


Cite this: *RSC Adv.*, 2023, 13, 4669

Aza-Michael promoted glycoconjugation of PETIM dendrimers and selectivity in mycobacterial growth inhibitions†

Biswajit Sarkar,^a Avishek Mahapa,^{*bc} Kalyan Dey,^a Rakshit Manhas,^c Dipankar Chatterji^{*b} and Narayanaswamy Jayaraman ^{*a}

The benign nature of aza-Michael addition reaction in aqueous solutions is demonstrated herein to conduct a direct glycoconjugation of amine-terminated poly(ether imine) (PETIM) dendrimers. Zero to three generations of dendrimers, possessing up to 16 amine functionalities at their peripheries, undergo aza-Michael reaction with unsaturated sugar vinyl sulfoxide in aq. MeOH solutions and afford the corresponding dendrimers modified with multiple glycosyl moieties at the periphery. First order kinetics of the glycoconjugation is monitored at varying temperatures and the rate constants are observed to be 60–508 s^{−1}, for zero and first generation dendrimers. The antibacterial effects of amine-terminated dendrimers and the corresponding glycoconjugates are studied across Gram-positive, Gram-negative and acid-fast bacteria. Among the species, *M. smegmatis* and *M. tuberculosis* showed the greatest growth inhibition effect at micromolar concentrations, for the native amine-terminated and the corresponding glycoconjugated dendrimers. Quantitative assays are performed to adjudicate the inhibition efficacies of dendrimers and the glycoconjugates. Selectivity to inhibit *M. smegmatis* and *M. tuberculosis* growth, and minimal effects on other bacterial species by dendrimers and glycoconjugates are emphasized.

Received 23rd December 2022
Accepted 25th January 2023

DOI: 10.1039/d2ra08196b

rsc.li/rsc-advances

Introduction

Many biological processes depend on the post-translational modifications of proteins and lipids with carbohydrate moieties.^{1–3} Cell surface modifications,⁴ assembly of *N*-glycoproteins,⁵ evolution of glycoarrays,⁶ and development of drugs and vaccines are enabled by glycoconjugations with appropriate sugar moieties.^{7–11} As a result, robust conjugation methodologies have been developed, which include amidations, thiol-ene, urea formations, Michael addition, click reactions and more.^{12–21} In these instances, a reactive moiety is tethered to the sugar derivative so as to enable glycoconjugation with the chosen receptors. We demonstrated recently a glycoconjugation methodology which relies on the aza-Michael addition reaction of a number of oxygen, nitrogen, carbon and sulfur nucleophiles with a sugar vinyl sulfoxide under aqueous conditions.²² The reactions are conducted under aqueous milieu, as a result, glycoconjugation on to proteins becomes feasible. The side

chain amine functionality of lysine residues in proteins reacts with sugar vinyl sulfoxide in such glycoconjugations. In advancing further, the utility of the methodology for glycoconjugation of dendritic macromolecules is undertaken herein.

Synthesis of monodispersed dendritic macromolecules requires tight control over their molecular weights and the number of chain ends.^{23–25} Interfacing dendrimers with diverse biological, chemical and materials entities is one of the major advancements in contemporary macromolecular research.^{26–29} Carbohydrate-binding proteins, namely, lectins and antibodies are the first point of attachment of carbohydrate ligands on to biologically-relevant receptors.³⁰ Following these clues, intense efforts were undertaken in the glycoconjugation of dendrimers. In this context, multitude of developments also include exploiting efficacies of glycodendrimers to study multivalent carbohydrate-protein interactions, inhibitions of enzyme functions, in gene and drug delivery applications.^{31,32}

An area of intense investigations relates to the role of the cationic dendrimers as a new class of antibacterial agents.^{33–45} When a microbial infection is treated with traditional antibiotics at sub-lethal concentrations or for a long time, it inversely induces resistance to the pathogens, prompting efforts to mitigate the multidrug-resistant strains. Extensively drug-resistant *M. tuberculosis*, Methicillin-resistant *S. aureus*, beta-lactamase-producing *E. coli* and *P. aeruginosa* are prominent among multidrug-resistant (MDR) strains, associated with healthcare infections. Effective antibacterial regimens become

^aDepartment of Organic Chemistry, Indian Institute of Science, Bangalore 560 012, India. E-mail: jayaraman@iisc.ac.in

^bMolecular Biophysics Unit, Indian Institute of Science, Bangalore 560 012, India. E-mail: dipankar@iisc.ac.in

^cInfectious Disease Department, CSIR-Indian Institute of Integrative Medicine, Jammu-180001, India. E-mail: avishek.mahapa@gmail.com

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2ra08196b>


an important query, in efforts to address such disease-causing bacterial infections to hosts.

Studies of poly(lysine),^{43,44} poly(amido amine),⁴⁵ poly(propylene imine),^{39,40} poly(phosphane)³⁵ and carbosilane dendrimers³⁸ show that the antibacterial properties of these dendrimers arise due to the cationic sites present within the structure. The antibacterial efficacies of such dendrimer also enhance further upon conjugation with small molecule inhibitors. The series of poly(ether imine) (PETIM) dendrimers were assessed earlier for their cytotoxicity and biological effects, in mammalian cell cultures, from which non-toxicity profiles were observed for the dendrimers possessing hydroxyl functionalities at their peripheries.^{46–49} This series of dendrimers also possesses varied functional moieties, namely, amines and carboxylic acids at the peripheries. The amine functionalities afford cationic sites to the dendrimer, suitable to interfere with the cell wall structures and functions. Specific relevance herein is the membrane cell wall biosynthesis in a bacterium. Controlling the biosynthesis of cell wall components assumes a greater significance to mitigate bacterial pathogenicity to host cells. The presence of multiple chain ends in dendritic macromolecules attracted efforts to identify potent antimicrobial properties, either in the native or chain end-functionalized forms.⁵⁰ The clustering of functional groups present at the peripheries of dendrimers leads to specific and non-specific interactions of the designed dendrimer agents with the microbial targets. The dendrimers with quaternary ammonium groups are effective against Gram-negative *E. coli* and Gram-positive *S. aureus* strains. Concerning dendrimers functionalized with sugars, it was demonstrated early by Magnusson and co-workers that glycoconjugation of dendrimers with di- to tetravalent galabiosides led to more effective hemagglutination inhibition of *S. suis* than the monomer galabioside.⁵¹ PAMAM dendrimers are by far the most studied and are known to engage in membrane disruption against Gram-positive and Gram-negative pathogens against opportunistic pathogens.⁵² An increasing interest in the antimicrobial properties of dendrimers arises in order to develop more effective dendrimer-based antimicrobial agents against MDR strains.

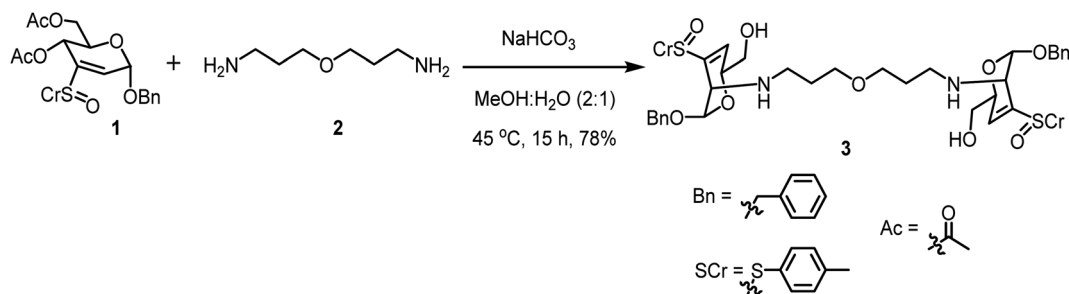
We report herein a glycoconjugation method, where the modified sugar moieties are covalently-linked to amine functionalities present at the peripheries of zero to three generations of PETIM dendrimers. Following synthesis, the native,

amine-terminated dendrimers and the corresponding glycoconjugates are studied in order to evaluate their efficacies on the growth of varied planktonic bacterial strains. Details are presented herein.

Results

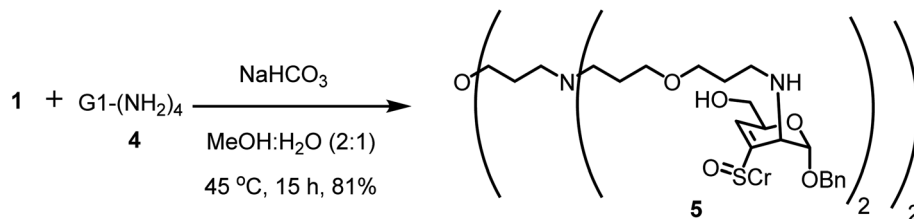
Synthesis

One-step glycoconjugation of amine-terminated PETIM dendrimers⁵³ in an aqueous medium, namely, aza-Michael addition, is described herein. Zero, first, second and third generation dendrimers possess 2, 4, 8 and 16 peripheral amine moieties, respectively, and the amine functionality acts as the aza-Michael donor. Benzyl 4,6-di-*O*-acetyl-2,3-dideoxy-3-(*p*-tolylsulfinyl)- α -D-erythro-hex-2-enopyranoside (**1**), possessing 2,3-unsaturated vinyl sulfoxide functionality acted as the Michael acceptor. The reaction of sugar vinyl sulfoxide **1** (ref. 54) with the bis-amine **2** (ref. 55) is shown in Scheme 1. The reaction was conducted in an aqueous solution, under basic condition (pH \sim 7.5–8) and the reaction afforded the bis-conjugated product **3**. Three molar equivalents of sulfoxide **1** were used to one molar equivalent of acceptor **2** and the reaction occurring at 45 °C was identified to be optimal for the glycoconjugation. Insolubility of **1** in water necessitated MeOH for solubilization and the reaction was conducted at 45 °C. Following column purification of the reaction mixture, bis-glycoconjugate **3** was secured in a good yield. As observed earlier, an addition–elimination occurred to afford the 3,4-unsaturated vinyl sulfoxide. The basic pH of the reaction also led to hydrolysis of the acetate moiety, leading to product **3** with a primary hydroxyl group. The addition occurred from the axial face to afford the *manno*-configured product, as observed generally for the primary amine additions.⁵⁴ Olefinic proton at C-2 carbon of **1** at 6.80 ppm disappeared and the olefinic proton at C-4 of **3** appeared at 6.59 ppm. Further, the anomeric H-1 in **1** appearing at 5.4 ppm up-field shifted to 5.10 ppm in **3**, the $J_{H-1-H-2}$ was 2.6 Hz, implying a *trans*-diequatorial disposition of H-1–H-2 protons in **3**. The dendrimer methylene protons in **3** appeared at 3.32, 2.44 and 1.55 ppm and shifted up-field by \sim 0.2–0.3 ppm after conjugation with sulfoxide **1**. The anomeric resonance shifted down-field to 94.2 ppm for product **3**, whereas that for vinyl sulfoxide **1** appeared at 93.7 ppm and 93.5 ppm. The methylene carbon adjacent to amine moiety in **3** appeared at 54.7 ppm, that for bis-amine **1**



Scheme 1 Reaction of sugar vinyl sulfoxide **1** with bis-amine **2**.



Scheme 2 Glycoconjugation of G1-(NH₂)₄ dendrimer with vinyl sulfoxide 1.

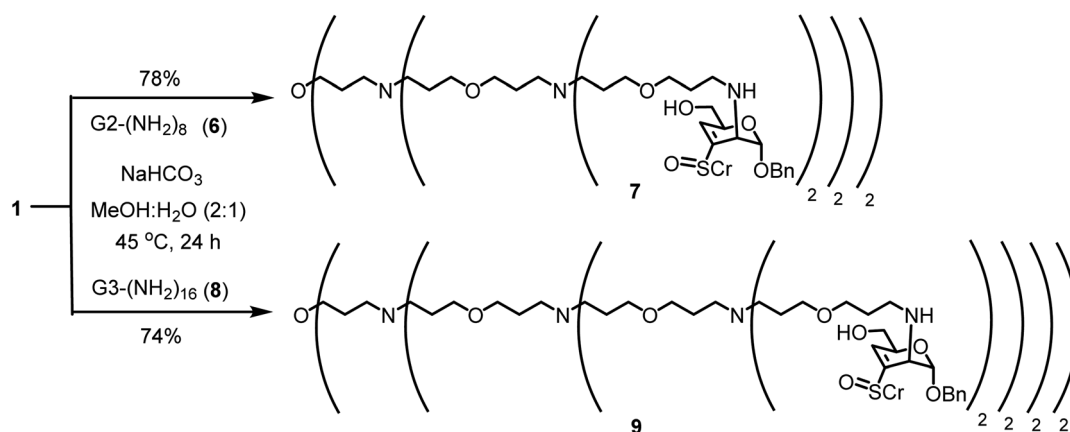
appeared at 47.5 ppm. The mass spectrum of 3 showed molecular ion peak as $[M + H]^+$ species, appearing at m/z 845.3532 (calcd mass m/z 845.3505).

First generation PETIM dendrimer 4 (ref. 53) with 4 peripheral amine moieties was subjected to glycoconjugation with vinyl sulfoxide 1 (Scheme 2). The reaction was conducted with one-and-half molar equivalent of 1 per amine moiety in tetraamine 4. Following reaction over 15 h, the solution was evaporated, suspended in water and triturated with EtOAc several times in order to remove excess of vinyl sulfoxide. The product thus obtained was analysed to determine the extent of conjugation of sugar moiety with the dendrimer 5. MALDI-TOF mass analysis revealed the molecular ion peak as $[M + 2H]^+$ adduct at m/z 2018.380 (calcd m/z 2018.9737), corresponding to complete conjugation at all the available primary amine moieties (Fig. S8†). In addition, multiple peaks due to the presence of acetate moiety at C-6 carbon were also seen. Further the conjugation was ascertained by comparison of the resonances of aromatic moiety with methylene moiety adjacent to the amine functionality in the 1H NMR spectrum of 5. The NMR spectra were recorded in CD₃OD. The resonances appeared broad and in sets of resonances, presumably due to sulfoxide chirality and the resulting diastereomeric nature of the product. The methylene moiety adjacent to primary amine in 4 appeared 2.79 ppm. Whereas in the glycoconjugated product 5, methylene moieties adjacent to the amine sites appeared at 2.67–3.00 ppm. The H-4 olefinic proton and the H-1 proton of 5 were found to resonate at 6.54 and 5.13 ppm, respectively, in 1H NMR spectra. The peak corresponding to anomeric carbon resonated

at 95.3 ppm in ^{13}C NMR spectra. These newly observed peaks in 1H and ^{13}C NMR spectroscopy ascertained the formation of conjugated dendrimer 5.

Glycoconjugations of G2-(NH₂)₈ and G3-(NH₂)₁₆,⁵³ possessing 8 and 16 amine moieties, respectively, were conducted (Scheme 3). The reactions were conducted in aq. MeOH for 24 h at 45 °C, in both the reactions. Removal of solvents and trituration with EtOAc afforded glycoconjugates 7 and 9, in 78% and 74%, respectively.

NMR spectra of these conjugated products 7 and 9 were recorded in CD₃OD and the resonances appeared broad. Major observations were that: (i) the acetate moiety at C-6 position was susceptible to hydrolysis; (ii) with an exponential increase in the molecular weights of glycoconjugates 7 and 9 without acetate moiety, 4362.19 and 9052.65 g mol⁻¹, respectively, gradual line-broadening and reduced signal-to-noise ratio of the peaks occurred in 1H and ^{13}C NMR spectra and (iii) mass spectra of these derivatives could not be secured by ESI-MS and MALDI-TOF mass spectrometries. Upon conjugation, the H-4, H-1 protons in 7 and 9 appeared at 6.59, 5.30 and 6.60, 5.19 ppm in 1H NMR spectra, respectively. Whereas, the anomeric carbon appeared at 93.9 and 95.4 ppm in ^{13}C NMR spectra, respectively. For these higher generations of dendrimers, the extent of conjugation was an average, calculated from the integral values of 1H NMR spectra. In the case of second-generation amine 6, the methylene moieties adjacent to amines appeared at 2.94 and 2.60 ppm, as apparent triplets. For the conjugation product 7, these and tolyl-CH₃ protons appeared 2.21–2.85 ppm, as broad resonances. The aryl moieties of sugar component



Scheme 3 Synthesis of second and third generation sugar vinyl sulfoxide conjugates 7 and 9.

Table 1 Kinetic constant for the glycoconjugation of PETIM dendrimers with sugar vinyl sulfoxide 1

Dendrimer generation	Kinetic constant (s^{-1})		
	25 °C	35 °C	45 °C
G0 (2)	61.2	158.4	284.4
G1 (4)	144.1	295.2	507.6

appeared at 7.38–7.48 ppm and 7.68–7.73 ppm. Integration of these aryl moiety protons with that of the methylene moieties adjacent to amine sites and tolyl-CH₃ protons enabled to adjudge the extent of conjugation. In this manner, a conjugation of 7 out of 8 sugar moieties was adjudged for the complete structure as given for 7 in Scheme 3. A similar analysis of the ¹H NMR spectrum of product 9 was also undertaken. In third-generation dendrimer 8, methylene protons adjacent to amine sites appeared at 3.03 and 2.54 ppm, as broad peaks. A comparison of the integrations of these and tolyl-CH₃ protons, 2.19–2.71 ppm and the aromatic moieties, 7.25–7.42 and 7.66–7.74, in 9 suggested a conjugation of 14 sugar moieties for the complete structure as given in Scheme 3. This incomplete conjugation also appeared to cause reduced resolutions of the NMR spectra in these glycoconjugates.

Determination of the kinetic constants

The kinetics of the conjugation was monitored, by ¹H NMR spectroscopy, in a mixture of CD₃OD and D₂O. Disappearance of the methylene protons adjacent to primary amine of dendrimer scaffold was chosen to monitor the conjugation, occurring at 25, 35 and 45 °C, over 7 h. The conjugation led to a reduction in the intensity of this methylene resonance at ~2.67 ppm in bis-

amine 2, adjacent to primary amine moiety (–CH₂NH₂). Conjugation of G1 generation dendrimer also led to a reduction of the intensity of resonance at ~2.56 ppm, corresponding to –CH₂NH₂ moiety. Due to overlapping resonances, kinetics of the conjugations in the case of G2 and G3 generation could not be conducted. ¹H NMR data were collected for up to 7 h and the time course analysis was fit to a first-order kinetic equation and the rate constants of conjugation were calculated (Table 1). Rate of the reaction increased approximately twice for each 10 °C raise in the temperature. The rate constant was found to be higher for the first (4) generation dendrimer than bis-amine 2 (Table 1).

Bacterial growth assays

In the initial screening, the antibacterial efficacies of amine-terminated dendrimers and the corresponding glycoconjugates (25 μg mL^{−1}) were evaluated on the growth of *E. coli* DH5α (Gram-negative), *P. aeruginosa* (Schroeter) Migula 27853 (Gram-negative), *S. aureus* Rosenbach 25923 (Gram-positive) and *M. smegmatis* mc² 155 (acid fast). Kanamycin (Kan) was used as positive antibacterial control for the assays. The bacterial growth profiles are shown in Fig. 1, the corresponding growth inhibition percentages in the presence and absence of dendrimers are given in Table 2. The untreated cells were considered as negative control (NC). The growth assays yielded a generation time of ~40 min, 90 min, 51 min. and 4.8 h for *E. coli*, *P. aeruginosa*, *S. aureus* and *M. smegmatis*, respectively, without the presence of the dendrimer agents.

As seen in Fig. 1a and b, the growth profiles of *E. coli* and *P. aeruginosa* encountered little or no inhibition of the bacterial growth in the presence of dendrimers, the maximum inhibition percentages were 22% and 6% for the conjugate 9, respectively.

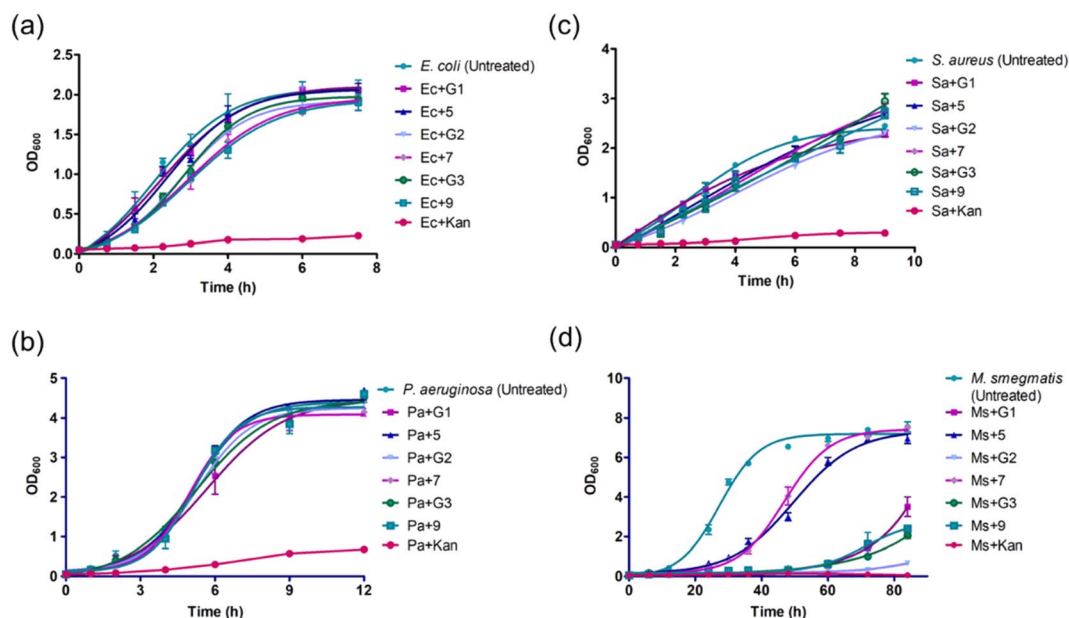


Fig. 1 Bacterial growth observed in the presence of different native and glycoconjugated dendrimer generations: (a) *E. coli* DH5α; (b) *P. aeruginosa*; (c) *S. aureus* and (d) *M. smegmatis* mc² 155.



The native un-conjugated G-2 dendrimer showed 32% growth inhibition of Gram-positive *S. aureus* (Fig. 1c). Further 10–12% growth inhibitions were observed in the presence of remaining native and glycoconjugated dendrimers, in comparison to untreated control. Among the strains, *M. smegmatis* was observed to undergo a significant growth reduction. G-1: 91.2%; 5: 43.2%; G-2: 43.8%; 7: 38%; G-3: 92.2% and 9: 90.6% (Fig. 1d), as compared to the untreated control. Also, the doubling times of *M. smegmatis* cells increased significantly: G-1: 12.5 h; 5: 2.1 h; G-2: 17.7 h; 7: 3.7 h; G-3: 31 h and 9: 7.4 h (Fig. 1d).

The above observations reveal that the dendrimer agents are effective to inhibit the growth of mycobacterium strains and, to some extent, Gram-positive *S. aureus*. The native and glycoconjugated dendrimers did not act on Gram-negative *E. coli* or *P. aeruginosa* strains, even when cationic amine sites constitute the dendrimers. A defining feature of the Gram-positive and Gram-negative strains is the altered cell wall structure compositional features. We presume that the cell surface-active dendrimer agents might differentiate these features, leading to the most effects of the dendrimer agents on acid-fast *M. smegmatis*, followed by Gram-positive *S. aureus* and minimal effects on the Gram-negative *P. aeruginosa* and *E. coli* strains.

MIC values determination for *M. smegmatis* and *M. tuberculosis*

Due to the selectivity of the dendrimers towards mycobacterial growth inhibition, further evaluation of the MIC value was conducted on *M. smegmatis* and, importantly, *M. tuberculosis* H37Rv cells and the results are presented in Table 3. The MIC values found for *M. smegmatis* were 50 $\mu\text{g mL}^{-1}$ for G1, G2 and G3 generations of amine terminated PETIM dendrimers. Whereas, the same for *M. tuberculosis* H37Rv cells were 100 $\mu\text{g mL}^{-1}$, 100 $\mu\text{g mL}^{-1}$ and 200 $\mu\text{g mL}^{-1}$, respectively. The glycoconjugated dendrimer generations G1 and G2, the MIC values were found 100 $\mu\text{g mL}^{-1}$ and 200 $\mu\text{g mL}^{-1}$, in each case for *M. smegmatis* and *M. tuberculosis* H37Rv, respectively. These values reiterated that the amine terminated dendrimers are relatively more efficient in inhibiting mycobacterial growth than the glycoconjugated dendrimers. When MIC of *M. smegmatis* was evaluated against third generation glycoconjugate 9, twice the concentration as that of the lower generation dendrimer glycoconjugates, implying that lower generation dendrimer glycoconjugates surpass inhibition efficiencies as compared to the high generation glycoconjugates. Bacterial strains *S. aureus*, *E.*

Table 3 MIC values of the PETIM dendrimers on *M. smegmatis* and *M. tuberculosis*^a

Compound	<i>M. smegmatis</i> MIC value ($\mu\text{g mL}^{-1}$)	<i>M. tuberculosis</i> MIC value ($\mu\text{g mL}^{-1}$)
G1	50	100
5	100	200
G-2	50	100
7	100	200
G3	50	200
9	200	NA

^a NA: MIC value is not within the limit.

coli and *P. aeruginosa* did not show growth inhibitions in the presence of the dendrimers and the corresponding glycoconjugates.

Discussion

Dendrimers with cationic sites at the peripheries are known to act as bactericidal agents, as a result of ionic interactions with anionic bacterial cell membranes, that lead to penetration and permeability of the exogenous dendrimers into the cellular plasma environment and the damage of the intracellular components.^{45,50} Carbohydrate-functionalized dendrimers were studied, specifically as anti-adhesive agents that block the binding of the host cell surface glycans with lectins present at the outer membranes of a bacterium.^{56,57} Among the most studied are the polycationic poly(amido amine) (PAMAM) dendrimers that show anti-bacterial effects on Gram-positive and Gram-negative bacteria, in low $\mu\text{g mL}^{-1}$ concentrations. Such bactericidal effects of the amine-terminated PAMAM dendrimers were off-set when the peripheral functionalities were modified with poly(ethylene glycol) moieties.⁵⁸ Similar bactericidal effects were established to poly(propylene imine) (PPI) and carbosilane dendrimers, possessing amine functionalities at their peripheries.^{59,60} Further, the antibacterial efficacies of maltose sugar-functionalized fourth generation PPI dendrimers were demonstrated in low micromolar concentrations against Gram-positive and Gram-negative bacteria.^{39,40} Bare PPI dendrimer, without the sugar functionalization, was found to be more cytotoxic than the sugar functionalized PPI dendrimer. Apoptosis, coupled with oxidative stress, of cells was found to be major causes for the antibacterial activity. The maltose-

Table 2 Percentage bacterial growth in the presence of native and glyco-conjugated dendrimer generations G-1 to G-3

	<i>M. smegmatis</i>	<i>E. coli</i>	<i>P. aeruginosa</i>	<i>S. aureus</i>
Untreated	100.0	100.0	100.0	100.0
G-1	8.8 \pm 1.5	97.1 \pm 3.4	91.0 \pm 1.9	91.9 \pm 2.7
5	56.8 \pm 0.23	95.3 \pm 3.4	99.5 \pm 5.6	88.1 \pm 4.4
G-2	4.0 \pm 0.08	85.1 \pm 2.9	94.5 \pm 2.8	67.2 \pm 4.7
7	62.0 \pm 2.7	80.0 \pm 3.4	95.5 \pm 0.2	82.8 \pm 3.6
G-3	7.8 \pm 0.24	87.0 \pm 1.1	95.3 \pm 0.5	81.8 \pm 2.4
9	9.4 \pm 2.9	78.0 \pm 3.3	95.7 \pm 0.07	81.1 \pm 0.7



functionalized fourth generation PPI dendrimer was found to disrupt outer membrane of bacterial cells, thereby permitting muralytic enzyme, endolysin, to penetrate and degrade peptidoglycans, leading to lysis of bacterial cells.⁴⁰

One of the well-studied polycationic macromolecules are the polylysine glycoconjugates that act on both the Gram-positive and Gram-negative bacterial cell membranes and rupture membrane structures, leading to cytoplasm damage and bacterial cell lysis.⁴³ The selectivity of the cell lysis by the polylysine glycoconjugate was found to be about half that of the native polylysine, even when the microbial targeting was superior with the polymer glycoconjugate. Reduction in the cationic sites density in the polymer glycoconjugate was proposed to cause reduced cell lysis, related with the increased minimum inhibitory concentration required to affect the bacterial growth. Important outcomes of our present study are that (i) the native dendrimers are more effective to inhibit the growth Gram-positive and acid-fast bacteria and very minimal or none of the inhibition effect on Gram-negative bacteria; (ii) the native dendrimers with dense cationic sites are far more mycobacterial growth inhibitors than the corresponding glycoconjugates and (iii) the higher generation native dendrimers are far more effective inhibitors than the glycoconjugates. The pattern reiterates that amine-rich dendrimers are by-far the efficient inhibitors and the corresponding glycoconjugations compensate the cationic sites induced growth inhibitions and reduce the inhibitory efficacies.

The selectivity to inhibit the growth of mycobacterial strains *M. smegmatis* and *M. tuberculosis* is revealing. We presume that the cell surface composition of various components is likely the origin of this selectivity. The bacterial outer layer is a complex, multi-layered arrangement of biological macromolecules. The organization and the components of the bacterial cell wall differ across bacterial species. The cell wall structures of *E. coli* and *P. aeruginosa*, used as representatives from Gram-negative bacteria, Gram-positive *S. aureus*, *M. tuberculosis* and acid-fast *M. smegmatis* are categorized broadly.^{61,62} The Gram-negative bacterial cell wall consists of layers of the inner membrane, the peptidoglycan and the outer membrane. The outer membrane is a bilayer of lipid, mainly composed of lipopolysaccharide. The Gram-positive bacteria lacks such an outer membrane. Repeating *N*-acetyl glucosamine and *N*-acetyl muramic acid disaccharide, along with cross-linked pentapeptide side chains, constituted the peptidoglycan layer. This peptidoglycan layer in Gram-positive bacteria is more dense than Gram-negative bacteria. Further, the Gram-positive bacteria possesses long anionic polymers called teichoic acids. On the other hand, the mycobacterial cell wall is different, called myco-membrane, composed of mycolic acids (MA). This outer membrane is critical for mycobacterial physiology and protects mycobacteria from different stress. This lipid layer also provides high resistance to many broad spectra of antibiotics. The cell wall is orders of magnitude less permeable, than the most resistant Gram-negative bacteria *E. coli* and *P. aeruginosa*. The lipids content in the mycobacterial cell wall is 60%, which is threefold higher than most Gram-negative bacteria. The mycolic acids in the mycobacterial cell wall are covalently linked with cell wall

polysaccharides, as in lipomannan and lipoarabinomannan.⁶³ This unique architecture and higher acidic moiety in mycobacteria provide negative charge to the mycobacterium cell wall compared. We presume that the un-modified cationic dendrimers interact efficiently with the anionic mycobacterial cell wall, when compared to other bacterial strains. This interaction at the mycobacterial cell wall plays an important role in the mycobacterial growth inhibition by dendrimers.

Conclusion

A benign glycoconjugation of amine-rich PETIM dendrimers of 0–3 generations, possessing 2–16 amine functionalities at the peripheries, is conducted with sugar moieties in aqueous MeOH solutions, by an aza-Michael addition reaction of sugar vinyl sulfoxide. The conjugation is observed to be faster in the case of the first generation dendrimer, than zero generation bis-amine derivative. The glycoconjugated dendrimers were studied for their antibacterial properties against Gram-positive, Gram-negative and acid-fast bacteria. Among the bacteria, the growths of *M. smegmatis* and *M. tuberculosis* are inhibited profoundly by the native, unmodified and glycoconjugated dendrimers, whereas Gram-negative bacterial strains are not affected. Results of this study show that the dendrimers inhibit the mycobacterial growth specifically and such a specificity opens up leads for the development of target-driven dendrimer-based anti-mycobacterial agents.

Experimental section

G0-Glycoconjugate (3)

A mixture of PETIM G0-(NH₂)₂ **2** (4 mg, 0.03 mmol) and sugar vinyl sulfoxide **1** (41 mg, 0.09 mmol) in MeOH/H₂O (2 : 1) (1.5 mL) was added with NaHCO₃ (2 mg) and stirred at 45 °C for 15 h. The crude reaction mixture was evaporated and purified using column chromatography (SiO₂, 100–200 mesh) (20% MeOH/CHCl₃) to afford compound **3**, as a gum (20 mg, 78%). *R*_f = 0.2 (20% MeOH/CHCl₃); ¹H NMR (400 MHz, CDCl₃): δ 7.63–7.59 (m, 4H), 7.36–7.22 (m, 14H), 6.68 (d, *J* = 16 Hz, 0.5H), 6.59 (app. s, 1.5H), 5.15–5.10 (m, 2H), 4.87–4.81 (m, 2H), 4.63–4.51 (m, 4H), 3.85–3.75 (m, 4H), 3.34–3.32 (m, 4H), 2.92 (s, 2H), 2.45–2.41 (m, 4H), 2.39 (s, 6H), 2.11–2.09 (m, 4H), 1.57–1.52 (m, 4H); ¹³C NMR (CDCl₃, 100 MHz) δ 143.8, 141.9, 139.5, 137.3, 130.2, 130.1, 129.9, 128.7, 128.6, 128.3, 128.2, 128.1, 128.0, 126.7, 125.8, 94.25, 71.2, 69.9, 69.1, 64.6, 54.73, 44.3, 30.5, 29.8, 21.6 and 14.2. ESI-MS *m/z*: [M + Na]⁺ calcd For C₄₆H₅₆N₂O₉S₂H, 845.3505; found: 845.3532.

G1-Glycoconjugate (5)

A mixture of PETIM G1-(NH₂)₄ **4** (8 mg, 0.014 mmol) and sugar vinyl sulfoxide **1** (38.5 mg, 0.084 mmol) in MeOH/H₂O (2 : 1) (1.5 mL) was added with NaHCO₃ (2 mg) and stirred at 45 °C for 15 h. The crude reaction mixture was evaporated, resuspended in H₂O (2 mL) and followed by washing with ethyl acetate (10 × 2 mL). The aqueous layer was concentrated *in vacuo* to afford compound **5**, as a yellowish white gum (22 mg, 81%). ¹H NMR



(500 MHz, CD₃OD): δ 7.69–7.60 (m, 13H), 7.33–7.20 (m, 23H), 6.69–6.54 (m, 4H), 5.13 (b, 4H), 4.61–4.44 (m, 8H), 3.71 (b, 7H), 3.63 (m, 7H), 3.45 (br. s, 21H), 3.00–2.67 (m, 9H), 2.38–2.34 (m, 24H), 1.77–1.47 (m, 19H). ¹³C NMR (CD₃OD, 125 MHz) δ 138.9, 131.1, 129.8, 129.6, 129.1, 128.2, 127.9, 127.0, 95.3, 72.4, 71.2, 70.8, 69.6, 64.8, 61.0, 51.9, 37.7, 31.0, 28.4, 26.8, 23.9, 23.7, 21.5, 21.3, 18.7. MALDI-MS m/z : [M + Na]⁺ calcd for C₁₁₀H₁₄₈N₆O₂₁S₄H₂, 2018.9737; found: 2018.330.

G2-Glycoconjugate (7)

A mixture of PETIM G2-(NH₂)₈ **6** (10 mg, 0.007 mmol) and sugar vinyl sulfoxide **1** (38.5 mg, 0.084 mmol) in MeOH/H₂O (2 : 1) (1.5 mL) was added with NaHCO₃ (2 mg) and stirred at 45 °C for 24 h. The crude reaction mixture was evaporated, resuspended in H₂O (2 mL) and followed by washing with ethyl acetate (10 × 2 mL). The aqueous layer was concentrated *in vacuo* to afford compound **7**, as a yellowish white gum (18 mg, 78%). ¹H NMR (500 MHz, CD₃OD): δ 7.73–7.68 (m, 18H), 7.48–7.38 (m, 51H), 6.81 (app. s., 2H), 6.59 (app. d, J = 8 Hz, 6H), 5.36–5.26 (m, 7H), 4.18–4.17 (m, 2H), 4.93–4.85 (m, 5H), 3.74 (s, 11H), 3.63 (s, 97H), 3.39 (s, 3H), 3.20–3.13 (m, 12H), 2.85–2.76 (m, 6H), 2.61–2.55 (m, 2H), 2.45–2.42 (m, 54H), 2.26–2.21 (m, 1H), 2.20 (s, 9H), 1.80 (s, 14H), 1.58–1.27 (br. m, 14H), 1.11–0.94 (m, 14H). ¹³C NMR (CD₃OD, 125 MHz) δ 169.1, 147.5, 142.1, 141.3, 140.4, 130.1, 128.9, 128.5, 128.1, 127.9, 127.5, 127.4, 126.0, 125.6, 93.9, 72.9, 72.7, 70.0, 68.7, 68.3, 61.4, 60.2, 59.1, 38.2, 31.8, 29.8, 29.5, 26.6, 22.9, 20.7, 20.2, 20.0.

G3-Glycoconjugate (9)

A mixture of PETIM G3-(NH₂)₁₆ **8** (10 mg, 0.003 mmol) and sugar vinyl sulfoxide **1** (33 mg, 0.072 mmol) in aq. MeOH/H₂O (2 : 1) (1.5 mL) was added with NaHCO₃ (2 mg) and stirred at 45 °C for 24 h. The crude reaction mixture was evaporated, resuspended in H₂O (2 mL) and followed by washing with ethyl acetate (10 × 2 mL). The aqueous layer was concentrated *in vacuo* to afford compound **9**, as a yellowish white gum (20 mg, 74%). ¹H NMR (400 MHz, CD₃OD): δ 7.74–7.66 (m, 57H), 7.42–7.25 (m, 114H), 6.75–6.60 (m, 15H), 5.19 (b, 16H), 4.67–4.58 (m, 15H), 4.50 (br. s, 15H), 3.77–3.68 (m, 25H), 3.54 (br. s, 121H), 3.36–3.33 (m, 9H), 3.09–3.06 (m, 138H), 2.64–2.61 (m, 8H), 2.46–2.40 (m, 150H), 2.25–2.19 (m, 17H), 1.90–1.87 (m, 62H). ¹³C NMR (CD₃OD, 100 MHz) δ 138.9, 131.1, 129.8, 129.6, 129.2, 129.1, 128.2, 126.9, 95.4, 72.4, 70.8, 69.3, 67.9, 64.8, 60.3, 59.3, 55.9, 52.8, 52.0, 39.0, 30.9, 28.7, 26.5, 23.9, 21.5, 21.3.

Biological studies

The antibacterial efficacies of amine-terminated dendrimers and the corresponding glycoconjugates were evaluated on the growth of *E. coli* DH5 α (Gram-negative), *P. aeruginosa* (Schroeter) Migula 27853 (Gram-negative), *S. aureus* Rosenbach 25923 (Gram-positive) and *M. smegmatis* mc² 155 (acid fast bacilli). All the bacterial strains were maintained in specific growth medium. *P. aeruginosa* and *S. aureus* were cultivated in Mueller-Hinton (MH) and Tryptic Soy Broth (TSB), respectively. *E. coli*, *P. aeruginosa* and *S. aureus* strains were grown in LB (Luria-Bertani) broth, whereas *M. smegmatis* and *M. tuberculosis*

H37Rv strains was grown in MB7H9 medium (Difco) and supplemented with glucose (2%), Tween 80 (0.05%) and 10% OADC only for *M. tuberculosis* at 37 °C. Kanamycin (25 μ g mL⁻¹) was used as positive control in the bacterial growth experiments. For growth assay, each compound (25 μ g mL⁻¹) was added at 0 h and the growth of cells was maintained under shaking condition at 37 °C. The bacterial growth was measured by optical density (OD) changes of the cultures at 600 nm. OD values for a specific time point were plotted and analyzed by Graph Pad Prism software. The growth inhibition assays for all bacterial species were performed three independent triplicates. Growth medium without any dendrimer compounds was taken as the positive control in all the experiments.

Minimum inhibitory concentration (MIC) estimation of the dendrimers

Determination of the MIC of *M. smegmatis* and *M. tuberculosis* H37Rv was confirmed by cell viability study using 96-well Microplate Alamar Blue Assay (MABA). The water solubility, permeability inside cell, low toxicity towards bacterial cell and stability in cell culture makes it perfect as visual indicator for the analysis of viability of the bacterial cell culture. Resazurin is a blue non-fluorescent dye, turns fluorescent pink dye resorufin when reduced. So, the reduction of resazurin (blue, non-fluorescent) to resorufin (pink, fluorescent) indicates metabolic impairment of the cells. Thus, the visual change of the colour is used for the assay to determine the minimum inhibitory concentration (MIC). The determination of MIC for PETIM and sugar vinyl sulfoxide modified PETIM was conducted by 96-well Microplate Alamar Blue Assay. The protocol of the assay is well established and commonly as resazurin microtiter assay (REMA) method.^{64–66} Briefly, *M. smegmatis* and *M. tuberculosis* H37Rv cells (at 0.05 OD) was taken in a 96-well flat-bottom microtiter plate, where PETIM and modified PETIM dendrimers were added to maintain a final volume of 200 μ L. The starting concentrations of the dendrimers were 200 μ g mL⁻¹ and 400 μ g mL⁻¹, for *M. smegmatis* and *M. tuberculosis* H37Rv, respectively. The compounds were diluted serially up to concentrations 0.39 μ g mL⁻¹. The cultures were allowed to grow by incubating for 48 h and 6 days for *M. smegmatis* and *M. tuberculosis* H37Rv, respectively, at 37 °C. After that, resazurin solution (0.04%) was added to the culture and incubated further for another 4 h for *M. smegmatis* and 24 h for *M. tuberculosis* H37Rv cells. The colour remained blue if the growth of the bacteria cell was inhibited. A negative control with no dendrimers added was used to compare with the dendrimer treated bacteria culture. The experiments were conducted in triplicate.

Author contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We are grateful to Department of Biotechnology, New Delhi, for a research support (Grant Number BT/PR33123/MED/29/1497/2020). B. S. and K. D. are grateful to IISc, Bangalore, CSIR, New Delhi, for research fellowships. AM acknowledge fellowship support from the DBT-Research Associateship (RA) Programme in Life Sciences. We thank Prof. Jayanta Chatterji, IISc, Bangalore, for providing *S. aureus* and *P. aeruginosa* strains.

References

- 1 K. Ohtsubo and J. D. Marth, *Cell*, 2006, **126**, 855.
- 2 C. Reily, T. J. Stewart, M. B. Renfrow and J. Novak, *Nat. Rev. Nephrol.*, 2019, **15**, 346.
- 3 K. Handa and S.-i. Hakomori, *Glycoconjugate J.*, 2017, **34**, 693.
- 4 J. M. Baskina and C. R. Bertozzi, *QSAR Comb. Sci.*, 2007, **26**, 1211.
- 5 C. Unverzagt and Y. Kajihara, *Chem. Soc. Rev.*, 2013, **42**, 4408.
- 6 J. W. Wehner, M. Hartmann and T. K. Lindhorst, *Carbohydr. Res.*, 2013, **371**, 22.
- 7 R. D. Astruc and D. R. Burton, *Nat. Rev. Drug Discovery*, 2010, **9**, 308.
- 8 S. Feng, C. Xiong, S. Wang, Z. Guo and G. Gu, *ACS Infect. Dis.*, 2019, **5**, 1423.
- 9 B. Liet, E. Laigre, D. Goyard, B. Todaro, C. Tiertant, D. Boturyn, N. Berthet and O. Renaudet, *Chem.-Eur. J.*, 2019, **25**, 15508.
- 10 N. T. Supekar, V. Lakshminarayanan, C. J. Capicciotti, A. Sirohiwal, C. S. Madsen, M. A. Wolfert, P. A. Cohen, S. J. Gendler and G.-J. Boons, *ChemBioChem*, 2018, **19**, 121.
- 11 K. Streichert, C. Seitz, E. Hoffmann, I. Boos, W. Jelkmann, T. Brunner, C. Unverzagt and M. Rubini, *ChemBioChem*, 2019, **20**, 1914.
- 12 B. Sarkar and N. Jayaraman, *Front. Chem.*, 2020, **8**(1–29), 570185.
- 13 G. C. Daskhan, H.-T. T. Tran, P. J. Meloncelli, T. L. Lowary, L. J. West and C. W. Cairo, *Bioconjugate Chem.*, 2018, **29**, 343.
- 14 M. G. García, J. M. Benito, A. P. Butera, C. O. Mellet, J. M. G. Fernández and J. L. J. Blanco, *J. Org. Chem.*, 2012, **77**, 1273.
- 15 M. Wolfenden, J. Cousin, P. N. Makker, A. Raz and M. J. Cloninger, *Molecules*, 2015, **20**, 7059.
- 16 C. Fessele, S. Wachtler, V. Chandrasekaran, C. Stiller, T. K. Lindhorst and A. Krueger, *Eur. J. Org. Chem.*, 2015, 5519.
- 17 P. Wu, M. Malkoch, J. N. Hunt, R. Vestberg, E. Kaltgrad, M. G. Finn, V. V. Fokin, K. B. Sharpless and C. J. Hawker, *Chem. Commun.*, 2005, 5775.
- 18 M. G. Cuesta, D. Goyard, E. Nanba, K. Higaki, J. M. G. Fernández, O. Renaudet and C. O. Mellet, *Chem. Commun.*, 2019, **55**, 12845.
- 19 L. G. Weaver, Y. Singh, G. Vamvounis, M. F. Wyatt, P. L. Burn and J. T. Blanchfield, *Polym. Chem.*, 2014, **5**, 1173.
- 20 X. Li, T. Fang and G.-J. Boons, *Angew. Chem., Int. Ed.*, 2014, **53**, 7179.
- 21 S. Munneke, E. M. Dangerfield, B. L. Stocker and M. S. M. Timmer, *Glycoconjugate J.*, 2017, **34**, 633.
- 22 B. Sarkar, A. Mahapa, D. Chatterji and N. Jayaraman, *Biochemistry*, 2019, **58**, 3561.
- 23 D. Astruc, L. Liang, A. Rapakousiou and J. Ruiz, *Acc. Chem. Res.*, 2012, **45**, 630.
- 24 D. A. Tomalia, J. B. Christensen and U. Boas, in *Dendrimers, Dendrons and Dendritic Polymers*, Cambridge University Press, Cambridge, UK, 2012.
- 25 R. K. O'Reilly, C. J. Hawker and K. L. Wooley, *Chem. Soc. Rev.*, 2006, **35**, 1068.
- 26 S. Svenson and D. A. Tomalia, *Adv. Drug Delivery Rev.*, 2012, **64**, 102.
- 27 D. Astruc, E. Boisselier and C. Ornelas, *Chem. Rev.*, 2010, **110**, 1857.
- 28 L. Röglin, E. H. M. Lempens and E. W. Meijer, *Angew. Chem., Int. Ed.*, 2011, **50**, 102.
- 29 A.-M. Caminade, A. Ouali, R. Laurent, C.-O. Turrin and J.-P. Majoral, *Chem. Soc. Rev.*, 2015, **44**, 3890.
- 30 N. Jayaraman, *Chem. Soc. Rev.*, 2009, **38**, 3463.
- 31 S. Mignani, J. Rodrigues, R. Roy, X. Shi, V. Ceña, S. E. Kazzouli and J.-P. Majoral, *Drug Discovery Today*, 2019, **24**, 1184.
- 32 L. I. F. Moura, A. Malfanti, C. Peres, A. I. Matos, E. Guegain, V. Sainz, M. Zloh, M. J. Vicent and H. F. Florindo, *Mater. Horiz.*, 2019, **6**, 1956.
- 33 M. L. Wolfenden and M. J. Cloninger, *Bioconjugate Chem.*, 2006, **17**, 958.
- 34 C. Sehad, T. C. Shiao, L. M. Sallam, A. Azzouz and R. Roy, *Molecules*, 2018, **23**, 1890.
- 35 S. Mignani, V. D. Tripathi, D. Soam, R. P. Tripathi, S. Das, S. Singh, R. Gandikota, R. Laurent, A. Karpus, A.-M. Caminade, A. Steinmetz, A. Dasgupta, K. K. Srivastava and J.-P. Majoral, *Biomacromolecules*, 2021, **22**, 2659.
- 36 A. Castonguay, E. Ladd, T. G. M. van de Ven and A. Kakkar, *New J. Chem.*, 2012, **36**, 199.
- 37 M. A. Mintzer, E. L. Dane, G. A. O'Toole and M. W. Grinstaff, *Mol. Pharm.*, 2012, **9**, 342.
- 38 I. H. Bermejo, J. M. H. Ros, L. S. García, M. Maly, C. V. Expósito, J. Soliveri, F. J. de la Mata, J. L. C. Patiño, J.-P. Serrano, J. S. Nieves and R. Gómez, *Eur. Polym. J.*, 2018, **101**, 159.
- 39 A. Felczak, N. Wrońska, A. Janaszewska, B. Klajnert, M. Bryszewska, D. Appelhans, B. Voit, S. Rózska and K. Lisowska, *New J. Chem.*, 2012, **36**, 2215.
- 40 K. Ciepluch, B. Maciejewska, K. Gałczyńska, D. K. Ciepluch, M. Bryszewska, D. Appelhans, Z. D. Kawa and M. Arabski, *Bioorg. Chem.*, 2019, **91**(1–6), 103121.
- 41 S. Alfei and A. M. Schito, *Nanomaterials*, 2020, **10**(1–52), 2022.
- 42 M. Á. Ortega, A. G. Merino, O. F. Martínez, J. R. Ruiz, L. Pekarek, L. G. Guijarro, N. G. Honduvilla, M. A. Mon,



- J. Buján and S. G. Gallego, *Pharmaceutics*, 2020, **12**(1–27), 874.
- 43 D. Alkekhia and A. Shukla, *J. Biomed. Mater. Res., Part A*, 2019, **107A**, 1324.
- 44 D. Pranantyo, L. Q. Xu, Z. Hou, E.-T. Kang and M. B.-C. Park, *Polym. Chem.*, 2017, **8**(1–10), 3364.
- 45 B. Wang, R. S. Navath, A. R. Menjoge, B. Balakrishnan, R. Bellair, H. Dai, R. Romero, S. Kannan and R. M. Kannan, *Int. J. Pharm.*, 2010, **395**, 298.
- 46 T. R. Krishna, S. Jain, U. S. Tatu and N. Jayaraman, *Tetrahedron*, 2005, **61**, 4281.
- 47 U. P. Thankappan, S. N. Madhusudana, A. Desai, G. Jayamurugan, Y. B. R. D. Rajesh and N. Jayaraman, *Bioconjugate Chem.*, 2011, **22**, 115.
- 48 A. Lakshminarayanan, V. K. Ravi, R. Tatineni, Y. B. R. D. Rajesh, V. Maingi, K. S. Vasu, N. Madhusudhan, P. K. Maiti, A. K. Sood, S. Das and N. Jayaraman, *Bioconjugate Chem.*, 2013, **24**, 1612.
- 49 A. Lakshminarayanan, B. U. Reddy, N. Raghav, V. K. Ravi, A. Kumar, P. K. Maiti, A. K. Sood, N. Jayaraman and S. Das, *Nanoscale*, 2015, **7**, 16921.
- 50 C. Z. Chen and S. L. Cooper, *Adv. Mater.*, 2000, **12**, 843.
- 51 H. C. Hansen, S. Haataja, H. J. Finne and G. Magnusson, *J. Am. Chem. Soc.*, 1997, **119**, 6974.
- 52 A. M. Holmes, J. R. Heylings, K. W. Wan and G. P. Moss, *Int. J. Antimicrob. Agents*, 2019, **53**, 500.
- 53 T. R. Krishna and N. Jayaraman, *J. Org. Chem.*, 2003, **68**, 9694.
- 54 A. Mukherjee and N. Jayaraman, *Tetrahedron*, 2012, **68**, 8746.
- 55 A. Mukherjee and N. Jayaraman, *Carbohydr. Res.*, 2013, **380**, 51.
- 56 M. Touaibia and R. Roy, *Mini-Rev. Med. Chem.*, 2007, **7**, 1270.
- 57 J. Rojo and R. Delgado, *Anti-Infect. Agents Med. Chem.*, 2007, **6**, 151.
- 58 A. I. Lopez, R. Y. Reins, A. M. McDermott, B. W. Trautner and C. Cai, *Mol. Biosyst.*, 2009, **5**, 1148.
- 59 C. Z. Chen, S. L. Cooper and N. C. B. Tan, *Chem. Commun.*, 1999, 1585.
- 60 P. Ortega, J. L. C. Patino, M. A. M. Fernandez, J. Soliveri, R. Gomez and F. J. de la Mata, *Org. Biomol. Chem.*, 2008, **6**, 3264.
- 61 L. J. Alderwick, J. Harrison, G. S. Lloyd and H. L. Birch, *Cold Spring Harbor Perspect. Med.*, 2015, **5**(1–15), a021113.
- 62 T. J. Silhavy, D. Kahne and S. Walker, *Cold Spring Harbor Perspect. Biol.*, 2010, **2**(1–16), a000414.
- 63 A. K. Mishra, N. N. Driessen, B. J. Appelmek and G. S. Besra, *FEMS Microbiol. Rev.*, 2011, **35**, 1126.
- 64 S. N. Rampersad, *Sensors*, 2012, **12**, 12347.
- 65 K. R. Gupta, S. Kasetty and D. Chatterji, *Appl. Environ. Microbiol.*, 2015, **81**, 2571.
- 66 M. Elshikh, S. Ahmed, S. Funston, P. Dunlop, M. McGaw, R. Marchant and I. M. Banat, *Biotechnol. Lett.*, 2016, **38**, 1015.

