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# Silver ions blocking crystallization of guanosine-based hydrogel for potential antimicrobial applications†

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In this work, the detailed crystallization process of 2'-deoxy-2'-fluoroguanosine (<sup>F</sup>G<sub>d</sub>) hydrogel has been studied using single crystal X-ray diffraction, variable-temperature nuclear magnetic resonance (VT-NMR), and scanning electron microscopy (SEM). Both solid and solution results indicated that the K<sup>+</sup>-mediated G-quartet structures were unstable and easily resulted in the breakdown of the hydrogel to form linear ribbon structures by forming mimic reverse Watson–Crick base pairs between the two faces with an intermolecular hydrogen-bond (N10H–O11). Accordingly, Ag<sup>+</sup> was introduced to block the crystallization of <sup>F</sup>G<sub>d</sub> to form long lifetime stable supramolecular hydrogel (>6 months) and possible silver-ions-mediated base pair motifs were suggested *via* NMR, UV, and mass spectroscopy (MS) in combination with powder X-ray diffraction (PXRD) and circular dichroism spectroscopy (CD). Furthermore, <sup>F</sup>G<sub>d</sub>Ag hydrogel exhibited low toxicity for normal oral keratinocyte cells (NOK-SI) and good antibacterial activities for *Fusobacterium nucleatum* *in vitro*.

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## Introduction

Supramolecular hydrogel self-assembled from low-molecular-weight gelators have gathered increasing attention over the past few decades owing to their potential applications in various fields including tissue scaffolding, drug delivery system, and functional biomaterials.<sup>1–13</sup> Among the five natural nucleosides, guanosine and its derivatives are of special interest as their self-complementary hydrogen-bond edges and aromatic surfaces, which promote them to further self-associate into highly ordered structures such as dimers, ribbons, and macrocyclic tetramers and even form three-dimensional (3D) gels containing certain alkali metal ions in aqueous or organic media.<sup>14–19</sup> Although there has been a long history of guanosine-derivative-based hydrogels since Bang demonstrated the formation of gels with guanylic acid in 1910,<sup>19–24</sup> a major limitation of guanosine-based hydrogels was their poor lifetime stability owing to their crystallization within hours at room temperature, which hinders their latent applications in biological or pharmaceutical fields.<sup>24</sup> To overcome the above limitation, guanosine

derivatives,<sup>25–34</sup> such as 5'-hydrazide-guanosine, 8-methoxy-2',3',5'-tri-*O*-acetylguanosine, and guanosine-borate esters, have been developed to improve the lifetime stability of hydrogels.

However, to the best of our knowledge, the crystallization process of guanosine-based hydrogels has not been explored yet. Thus, determining their delicate structural balance between gelation and crystallization is critical, not only to gain more insight into the reasons for crystallization and further understand how to design new guanosine-based gelators but also to tune their lifetime stability and mechanical properties to develop multifunction supramolecular materials in the future. Inspired by this, the crystallization process of 2'-deoxy-2'-fluoroguanosine (<sup>F</sup>G<sub>d</sub>) hydrogel has been studied in this work. <sup>F</sup>G<sub>d</sub> has the same base and similar sugar moiety as guanosine. It was first synthesized by J. Imura in 1981 and thereafter, it was observed to exhibit significant anti-virus activities for influenza A and B, herpes simplex virus (HSV), and H7N1.<sup>35–39</sup> Furthermore, Christopher J. L. suggested that <sup>F</sup>G<sub>d</sub> is a powerful anti-favoring tool to manipulate G-quadruplex polymorphism and folding topology.<sup>40</sup> Recently, the gel properties of <sup>F</sup>G<sub>d</sub> have been investigated by our group and the results indicated that <sup>F</sup>G<sub>d</sub> forms a transparent hydrogel only in the presence of K<sup>+</sup>.<sup>41</sup> In this work, the detailed crystallization process of <sup>F</sup>G<sub>d</sub> hydrogel was first studied using single crystal X-ray diffraction, VT NMR, SEM, PXRD, and CD. Based on the above information, we observed that silver ions could not only block the crystallization of <sup>F</sup>G<sub>d</sub> but also induce <sup>F</sup>G<sub>d</sub> to form a long-term stable supramolecular hydrogel with favorable antimicrobial activities (Fig. 1).

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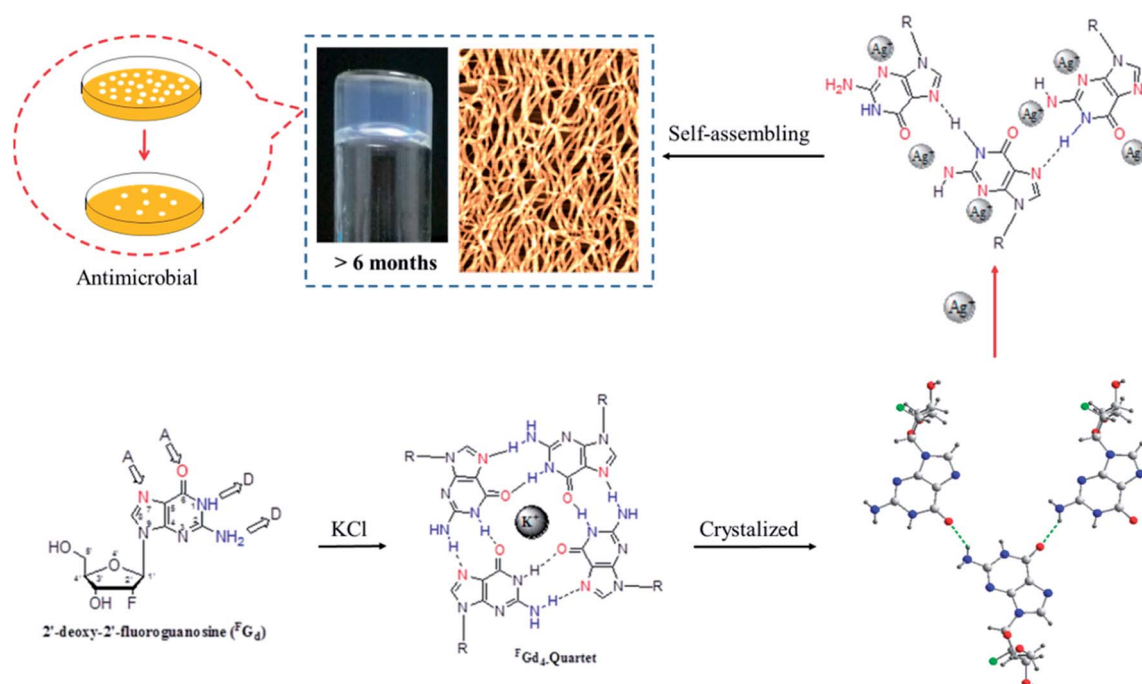


Fig. 1 Schematic illustration of a long lifetime stable supramolecular hydrogel formed when silver ions were used to block the crystallization of  $^F\text{G}_d$ . The hydrogel further showed highly antimicrobial activities.

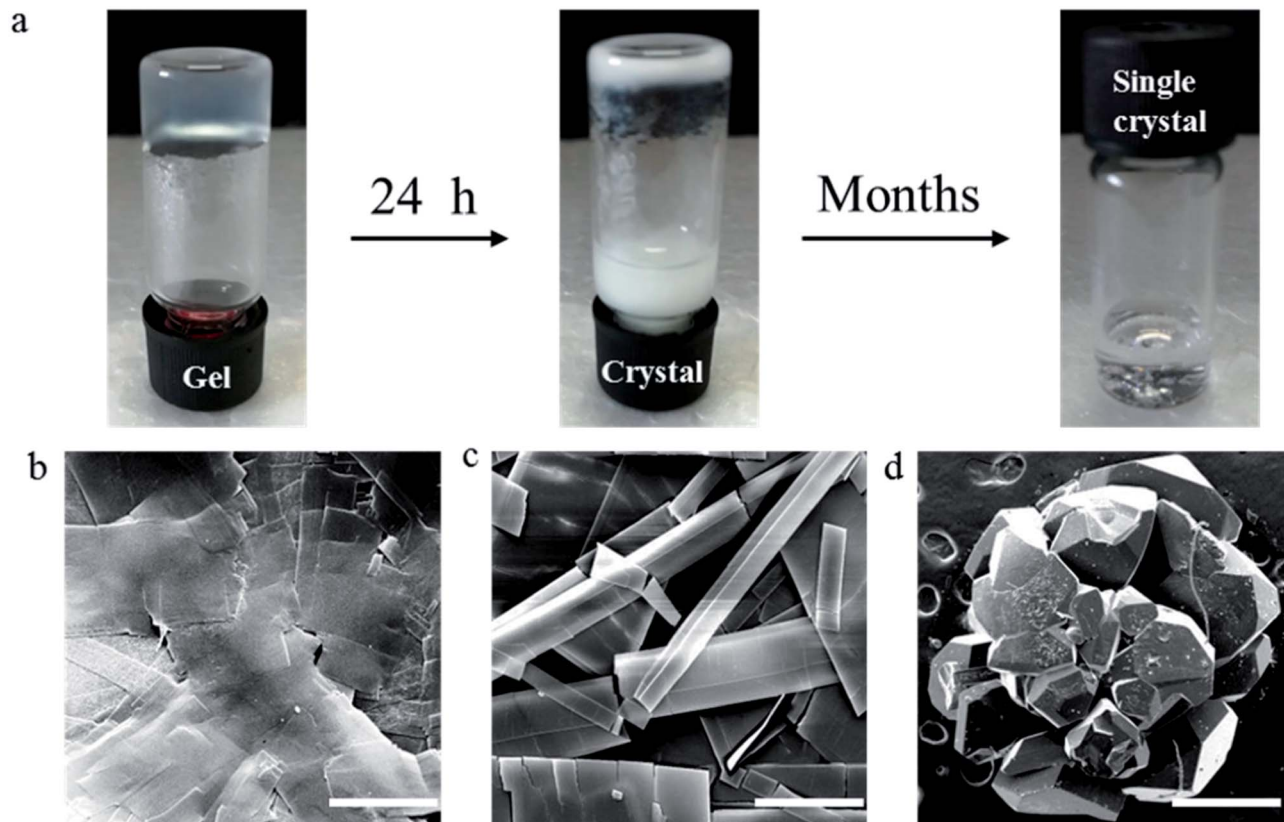


Fig. 2 (a) Images of the crystal process of  $^F\text{G}_d$  hydrogel at a concentration of 1.4 mg per 100  $\mu\text{L}$  of 0.2 M KCl solutions. (b–d) SEM images of the xerogels, crystal, and single crystal prepared from  $^F\text{G}_d$ . Scale bars: 10  $\mu\text{m}$  (b), 25  $\mu\text{m}$  (c), and 0.5 mm (d).



## Result and discussion

Recently, we reported the gel properties of  $^F\text{G}_d$  and the results indicated that  $^F\text{G}_d$  forms a transparent gel in the presence of  $\text{K}^+$  and crystallizes in the presence of  $\text{Li}^+$ ,  $\text{Na}^+$ ,  $\text{Rb}^+$ , and  $\text{Cs}^+$ .<sup>41</sup> The possible self-assembling process of  $^F\text{G}_d$  hydrogel is shown in Fig. S1a† according to the previous guanosine-based hydrogel. To obtain an insight into the microstructure of the hydrogel formed by  $^F\text{G}_d$ , SEM was performed on xerogels containing KCl at a lower concentration to avoid the negative impact caused by the crystallization of salt. 3D networks with porous-like structures consisting of intensive slices were observed and the diameter of the hole was approximately 150  $\mu\text{m}$  (see ESI, Fig. S1b and c†). Accordingly, atomic force microscope (AFM) measurements were carried out and a similar slice structure was obtained. Notably, some flexible fibers with a diameter of about 10–20 nm were observed on the surface of slices. These fibers inter-tangled with each other to form highly ordered slice

structures and thereafter formed 3D hydrogel networks. Nevertheless,  $^F\text{G}_d$  hydrogel was easily crystallized within hours similar to other guanosine-based hydrogels. Recent studies gave much evidence to explore the properties of gel to crystal transition for peptide, amino acid and sugar derivatives.<sup>42–45</sup> However, there is no literature to report the detail crystallization process of guanosine-based hydrogels. To find out and understand the reason for the crystallization of  $^F\text{G}_d$  hydrogel, the visualize dynamic crystal process of  $^F\text{G}_d$  hydrogel has been investigated as shown in Fig. 2a. First,  $^F\text{G}_d$  formed a transparent gel at the concentration of 1.4 mg per 100  $\mu\text{L}$  of 0.2 M KCl solution. Subsequently, the nucleoside  $^F\text{G}_d$  began to crystallize and the hydrogel broke down after 24 h. Eventually, we got the single crystal of  $^F\text{G}_d$  in the above solution after several months, which was, to the best of our knowledge, the first free guanosine-based nucleoside crystal formed from hydrogel and would provide us more detailed information about the reason for crystallization at molecular level. Meanwhile, the crystal

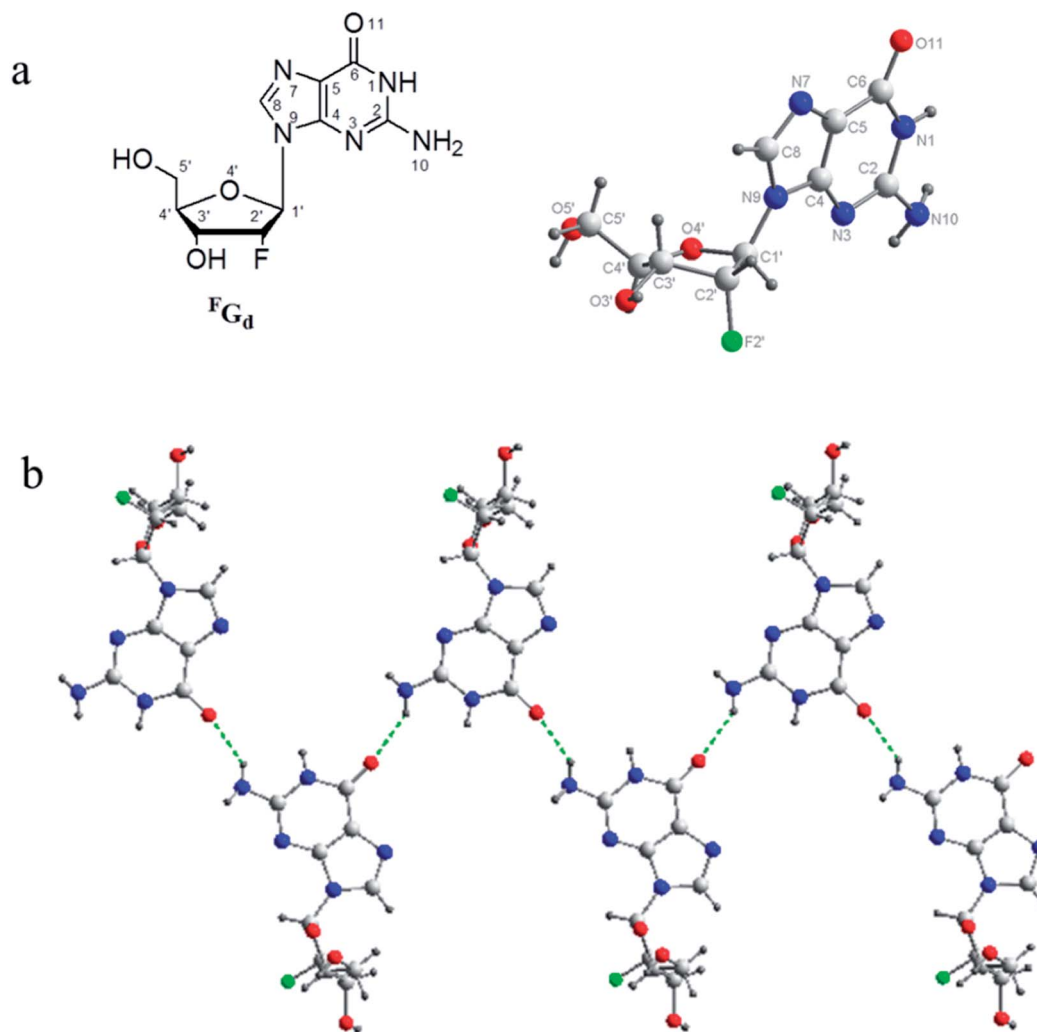


Fig. 3 (a) Molecular structure with systematic numbering and single crystal structure of  $^F\text{G}_d$ , which adopted an *anti*-conformation with an N-type (*3'*-*endo*) sugar pucker and 5'-OH at ap position. (b) A detailed view of the mimic reverse Watson–Crick base pairs in the solid state of  $^F\text{G}_d$  and the repeated hydrogen bonds unit in the entire assembly. Atoms were coded as follows: red, oxygen; blue, nitrogen; gray, carbon; green, fluorine; black, hydrogen.

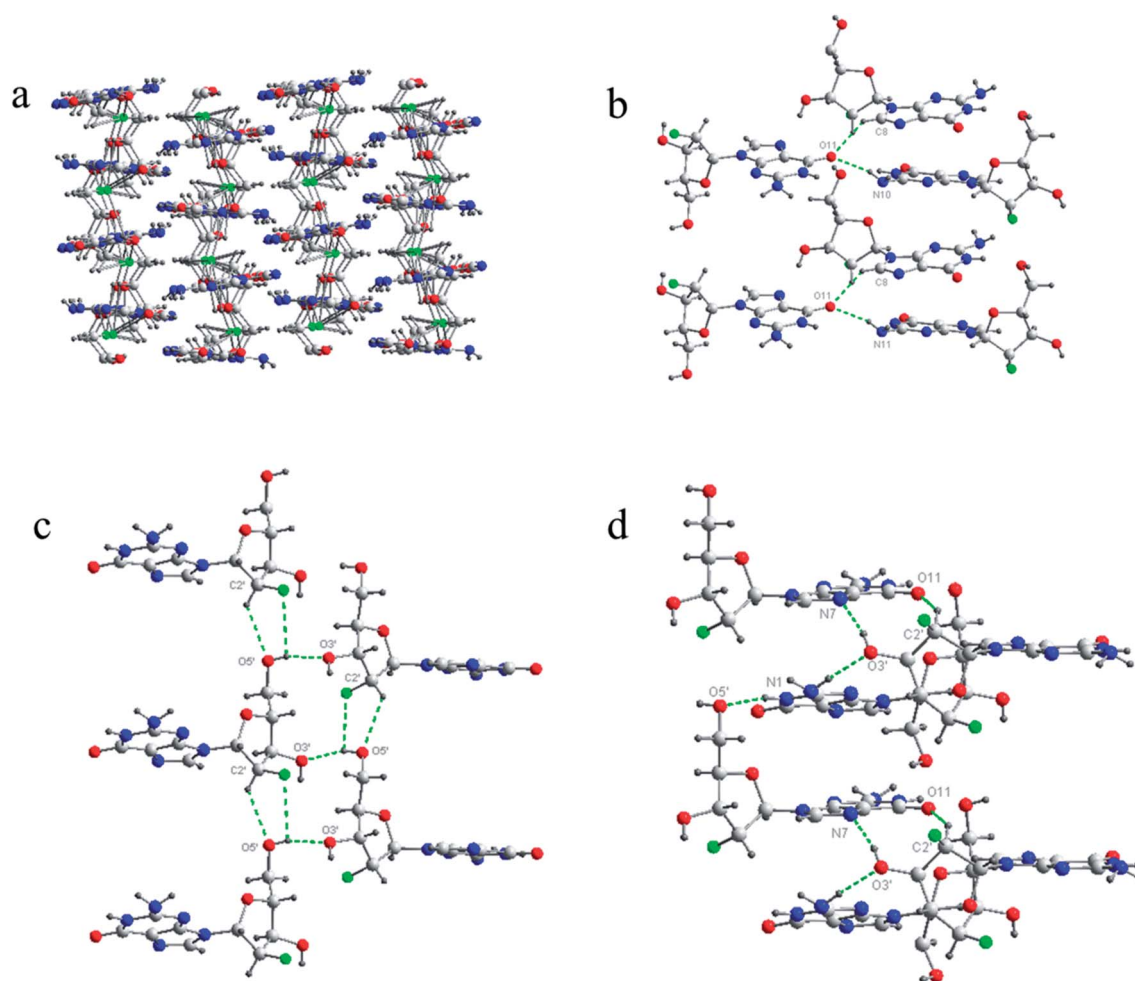


process was also monitored by SEM at micro level (Fig. 2b). The results indicated that the micron-scale slice-like structures gradually grew to piece-like and even colorless millimeter-scale granule-like structures while the gel transformed to a single crystal.

Furthermore, to get more detail information about the crystal from the hydrogel in molecular level, the single crystal structure of  ${}^F\text{G}_d$  has been analyzed carefully from three levels: monomer molecular, base pair motif and three-dimension structure. Firstly, the monomer molecular structure with systematic numbering and the single crystal structure of  ${}^F\text{G}_d$  were shown in Fig. 3. The *anti/syn* conformation of the glycosyl bond was paramount to the canonical purine or pyrimidine nucleosides, which could be defined by the torsion angle  $\chi$  ( $\text{O4}'\text{-C1}'\text{-N9-C4}$ ). The torsion angles  $\chi$  of  ${}^F\text{G}_d$  were measured to be  $-93.15(33)^\circ$ , indicating that it adopted a normal *anti* conformation. Sugar pucker is another significant conformational parameter defined by the pseudorotation phase angle ( $P$ ) and maximum pucker amplitude ( $\tau_m$ ). Two ranges of pseudorotation phase angle were initially observed in natural and

synthetic nucleosides:  $\text{C3}'\text{-endo}$  with  $0^\circ \leq P \leq 36^\circ$  (North) or  $\text{C2}'\text{-endo}$  with  $144^\circ \leq P \leq 180^\circ$  (South).  ${}^F\text{G}_d$  adopted a typical N-type conformation with a twist of  $\text{C3}'\text{-endo}$  ( ${}^3T_2$ ,  $N$ ,  $P = 15.6(4)^\circ$ ,  $\tau_m = 33.6(7)^\circ$ ). Moreover, it was also a crucial structural parameter for the orientation of the 5'-hydroxyl group defined by the torsion angle  $\gamma$  ( $\text{O5}'\text{-C5}'\text{-C4}'\text{-O4}'$ ) relative to the sugar ring. For  ${}^F\text{G}_d$ , it was in the *ap* (*gauche, trans*) range with the  $\text{O5B}'\text{-OH}$  at the axial position and pointing outside the sugar ring ( $\gamma = 61.98(33)^\circ$ ). Secondly, as  ${}^F\text{G}_d$  has the same base moiety as guanosine, which has three faces with hydrogen bonding donors and acceptors, it can self-assemble into complex and unique supramolecular structures, such as dimers, ribbons, and tetramers. In this case, the aforementioned structures were not obtained as the hydrogels broke down and it was only found that  ${}^F\text{G}_d$  could form mimic reverse Watson-Crick base pairs between two faces with an intermolecular hydrogen-bond ( $\text{N10H-O11}$ ), which extended to form a linear ribbon structure with sugar residues located on both sides.

Finally, based on the base pair motif, a multilayered structure was formed at the 3D supramolecular level with



**Fig. 4** Complicated hydrogen bond networks of  ${}^F\text{G}_d$ . (a) The overall multilayered supramolecular structure of  ${}^F\text{G}_d$ . (b) The interactions between bases. (c) The interactions between sugars. (d) The interactions between base and sugar. Nine hydrogen bond units repeated across the different layers in the entire assembly were highlighted in green color. Atoms were coded as follows: red, oxygen; blue, nitrogen; gray, carbon; green, fluorine; black, hydrogen.



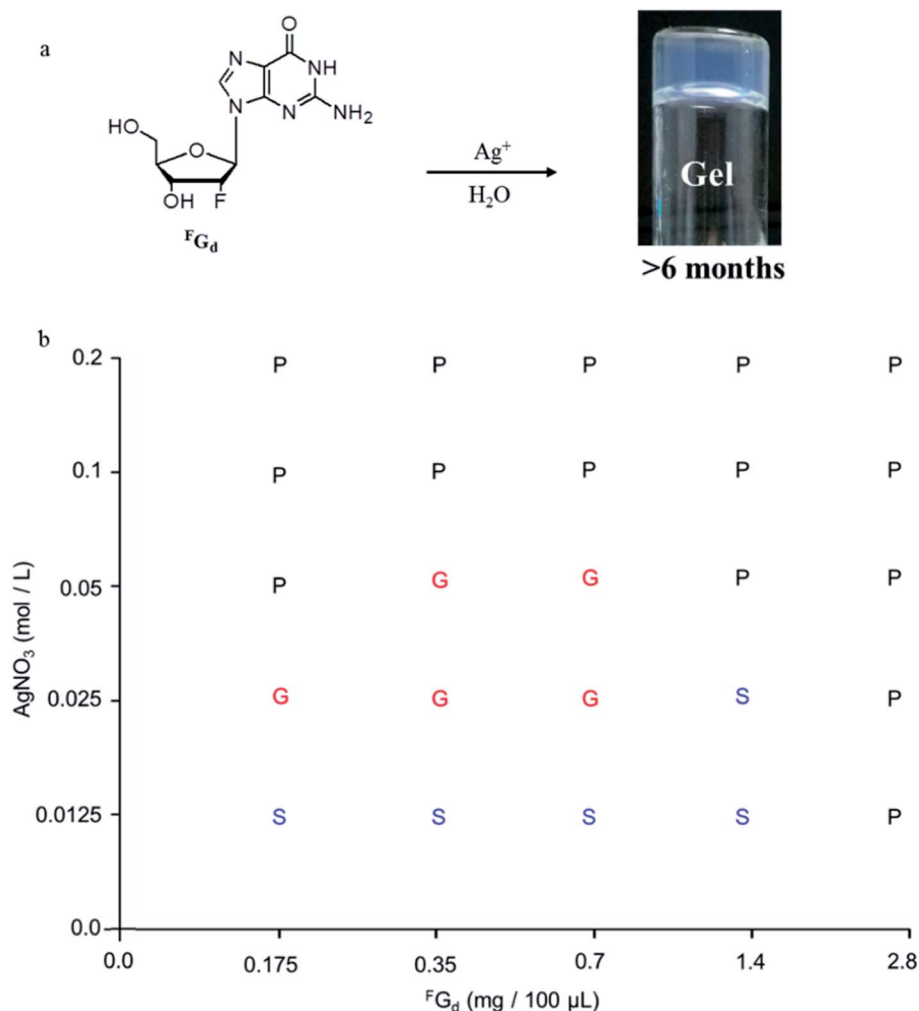


Fig. 5 Preparation and characterization of  ${}^F\text{G}_d\text{Ag}$  hydrogel in water. (a) The formation of a long-lasting stable supramolecular hydrogel *via* Ag<sup>+</sup>-mediated  ${}^F\text{G}_d$  in water at room temperature. (b) Phase diagram of gelation of  ${}^F\text{G}_d$  at different concentrations of aqueous AgNO<sub>3</sub>. G: gel, S: solution, P: precipitate.

complicated hydrogen bond networks including additional contributions from sugar residues (Fig. 4). To clearly display the hydrogen bond networks, the 3D supramolecular network was divided into three aspects: base–base, sugar–sugar, and base–sugar interactions of different layers. The interaction between base moieties is shown in Fig. 4b. O11 held three molecules from three neighboring layers together as a bridge with two hydrogen bonds, and it was connected to C8H of one molecule and N10H of another molecule. The sugar–sugar interactions are displayed in Fig. 4c. The C2'–H of one molecule was directly connected to the O5' of another molecule. 5'–OH linked together three molecules from three adjacent layers with two hydrogen bonds (2'F–5'OH–3'O). The interactions of base–sugar are shown in Fig. 4d. There were four hydrogen bonds between four molecules at three neighboring layers (5'O–N1H, N7–3'OH, 3'O–N10H, O11–C2'H). In general, a complicated hydrogen bond network formed with nine hydrogen bonds was repeated infinitely throughout the assembly process of the 3D multilayered supramolecular structure in the single-crystal state.

In order to explore their self-assembling properties in solution state, VT NMR spectrometry was used to investigate the formation of hydrogen bonds of  ${}^F\text{G}_d$ . As shown in Fig. S2,† the chemical shift of N10H atom of the amino group moved from  $\delta$  6.60 (298 K) to  $\delta$  6.46 ppm (338 K) with  $\Delta\delta = 0.14$  ppm, the chemical shift of 3'-OH atom of the hydroxyl group moved from  $\delta$  5.69 (298 K) to  $\delta$  5.53 ppm (338 K) with  $\Delta\delta = 0.16$  ppm, and the chemical shift of 5'-OH atom of the hydroxyl group moved from  $\delta$  5.15 (298 K) to  $\delta$  4.97 ppm (338 K) with  $\Delta\delta = 0.18$  ppm. These results indicated that these hydrogen atoms participated in the formation of intermolecular hydrogen bonds. Another hydrogen atom (N1H) from the imino group at  $\delta$  10.73 ppm was assigned as the intramolecular hydrogen bond as its chemical shift remained unchanged upon the increase in the temperature. To this end,  ${}^1\text{H}$ – ${}^1\text{H}$  nuclear overhauser effect (NOE) experiments was performed and they demonstrated that the C<sub>8</sub>–H $\cdots$ NH<sub>2</sub> and NH NOE signals (illustrated by the empty boxes, see ESI, Fig. S3†) were not observed, which was consistent with the solid state.



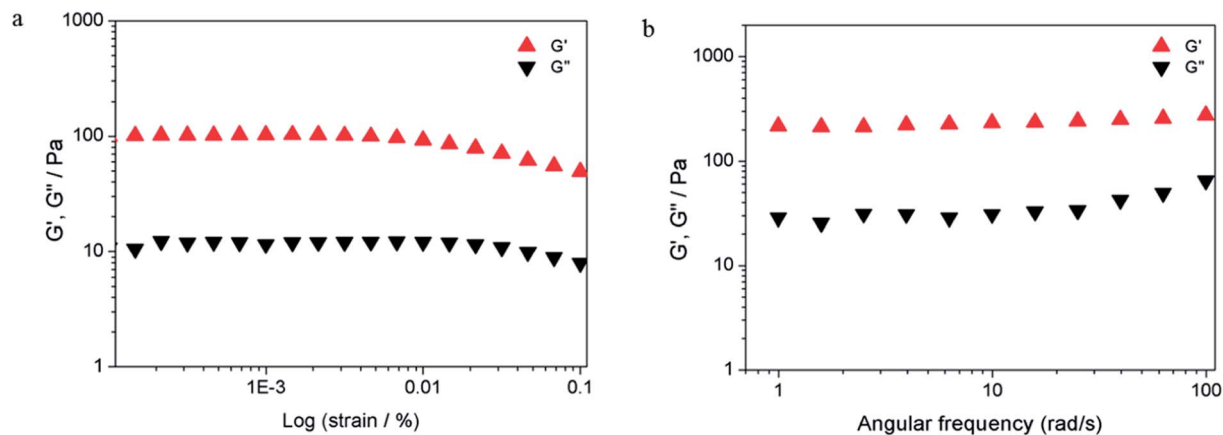


Fig. 6 Evaluation of  $G'$  and  $G''$  as a function of the strain (a) and frequency sweep (b). The compound was measured at the concentration of 0.7 mg per 100  $\mu\text{L}$  in 0.025 M  $\text{AgNO}_3$  solutions at 25  $^\circ\text{C}$ .

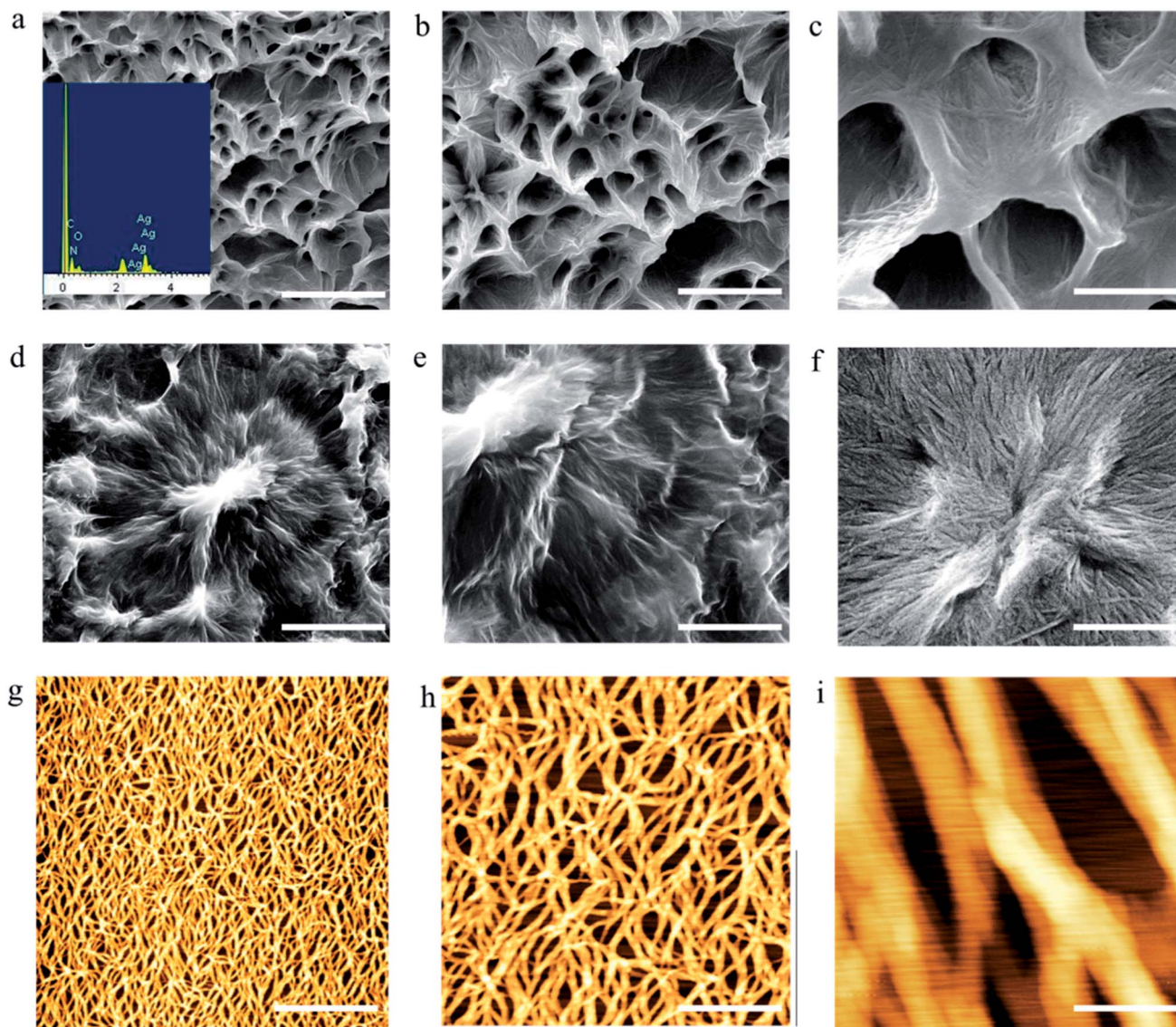


Fig. 7 Microstructures of  $\text{FGD}$  hydrogel at the concentration of 0.7% in 0.025 M  $\text{AgNO}_3$  solution. (a–f) SEM images of  $\text{FGD}$  hydrogel. (g–i) AFM images of  $\text{FGD}$  hydrogel. Scale bars: 5  $\mu\text{m}$  (a and d), 1.5  $\mu\text{m}$  (b and e), 500 nm (c and f), 1.5  $\mu\text{m}$  (g), 500 nm (h), and 50 nm (i).



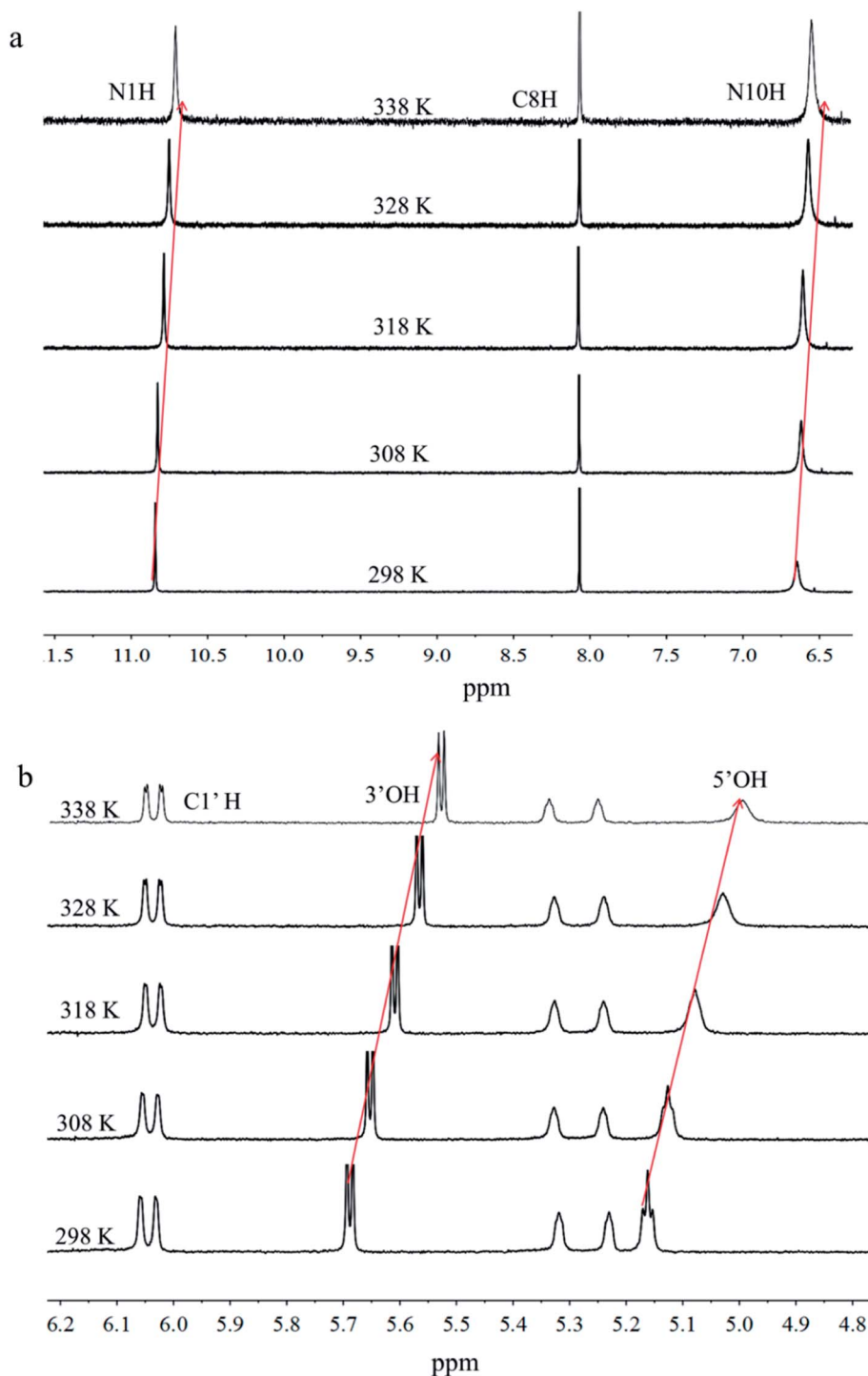


Fig. 8 VT  $^1\text{H}$  NMR spectra of  $^{\text{F}}\text{G}_d\text{Ag}$  recorded from 298 K to 338 K.

Both solid and solution results indicated that the  $\text{K}^+$ -introduced G-quartet structure was not stable and easily broke down in hydrogel and subsequently formed a linear ribbon structure with mimic reverse Watson–Crick base pairs between the two faces with an intermolecular hydrogen-bond (N10H–O11), which may be the reason for the propensity of guanosine-based hydrogel to crystallize over time. To block the crystallization of  $^{\text{F}}\text{G}_d$  hydrogel getting a long lifetime stable supramolecular

hydrogel, introducing stronger metal bonds would be a good choice. Previously, researchers determined that metal ions such as  $\text{Pt}^{2+}$ ,  $\text{Hg}^{2+}$ , and  $\text{Ag}^+$  tended to alter hydrogen bonding patterns between guanosine molecules by the coordination of the electron-rich nitrogen and oxygen groups around the purine ring to produce a range of metal ion-linked H-bonded architectures.<sup>34,46,47</sup> For example, Kraatz *et al.* reported that an  $\text{Ag}^+$  induced G gel was exploited for the light triggered *in situ*



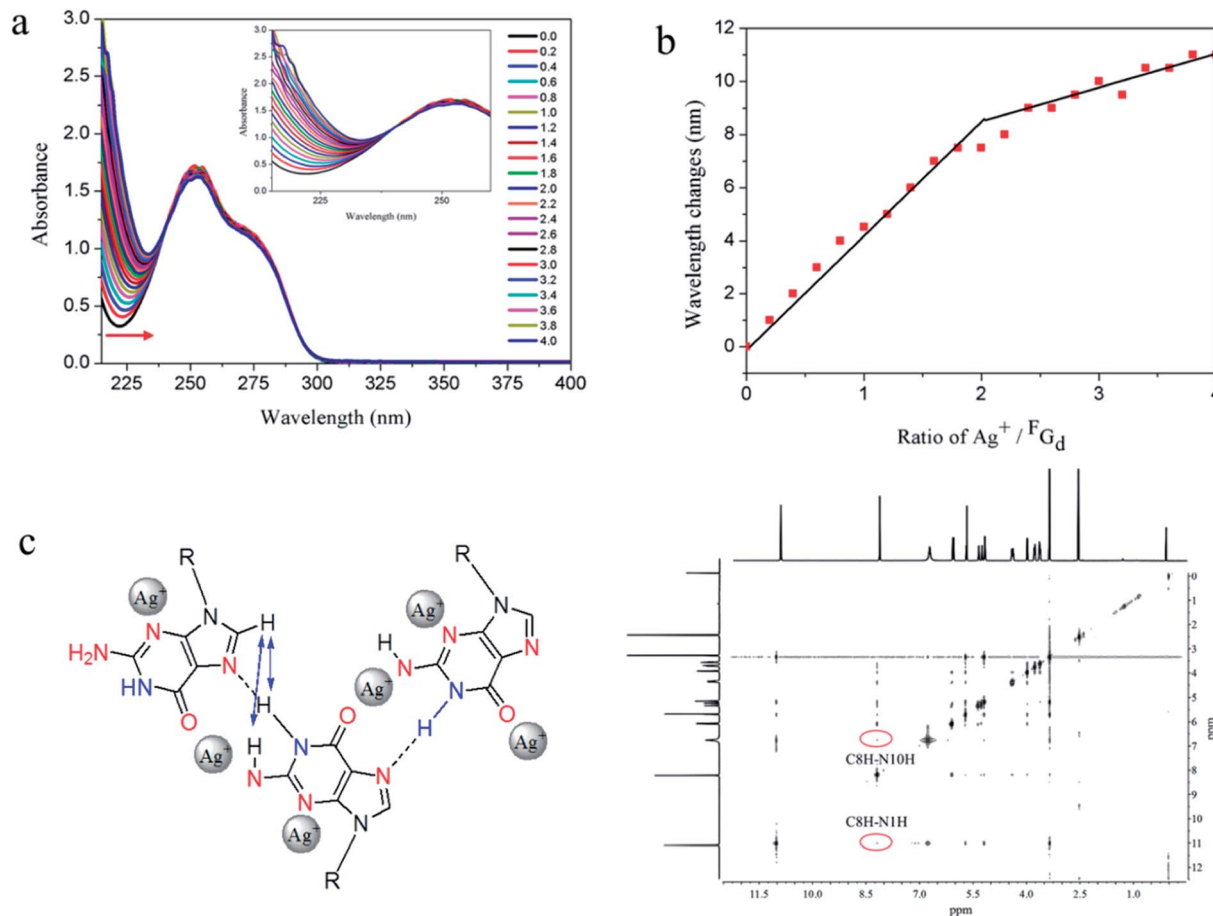


Fig. 9 Analysis and evaluation of possible silver-ions-mediated base pair in <sup>F</sup>G<sub>d</sub>Ag hydrogel. (a) UV spectrophotometric titration of 0.25 μM <sup>F</sup>G<sub>d</sub> with the increase in Ag<sup>+</sup> concentration measured in H<sub>2</sub>O. (b) Graphs of the ratio of silver ion/duplex versus absorbance changes measured at different wavelengths. (c) The proposed silver-mediated self-assembling structure of <sup>F</sup>G<sub>d</sub> and its <sup>1</sup>H-<sup>1</sup>H NOE spectra.

fabrication of uniform AgNPs within a gel to make a nano-bio hybrid material; Mann *et al.* found that supramolecular hydrogels produced by spontaneous self association of disodium guanosine 5'-monophosphate (Na<sub>2</sub>5'GMP) in the presence of Ag<sup>+</sup> ions. Inspired by this, Ag<sup>+</sup> was introduced in this work and the results showed that <sup>F</sup>G<sub>d</sub> can not only form hydrogel in the presence of silver ions but also demonstrate long lifetime stability (>6 months). Furthermore, <sup>F</sup>G<sub>d</sub> hydrogelation experiments were carried out at different concentrations of AgNO<sub>3</sub> and <sup>F</sup>G<sub>d</sub> (0.0125 M to 0.2 M AgNO<sub>3</sub> and 0.175 mg to 2.8 mg of <sup>F</sup>G<sub>d</sub> per 100 μL of solution). The phase diagram in Fig. 5 revealed that the formative qualification of the hydrogel. The concentration control between AgNO<sub>3</sub> and <sup>F</sup>G<sub>d</sub> was observed to play a critical role in the formation process of <sup>F</sup>G<sub>d</sub> hydrogel. Interestingly, <sup>F</sup>G<sub>d</sub> tended to form a hydrogel at low silver ion concentration, which suggested that the hydrogel had significant potential for biological or medical applications.

Furthermore, the mechanical properties of the hydrogel were analyzed in detail using rheological measurements. The storage modulus (measurement of elastic property) and loss modulus (measurement of fluidity) were expressed as  $G'$  and  $G''$ , respectively. As shown in Fig. 6, the typical elastic nature of <sup>F</sup>G<sub>d</sub> + Ag<sup>+</sup>

(<sup>F</sup>G<sub>d</sub>Ag) hydrogel was evident from the fact that  $G' > G''$  (solid-like behavior) *via* strain amplitudes ranging from 0.001 to 0.1% at 6.28 rad s<sup>-1</sup>. Moreover, the frequency sweep experiments of <sup>F</sup>G<sub>d</sub>Ag hydrogel were performed in the angular frequency range of 0.1–100 rad s<sup>-1</sup> under an initial strain of 0.1% and the results showed that <sup>F</sup>G<sub>d</sub>Ag hydrogel had a higher storage modulus  $G'$  than loss modulus  $G''$  over the entire applied frequency range, indicating that it exhibited a solid-like behavior.

To obtain microstructures of the <sup>F</sup>G<sub>d</sub>Ag hydrogel, we prepared xerogels for SEM observations (Fig. 7a–f). The hydrogels were prepared in a sample tube (0.7 w/v%) containing 0.025 M AgNO<sub>3</sub> and they were subsequently frozen. The frozen samples were lyophilized using a vacuum pump. Prior to examination, the xerogel was attached to the silica wafer and coated with a thin layer of gold. The SEM images revealed the three-dimensional uniform porous-like structures with a diameter of approximately 500 nm. Careful inspection of the images revealed that flexible fibers with a diameter of approximately 20–30 nm and several micrometers in length were formed spontaneously during the formation of hydrogel. The fibers intertwined with each other to form highly ordered film-like

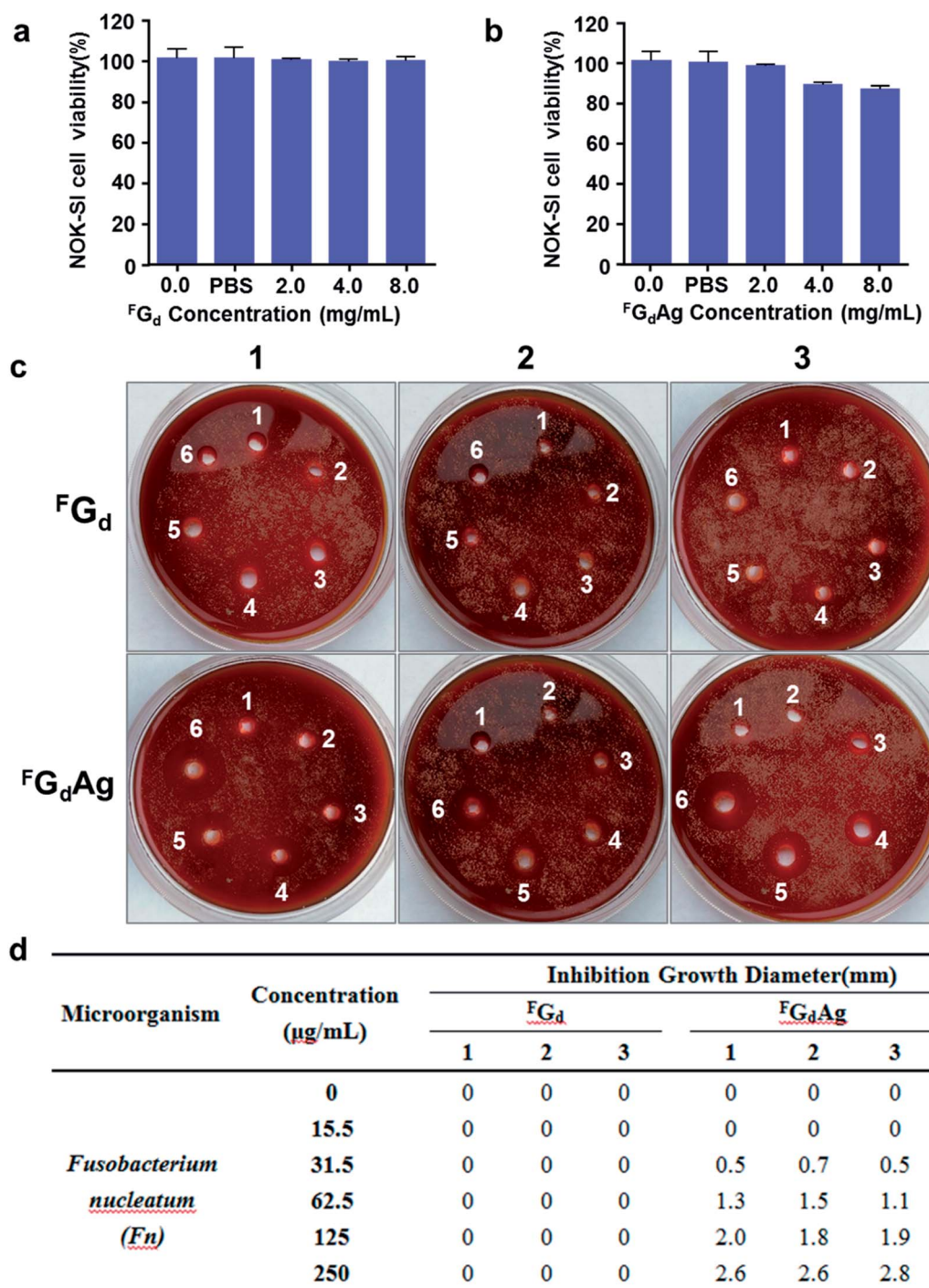




structures. Energy-dispersive X-ray spectroscopy analysis confirmed the presence of Ag in the fibers. AFM was carried out to further analyze and evaluate the structures of the hydrogel. Fig. 7g–i reveals the existence of an interconnected fibrous network with a diameter of approximately 20–30 nm, height of

6–8 nm, and length of several micrometers, which are consistent with the above results.

To explore more molecular-level evidences of the hydrogel, CD spectroscopy, PXRD, and NMR were carried out. Previous reports indicated that a planar quartet system show peaks at



**Fig. 10** Demonstration of low cytotoxicities and excellent antimicrobial efficiencies of  $\text{FG}_d$  and  $\text{FG}_d\text{Ag}$  *in vitro*. (a and b) CCK8 assay of the cytotoxicities of  $\text{FG}_d$  and  $\text{FG}_d\text{Ag}$  on immortalized normal oral keratinocyte cells (NOK-SI). The results represent mean  $\pm$  SD. (c) The antibacterial activities *in vitro* via evaluating the inhibition growth diameter of  $\text{FG}_d$  and  $\text{FG}_d\text{Ag}$  hydrogels against *Fusobacterium nucleatum*. In the sterilization experiment, (d) the concentrations of  $\text{FG}_d$  and  $\text{FG}_d\text{Ag}$  hydrogels were  $0 \mu\text{g mL}^{-1}$  (1);  $15.5 \mu\text{g mL}^{-1}$  (2);  $31.5 \mu\text{g mL}^{-1}$  (3);  $62.5 \mu\text{g mL}^{-1}$  (4);  $125 \mu\text{g mL}^{-1}$  (5) and  $250 \mu\text{g mL}^{-1}$  (6), respectively. The dates represent the growth inhibition diameter (in mm) of each sample in three independent experiments.



240 nm and 260 nm in CD spectrum.<sup>28</sup> Here, the <sup>F</sup>G<sub>d</sub>Ag hydrogel showed two negative peaks at 220 and 290 nm (see ESI, Fig. S4†), which indicated that the G-quartet structures were not observed in this hydrogel. As shown in Fig. S5,† the freeze-dried sample of <sup>F</sup>G<sub>d</sub>Ag hydrogel was studied using PXRD and it exhibited similar patterns for <sup>F</sup>G<sub>d</sub>Ag hydrogel and the crystal of <sup>F</sup>G<sub>d</sub> hydrogel in the presence of K<sup>+</sup>. All of them exhibited a significant peak at  $2\theta \approx 6.0^\circ$  and  $26.8^\circ$  ( $d = 14.5 \text{ \AA}$  and  $3.4 \text{ \AA}$ , respectively) consistent with the monomeric length and Pi–Pi stacking distance between two layers of G rings.<sup>31</sup> Subsequently, <sup>F</sup>G<sub>d</sub> and <sup>F</sup>G<sub>d</sub>Ag were characterized using <sup>1</sup>H NMR spectra at 298 K (see ESI, Fig. S6†). The results showed that N10H peak almost disappeared when Ag<sup>+</sup> was introduced to <sup>F</sup>G<sub>d</sub>, indicating that the amino group may participate in the silver base pair. VT NMR experiments were carried out to further investigate the formation of hydrogen bonds of <sup>F</sup>G<sub>d</sub>Ag. As shown in Fig. 8 and Table S1,† the chemical shift of N1H atom was from  $\delta$  10.84 (298 K) to  $\delta$  10.64 ppm (338 K) with  $\Delta\delta = 0.20$  ppm, the chemical shift of N10H atom of the amino group was from  $\delta$  6.65 (298 K) to  $\delta$  6.47 ppm (338 K) with  $\Delta\delta = 0.18$  ppm, and the peak began to increase in relative intensity when the temperature increased from 298 K to 338 K. The chemical shift of 3'-OH atom of the hydroxyl group moved from  $\delta$  5.69 (298 K) to  $\delta$  5.53 ppm (338 K) with  $\Delta\delta = 0.16$  ppm and the chemical shift of 5'-OH atom of the hydroxyl group moved from  $\delta$  5.16 (298 K) to  $\delta$  4.98 ppm (338 K) with  $\Delta\delta = 0.18$  ppm. These results demonstrated that these hydrogen atoms participated in the formation of intermolecular hydrogen bonds.

Stoichiometric titration experiments were performed to verify the formation of silver-mediated pairs and determine the amount of silver ions bound to the nucleoside <sup>F</sup>G<sub>d</sub>. As shown in Fig. 9, the changes in UV wavelengths were plotted *versus* the ratio of silver equivalents/<sup>F</sup>G<sub>d</sub>, suggesting that two silver ions were captured by one <sup>F</sup>G<sub>d</sub> (Fig. 9a and b). Accordingly, positive mode electrospray ionization-mass spectroscopy analysis was carried out and the corresponding silver complexes of 1 : 1 and 1 : 2 peaks at 475.1 and 624.1, respectively, were observed (see ESI, Fig. S7†), which provided powerful evidence further indicating that the two silver ions were captured by one <sup>F</sup>G<sub>d</sub>. Based on the above results, the possible silver-mediated base pair motifs are shown in Fig. 9c. To validate the assumption, <sup>1</sup>H-<sup>1</sup>H NOE experiments were carried out and the results suggested that C8-H produced a strong NOE at NH2a, NH2b, and NH simultaneously (Fig. 9c), which provided an unambiguous interpretation of the G-ribbon structures.

*Fusobacterium nucleatum* and *Porphyromonas gingivalis*, two Gram-negative anaerobes, are the most abundant microorganisms present in the oral cavity during periodontal disease.<sup>48,49</sup> In this study, the antibacterial activities of <sup>F</sup>G<sub>d</sub> and <sup>F</sup>G<sub>d</sub>Ag hydrogels *in vitro* were preliminarily evaluated. Initially, we assessed the biocompatibility of <sup>F</sup>G<sub>d</sub> (<sup>F</sup>G<sub>d</sub> dissolved in PBS) and <sup>F</sup>G<sub>d</sub>Ag hydrogels *in vitro*, the cell viability of the immortalized normal oral keratinocyte cells (NOK-SI) was assessed using CCK8 assay. As shown in Fig. 10, when the concentration of <sup>F</sup>G<sub>d</sub> and <sup>F</sup>G<sub>d</sub>Ag increased to  $2.0 \text{ mg mL}^{-1}$ , the viability of the NOK-SI cell line was higher than 97%. As the concentration of <sup>F</sup>G<sub>d</sub> and <sup>F</sup>G<sub>d</sub>Ag increased to  $8.0 \text{ mg mL}^{-1}$ , the viability of NOK-SI cell line was

higher than 86%. As expected, these results indicated that the <sup>F</sup>G<sub>d</sub> and <sup>F</sup>G<sub>d</sub>Ag hydrogels had only slight toxicity *in vitro* to be used as a safe biomaterial. Furthermore, the preliminary *in vitro* antimicrobial activities of <sup>F</sup>G<sub>d</sub> and <sup>F</sup>G<sub>d</sub>Ag hydrogels on *Fusobacterium nucleatum* and *Porphyromonas gingivalis* were evaluated by measuring the inhibition growth diameter and minimal bactericidal concentrations (MBCs). The results presented <sup>F</sup>G<sub>d</sub> and <sup>F</sup>G<sub>d</sub>Ag hydrogels had no obvious inhibition effect on *Porphyromonas gingivalis*. However, <sup>F</sup>G<sub>d</sub>Ag hydrogel exhibited excellent antibacterial activities for *Fusobacterium nucleatum* (MBC:  $31.25 \text{ \mu g mL}^{-1}$ ), compared to the control group, <sup>F</sup>G<sub>d</sub> hydrogel, which exhibited no antibacterial activities, silver ions as positive control (see ESI, Fig. S8†). In Fig. 10c and d, the diameters of *Fusobacterium nucleatum* inhibiting loops were approximately 0.57, 1.3, 1.9, 2.67 mm for <sup>F</sup>G<sub>d</sub>Ag hydrogel  $31.5$ ,  $62.5$ ,  $125$ , and  $250 \text{ \mu g mL}^{-1}$ , respectively. Therefore, these findings suggested that <sup>F</sup>G<sub>d</sub>Ag might be a possible biomaterial for antimicrobial medical applications.

## Conclusion

In summary, the detailed crystallization process of <sup>F</sup>G<sub>d</sub> hydrogel was studied using SEM. The results indicated that the micron-scale slice-like structure gradually grew to piece-like and even colorless millimeter-scale granule-like structure while the gel transformed to single crystal, which is, to the best of our knowledge, the first free guanosine-based nucleoside crystal formed from a hydrogel. The single crystal X-ray diffraction analysis indicated that <sup>F</sup>G<sub>d</sub> adopted an *anti*-conformation with an N-type (3'-*endo*) sugar puckering and 5'-OH at ap position, formed mimic reverses Watson–Crick base pairs with an intermolecular hydrogen-bond (N10H–O11), and had nine hydrogen bonds infinitely repeated in the entire assembly to form the 3D multilayered supramolecular structure in the solid state. These hydrogen bonds were also further investigated using VT NMR in solution state, and the results were consistent with the solid state. Then, Ag<sup>+</sup> was introduced to block its crystallization and form hydrogel with long lifetime stability (>6 months). Rheological measurements revealed that <sup>F</sup>G<sub>d</sub>Ag gel has a higher storage modulus  $G'$  than loss modulus  $G''$  over the entire applied frequency range (solid-like behavior). SEM and AFM studies demonstrated that flexible fibers with a diameter of approximately 20–30 nm, height measurements of approximately 7 nm, and length of several micrometers were formed spontaneously during the formation of <sup>F</sup>G<sub>d</sub>Ag gel whereas <sup>F</sup>G<sub>d</sub> formed a slice structure. Possible silver-mediated base pair motifs were suggested using PXRD, CD NMR, UV, and MS. Finally, <sup>F</sup>G<sub>d</sub>Ag hydrogel exhibited low toxicity for NOK-SI cell and good antibacterial activities for *Fusobacterium nucleatum in vitro*, whereas <sup>F</sup>G<sub>d</sub> exhibited no antibacterial activity. The above results suggested that <sup>F</sup>G<sub>d</sub>Ag might be useful for future applications in the field of antimicrobial medicines or drug delivery.

## Conflicts of interest

The authors declare no conflict of interest.



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