as a guide to develop suitable additives for polyacrylamide, providing a useful idea for the design of high temperature materials for physical blending.

2. Results and Discussion
2.1. Molecular Dynamics

Figure 1 shows the synthetic process and chemical structures of three different types of polyacrylamides, as well as the structures of poloxamers. The structures of CPAM, APAM, NPAM, and poloxamer can be verified in the $^1$H NMR spectrum (Figures S1–S4), which mainly show inactive hydrogens on methyl or methylene groups. To investigate possible interactions between such polymers, we first performed molecular dynamics (MD) simulations of simple NPAM and poloxamer [28–30]. Nonionic polyacrylamide (NPAM) has only -CO-NH$_2$ groups and no other ionic groups. The MD processes and interactions between the two polymers were mainly studied. In an aqueous solution, the molar ratio of NPAM to poloxamer is 1:1. Initially, 20 molecules were randomly contained in a cube length of 45 Å, equilibrated at 298.15 K for 1 ns, and finally MD simulations of 50 ns were performed. Eventually all molecules form different structures. After the simulation process, the molecular dynamics simulation plots at 10 ns, 35 ns, and 50 ns were extracted, and different colors were used to distinguish the two molecules (Figure 2A). It can also be seen
from the simulation results that an aggregated complex is formed between poloxamer and nonionic polyacrylamide, which is mainly due to the obvious hydrogen bonding between the hydroxyl, ether, and amide groups. Note that there are also obvious hydrogen bonds between the NPAM molecules. According to reports, multiple hydrogen bonds formed between polymers are beneficial to improving the stability of materials in most cases.

Figure 1. (A) Structure of anionic polyacrylamide (APAM), cationic polyacrylamide (CPAM), and nonionic polyacrylamide (NPAM). (B) Structure of Poloxamer. Among them, important monomers include acrylamide (AM), acrylic acid (AA), and acryloyloxyethyltrimethyl ammonium chloride (DAC).

Figure 2. (A) Molecular dynamics process of the interaction between NPAM and poloxamer. (B) Hydrogen bonding between two polymers.
In addition, MD simulations were performed on other composite systems under the same conditions (Figures S5 and S6). For anionic polyacrylamide (APAM), it has a -COO⁻ group and therefore has a pronounced negative charge (Figure 3A). For cationic polyacrylamide (CPAM), it has -N(CH₃)₃⁺ groups in cationic form on the branched chains, which have a pronounced positive charge (Figure 3B). Unlike the NPAM case, the binding of APAM and CPAM to poloxamer was not evident. APAM and poloxamer tend to disperse rather than aggregate. This may be due to the fact that the carboxyl groups on APAM are easily soluble in water [31,32], that is, tend to combine with water molecules, which instead reduces the complexation with poloxamer. For CPAM, there is a large branch on the molecular chain, which may be difficult to disperse due to the fact that the steric position can fix the poloxamer (Figure 3B) molecule [33,34]. From the results of the two simulations, poloxamer molecules still tend to form hydrogen bonds with amide groups on the polymer chain, rather than other anionic and cationic groups on the copolymer. Due to the relatively low content of amide groups in these two copolymers, it may lead to weaker hydrogen bonding with poloxamers.

![Molecular dynamics results of the interaction of two composite systems. (A) APAM and poloxamer. (B) CPAM and poloxamer.](image)

**Figure 3.** Molecular dynamics results of the interaction of two composite systems. (A) APAM and poloxamer. (B) CPAM and poloxamer.

### 2.2. Preparation and Characterization of Materials

Furthermore, the predicted results of molecular dynamics simulations are verified by actual experiments. For the convenience of the study, the ratio of polyacrylamide and poloxamer was fixed at 1:1, and the total content of both in composite systems was 10 wt%. Typically, the three low-concentration polyacrylamides do not gel at room temperature or high temperature. After adding poloxamer and mixing at room temperature to dissolve, CPAM/poloxamer and NPAM/poloxamer formed a more obvious self-supporting hydrogel (Figure 4A). In contrast, APAM/poloxamer was in the solution state. Figures 4B and 4C also provide the G' (storage modulus) and G'' (loss modulus) of CPAM/poloxamer and NPAM/poloxamer. Except for APAM/poloxamer (Figure S7), the G' values of the other blends were consistently greater than G'' over the entire range, indicating the elastic properties of the hydrogels [35]. In addition, it can also be found that the G' value of NPAM/poloxamer (135 Pa) was higher than that of CPAM/poloxamer (98 Pa), which means higher intensity [36,37].
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The effect of temperature on the viscosity of the three composite systems was also investigated. Meanwhile, three single polymer solutions with a concentration of 10 wt% were also prepared for comparison. According to reports, poloxamer is a classical temperature-sensitive copolymer, which produces a sol-gel phase transition with the change of temperature [22]. When the temperature increases, its viscosity first increases with the temperature and then slowly decreases (Figure S8). The viscosity of CPAM/poloxamer is relatively stable, which may benefit from the reduced temperature sensitivity due to the introduction of large cationic groups (Figure 5A). The viscosity of NPAM/poloxamer is also more stable over the entire temperature range (25–80 °C) compared to pure NPAM (Figure 5B). When the temperature was raised to 70 °C, the viscosity of the former dropped sharply to 12,800 mPa·S, but the latter remained at 25,150 mPa·S. Therefore, it can be pointed out that poloxamer significantly improves the thermal stability of the viscosity of NPAM at high temperature. As shown in Figure 5C, the addition of poloxamer did not significantly change the viscosity of APAM at high temperature, but instead caused a sharp drop. According to reports, it is known that poloxamer molecules are negatively charged due to the presence of ether groups (-O-) and hydroxyl groups (-OH). This leads to mutual repulsion with the similarly negatively charged APAM molecules, making it difficult for the two to bind and possibly reducing the viscosity of the latter.

The shear viscosities of the three complexes were also tested at continuously varying shear rates. In theory, composites tend to have higher viscosities than single polymers because crosslinking can restrict the movement of polymer chains. The viscosity of CPAM/poloxamer did not change much (Figure S9), and a very significant increase in the viscosity of NPAM/poloxamer was noted over the entire shear range (Figure S10). In addition, compared to the single APAM solution, the viscosity of APAM/poloxamer dropped sharply (Figure S11). The viscosity of all materials decreased with an increasing shear rate, indicating that they have polymer properties [38,39].
which may be the reason for the high temperature resistance of NPAM. Apparently, the SEM image of APAM, and (Figure 5F), but just some scattered and irregular flakes. It has been reported that the stronger hydrogen bonds [46–48].

The FTIR spectrum towards lower wavenumbers is reported to demonstrate the formation of increases, the characteristic peak shifts of O-H and C-O are more obvious. The shift in the broad peak was formed in the region near O-H. As the concentration of stretching bands of poloxamer shifted to 3445 and 1100 cm$^{-1}$ in Figure 6B, the characteristic O-H and C-O stretching bands of poloxamer are shown at the concentrations of 4, 8, and 12 wt%, respectively. To further elucidate the interactions of NPAM, observations suggest that at the beginning of NPAM, molecules and poloxamer chains cross-link with each other and form an interpenetrating network (Figure 5E) [10,42,43]. Apparently, the SEM image of APAM/poloxamer does not show obvious micropores (Figure 5F), but just some scattered and irregular flakes. It has been reported that the complete interpenetrating network structure is beneficial to the stability of the material, which may be the reason for the high temperature resistance of NPAM/poloxamer.

2.3. The Formation Mechanism of Hydrogels

Due to the excellent thermal stability of NPAM/poloxamer hydrogel, a more comprehensive characterization was performed. In fact, the hydrogel still had some fluidity at low concentration (4 wt%), while at high concentration (12 wt%) it was easy to generate bubbles and led to opacity (Figure 6A). It is speculated that the optimal concentration of the composite system may be between 6 and 11 wt%. For convenience, NPAM/poloxamer-4, NPAM/poloxamer-8, and NPAM/poloxamer-12 were used to represent the hydrogels at the concentrations of 4, 8, and 12 wt%, respectively. To further elucidate the interactions of mixtures, Fourier transform infrared spectroscopy (FTIR) was performed. As can be seen in Figure 6B, the characteristic O-H and C-O stretching bands of poloxamer are shown at 3450 and 1105 cm$^{-1}$, respectively [44–46]. After blending with NPAM, the characteristic stretching bands of poloxamer shifted to 3445 and 1100 cm$^{-1}$, respectively, and a distinct broad peak was formed in the region near O-H. As the concentration of NPAM/poloxamer increases, the characteristic peak shifts of O-H and C-O are more obvious. The shift in the FTIR spectrum towards lower wavenumbers is reported to demonstrate the formation of stronger hydrogen bonds [46–48].
In addition, hydrogel concentrations may affect the structure of the composite system, and SEM was used to study the composite system at different concentrations. As shown in Figure 6C, the lyophilized NPAM/poloxamer-4 sample has an irregular multilayered scale structure with insignificant micropores. With increasing concentration, the NPAM/poloxamer-8 hydrogel showed continuous pore walls and regular micropore, indicating the formation of a denser interpenetrating network structure. Similarly, NPAM/poloxamer-12 hydrogel has a distinct interpenetrating network structure, but also has a ragged pore structure which may be caused by inhomogeneous mixing. These results provide additional evidence for the formation of hydrogels and also demonstrate a stronger interaction between poloxamer and NPAM. Furthermore, this seems to validate the results of the above molecular dynamics simulations where the amide groups (−CO-NH$_2$) of NPAM have strong interactions with the ether groups (−O-) and (−OH) of poloxamer, mainly in the form of hydrogen bonds.

2.4. Thermosensitivity of Hydrogels

Since the hydrogen bonding between the poloxamer and the nonionic polyacrylamide is crucial for gel formation, the concentration has a crucial effect on the performance of the hydrogel. Therefore, we investigated the stability of hydrogels with different concentrations at high temperature. Figure 7A is the change curve of the three concentrations of the NPAM/poloxamer composite system with the increase of temperature. It can be seen from the figure that with an increase in concentration, the viscosity of the composite system tends to be stable in the process of increasing the temperature. For NPAM/poloxamer-4 hydrogel, the viscosity is stable between 8149–32,500 mPa·S in the range of 50–80 °C, and the variation range is relatively large. Obviously, the viscosity of the hydrogel further increased with increasing concentration. Note that at 75 °C, the viscosity of the NPAM/poloxamer-4 hydrogel is 11,000 mPa·S compared to 23,700 mPa·S for NPAM/poloxamer-8, which is an improvement of approximately two-fold. However, the viscosity of NPAM/poloxamer-12 is only 28,200 mPa·S at the same temperature, and there is no obvious increase. This shows...
that after the concentration reaches a certain level, the maximum viscosity of the hydrogel at high temperature has reached saturation.

Figure 7. (A) Viscosity of NPAM/poloxamer hydrogels at different concentrations. Temperature sweep tests of (B) NPAM/poloxamer-4, (C) NPAM/poloxamer-8, and (D) NPAM/poloxamer-12 hydrogels. (E) Pictures of NPAM/poloxamer-8 hydrogel heated to 80 °C and cooled to 25 °C.

The temperature sweep data of $G'$ (storage modulus) and $G''$ (loss modulus) of the composite system were further investigated. As shown in Figure 7B, although the $G'$ of the NPAM/poloxamer-4 hydrogel is always higher than $G''$, the decreasing trend is very obvious, which indicates the instability of the hydrogel structure [30,49]. In addition, the $G'$ of NPAM/poloxamer-8 (Figure 7C) and NPAM/poloxamer-12 (Figure 7D) showed similar variation patterns in the whole temperature range, and the magnitude of decrease was much smaller. In addition, it can also be seen from the data that the viscosity of all hydrogels shows a slow increase between 25 and 35 °C, and then decreases. In the early stage of the temperature increase, the poloxamer gelled and caused the viscosity to increase (Figure S12). This might also enhance the hydrogen bonding with NPAM molecules, thereby improving the stability of the solution at high temperature. The stabilization of the hydrogel may also benefit from the complete interpenetrating network formed by multiple hydrogen bonds which can greatly enhance the strength of the structure [50,51].

As the temperature continued to increase, the hydrogen bond between the polyacrylamide and poloxamer in the composite system would become weaker, which eventually led to a decrease in viscosity.
It is worth noting that high temperature also has an effect on the appearance of hydrogels. It can be seen that NPAM/poloxamer-8 hydrogel appeared with obvious turbidity at 80 °C and gradually clarified after lowering the temperature (Figure 7E). When returned to a room temperature of 25 °C, the composite system returned to its original clear and transparent state. It has been reported that poloxamer will appear cloudy and gelatinized at high temperature [52,53], which may be beneficial to the stabilization of NPAM/poloxamer. Therefore, this facilely prepared blended hydrogel is expected to be used to improve the application range of polyacrylamide, especially in high temperature environments.

3. Materials and Methods

3.1. All-Atom Molecular Dynamics (AAMD)

Fragment structures of poloxamers and three polyacrylamides (NPAM, CPAM, and APAM) were established by Guassian16. Afterwards, geometry optimization was performed by the B3LYP-D3 method and 6–31G (d, p) basis set. Atomic and molecular dynamics simulations were performed in the GROMACS (version 2020.6) simulation package using the general Amber force field (GAFF) combined with the TIP3P water model [54]. The ratio of poloxamers to NPAM was 1:1, and all systems were constructed by randomly placing 50 molecules into cubic boxes of approximately 6 nm. After solvation with water molecules, molecular dynamics simulations in the isothermal and isobaric (NPT) ensemble after thousands of steps of energy minimization were performed using Berendsen’s method at 298 K, 1 atm, and 50 ns. Structural pictures at 0 ns, 10 ns, 35 ns, and 50 ns were extracted to observe the interactions between molecules. The cutoff length for non-bonded interactions is 1.2 nm, and the long-range electrostatic interactions use the particle grid Ewald method with a Fourier spacing of 0.1 nm. Constraint of all covalent bonds to hydrogen atoms used the LINCS algorithm. CPAM and APAM were simulated using the same method. The final result was displayed by VMD 1.9.3 software [55], while the molecular structure was displayed by open-source PYMOL 2.3.0.

3.2. Materials

Acrylamide (A.R, 99.0%), acrylic acid (A.R, 99%), acryloyloxyethyltrimethyl ammonium chloride (A.R, 99%), poloxamer (A.R.), and other chemicals were purchased from Aladdin Reagent Co., Ltd. (Shanghai, China). All chemicals were used without purification, and deionized water was used for all experiments.

3.3. Synthesis of Nonionic Polyacrylamide (NPAM)

To begin with, 2 g of acrylamide was added to enough deionized water, stirred slowly until the acrylamide was completely dissolved. The pH was adjusted to about 4, stirred again and passed through N₂ gas to remove air. Subsequently, amine persulfate-sodium bisulfite was added as an initiator, and the resulting mixed solution was reacted at 25 °C for 24 h. The product was then precipitated with absolute ethanol and washed 3 times. Finally, a white powdery NPAM sample was obtained by freeze-drying for 24 h.

3.4. Synthesis of Anionic Polyacrylamide (APAM)

A mixed solution prepared from 2 g of acrylamide and 2 g of acrylic acid was adjusted to pH 9.0 with NaOH, and N₂ gas was introduced to remove air. The next steps were as above.

3.5. Synthesis of Cationic Polyacrylamide (NPAM)

A mixed solution of 2 g of acrylamide and 2 g of acryloyloxyethyltrimethyl ammonium chloride was prepared, and N₂ gas was introduced to remove air. The next steps were as above.
3.6. Preparation of Hydrogels

The obtained polymers were freeze-dried to obtain powder products. Poloxamer solution (5 wt%) was obtained by dissolving poloxamer powder into deionized water at a temperature of 50 °C for several minutes. Then, nonionic polyacrylamide (5 wt%) was added and stirring was continued for a few minutes to finally obtain a 10 wt% NPAM/poloxamer mixture. After sonication to remove air bubbles, the product was freeze-dried to obtain a powder. Cationic polyacrylamide and anionic polyacrylamide were prepared using the same method.

3.7. Characterization

$^1$H NMR spectra were recorded by using an AMX-400 (Bruker, Switzerland). The freeze-dried samples were dissolved in D$_2$O and scanned under a temperature of 300 K. The microstructure images of NPAM/poloxamer hydrogels were characterized by a scanning electron microscope (SEM, JEOL JSM-6701F, Japan, Tokyo). FT-IR spectra of samples were obtained with a Perkin Elmer Spectrum One instrument (Waltham, MA, USA) and the scanning range was 4000 cm$^{-1}$–400 cm$^{-1}$. The test samples were prepared by mixing freeze-dried hydrogel powder after 48 h of vacuum drying with KBr crystals baked by infrared light.

3.8. Rheological Mechanical

The rheological behavior of NPAM/poloxamer samples was analyzed using a modular compact rheometer (Anton Paar, MCR 302). Time sweep measurements were performed at 0.1% strain, 1 Hz frequency, and the time was held at 20 min. Frequency ($\omega$) sweep tests were performed in the range 0.1–100 rad/s and the temperature was fixed at 25 °C. The temperature sweep test (T) was performed in the range of 25–80 °C with $\omega$ fixed at 0.5 rad/s. The strain ($\gamma$) was fixed at 0.5% in all the above tests.

4. Conclusions

In this study, a polyacrylamide/poloxamer hydrogel with an interpenetrating network was successfully constructed using a hybrid modification strategy. To study the interactions between polymers, molecular simulations were used to study the formation of three polyacrylamides and poloxamers, which were anionic polyacrylamide, cationic polyacrylamide, and nonionic polyacrylamide. The simulation results predict strong hydrogen bonding between the amide group on the polyacrylamide and the oxygen-containing groups (ether and hydroxyl) on the poloxamer, which may be the point of maintaining stability at high temperature.

Subsequent experiments tested the properties of the three complexes and verified the effect of poloxamers on polyacrylamides within different groups. The results of this experiment also showed that the addition of poloxamers significantly increased the viscosity and stability at high temperature of the nonionic polyacrylamide. Through a series of characterizations, it was concluded that the prepared nonionic polyacrylamide/poloxamer composite system formed an interpenetrating three-dimensional network with a regular porous structure which could form a self-supporting hydrogel at room temperature. In addition, the experiments further proved that when the ratio of the two polymers was 1:1, the hydrogel with a content of 8 wt% already had excellent high temperature resistance properties.

Considering the results obtained in this study, it can be emphasized that nonionic polycrylamides based on non-toxic and harmless poloxamers are very suitable for use as additives, resulting in higher viscosity and more remarkable stability. Since this material is easy to prepare and suitable for large-scale production, it has great application potential in petroleum, water treatment, and other fields. Furthermore, the introduction of molecular dynamics simulations also provides a simple and practical strategy for designing high temperature-resistant acrylicamide polymers.
Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/molecules27165326/s1, Figure S1: The full $^1$H NMR spectrum of the APAM; Figure S2: The full $^1$H NMR spectrum of the NPAM; Figure S3: The full $^1$H NMR spectrum of the CPAM; Figure S4: The full $^1$H NMR spectrum of the poloxamer; Figure S5: Molecular dynamics process of the interaction between APAM and poloxamer; Figure S6: Molecular dynamics process of the interaction between CPAM and poloxamer; Figure S7: $G'$ and $G''$ values of CPAM/poloxamer composite system; Figure S8: Viscosity of poloxamer (mass fraction: 10 wt%) in the temperature range of 25–80 °C; Figure S9: Viscosity of CPAM and CPAM/poloxamer; Figure S10: Viscosity of NPAM and NPAM/poloxamer; Figure S11: Viscosity of APAM and APAM/poloxamer; Figure S12: Viscosity of poloxamer (mass fraction: 4 wt%) in the temperature range of 25–80 °C.

Author Contributions: X.S. participated in the conception and design of the study; G.L. carried out the material characterization and simulation calculations; J.W. and X.S. carried out cytotoxicity experiments, antibacterial experiments, and analyzed the data; J.Z. and S.S. contributed to the analysis of the data; Q.L. and Y.Z. contributed to the design of study and revised the manuscript. All authors have read and agreed to the published version of the manuscript.

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References

1. Jiang, F.; Chi, Z.; Ding, Y.; Quan, M.; Tian, Y.; Shi, J.; Song, F.; Liu, C. Wound Dressing Hydrogel of Enteromorpha prolifera Polysaccharide-Polyacrylamide Composite: A Facile Transformation of Marine Blooming into Biomedical Material. *ACS Appl. Mater. Interfaces* **2021**, *13*, 14350–14352. [CrossRef] [PubMed]

2. Kawai, R.; Tanaka, H.; Matsubara, S.; Ida, S.; Uchida, M.; Okumura, D. Implicit rule on the elastic function of a swollen polyacrylamide hydrogel. *Soft Matter* **2021**, *17*, 4979–4988. [CrossRef]

3. Abbasi, S.; Yousefi, G.; Tamaddon, A.M.; Firuzi, O. Paclitaxel-loaded polypeptide-polyacrylamide nanomicelles overcome drug-resistance by enhancing lysosomal membrane permeability and inducing apoptosis. *J. Biomed. Mater. Res. A* **2021**, *109*, 18–30. [CrossRef] [PubMed]

4. Daliri, K.; Pfannkuche, K.; Garipcan, B. Effects of physicochemical properties of polyacrylamide (PAA) and (polydimethylsiloxane) PDMS on cardiac cell behavior. *Soft Matter* **2021**, *17*, 1156–1172. [CrossRef] [PubMed]

5. Perez-Calixto, D.; Amat-Shapiro, S.; Zamarron-Hernandez, D.; Vazquez-Victorio, G.; Puech, P.H.; Hautefeuille, M. Determination by Relaxation Tests of the Mechanical Properties of Soft Polyacrylamide Gels Made for Mechanobiology Studies. *Polymers* **2021**, *13*, 629. [CrossRef]

6. Scalet, J.M.; Suekama, T.C.; Jeong, J.; Gehrke, S.H. Enhanced Mechanical Properties by Ionomeric Complexation in Interpenetrating Network Hydrogels of Hydrolyzed Poly (N-vinyl Formamide) and Polyacrylamide. *Gels* **2021**, *7*, 80. [CrossRef]

7. Song, T.; Li, S.; Yin, Z.; Bao, M.; Lu, J.; Li, Y. Hydrolyzed polyacrylamide-containing wastewater treatment using ozone reactor-upflow anaerobic sludge blanket reactor-aerobic biofilm reactor multistage treatment system. *Environ. Pollut.* **2021**, *269*, 116111. [CrossRef]

8. Werber, J.R.; Peterson, C.; Van Zee, N.J.; Hillmyer, M.A. Functionalized Polymersomes from a Polyisoprene-Activated Polyacrylamide Precursor. *Langmuir* **2021**, *37*, 490–498. [CrossRef]

9. Zhang, Z.; Lin, T.; Li, S.; Chen, X.; Que, X.; Sheng, L.; Hu, Y.; Peng, J.; Ma, H.; Li, J.; et al. Polyacrylamide/Copper-Alginate Double Network Hydrogel Electrolyte with Excellent Mechanical Properties and Strain-Sensitivity. *Macromol. Biosci.* **2022**, *22*, 2100361. [CrossRef]

10. Zhao, D.; Feng, M.; Zhang, L.; He, B.; Chen, X.; Sun, J. Facile synthesis of self-healing and layered sodium alginate/polyacrylamide hydrogel promoted by dynamic hydrogen bond. *Carbohydr. Polym.* **2021**, *256*, 117580. [CrossRef]
11. Jayaramudu, T.; Ko, H.U.; Kim, H.C.; Kim, J.W.; Kim, J. Swelling Behavior of Polyacrylamide-Cellulose Nanocrystal Hydrogels: Swelling Kinetics, Temperature, and pH Effects. *Materials* 2019, 12, 2080. [CrossRef] [PubMed]

12. Li, Y.; Gong, Q.; Liu, X.; Xia, Z.; Yang, Y.; Chen, C.; Qian, C. Wide temperature-tolerant polyaniline/cellulose/polyacrylamide hydrogels for high-performance supercapacitors and motion sensors. *Carbohydr. Polym.* 2021, 267, 118207. [CrossRef]

13. Mirzaie Yegane, M.; Hashemi, F.; Vercauteren, F.; Meulendijks, N.; Gharbi, R.; Boukany, P.E.; Zitha, P. Rheological response of a modified polycrylamide-silica nanoparticles hybrid at high salinity and temperature. *Soft Matter* 2020, 16, 10198–10210. [CrossRef] [PubMed]

14. Bulut, E.; Turhan, Y. Synthesis and characterization of temperature-sensitive microspheres based on acrylamide grafted hydroxypropyl cellulose and chitosan for the controlled release of amoxicillin trihydrate. *Int. J. Biol. Macromol.* 2021, 191, 1191–1203. [CrossRef]

15. Keratimanoch, S.; Takahashi, K.; Kuda, T.; Okazaki, E.; Geng, J.T.; Osako, K. Effects of tyndallization temperature on the sterility and quality of kamaboko. *Food Chem.* 2022, 366, 130692. [CrossRef] [PubMed]

16. Yin, C.; Lan, J.; Wang, X.; Zhang, Y.; Ran, R.; Shi, L.Y. Shape-Stable Hydrated Salts/Polyacrylamide Phase-Change Organohydrogels for Smart Temperature Management. *ACS Appl. Mater. Interfaces* 2021, 13, 21810–21821. [CrossRef]

17. Zhang, H.; Liu, Z.; Mai, J.; Wang, N.; Liu, H.; Zhong, J.; Mai, X. A Smart Design Strategy for Super-Elastic Hydrogel with Long-Term Moisture, Extreme Temperature Resistance, and Non-Flammability. *Adv. Sci.* 2021, 8, 2100320. [CrossRef]

18. Zhang, S.; Peng, B.; Wang, W. Temperature-Responsive Hydrogel Carrier for Reducing Adsorption Loss of Petroleum Sulfonates. *Langmuir* 2021, 37, 9809–9816. [CrossRef]

19. Zhang, J.; Zeng, L.; Qiao, Z.; Wang, J.; Jiang, X.; Zhang, Y.S.; Yang, H. Functionalizing Double-Network Hydrogels for Applications in Remote Actuation and in Low-Temperature Strain Sensing. *ACS Appl. Mater. Interfaces* 2020, 12, 30247–30258. [CrossRef]

20. Pi, K.; Liu, J.; Van Cappellen, P. A DNA-based biosensor for aqueous Hg(II): Performance under variable pH, temperature and competing ligand composition. *J. Hazard. Mater.* 2020, 385, 121572. [CrossRef]

21. Yin, B.; Liang, R.; Liang, X.; Fu, D.; Wang, L.; Sun, G. Construction of Stable Wide-Temperature-Range Proton Exchange Membranes by Incorporating a Carbonized Metal-Organic Frame into Polybenzimidazoles and Polyacrylamide Hydrogels. *Small* 2021, 17, 2103214. [CrossRef] [PubMed]

22. Colly, A.; Marquette, C.; Courtial, E.J. Poloxamer/Poly(ethylene glycol) Self-Healing Hydrogel for High-Precision Freeform Reversible Embedding of Suspended Hydrogel. *Langmuir* 2021, 37, 4154–4162. [CrossRef]

23. Feng, P.; Qiu, H.; Luo, Y.; Hu, J.; Cao, Y.; Pang, Q.; Mou, X.; Hou, R.; Hou, W.; Zhu, Y. Development of Poloxamer Hydrogels Containing Antibacterial Guanidine-Based Polymers for Healing of Full-Thickness Skin Wound. *ACS Biomater. Sci. Eng.* 2021, 7, 4537–4568. [CrossRef]

24. Rodrigues Sousa, H.; Lima, I.S.; Neris, L.M.L.; Silva, A.S.; Santos Nascimento, A.M.S.; Araujo, F.P.; Ratke, R.F.; Silva, D.A.; Osajima, J.A.; Bezerra, L.R.; et al. Superabsorbent Hydrogels Based to Polyacrylamide/Cashew Tree Gum for the Controlled Release of Water and Plant Nutrients. *Molecules* 2021, 26, 2680. [CrossRef] [PubMed]

25. Azad, M.S.; Trivedi, J.J. Synergistic Behavior of Anionic Surfactants and Hydrolyzed Polyacrylamide under an Extensional Field: Effect of Hydrophobicity. *Langmuir* 2021, 37, 13645–13653. [CrossRef]

26. Tian, G.; Zhong, X.; Wu, X.; Wang, Z. Self-Assembly Preparation of Nano-Lignin/Cationic Polyacrylamide Complexes. *Polymers* 2021, 13, 1726. [CrossRef] [PubMed]

27. Du, P.; Li, X.; Yang, Y.; Fan, X.; Fang, Z.; Zhou, Z. Enhanced coagulation by two-stage alum addition: The role of solution pH, floe breakage and assistant of non-ionic polyacrylamide. *Environ. Technol.* 2021, 42, 4456–4465. [CrossRef]

28. Mercado-Montijo, J.; Anstine, D.M.; Rukmani, S.J.; Colina, C.M.; Andrew, J.S. PEGDA hydrogel structure from semi-dilute concentrations: Insights from experiments and molecular simulations. *Soft Matter* 2022, 18, 3565–3574. [CrossRef]

29. Wu, B.; Zhao, S.; Yang, X.; Zhou, L.; Ma, Y.; Zhang, H.; Li, W.; Wang, H. Biomimetic Heterodimerization of Tetrapeptides to Generate Liquid Crystalline Hydrogel in A Two-Component System. *ACS Nano* 2022, 16, 4126–4138. [CrossRef]

30. Sun, Y.; Li, X.; Zhao, M.; Chen, Y.; Xu, Y.; Wang, K.; Bian, S.; Jiang, Q.; Fan, Y.; Zhang, X. Bioinspired supramolecular nanofiber hydrogel through self-assembly of biphenyl-tripeptide for tissue engineering. *Bioact. Mater.* 2022, 8, 396–408. [CrossRef]

31. Binaeian, E.; Babae Zadvarzi, S.; Yuan, D. Anionic dye uptake via composite using chitosan-polyacrylamide hydrogel as matrix containing TiO2 nanoparticles; comprehensive adsorption studies. *Int. J. Biol. Macromol.* 2020, 162, 150–162. [CrossRef] [PubMed]

32. Moghaddam, R.H.; Dadfarinia, S.; Shabani, A.M.H.; Tavakol, M. Synthesis of composite hydrogel of glutamic acid, gum tragacanth, and anionic polyacrylamide by electron beam irradiation for uranium (VI) removal from aqueous samples: Equilibrium, kinetics, and thermodynamic studies. *Carbohydr. Polym.* 2019, 206, 352–361. [CrossRef] [PubMed]

33. Wang, B.; Dai, L.; Hunter, L.A.; Zhang, L.; Yang, G.; Chen, J.; Zhang, X.; He, Z.; Ni, Y. A multifunctional nanocellulose-based hydrogel for strain sensing and self-powering applications. *Carbohydr. Polym.* 2021, 268, 118210. [CrossRef]

34. Zhang, S.; Kieffer, S.J.; Alleyne, A.G.; Braun, P.V. Directed Molecular Collection by E-Jet Printed Microscale Chemical Potential Wells in Hydrogel Films. *Adv. Mater.* 2018, 30, 1803140. [CrossRef] [PubMed]

35. Song, X.W.; He, T.; Qi, Y.J.; Liu, Y.L.; Wu, H.Q.; Liu, C.T.; Zhang, Y. Properties of Cell-Compatible Poly(vinyl alcohol) Hydrogels Cross-Linked with Hydrophobic Lutecin. *ACS Appl. Polym. Mater.* 2021, 3, 3019–3027. [CrossRef]

36. Azizighannad, S.; Wang, Z.; Siddiqui, Z.; Kumar, V.; Mitra, S. Nano Carbon Doped Polyacrylamide Gel Electrolytes for High Performance Supercapacitors. *Molecules* 2022, 26, 2631. [CrossRef]
37. Song, X.W.; Zhang, Z.Q.; Shen, Z.L.; Zheng, J.; Liu, X.; Ni, Y.Q.; Quan, J.; Li, X.R.; Hu, G.; Zhang, Y. Facile Preparation of Drug-Releasing Supramolecular Hydrogel for Preventing Postoperative Peritoneal Adhesion. *ACS Appl. Mater. Inter.* 2021, 13, 56881–56891. [CrossRef]

38. Agasty, A.; Wisniewska, A.; Kalwarczyk, T.; Koynow, K.; Holyst, R. Macroscopic Viscosity of Polymer Solutions from the Nanoscale Analysis. *ACS Appl. Poly. Mater.* 2021, 3, 2813–2822. [CrossRef]

39. Al-Saleh, S.; Alateeq, A.; Alshaya, A.H.; Al-Qahimi, A.S.; Tulfah, H.I.; Binhasan, M.; Shabib, S.; Farooq, I.; Vohra, F.; Abduljabbar, T. Influence of TiO$_2$ and ZrO$_2$ Nanoparticles on Adhesive Bond Strength and Viscosity of Dentin Polymer: A Physical and Chemical Evaluation. *Polymers* 2021, 13, 3794. [CrossRef]

40. Liao, X.; Song, X.; Li, J.; Li, L.; Fan, X.; Qin, Q.; Zhong, C.; Yang, P.; Zhan, J.; Cai, Y. An injectable co-assembled hydrogel blocks reactive oxygen species and inflammation cycle resisting myocardial ischemia-reperfusion injury. *Acta Biomater.* 2022, 149, 82–95. [CrossRef]

41. Zheng, J.; Fan, R.; Wu, H.; Yao, H.; Yan, Y.; Liu, J.; Ran, L.; Sun, Z.; Yi, L.; Dang, L.; et al. Directed self-assembly of herbal small molecules into sustained release hydrogels for treating neural inflammation. *Nat. Commun.* 2019, 10, 1604. [CrossRef] [PubMed]

42. Yue, Y.; Wang, X.; Wu, Q.; Han, J.; Jiang, J. Assembly of Polyacrylamide- Sodium Alginate-Based Organic-Inorganic Hydrogel with Mechanical and Adsorption Properties. *Polymers* 2019, 11, 1239. [CrossRef] [PubMed]

43. Zhao, Y.; Chen, G.; Sun, M.; Jin, Z.; Gao, C. Study on preparation of the pH sensitive hydroxyethyl chitin/poly (acrylic acid) hydrogel and its drug release property. *Sheng Wu Yi Xue Gong Cheng Xue Za Zhi* 2006, 23, 338–341.

44. Anwer, M.K.; Iqbal, M.; Ahmed, M.M.; Al-Saleh, S.; Alates, R.; Ansari, M.N.; Ezzeldin, E.; Khalil, N.Y.; Ali, R. Improving the Solubilization and Bioavailability of Arbidol Hydrochloride by the Preparation of Binary and Ternary beta-Cyclodextrin Complexes with Poloxamer 188. *Pharmaceuticals* 2021, 14, 411. [CrossRef] [PubMed]

45. Constantin, M.; Cosman, B.; Berea, M.; Ailiesei, G.L.; Fundueanu, G. Thermosensitive Poloxamer-graft-Carboxymethyl Pullulan: A Potential Injectable Hydrogel for Drug Delivery. *Polymers* 2021, 13, 3025. [CrossRef]

46. Zafar, A.; Alruwaili, N.K.; Imam, S.S.; Alsaidan, O.A.; Alsaidan, O.A.; Alkholifi, F.K.; Alharbi, K.S.; Mostafa, E.M.; Alanazi, A.S.; Gilani, S.J.; Musa, A.; et al. Formulation of Genistein-HP beta Cyclodextrin-Poloxamer 188 Ternary Inclusion Complex: Solubility to Cytotoxicity Assessment. *Pharmaceutics* 2021, 13, 1997. [CrossRef]

47. Su, G.; Jia, L.; Zhang, X.; Zhang, Y.; Deng, P.; Zhou, T. Exploration of the unusual two-step volume phase transition of the poly(N-vinylcaprolactam-co-hydroxyethyl methacrylate) hydrogel. *Phys. Chem. Chem. Phys.* 2018, 20, 23013–23024. [CrossRef] [PubMed]

48. Su, G.; Zhou, T.; Zhang, Y.; Liu, X.; Zhang, A. Microdynamics mechanism of D2O absorption of the poly(2-hydroxyethyl methacrylate)-based contact lens hydrogel studied by two-dimensional correlation ATR-FTIR spectroscopy. *Soft Matter* 2016, 12, 1145–1157. [CrossRef]

49. Du, W.; Deng, A.; Guo, J.; Chen, J.; Li, H.; Gao, Y. An injectable self-healing hydrogel-cellulose nanocrystals conjugate with excellent mechanical strength and good biocompatibility. *Carbohydr. Polym.* 2019, 223, 115084. [CrossRef]

50. Ding, H.; Liang, X.; Wang, Q.; Wang, M.; Li, Z.; Sun, G. A semi-inter-penetrating network ionic composite hydrogel with low modulus, fast self-recoverability and high conductivity as flexible sensor. *Carbohydr. Polym.* 2020, 248, 116797. [CrossRef]

51. Hua, L.; Xie, M.; Jian, Y.; Wu, B.; Chen, C.; Zhao, C. Multiple-Responsive and Amphibious Hydrogel Actuator Based on Asymmetric UCST-Type Volume Phase Transition. *ACS Appl. Mater. Interfaces* 2019, 11, 43641–43648. [CrossRef] [PubMed]

52. Alhakamy, N.A.; Hosny, K.M.; Rizg, W.Y.; Shmawbi, B.A.; Badr, M.Y.; Safi, A.Y.; Murshid, S.S.A. Development and Optimization of Hyaluronic Acid-Poloxamer In-Situ Gel Loaded with Voriconazole Cubosomes for Enhancement of Activity against Ocular Fungal Infection. *Gels* 2022, 8, 241. [CrossRef] [PubMed]

53. Ruan, X.; Hu, J.; Lu, L.; Wang, Y.; Tang, C.; Liu, F.; Gao, X.; Zhang, L.; Wu, H.; Huang, X.; et al. Poloxamer 407/188 Binary Thermosensitive Gel as a Moxidectin Delivery System: In Vitro Release and In Vivo Evaluation. *Molecules* 2022, 27, 3063. [CrossRef]

54. Hess, B.; Kutzner, C.; van der Spoel, D.; Lindahl, E. GROMACS 4: Algorithms for highly efficient, load-balanced, and scalable molecular simulation. *J. Chem. Theory Comput.* 2008, 4, 435–447. [CrossRef] [PubMed]

55. Humphrey, W.; Dalke, A.; Schulten, K. VMD: Visual molecular dynamics. *J. Mol. Graph.* 1996, 14, 33–38. [CrossRef]