

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)

Cite this: *Polym. Chem.*, 2022, **13**, 5234

Effective pH-regulated release of covalently conjugated antibiotics from antibacterial hydrogels

Rita Das Mahapatra,^{†a} Ara Jo,^{†b} Kusuma Betha Cahaya Imani,^a Jin-Woong Chung^{*c} and Jinhwan Yoon ^{*a}

This paper describes the development of a pH-sensitive ampicillin-loaded double-network hydrogel for the local delivery of antibiotics against Gram-positive bacteria. Covalent polymerization of a cationic monomer constituted the first network, whereas imine bond formation between the aldehyde end group of Pluronic F127 (F127-CHO) and the amine of polyethyleneimine constituted the second network. The hydrogel was loaded with the antibiotic ampicillin. Ampicillin was conjugated by forming a reversible imine bond with F127-CHO, which allowed its sustained release over an extended period of time. The antibiotic's release was dependent on the initial amount of drug conjugated in the hydrogel and was greater at pH 5 than at pH 7. The hydrogel displayed excellent biocompatibility and antimicrobial activity against Gram-positive bacteria *in vitro*. Therefore, in the future, the proposed hydrogel could be used for *in situ* delivery of antibiotics in post-operative patients.

Received 20th April 2022,
Accepted 19th August 2022

DOI: 10.1039/d2py00505k

rsc.li/polymers

Introduction

Combating bacterial infections in vascular tissues (*e.g.*, bone infection) and necrotic tissues (*e.g.*, surgical site infections) is extremely challenging owing to the limited penetration of antibacterial drugs into the infection sites.^{1–3} Generally, treatment procedures include prolonged courses of antibacterial drugs administered orally or through intravenous injection over periods of weeks or months.⁴ Nevertheless, severe whole-body infection known as sepsis can emerge due to lack of effective antibiotic treatment at the infection site.^{5,6} Owing to the occurrence of bacterial resistance at the target site, a higher dose of antibiotic is sometimes required; however, this can have lethal side effects.^{7,8} Therefore, the development of biocompatible biomaterials with antibacterial properties could dramatically reduce the side effects of excessive drug loads at the target site by enabling instead their slow and sustained release. To this end, antibacterial hydrogels offer several advantages as their

high-viscosity matrix allows the controlled and local release of therapeutics.^{9–26} Moreover, hydrogels are characterized by elevated biocompatibility, soft mechanical properties, high water content, and high capacity for absorption. In general, antibacterial hydrogels are fabricated from polymers with antibacterial properties,^{27,28} through incorporation of specific drugs, metal nanoparticles or a combination of the two.^{4,29,30} While conjugation of antibiotics in the hydrogel matrix is highly effective against bacteria, the rapid release of antibiotics limits their long-term effectiveness.

Various strategies have been proposed for the long-lasting, sustained release of antibiotics, including manipulation of gel porosity and microstructure for slow diffusion,²⁰ slow degradation for hydrogel matrix-assisted release,²¹ squeezing of antibiotics by a stimuli-responsive volume change of the hydrogels,^{22,23} and covalent conjugation of the hydrogel network with degradable linkers.^{24–26} Covalent conjugation of antibiotics in hydrogels has received growing attention owing to its long-lasting and controlled release of antibiotic through the dissociation of covalent bonds at a specific pH. Haldar *et al.* developed vancomycin-loaded injectable hydrogels to deliver antibiotics locally.⁴ The hydrogel was developed by mixing polydextran aldehyde with amine-functionalized chitosan through reversible imine bond formation. Vancomycin was conjugated covalently with the hydrogel network and showed sustained release over an extended period in a pH-dependent manner. Stewart *et al.* reported the sustained release of tobramycin, a polycationic aminoglycoside anti-

^aGraduate Department of Chemical Materials, Institute for Plastic Information and Energy Materials, Sustainable Utilization of Photovoltaic Energy Research Center, Pusan National University, Busan, 46241, Republic of Korea.

E-mail: jinhwan@pusan.ac.kr

^bDepartment of Molecular Biology and Immunology, College of Medicine, Kosin University, Busan, 49267, Republic of Korea

^cDepartment of Biomedical Science, Dong-A University, 37 Nakdong-Daero 550 beon-gil, Saha-gu, Busan, 49315, Republic of Korea. E-mail: jwchung@dau.ac.kr

[†]These authors contributed equally to this work.

biotic, from a hydrogel over a period of 50 days at pH 6–8.¹⁹ Because infection sites are generally acidic in nature,⁴ efficient antimicrobial hydrogels should be designed for the controlled release of antibiotics under acidic conditions. While there is a previous report regarding the pH-dependent release of antibiotics for only injectable hydrogels,⁴ to the best of our knowledge, no one reports mechanical robust double network hydrogel which show the controlled release of antibiotics in the acidic pH.

The present study aimed to develop efficient, long-lasting antibacterial hydrogels, whose mechanical robustness was based on a double-network structure that enabled the covalent conjugation of antibiotics. The first network originated from the covalent polymerization of positively charged monomers, whereas the second network was attributed to imine bond formation between the aldehyde end of modified Pluronic F127 (F127-CHO) and the primary amine of polyethyleneimine (PEI). Accordingly, the first network maintained the mechanical properties of the hydrogel over time, which is particularly important in infections associated with wound healing, whereas the second network helped release the antibiotic at the target site in a pH-dependent manner. Ampicillin (Amp), a highly effective broad-spectrum antibiotic, was selected for this study. Because it contains a free primary amine group, it can be covalently conjugated in the hydrogel matrix through imine bond formation. Covalent conjugation allowed the sustained release of the antibiotic in a pH-dependent manner and was highly effective at killing bacteria *in vitro*.

Experimental

Materials

Acrylamide (AAM) and *N,N'*-methylenebisacrylamide (BisAA) were purchased from Bio Basic Inc. (Markham, ON, Canada). All other chemicals, including PEI ($M_w = 25\,000\text{ g mol}^{-1}$), Pluronic F127 ($M_w = 12\,600\text{ g mol}^{-1}$), ammonium persulfate (APS), *N,N,N',N'*-tetramethylethylenediamine (TEMED), and 2-(dimethylamino)ethyl methacrylate (DMAEMA), were purchased from Sigma-Aldrich (St. Louis, MO, USA).

Synthesis of cationic dimethylamino methacrylate monomer (DMA-6)

Briefly, 1 g DMAEMA (1 eq., 6.36 mmol) in 3.49 mL 1-chlorohexane (4 eq.) was mixed with 5 mL dimethyl formamide (DMF) in a single-neck round-bottom flask. The reaction mixture was refluxed for 24 h at 160 °C. The reaction was stopped, and once the mixture cooled to room temperature, ethyl acetate was added to obtain a white precipitate. The precipitate was collected *via* filtration and washed with hexane three times to remove unreacted 1-chlorohexane. The product was then transferred to a vacuum oven and dried.

The chemical shifts (δ) obtained from the proton nuclear magnetic resonance (¹H NMR) spectrum (400 MHz, CDCl₃) of the synthesized DMA-6 was as follows: 0.86 ppm (3H, CH₃CH₂CH₂CH₂), 1.33 ppm (6H, CH₃CH₂CH₂CH₂), 1.74 ppm (2H, CH₂CH₂N(CH₃)₂), 1.96 ppm (3H, CH₃C=CH₂), 3.59 ppm (6H, N(CH₃)₂), 3.63 ppm (2H, CH₂CH₂N(CH₃)₂), 4.16 ppm (2H,

N(CH₃)₂CH₂CH₂CO), 4.64 ppm (2H, N(CH₃)₂CH₂CH₂CO), as well as 5.66 and 6.13 ppm (2H, CH₂=C(CH₃)).

Synthesis of F127-CHO

First, 6 g Pluronic F127 was dissolved in 40 mL dry dichloromethane (DCM) in a two-neck round-bottom flask under a N₂ atmosphere. Then, an excess of triethylamine (10-fold molar ratio) was added to the mixture, and the temperature of the flask was brought to 0–5 °C using an ice bath. After 10 min, methanesulfonyl chloride (four-fold molar ratio) dissolved in a small amount of dry DCM was added in a dropwise fashion. The ice bath was removed, and the mixture was stirred at room temperature for an additional 24 h. Next, the reaction mixture was washed with water five times to remove side products, and the organic fraction was collected and concentrated under vacuum. The concentrated mixture was precipitated over chilled diethyl ether; the product was collected by centrifugation and then dried in vacuum for 1 day at 40 °C.

The obtained white product was dissolved in dry DMF with K₂CO₃ (5-fold molar ratio) and 4-hydroxybenzaldehyde (three-fold molar ratio). The reaction mixture was refluxed at 80 °C for 48 h and washed with water and DCM five times to remove side products, and the organic fraction was collected and concentrated under vacuum. This concentrated mixture was then precipitated over chilled diethyl ether, and the product was collected by centrifugation. Based on ¹H NMR spectra (400 MHz, CDCl₃), the degree of end-functionalization was calculated to be greater than 90%: $\delta = 9.88\text{ ppm}$ (2H, CHO), $\delta = 7.81\text{ ppm}$ (4H, aromatic ring), $\delta = 7.03\text{ ppm}$ (4H, aromatic ring), and $\delta = 1.14\text{ ppm}$ (183H, PPG methyl protons).

Preparation of hydrogels

To prepare hydrogels, F127-CHO (55 mg mL⁻¹) and BisAA (5.88 mg mL⁻¹) were dissolved in buffer at pH 7. Separately, AAm (54 mg mL⁻¹), DMA-6 (63 mg mL⁻¹), and PEI (17.64 mg mL⁻¹) were also mixed in buffer at pH 7. Both solutions were degassed to remove any dissolved oxygen and then mixed under N₂ atmosphere with concomitant addition of APS and TEMED. The solution was quickly transferred into a mold and kept under N₂ atmosphere for 1 h to obtain transparent hydrogels. To prepare the ampicillin-loaded hydrogels, ampicillin at different concentrations (5 or 10 mg mL⁻¹) was mixed with F127-CHO (F127-CHO : AMP = 1 : 3 for 5 mg AMP and 1 : 6 for 10 mg AMP), and the solution was left for 15 min at 37 °C to ensure complete conjugation of ampicillin with F127-CHO. After gel formation, the hydrogel was immersed in water for 3–4 days to remove any unreacted monomers.

Characterization

The equilibrium swelling ratio (SR) was measured for lyophilized hydrogel and determined by the following equation:

$$SR = m_{eq} - m_0 / m_0,$$

where m_{eq} and m_0 are mass at equilibrium and initial state, respectively.

Fourier-transform infrared (FT-IR) spectra of the hydrogels were obtained using an Agilent Cary 630 FT-IR spectrometer (Agilent Technologies, Santa Clara, CA, USA). The samples were freeze-dried prior to the measurement. NMR measurements were conducted in CDCl_3 using a Bruker Avance Neo 500 MHz spectrometer (Bruker, Billerica, MA, USA). Field emission scanning electron microscopy (FE-SEM) images of the dried hydrogels were captured using a Sigma microscope (Zeiss, Oberkochen, Germany). The samples were sputtered with gold, and images were captured. A rotational rheometer AR G2 (TA instrument, New Castle, USA) having a parallel plate geometry with a diameter and gap of 20 mm 0.55 mm respectively was used to perform the rheological measurements. A solvent trap was used to prevent water evaporation during rheological measurements.

Agar diffusion test

Antimicrobial activity was tested in accordance with the Clinical Laboratory Standards Institute manual using disk diffusion and growth curve measurements. *Bacillus subtilis* Gram-positive bacteria were grown on cation-adjusted Mueller-Hinton broth (CAMHB) agar, from which bacterial colonies were selected using a sterile inoculating loop and suspended in 10 mL sterile broth medium for overnight incubation at 35 °C. On the next day, when the cultures' optical density at 600 nm (OD_{600}) reached 0.4–0.5, bacteria were spread on dried agar plates. Hydrogels were placed on the surface of the agar plates, and they were incubated for 24 h at 35 °C. Antibacterial activity was measured by evaluating the bacterial growth inhibition zone around the hydrogel.

Bacterial growth curve and colony forming unit (CFU) determination

Bacteria were grown in 100 mL CAMHB medium at 35 °C starting at $\text{OD}_{600} = 0.1$. The inoculum density was determined every 3 h for a total of 24 h by measuring OD_{600} using a UV/visible spectrophotometer. All experiments were performed in triplicates.

For CFU determination, bacteria were cultured overnight at 35 °C in a continuous shaking incubator at 180 rpm. The next day, hydrogel samples were added at a ratio of 1 g per 50 mL bacterial solution. Samples were incubated at 35 °C with shaking for 24 h. Every 3 h, OD_{600} was measured using a microplate reader. The bacterial solution (100 μL) was diluted with 900 μL culture medium in serial 10-fold dilutions from 10^{-1} to 10^{-7} . The diluted bacterial suspension (100 μL) was spread onto broth agar medium, and colonies were counted after incubation for 24 h at 35 °C. The following equation was used: $\text{CFU/mL} = (\text{number of colonies/dilution factor})/\text{volume of culture solution}$. The dilution factor was calculated as the ratio of the bacterial suspension's volume after dilution vs. its volume before dilution.

Cell culture and proliferation assay

NIH3T3 mouse embryonic fibroblasts, human dermal fibroblasts, and newborn human foreskin fibroblasts were pur-

chased from the American Type Culture Collection (Manassas, VA, USA). All cell lines were cultured in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum and 1% penicillin/streptomycin (100 U mL^{-1}) and maintained at 37 °C in a humidified atmosphere containing 5% CO_2 .

The cytotoxicity of the hydrogel was measured using the methyl thiazolyl tetrazolium (MTT) cell proliferation assay. The hydrogel was under the same conditions as those used for the disk diffusion experiments. Cells were seeded in 12-well plates at a density of 1×10^5 cells per well. The following day, the hydrogel was added to the medium, and cells were cultured for 24 h at 37 °C in a humidified atmosphere in a CO_2 incubator. After incubation, the cells were observed under a microscope, the hydrogel was removed, the medium was replaced with 10 μL MTT solution at 5 mg mL^{-1} in $1 \times$ phosphate-buffered saline, and the cells were incubated for 4 h at 37 °C. MTT-formazan was dissolved in 100 μL dimethyl sulfoxide, and absorbance at 550 nm (A_{550}) was measured using a microplate reader. Cell viability (%) was calculated as $(A_{550} \text{ of treated cells}/A_{550} \text{ untreated cells}) \times 100$.

Results and discussion

Fig. 1 illustrates the synthetic procedure for the various components of double-network antibacterial hydrogels. The resulting hydrogel comprised a positively charged cross-linked polymer chain as the first network and covalently-linked Pluronic F127 and PEI as the second network. The positively charged DMA-6 monomer was chosen because of its inherent antibacterial properties, whereas F127-CHO and PEI were selected because of their biocompatibility and amine functionality, respectively. Branched PEI has been applied in biomedicine as a nanoparticle coating for drug and gene delivery.²⁷ As PEI contains many free amines that remain in the protonated form at physiological pH, it can bind to the negatively charged bacterial cell wall, resulting in cell membrane disruption and lysis.

DMA-6 was synthesized by reacting DMAEMA with 1-chlorohexane under reflux condition for 24 h in DMF (Fig. 1a). F127-CHO was synthesized from commercially available Pluronic F127 using a two-step method (Fig. 1b). First, Pluronic F127 was treated with methanesulfonyl chloride, followed by reaction with 4-hydroxybenzaldehyde in DMF. Second, it was reacted with 4-hydroxybenzaldehyde in dry DMF in the presence of K_2CO_3 .

The double-network resulted from the covalent polymerization of DMA-6, AAm, and BisAA as the first network and imine bond formation between the aldehyde of F127-CHO with primary amines of PEI as the second network (Fig. 1c). The total polymer concentration was 30% (w/w), and the first to second network ratio was 70 : 30. To prepare the hydrogels, a solution containing DMA-6, AAm, and PEI was mixed with a second solution containing F127-CHO and BisAA. After APS and TEMED were added to initiate free radical polymerization, the mixture was transferred to a mold and maintained in a N_2

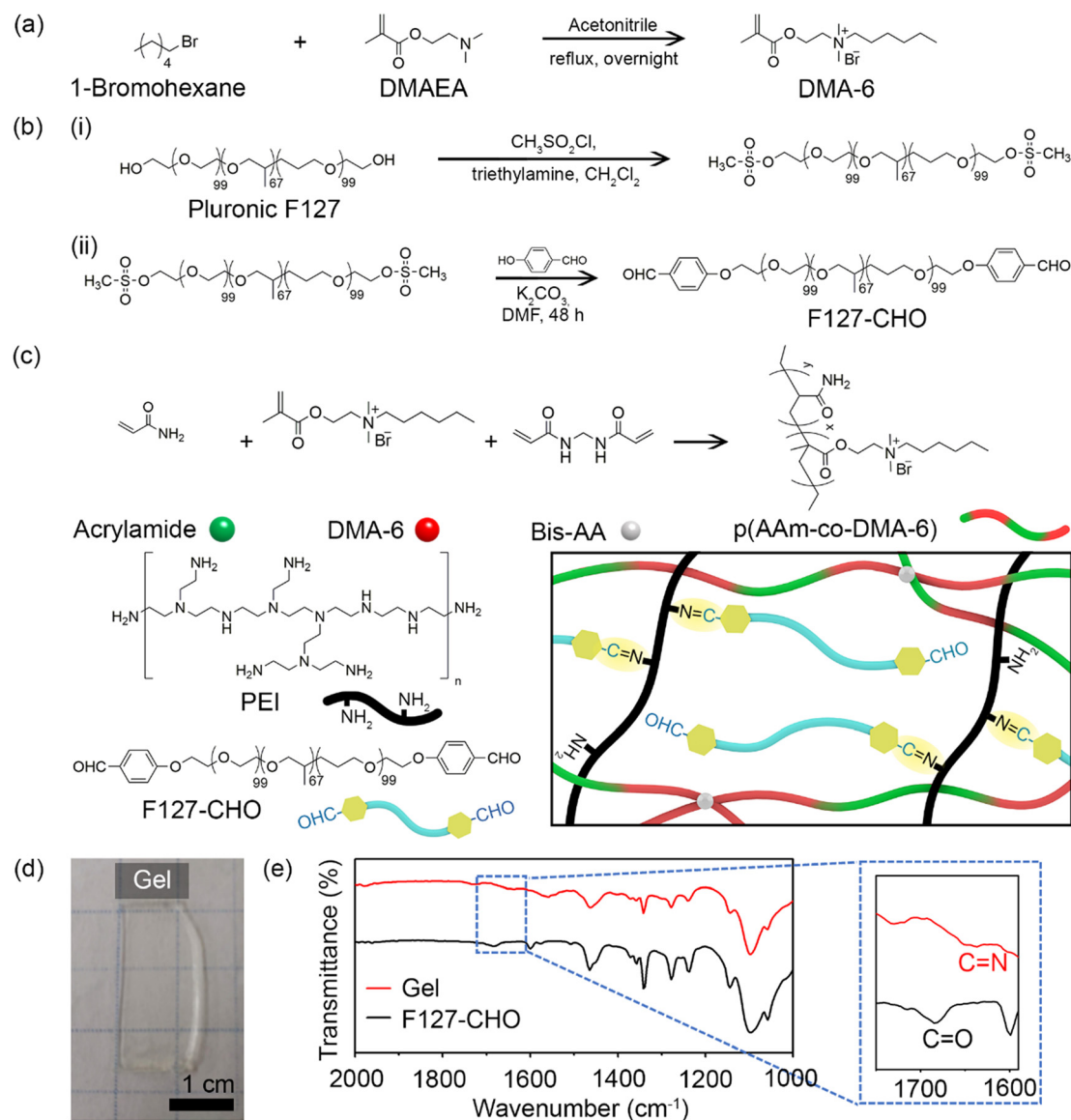


Fig. 1 Schematic representation of the synthesis of (a) DMA-6 and (b) F127-CHO. (c) Illustration of the double-network structure of the gel. (d) Photograph of the prepared gel. (e) FT-IR spectra of the gel and F127-CHO.

atmosphere for 1 h until optically transparent hydrogels were obtained (Fig. 1d). Because DMA-6 alone formed a weak hydrogel, 8% (mol mol⁻¹) AAm was introduced to fabricate a much stronger hydrogel capable of maintaining structural integrity for several months. The swelling ratio of the hydrogel was measured to be 5.6.

To confirm imine bond formation between the aldehyde and amine groups, the F127-CHO solution and hydrogel were analyzed by FT-IR. As can be seen in Fig. 1e, the F127-CHO solution showed a peak at 1684 cm^{-1} , corresponding to the C=O in aldehyde group, but was shifted to 1650 cm^{-1} after gel formation with PEI. The peak at 1650 cm^{-1} corresponded to the C=N imine bond formed during hydrogel cross-linking.

As the final goal of this study was to develop effective long-lasting hydrogels with antibacterial properties, the free

primary amine group of ampicillin was allowed to form a reversible covalent imine bond with F127-CHO. This bond led to covalent conjugation of the drug in the gel network. Two different ampicillin concentrations (5 and 10 mg mL⁻¹) were used, and ampicillin was dissolved in DMA-6 during hydrogel formation.

To form the covalent imine conjugate, F127-CHO (50 mg mL⁻¹) and ampicillin (5 mg mL⁻¹) were mixed and incubated at 37 °C for 1 h. The solution was then dialyzed for 24 h at 4 °C in deionized water, with water changes at regular intervals to remove any non-conjugated ampicillin. The polymer-antibiotic solution was then lyophilized, and ¹H NMR spectra were recorded with D₂O to confirm the covalent coupling of ampicillin with F127-CHO. The ¹H NMR spectrum of ampicillin was recorded for comparison, revealing that the peaks in the aro-

matic region (5.0–8.0 ppm) exhibited by the polymer–drug corresponded to ampicillin peaks along with other protons from F127-CHO (Fig. 2a). This result clearly indicated that ampicillin was bound to F127-CHO *via* imine conjugation. The covalent conjugation of ampicillin with the hydrogel network was further confirmed by FT-IR spectra, which revealed a stronger peak at 1652 cm^{-1} (C=N stretching) for the drug-polymer (Fig. 2b).

Given that some double-network hydrogels are chemically cross-linked, permanent damage to the network causes irreversible softening of the gels and poor fatigue resistance.^{28,29,31} To overcome this issue, non-covalent, reversible bonds are introduced. Here, a reversible imine bond was introduced in the second network, and the viscoelastic properties of the hydrogels were determined by rheological experiments.

As shown in Fig. 2c, the hydrogel without ampicillin exhibited a G' value of $750 \pm 50\text{ Pa}$; whereas addition of 5 and 10 mg ampicillin increased the G' value to $1100 \pm 100\text{ Pa}$ and $2000 \pm 180\text{ Pa}$, respectively. Similarly, measurements of the storage modulus (G') and loss modulus (G'') indicated that the G' and G'' values were higher for ampicillin-loaded hydrogels than unloaded hydrogels (Fig. 2c). This behavior can be attributed to the increased density in ampicillin-loaded hydrogels. Notably, the G' value was much higher than the G'' value throughout the frequency range ($1\text{--}100\text{ rad s}^{-1}$) for all hydrogel formulations, indicating strong hydrogel formation (Fig. 2d). These results clearly suggest that the hydrogel network tends to become stronger with increasing ampicillin concentration. Moreover, ampicillin owing to the presence of amide and carboxylic acid functionality in its chemical structure can

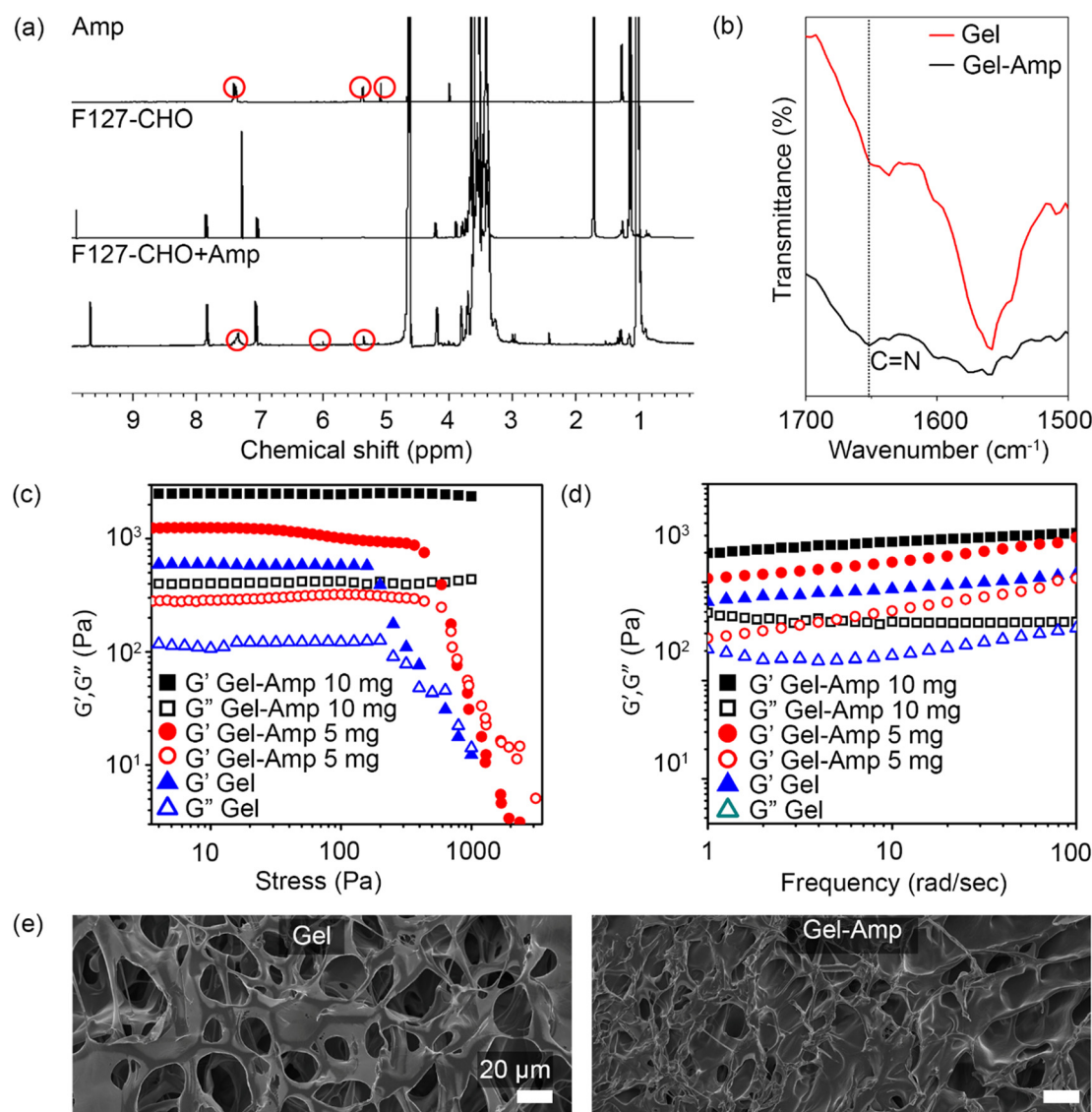


Fig. 2 (a) $^1\text{H-NMR}$ spectra of free ampicillin (AMP), F127-CHO and ampicillin-contained F127-CHO recorded using D_2O as the solvent. (b) FT-IR spectra of the hydrogel and ampicillin-containing hydrogel. (c, d) Storage modulus (G') and loss modulus (G'') of hydrogels as a function of (c) mechanical stress and (d) frequency. (e) SEM images of hydrogel and ampicillin-containing hydrogel.

undergo H-bonding bonding interactions with the hydrogel network.^{32,33} This additional H-bonding interactions can affect the overall mechanical properties of the hydrogels.

The surface morphology of hydrogels with and without ampicillin was observed by FE-SEM. Following air-drying, the hydrogels without ampicillin showed a porous structure with large pores, which became smaller and denser in the case of ampicillin-loaded hydrogels (Fig. 2e). This denser network can be attributed to the conjugation between the free amines of ampicillin and the aldehyde of F127-CHO, explaining also the results obtained in the rheological experiments, whereby hydrogels became stronger as the amount of ampicillin was increased. Overall, these findings clearly indicated that ampicillin was successfully conjugated in the hydrogel network.

As previously mentioned, the antibiotic was covalently bound to the hydrogel network to allow its release over time and thereby, ensure long-lasting antibacterial activity. As illustrated in Fig. 3a, in an acidic medium (pH 5), the imine bonds between ampicillin and F127-CHO, as well as between PEI and F127-CHO, broke down, resulting in antibiotic release.

Moreover, this is to notify that the pK_a values for carboxylic acid and the amines are 3.24 and 7.44 respectively hence, ampicillin will exist in zwitterionic state in the pH range of pH 3.0 and 7.5. Hence, the drug release would remain same in our experimental pH values.³⁴

To attain the controlled release of antibiotic under acidic conditions, we determined the release kinetics of the prepared hydrogels loaded with different amounts of drug (5 and 10 mg) and at pH 5 and pH 7. Hydrogels (0.4 mL) were placed in Eppendorf tubes and covered with 1 mL buffer of different pH.

The hydrogels with the buffer were incubated at 37 °C with constant shaking for 24 h, after which the upper part of the buffer was collected for UV-visible spectroscopy measurements and replaced with fresh buffer. This process was repeated every day for 7 days, and the amount of ampicillin released was quantified based on absorbance at 257 nm.

As shown in Fig. 3b, ampicillin was successfully released from the hydrogels for over 7 days in both pH groups. Ampicillin release was rapid at the initial stage and slowed down over time for all samples in a manner that depended on

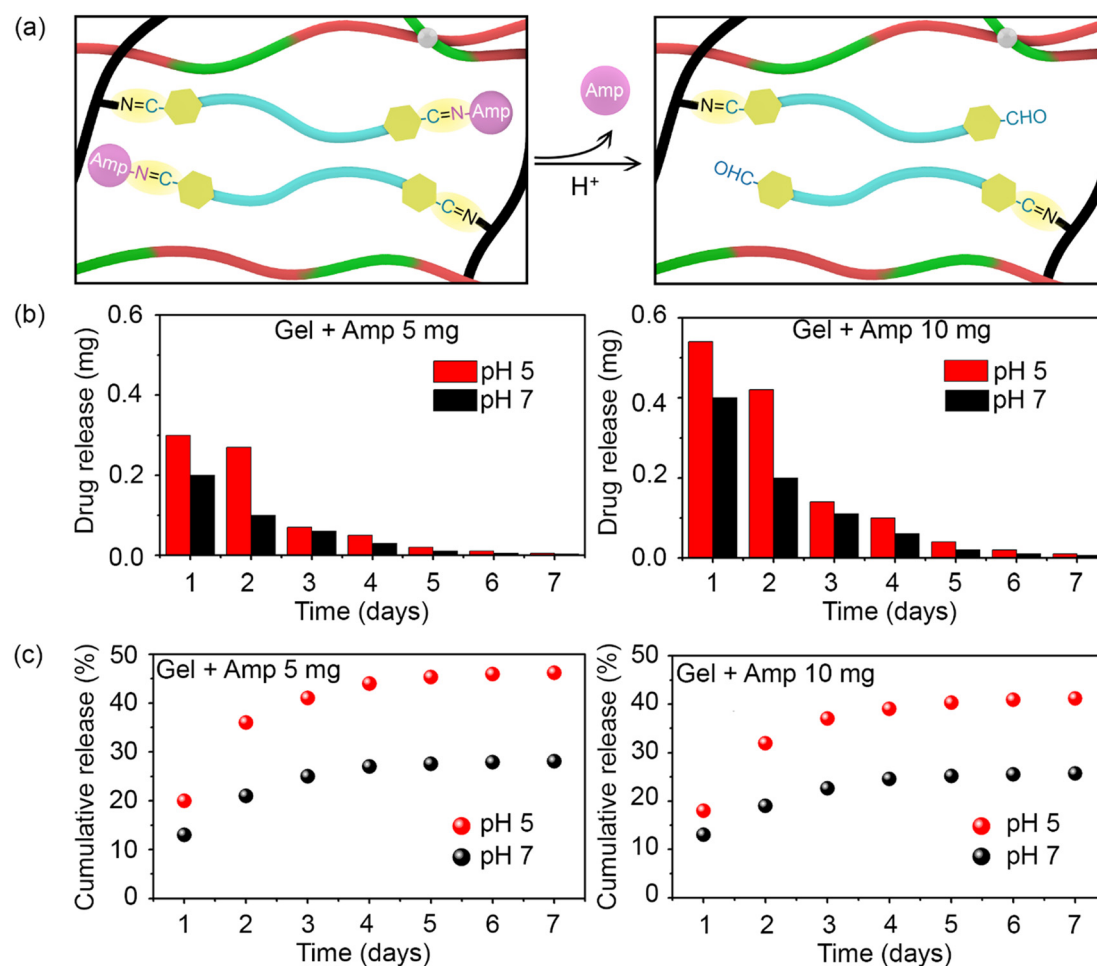


Fig. 3 (a) Illustration of ampicillin release from encapsulation in the hydrogel network upon acidification of the medium. (b) Ampicillin release rate from the hydrogel with respect to antibiotic concentration and medium pH. (c) Cumulative release of ampicillin from the hydrogel with respect to antibiotic concentration and medium pH.

the initial ampicillin concentration and medium pH. The release was always higher at pH 5 than at pH 7 because the acidic buffer broke the imine bonds between the antibiotic and F127-CHO in the hydrogel network. Interestingly, the ampicillin release rate was much higher during the first and second days, tapering off after that at both pH values.

Based on the detected amount of ampicillin, its cumulative release was plotted. A higher cumulative release was measured at pH 5 than at pH 7 for both concentrations of ampicillin (Fig. 3c). Over 7 days, almost 45% and 40% of overall ampicillin present in the hydrogels at 10 mg and 5 mg, respectively, was released. In comparison, only half this amount of drug was released at pH 7. These results showcase how hydrogels in which ampicillin is conjugated through imine bonds can be

used for the controlled and prolonged release of antibiotic under acidic conditions.

To confirm the antibacterial activity of the prepared hydrogels, an agar diffusion test was carried out with the Gram-positive bacterium *B. subtilis*. When material with antibacterial activity is placed on an agar-grown bacterial culture, bacterial growth is inhibited by the diffused antibiotic. The antibiotic activity was determined by measuring the diameter of the inhibition zone around the hydrogel. Unloaded hydrogels and hydrogels loaded with 5 and 10 mg ampicillin produced clear zones of 1.3, 2.6, and 4.3 mm in diameter, respectively (Fig. 4a). Even though a minor inhibitory effect was detected for the unloaded hydrogel, the antibacterial effect was more pronounced at a higher ampicillin concentration.

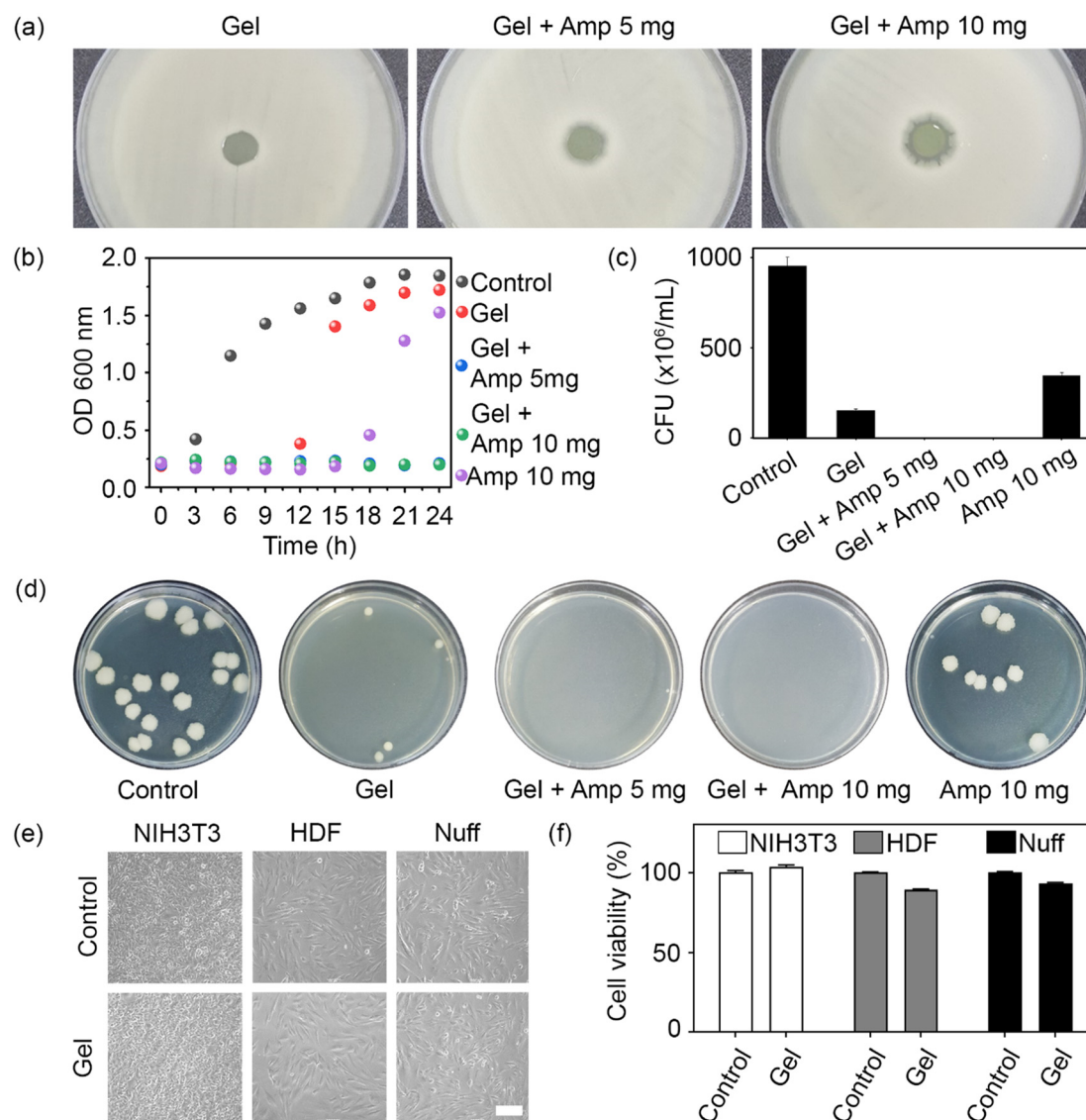


Fig. 4 (a) Inhibition zone test using *B. subtilis* on agar plates; hydrogels with or without ampicillin were placed in the center of the bacterial culture, and the inhibitory halo was quantified. (b) Growth of *B. subtilis* in the presence of hydrogels containing ampicillin or not over a period of 24 h. (c) CFU measurement after incubation of *B. subtilis* with the prepared gels for 24 h. (d) Representative images of surviving *B. subtilis* colonies on agar plates. (e) Morphology of NIH3T3 mouse fibroblasts, human dermal fibroblasts (HDF), and newborn human foreskin fibroblasts (Nuff) exposed to the prepared hydrogels for 24 h. Scale bar = 200 μ m. (f) Cytotoxicity of the prepared hydrogels toward the three types of fibroblasts.

To assess the antibacterial activity of the prepared hydrogels in a three-dimensional environment, the number of live cells in a bacterial population exposed to the loaded or unloaded hydrogels over 24 h was recorded, both in the form of OD₆₀₀ measurements (Fig. 4b) and CFU (Fig. 4c). While ampicillin alone (positive control) retarded bacterial growth for 15 h, no cell growth was observed in the ampicillin-loaded hydrogel groups over at least 24 h owing to the long-lasting antibacterial activity of the hydrogel (Fig. 4b). This finding was consistent with the outcome of the agar diffusion test. Similarly, CFU counts confirmed the absence of live bacteria when exposed to hydrogels loaded with either 5 or 10 mg ampicillin (Fig. 4c). It should be noted that the number of bacteria in the unloaded hydrogel group was lower than in the ampicillin control, suggesting that the positively charged chain and PEI backbone conferred some basic antibacterial activity to the hydrogel network. Therefore, a synergistic antibacterial effect was attributed to the chemical structure of the hydrogel and the loaded antibiotic, which endowed the prepared hydrogels with superior antimicrobial activity.

To evaluate the biocompatibility of the unloaded hydrogel, mouse and human fibroblasts were cultured with the hydrogel for 24 h. No morphological differences were observed between the control cells and those exposed to unloaded hydrogels (Fig. 4e). Finally, an MTT assay was performed to determine the cytotoxicity of the hydrogels toward mouse and human fibroblasts. The unloaded hydrogel did not exhibit any cytotoxicity toward the tested cell lines (Fig. 4f). Taken together, these results show that the unloaded hydrogel had a negative effect only on bacteria but displayed an otherwise excellent biocompatibility toward mammalian cells.

Conclusions

In summary, the present study describes the development of antibiotic-loaded antimicrobial hydrogels based on a double-network structure. The hydrogels integrated a first network composed of a positively charged monomer obtained *via* free radical polymerization, with a second network generated *via* a reversible imine bond between the free amines of PEI and the aldehyde of F127-CHO. The antibiotic ampicillin was loaded on the hydrogel through reversible imine bond formation. This pH-sensitive bond enabled the effective long-lasting release of antibiotic in an acidic environment. Importantly, the prepared hydrogels showed excellent antimicrobial activity against Gram-positive bacteria, while at the same time allowing for strong biocompatibility toward mammalian cells. Hence, the proposed hydrogels promise materials with long-lasting antibiotic activity and mechanical integrity.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This study was supported by the Ministry of Science and ICT, Ministry of Education, and the National Research Foundation of Korea through the Basic Science Research Program (grants 2019R11A2A01040856 and 2022R1A2C2008256).

References

- 1 E. S. R. Darley and A. P. MacGowan, *J. Antimicrob. Chemother.*, 2004, **53**, 928–935.
- 2 R. L. Smith, D. J. Schurman, G. Kajiyama, M. Mell and E. Gilkerson, *J. Bone Joint Surg. Am.*, 1987, **69**, 1063–1068.
- 3 C. D. Owens and K. Stoessel, *J. Hosp. Infect.*, 2008, **70**, 3–10.
- 4 J. Hoque, B. Bhattacharjee, R. G. Prakash, K. Paramanandham and J. Haldar, *Biomacromolecules*, 2018, **19**, 267–278.
- 5 D. E. Reichman and J. A. Greenberg, *Rev. Obstet. Gynecol.*, 2009, **2**, 212–221.
- 6 L. Hall-Stoodley, J. W. Costerton and P. Stoodley, *Nat. Rev. Microbiol.*, 2004, **2**, 95–108.
- 7 R. S. Howell-Jones, M. J. Wilson, K. E. Hill, A. J. Howard, P. E. Price and D. W. Thomas, *J. Antimicrob. Chemother.*, 2005, **55**, 143–149.
- 8 D. Stengel, K. Bauwens, J. Sehouli, A. Ekkernkamp and F. Porzolt, *Lancet Infect. Dis.*, 2001, **1**, 175–188.
- 9 J. H. Ryu, Y. Lee, W. H. Kong, T. G. Kim, T. G. Park and H. Lee, *Biomacromolecules*, 2011, **12**, 2653–2659.
- 10 Y. Li, K. Fukushima, D. J. Coady, A. C. Engler, S. Liu, Y. Huang, J. S. Cho, Y. Guo, L. S. Miller, J. P. K. Tan, P. L. R. Ee, W. Fan, Y. Y. Yang and J. L. Hedrick, *Angew. Chem., Int. Ed.*, 2013, **52**, 674–678.
- 11 K. Vulic and M. S. Shoichet, *J. Am. Chem. Soc.*, 2012, **134**, 882–885.
- 12 S. P. Hudson, R. Langer, G. R. Fink and D. S. Kohane, *Biomaterials*, 2010, **31**, 1444–1452.
- 13 J. Guo, W. Wang, J. Hu, D. Xie, E. Gerhard, M. Nisic, D. Shan, G. Qian, S. Zheng and J. Yang, *Biomaterials*, 2016, **85**, 204–217.
- 14 Y. Zhang, J. Zhang, M. Chen, H. Gong, S. Thamphiwatana, L. Eckmann, W. Gao and L. Zhang, *ACS Appl. Mater. Interfaces*, 2016, **8**, 18367–18374.
- 15 I. Stefanov, S. Pérez-Rafael, J. Hoyo, J. Cailloux, O. O. S. Pérez, D. Hinojosa-Caballero and T. Tzanov, *Biomacromolecules*, 2017, **18**, 1544–1555.
- 16 S. Q. Liu, C. Yang, Y. Huang, X. Ding, Y. Li, W. M. Fan, J. L. Hedrick and Y.-Y. Yang, *Adv. Mater.*, 2012, **24**, 6484–6489.
- 17 M. C. Giano, Z. Ibrahim, S. H. Medina, K. A. Sarhane, J. M. Christensen, Y. Yamada, G. Brandacher and J. P. Schneider, *Nat. Commun.*, 2014, **5**, 4095.
- 18 D. A. Salick, J. K. Kretsinger, D. J. Pochan and J. P. Schneider, *J. Am. Chem. Soc.*, 2007, **129**, 14793–14799.
- 19 D. D. Lane, A. K. Fessler, S. Goo, D. L. Williams and R. J. Stewart, *Acta Biomater.*, 2017, **50**, 484–492.

- 20 T. R. Hoare and D. S. Kohane, *Polymer*, 2008, **49**, 1993–2007.
- 21 C. S. Kwok, T. A. Horbett and B. D. Ratner, *J. Controlled Release*, 1999, **62**, 301–311.
- 22 W.-C. Liao, S. Lilienthal, J. S. Kahn, M. Riutin, Y. S. Sohn, R. Nechushtai and I. Willner, *Chem. Sci.*, 2017, **8**, 3362–3373.
- 23 H. Liu, X. Shi, D. Wu, F. K. Khshen, L. Deng, A. Dong and J. Zhang, *ACS Appl. Mater. Interfaces*, 2019, **11**, 19700–19711.
- 24 R. G. Schoenmakers, P. Van De Wetering, D. L. Elbert and J. A. Hubbell, *J. Controlled Release*, 2004, **95**, 291–300.
- 25 Y. Yeo, T. Ito, E. Bellas, C. B. Highley, R. Marini and D. S. Kohane, *Ann. Surg.*, 2007, **245**, 819–824.
- 26 K. Nakamae, T. Nishino, K. Kato, T. Miyata and A. S. Hoffman, *J. Biomater. Sci. Polym. Ed.*, 2004, **15**, 1435–1446.
- 27 H. Du, G. Zha, L. Gao, H. Wang, X. Li, Z. Shen and W. Zhu, *Polym. Chem.*, 2014, **5**, 4002–4008.
- 28 C. J. Waschinski, S. Barnert, A. Theobald, R. Schubert, F. Kleinschmidt, A. Hoffmann, K. Saalwächter and J. C. Tiller, *Biomacromolecules*, 2008, **9**, 1764–1771.
- 29 A. GhavamiNejad, C. H. Park and C. S. Kim, *Biomacromolecules*, 2016, **17**, 1213–1223.
- 30 V. W. L. Ng, J. M. W. Chan, H. Sardon, R. J. Ono, J. M. García, Y. Y. Yang and J. L. Hedrick, *Adv. Drug Delivery Rev.*, 2014, **78**, 46–62.
- 31 C. Maiti, K. B. C. Imani and J. Yoon, *ChemPlusChem*, 2021, **86**, 601–611.
- 32 M. Catauro, M. G. Raucci, D. de Marco and L. Ambrosio, *J. Biomed. Mater. Res., Part A*, 2006, **77A**, 340–350.
- 33 E. Khan, A. Shukla, A. Srivastava, Shweta and P. Tandon, *New J. Chem.*, 2015, **39**, 9800–9812.
- 34 G. Mallocci, G. Serra, A. Bosin and A. V. Vargiu, *Computation*, 2016, **4**, 5.