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## Heparan sulfate glycomimetics via iterative assembly of “clickable” disaccharides†

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Heparan sulfate (HS) glycosaminoglycans are widely expressed on the mammalian cell surfaces and extracellular matrices and play important roles in a variety of cell functions. Studies on the structure–activity relationships of HS have long been hampered by the challenges in obtaining chemically defined HS structures with unique sulfation patterns. Here, we report a new approach to HS glycomimetics based on iterative assembly of clickable disaccharide building blocks that mimic the disaccharide repeating units of native HS. Variably sulfated clickable disaccharides were readily assembled into a library of mass spec–sequenceable HS-mimetic oligomers with defined sulfation patterns by solution-phase iterative syntheses. Microarray and surface plasmon resonance (SPR) binding assays corroborated molecular dynamics (MD) simulations and confirmed that these HS-mimetic oligomers bind protein fibroblast growth factor 2 (FGF2) in a sulfation-dependent manner consistent with that of the native HS. This work established a general approach to HS glycomimetics that can potentially serve as alternatives to native HS in both fundamental research and disease models.

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## Introduction

As a major class of glycosaminoglycan polysaccharides, heparan sulfates (HS), are essential components of the cell surface glycocalyx and extracellular matrix that play important roles in cell proliferation, differentiation, migration, and cell–cell interaction.<sup>1</sup> HS consist of uronic acid-(1 → 4)-D-glucosamine disaccharide repeating units that are sulfated at various positions. Variable patterns of *N*-sulfate, *O*-sulfate and *N*-acetyl substitutions on the polymeric backbone of HS give rise to a high degree of sequence complexity, which in turn enables active modulation of the sulfate-dependent interactions of HS with a plethora of signaling proteins.<sup>2,3</sup> Many HS-binding proteins employ a distinct sequence of positively charged residues to engage with HS, and correspondingly, require specific sulfation

patterns of HS to achieve selectivity among the complex glycome.

To understand the structure–activity relationships of the sulfation patterns in HS, Seeberger,<sup>4,5</sup> Linhardt,<sup>6,7</sup> Liu,<sup>8,9</sup> Boons,<sup>10,11</sup> Weigel,<sup>12</sup> DeAngelis,<sup>13</sup> Hung,<sup>14,15</sup> Huang,<sup>16–18</sup> and others<sup>19–24</sup> have developed chemical and chemoenzymatic synthetic approaches to well-defined HS oligosaccharides that have significantly advanced our understanding of HS biology. Nevertheless, the chemical synthesis of HS oligosaccharides often requires exquisite synthetic route design and skillful carbohydrate chemistry manipulations, resulting in high production cost and scalability challenges. Chemoenzymatic synthesis has emerged as a scalable method to access HS oligosaccharides, but the requirements for engineered enzymes and large quantities of activated substrates often render the synthesis expensive and technically demanding.

Compared to the native HS structures, HS-mimetic oligomers and polymers require less efforts to synthesize and provide comparable activities in many functional assays, making them highly useful tools to study HS biology. For example, the work by Hsieh–Wilson<sup>25,26</sup> and Godula<sup>27,28</sup> demonstrated that the repeated display of sulfated disaccharides along a polymer backbone possesses activities similar to the native HS in anti-coagulation, stem cell differentiation, and chemokine response. However, most existing HS-mimetic polymers incorporate sulfated oligosaccharides as pendant groups and do not allow

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precision control of the sulfation patterns, limiting their utility in the investigation into functional roles of sulfation patterns in HS. Herein, we developed a series of “clickable” disaccharides that resemble the repeating unit of HS and can be rapidly assembled into sequence-defined glycomimetic oligomers *via* the copper(I)-catalyzed alkyne–azide cycloaddition (CuAAC) click chemistry (Fig. 1). These clickable disaccharides consist of an iduronic acid (IdoA)-(β1 → 4)-glucosamine (GlcN) core structure and an azide and a triisopropyl silyl (TIPS)-protected alkyne end groups, with various positions modified by the sulfate group (Fig. 1a). We have shown that a variety of HS-mimetic oligomer structures with defined sulfation patterns could be readily synthesized following efficient solution-phase iterative assembly and their sequences could be independently verified by tandem mass spectrometry (Fig. 1b). A microarray consisting of HS-mimetic oligomers with various sulfation patterns was fabricated to examine their ability to bind fibroblast growth factor-2 (FGF2), a model protein with well-established sulfation-dependent interactions with HS.<sup>29,30</sup> Molecular dynamics (MD) simulation of the glycomimetic oligomers bound to FGF2 corroborated the binding assay results, confirming that these HS-mimetic oligomers bind FGF2 in a sulfation-dependent fashion similar to the native HS oligosaccharides. Taken together, our studies suggest that the clickable disaccharides are facile tools for investigating the functional roles of sulfation in diverse physiological processes involving HS.

## Results and discussion

### Design and synthesis of clickable disaccharides

The development of chemically defined HS mimetics has been stymied by the difficulties in preparing large collections of these compounds with tunable chain length and precisely controlled

sulfation patterns. Inspired by the chemical synthesis of polypeptides<sup>31</sup> and nucleic acids<sup>32</sup> *via* solid-phase and solution-phase iterative synthesis, we envision that a synthetic approach to HS-mimetic oligomers could be realized by connecting modular disaccharide building blocks using the CuAAC click chemistry, due to its mild reaction condition, high coupling efficiency, and excellent tolerance to a broad scope of functionalities.<sup>33</sup> In particular, CuAAC coupling can be performed in the late stage of the synthesis with minimal impacts on the structural integrity of the oligomers generated from the iterative synthesis. To this end, we designed the partially protected IdoA-(β1 → 4)-GlcN disaccharides with sulfation at specific hydroxyl and amino groups and an azide and a TIPS-alkyne group at C1 and C4 positions, respectively (Fig. 1). Non-sulfated hydroxyl groups were protected by benzyl ether (Bn) groups. As such, these disaccharide building blocks could be iteratively assembled after the TIPS group is removed *via* tetra-*n*-butylammonium fluoride (TBAF) after each coupling step.

We began the synthesis of the “clickable” disaccharide building blocks using commercially available diacetone glucose and *N*-acetylglucosamine starting materials (Fig. 2). Specifically, 1,6-anhydro β-(L)-idopyranose was synthesized from diacetone glucose in six steps.<sup>34</sup> While our initial attempt to install triisopropyl silane (TIPS) protected alkyne using TIPS-protected propargyl bromide was unsuccessful due to ester migration, an alternative route from TIPS-protected propargylic trichloroacetimidate successfully afforded anhydrosugar **2**. Acetolytic ring opening of **2** provided the 1,6-diacetate, which was converted into thioglycoside **3** in a combined yield of 38% over 3 steps. After the removal of the C6 acetyl group, **4** was oxidized and esterified into thioglycoside methyl ester **5** as an iduronic acid donor in 57% overall yield over 3 steps. To access the glucosamine acceptors, glycosylation of (D)-*N*-acetylglucosamine **6** with the 2-chloroethanol followed by sequential acetylation and azidation yielded the 1-azidoethyl compound in 65% overall yield. The followed installation of 4,6-benzylidene acetal and benzyl ether furnished **7** in 77% yield. After the benzylation on C4 hydroxyl and the removal of the 4,6-benzylidene protection in **7**, the 6-hydroxyl was selectively protected by levulinic acid to afford *N*-acetylglucosamine acceptor **9** in 61% overall yield. To prepare the glucosamine acceptor for *N*-sulfation, we deacetylated **7** and protected the C2 amine with 2,2,2-trichloroethoxycarbonyl (Troc) group to arrive at **12** in 86% yield over two steps. Tin(II)-catalyzed benzylation of **12** under acidic conditions followed by the removal of 4,6-benzylidene protection produced the glucosamine acceptor **13** with Troc-protected amine in 31% overall yield. Lastly, glycosylation of acceptor **9** or **13** by thioglycoside donor **5** generated β-linked core disaccharide **10** and **14** in 65% and 76% yields, respectively.

The orthogonal protecting groups in the core disaccharide **10** provided the opportunity for regioselective deprotection and sulfation at various positions. Through protecting group manipulation and *O*-sulfation, non-sulfated disaccharide **16** and differentially sulfated disaccharides (**17–19**) were readily obtained. Similarly, trisulfated disaccharide building block **21** was generated from **20** *via* sequential deprotection and *O*- and

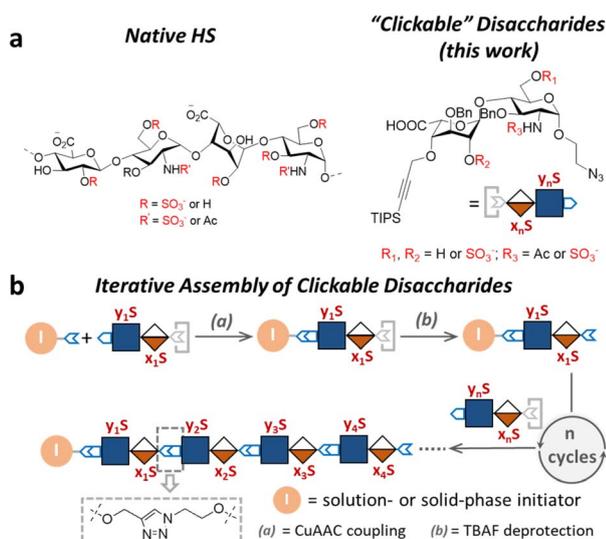
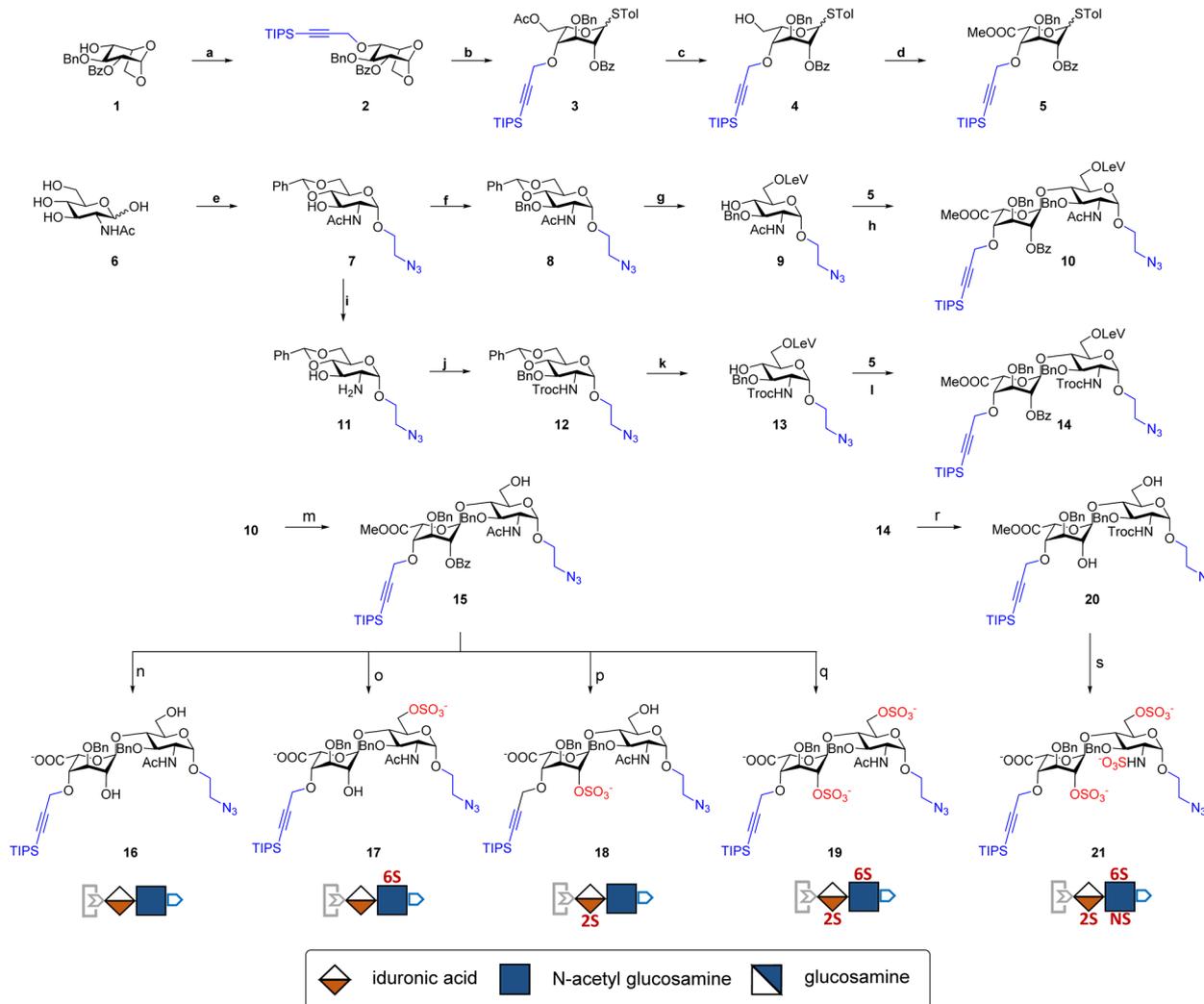


Fig. 1 Overview of this work. (a) Chemical structures of natural HS and the HS glycomimetics developed in this work. (b) The iterative assembly of the clickable disaccharides *via* CuAAC click chemistry.





**Fig. 2** Synthesis of clickable disaccharides. Reagents and conditions: (a) TIPSCHCCH<sub>2</sub>ONHCCl<sub>3</sub>, TfOH, molecular sieve 4 Å, 45 °C, 46%; (b) (i) Cu(OTf)<sub>2</sub>, acetic anhydride, (ii) TolSH, TfOH, DCM, 82% in two steps; (c) AcCl, DCM/MeOH, 92%; (d) (i) TEMPO, BAIB, DCM/H<sub>2</sub>O, (ii) Me<sub>2</sub>SO<sub>4</sub>, acetone, 0 °C, 62%; (e) (i) ClCH<sub>2</sub>CH<sub>2</sub>OH, AcCl, 70 °C, 65%, (ii) acetic anhydride, pyridine, quant, (iii) NaN<sub>3</sub>, DMF, 60 °C, quant, (iv) NaOMe, MeOH, quant, (v) benzaldehyde dimethyl acetal, TsOH, DMF, 65 °C, 77%; (f) BnBr, NaH, DMF, 90%; (g) (i) TFA, DCM/H<sub>2</sub>O, 30 °C, 74%, (ii) Lev acid, EDC, DMAP, DCM, 91%; (h) TIS, TfOH, molecular sieve 4 Å, 69%; (i) KOH, dioxane/2-methoxyethanol, 120 °C; (j) TrocCl, sat. NaHCO<sub>3</sub>, THF, 86% in two steps, (k) (i) BnOCNHCCl<sub>3</sub>, Sn(OTf)<sub>2</sub>, Et<sub>2</sub>O, molecular sieve 4 Å, 62%; (ii) TFA, DCM/H<sub>2</sub>O, 30 °C, 66%; (iii) Lev acid, EDC, DMAP, DCM, 76%; (l) TIS, TfOH, molecular sieve 4 Å, 76%; (m) hydrazine acetate, MeOH/THF, quant; (n) (i) LiOH, MeOH, 0 °C, (ii) NaOH, MeOH/H<sub>2</sub>O, 80% in 2 steps; (o) (i) SO<sub>3</sub>-Py, DMF, 87%; (ii) LiOH, MeOH/H<sub>2</sub>O, 0 °C, (iii) NaOH, MeOH/H<sub>2</sub>O, 93% in 2 steps; (p) (i) MeONa, MeOH, 88%, (ii) Lev acid, EDC, DMAP, DCM, 83%, (iii) SO<sub>3</sub>-Py, DMF, 77%, (iv) LiOH, NaOH, THF/H<sub>2</sub>O, 87%; (q) (i) MeONa, MeOH, 88%, (ii) SO<sub>3</sub>-Py, DMF, 90%, (iii) LiOH, MeOH/H<sub>2</sub>O, 93%; (r) (i) hydrazine acetate, THF/H<sub>2</sub>O, (ii) MeONa, MeOH, 66%; (s) (i) SO<sub>3</sub>-Py, DMF, 89%, (ii) LiOH, MeOH/H<sub>2</sub>O, (iii) SO<sub>3</sub>-Py, NaOH, TEA, DMF, 79% in two steps.

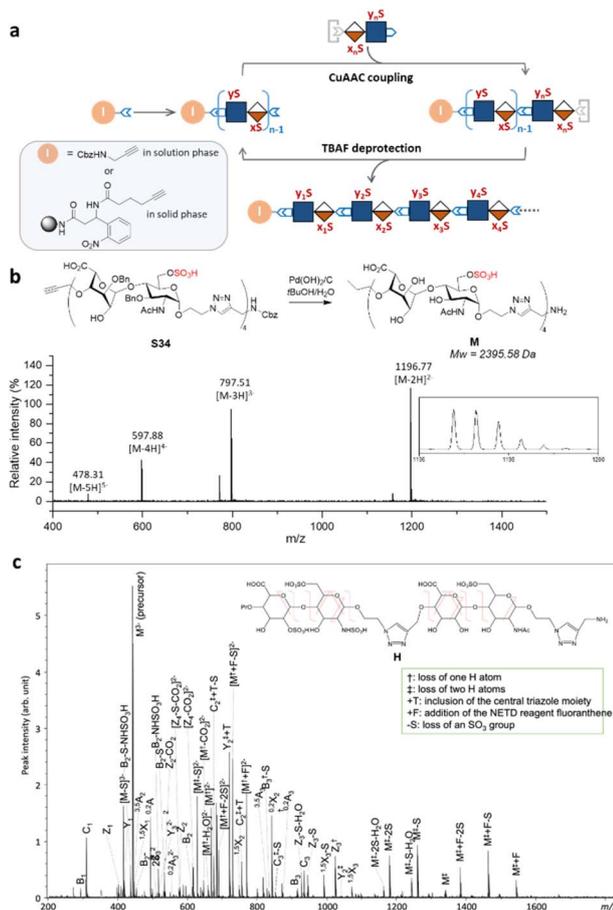
*N*-sulfation. Importantly, disaccharides **17–19** and **21** were confirmed to be compatible with CuAAC coupling and TBAF deprotection conditions (Fig. S1†). Taken together, variably sulfated clickable disaccharides were efficiently prepared and validated for an iterative coupling–deprotection procedure, making them promising building blocks for the synthesis of sequence-defined HS-mimetic oligomers.

### Iterative assembly of clickable disaccharides

Next, we explored the iterative assembly of the clickable disaccharides into sequence-defined HS-mimetic oligomers. Starting

from a benzyl chloroformate (Cbz)-protected propargyl amine initiator, successive CuAAC coupling and TBAF deprotection were performed in several iterations until the desired length of the oligomer was obtained (Fig. 3a). In each cycle, a unique disaccharide building block was incorporated to generate a sequence-defined oligomer with a specific sulfation pattern. The product of each coupling and deprotection step was isolated by flash silica chromatography and confirmed by liquid chromatography/electrospray ionization mass spectrometry (LC-ESI-MS) before the next cycle. Using 6-*O*-sulfated disaccharide **17** as a model building block, we constructed a model





**Fig. 3** Assembly and MS/MS sequencing of the HS-mimetic oligomer. (a) Solution-phase assembly of variably sulfated clickable disaccharides. (b) LC-ESI-MS analysis of a model HS-mimetic octasaccharide oligomer **M** after the hydrogenolysis of oligomer **S34** generated from solution-phase assembly. (c) NETD MS/MS sequencing of oligomer **H** allowed for confident assignment of the sulfation pattern.

octasaccharide **S34** in an overall yield of 16% in eight steps after isolation by semi-prep HPLC (80% average yield in each step) (Fig. S2†).

The Bn-protected oligomers from the solution-phase were deprotected *via* hydrogenolysis by Pd(OH)<sub>2</sub>/C in *tert*-butanol/water (1/1, v/v) in high efficiency, producing the HS-mimetic oligomers in excellent yields. For example, hydrogenolysis of **S34** afforded fully deprotected octasaccharide **M** in 70% yield (Fig. 3b). Following the synthetic procedures established in the model studies, we constructed a library of 14 HS-mimetic tetrasaccharides and octasaccharides with various sulfation patterns (**A–N**) (Fig. 4a).

The sequencing ability of the HS-mimetic oligomers is important for future combinatorial library synthesis and high-throughput screening. To this end, the sequence of the synthetic HS glycomimetics were independently verified by tandem mass spectrometry (MS/MS). The conventional collision-induced dissociation (CID) spectrum was dominated by sulfate loss and unable to produce informative tandem mass spectra for reliable sequencing due to the lability of the sulfate

group during collisional excitation (Fig. S3†). To overcome this challenge, negative electron transfer dissociation (NETD) was performed on the tetrasaccharide oligomer **H** (Fig. 3c).<sup>35,36</sup> Encouragingly, NETD generated complete series of glycosidic fragments, including all B-, C-, Y-, and Z-ions. Product ions at each cleavage site were detected without sulfate loss, allowing confident assignment of the sulfate modifications at the residue level. Localization of the sulfate modifications within each residue could also be achieved by identifying the mass difference between key cross-ring fragments and their corresponding glycosidic fragment. For example, the presence of a <sup>3,5</sup>A<sub>2</sub> ion (*m/z* 451.022) and a <sup>0,2</sup>X<sub>2</sub> ion (*m/z* 845.181) provides conclusive evidence for assignment of the 6-*O*-sulfation and *N*-sulfation at the second GlcN residue. The presence of the triazole linker does not appear to interfere with the structural characterization, as no fragmentation within the triazole moiety was observed in the NETD spectrum.

### Protein-binding studies

We used glycan microarray and surface plasmon resonance (SPR) to analyze the interactions between the HS-mimetic oligomers and FGF2. Glycan microarrays have become a key glycomics technology for systematically evaluating the structure-activity relationships of complex carbohydrates.<sup>37</sup> We fabricated microarrays of the HS-mimetic oligomers by reacting their terminal amino groups with *N*-hydroxysuccinimide (NHS)-activated glass slides (Fig. 4b). The microarray of HS-mimetic oligomers was incubated with FGF2 in the Tris buffer (20 mM Tris-HCl pH 7.4, 150 mM NaCl, 2 mM CaCl, 2 mM MgCl) containing Tween-20 (0.05%) and BSA (1%), and binding was indirectly observed using unlabeled anti-FGF2 antibody and a goat-anti-mouse IgG 635 detection antibody. Fluorescence signal at excitation wavelength 635 nm was measured and the average signal for replicate spots (*n* = 4) was calculated as the mean RFU. Encouragingly, the HS-mimetic oligomers exhibited sulfation-dependent binding to FGF2 similar to the native HS (Fig. 4c). It was observed that, at both 100 μM and 300 μM of HS-mimetic oligomers, FGF2 bound tetrasaccharide oligomers **G–K** comparably but to a greater extent than the oligomers with lower sulfation density. Specifically, the oligomers with *N*-sulfation, 2-*O*-, and 6-*O*-sulfation (*e.g.*, **J** and **K**) exhibited higher fluorescent intensity than those carrying only 2-*O*- or 6-*O*-sulfation (*e.g.*, **C**, **D**, **M**, or **N**) in the microarray experiment using fluorescently labeled FGF2. To rule out the avidity effect in which multiple surface-bound HS-mimetic oligomers in proximity bind to the same FGF2 protein, the FGF2-oligomer interactions were further investigated using an SPR competition assay (Fig. 4d). Biotinylated heparin was immobilized onto the streptavidin-functionalized SPR sensor chip, to which a solution containing a fixed concentration of FGF2 and various concentrations of a specific HS-mimetic oligomer was injected and allowed to flow through the sensor chip. The HS-mimetic oligomer competes with the immobilized heparin and inhibits its binding to FGF2, resulting in decreased responses in the SPR sensorgrams. The half maximal inhibitory concentration (IC<sub>50</sub>) of the HS-mimetic oligomers against the FGF2 binding by



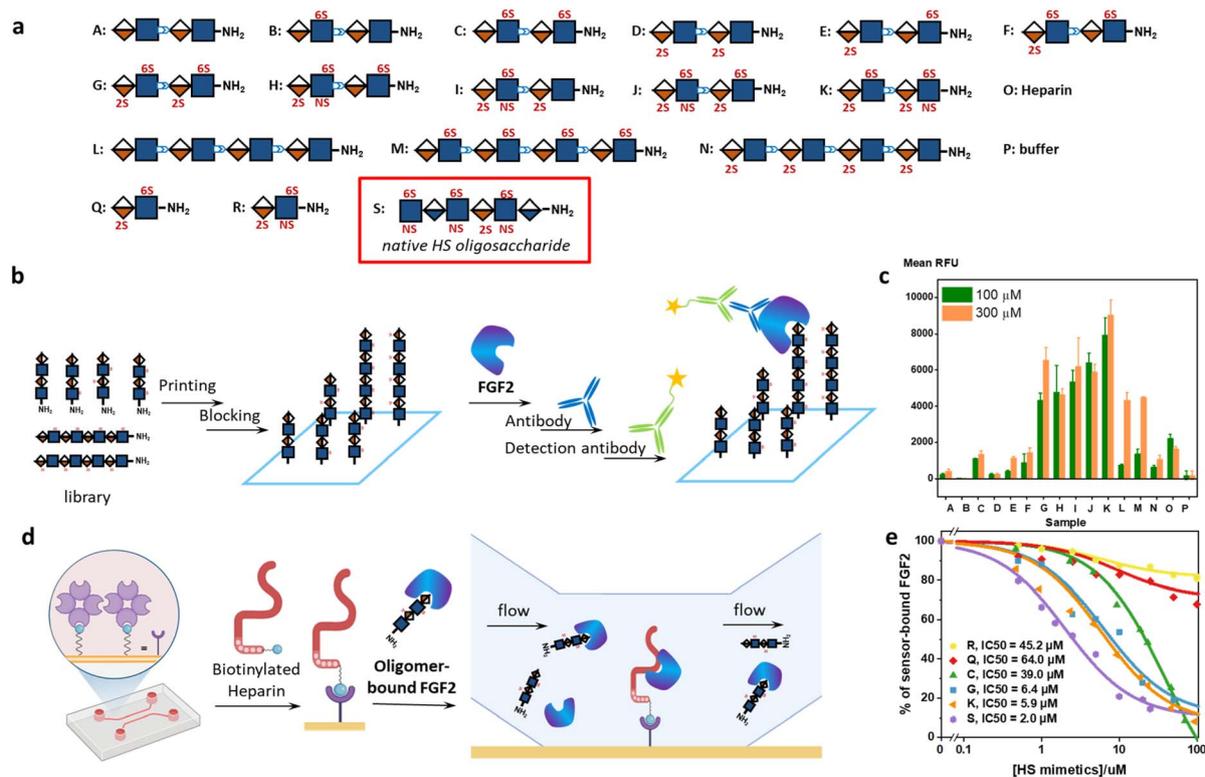


Fig. 4 Characterization of the interactions between the HS-mimetic oligomers and FGF2. (a) Sequences of the HS-mimetic oligomer library. A native HS hexasaccharide is used as a control in the SPR competitive binding assay. (b) Workflow of the glycan microarray experiment. (c) Binding profile of the HS-mimetic oligomer library microarray at 100  $\mu\text{M}$  and 300  $\mu\text{M}$  ( $n = 4$ ). The concentration of FGF2 was 5  $\mu\text{g mL}^{-1}$ . P: buffer only. (d) Workflow of the SPR experiment. (e) FGF2 binding of the HS-mimetic oligomers and native HS hexasaccharide in competition with the immobilized heparin in the SPR experiment. The results are fitted to the competitive inhibition model.

heparin could be determined by fitting the concentration-dependent SPR responses to a competitive inhibition model (see ESI Methods<sup>†</sup>).<sup>38</sup> Consistent with the microarray results, oligomers with higher sulfation, *e.g.*, **K** ( $\text{IC}_{50} = 5.9 \mu\text{M}$ ) and **G** ( $\text{IC}_{50} = 6.4 \mu\text{M}$ ), exhibited higher affinity to FGF2 than the ones with lower sulfation, *e.g.*, **C** ( $\text{IC}_{50} = 39.0 \mu\text{M}$ ) (Fig. 4e and S4<sup>†</sup>). Notably, the  $\text{IC}_{50}$  of **K** is only slightly lower than a native HS hexasaccharide **S**, GlcNS(6S)-GlcA-GlcNS(6S)-IdoA(2S)-GlcNS(6S)-GlcA ( $\text{IC}_{50} = 2.0 \mu\text{M}$ ). Furthermore, the  $\text{IC}_{50}$  value of **K** is also consistent with the FGF2 affinity of an analogous native IdoA-GlcNS(6S)-IdoA(2S)-GlcNS(6S) tetrasaccharide reported by Boons *et al.* ( $K_{\text{D}} = 1.2 \mu\text{M}$ ).<sup>10</sup> In contrast, disaccharide **Q** and **R** are insufficient to inhibit the FGF2 binding by heparin: not only their  $\text{IC}_{50}$  values are much higher ( $\text{IC}_{50} = 45.2 \mu\text{M}$  for trisulfated **R**;  $\text{IC}_{50} = 64.0 \mu\text{M}$  for disulfated **Q**), the maximal inhibition for **Q** and **R** were also lower than 30% of the total binding. Taken together, the structure-activity relationship demonstrated by the HS-mimetic oligomers in FGF2 recognition is consistent with that of the native HS oligosaccharides.<sup>10,39,40</sup>

### Molecular dynamics simulations

While the CuAAC coupling chemistry enables highly efficient iterative assembly in the solution phase, it involves incorporation of a non-natural triazole spacer between adjacent

disaccharide units. To study how the spacer in HS-mimetic oligomers affects their ability to replicate behavior of native HS, we performed molecular dynamics (MD) simulations of compound **K** (Fig. 5a) and compared it with its native counterpart, a IdoA(2S)-GlcNAc(6S)-IdoA(2S)-GlcNS(6S) tetrasaccharide. To model the compounds, we employed a combination of the GLYCAM 06j<sup>41,42</sup> and generalized Amber force fields<sup>43</sup> in the OpenMM software package<sup>44</sup> (refer to ESI<sup>†</sup> for details). We analyzed four major polymer properties to compare the behavior of HS mimetic with native HS: (1) glycosidic linkages; (2) ring-flipping states; (3) end-to-end distance; and (4) ion atmosphere. For this, we used the crystallographic conformation of an FGF2-bound HS tetramer (PDB: 1BFB<sup>45</sup>) as a point of reference. In addition, we performed simulations of FGF2 binding to demonstrate that mimetic HS can reproduce the sulfate-dependent binding behavior of the native polymer.

Rotation about glycosidic linkages are important for determining the long-chain conformation of HS polymers.<sup>46</sup> For **K**, we find that the flexibility of the GlcNAc(6S)-IdoA(2S) linkage is greatly enhanced due to the triazole spacer (Fig. S5a<sup>†</sup>). In contrast, conformations of adjacent linkages are identical between the native HS tetrasaccharide and **K** (Fig. S5b and c<sup>†</sup>), indicating they are unaffected by the spacer.

Ring flipping is also key for deciding HS conformation and can affect protein binding.<sup>47</sup> The ring of central IdoA(2S) has



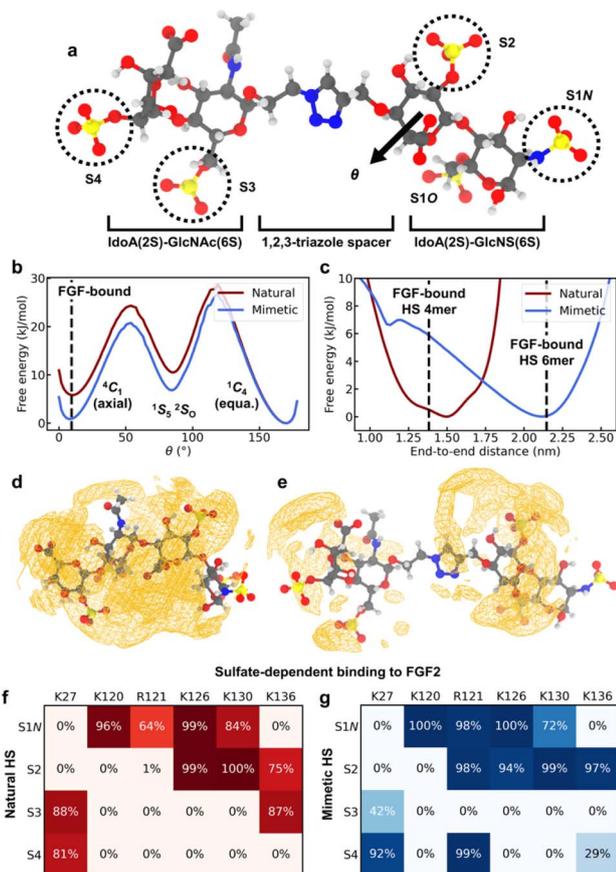


Fig. 5 MD simulation results. (a) Representative configuration of compound **K**, with the sulfate groups on each monosaccharide residue labeled. Atom color code: H – white, C – gray, N – blue, O – red, S – yellow. (b) Free energy of ring flipping for central IdoA(2S) (adjacent to spacer).  $\theta$  is the Cremer–Pople parameter<sup>50</sup> for monitoring ring flipping between chair conformers. The  ${}^1C_4$  conformer corresponds to 2-O-sulfate adopting the equatorial geometry, while  ${}^4C_1$  corresponds to the axial geometry. (c) Free energy of end-to-end distances, defined as the distance between C4 of terminal IdoA(2S) and C1 of terminal GlcNS(6S). Dashed lines give the crystallographic distances for FGF2-bound tetramer and hexamer structures (PDB: 1BFB and 1BFC). (d) Spatial distribution function (SDF) of  $\text{Na}^+$ , a representative counterion, around native HS. Orange wireframe shows the 1.2 M isosurface for  $\text{Na}^+$  concentration. (e) SDF of  $\text{Na}^+$  ions around compound **K**, with the 1.2 M isosurface shown. (f and g) Interactions between the sulfate groups of **K** or the native HS tetrasaccharide and the amino acids of the FGF2 heparin binding domain. (f) Probability for sulfate groups to be in contact with positively charged residues when the native HS tetrasaccharide binds to FGF2. A sulfate is defined to be in contact with a residue if the distance between any non-hydrogen atoms is less than 0.3 nm. (g) Probability of contact with positively charged residues when compound **K** binds to FGF2.

a thermodynamically preferred chair conformer ( ${}^1C_4$ ) (Fig. 5b), consistent with previous studies.<sup>47,48</sup> In **K**, the less favored chair conformer ( ${}^4C_1$ ) is stabilized by  $\sim 5$  kJ mol<sup>-1</sup> relative to the native HS tetrasaccharide. The lower energy of this conformer ( ${}^4C_1$ ) in the mimetic is due to its spacer, which creates a spatial buffer between residues and mitigates unfavorable interactions in the axial geometry. Based on the crystal structure (PDB: 1BFB) reported by Faham, *et al.*,<sup>45</sup> HS tetramer binds to FGF2 mainly

through two adjacent sulfate groups, one of which is on a central IdoA(2S) residue. Interestingly, in order for FGF2-binding to occur, IdoA(2S) must adopt the less favored ring conformer (Fig. 5b). Since the triazole spacer in mimetic HS stabilizes the less favored conformer, this suggests that the free-energy cost of binding due to ring flipping is lowered in the mimetic. Meanwhile, ring flipping for other residues is similar, in terms of free energy, between native HS tetrasaccharide and **K** (Fig. S6†).

The overall polymer conformation of HS can be characterized by measuring its end-to-end distance. The preferred end-to-end distance of native HS tetrasaccharide is 1.5 nm, close to the FGF2-bound distance (Fig. 5c). In contrast, the most favorable distance for **K** is 2.1 nm. However, the flexibility of the triazole spacer enables **K** to adopt folded conformations where its polymer length is close to the FGF2-bound distance, at a free-energy cost of  $\sim 6$  kJ mol<sup>-1</sup>. On the other hand, the preferred end-to-end distance of **K** is closest in length to HS hexamer (PDB: 1BFC<sup>45</sup>) when bound to FGF2 (Fig. 5c).

Protein binding in HS occurs through its sulfate groups,<sup>49</sup> which requires desolvation of counterions. We analyzed  $\text{Na}^+$  solvation around native HS tetrasaccharide and compound **K** through the potential of mean force (PMF). The PMF value at a given distance from a sulfate group is the free energy change associated with moving a  $\text{Na}^+$  ion from bulk solution to that distance. The PMF between sulfate groups and their surrounding  $\text{Na}^+$  solvation shell is greater in the native HS tetrasaccharide than in **K** (Fig. S7†). This suggests that desolvation of  $\text{Na}^+$  ions around sulfate groups can occur more easily in the mimetic than in native HS. This effect can be attributed to the greater polymer length, and hence smaller linear charge density, of **K**. Accordingly, we find the counterion density around most regions of compound **K** to be smaller than that around native HS (Fig. 5d and e). Furthermore, the average excess number<sup>51</sup> of  $\text{Na}^+$  ions surrounding native HS is greater than that around the mimetic by 0.4 (Fig. S7†). This is due to the stronger interaction of native HS tetrasaccharide, compared to **K**, with its surrounding  $\text{Na}^+$  ion environment.

Finally, we compared the sulfate-dependent binding behavior of native HS tetrasaccharide and **K** by performing representative simulations of these two oligomers binding to FGF2. Based on the crystal structure (PDB: 1BFB<sup>45</sup>), HS tetramer binds to the positively charged domain of FGF2 containing Lys120, Arg121, Lys126, and Lys130. For the native HS tetrasaccharide, we reproduced this interaction in our simulations (Fig. S8a†) and find that the polymer remains stably bound to FGF2 (Fig. S8c†). Specifically, sulfate groups S1N and S2 (Fig. 5a) show the most extensive binding to the positively charged domain (Fig. 5f and S8a†), consistent with the crystallographic binding configuration (Table S1†). Sulfates S3 and S4 (Fig. 5a) can also interact *via* Lys27 and Lys136, though to a lesser degree. Likewise, **K** binds to FGF2 stably (Fig. S8d†). This binding occurs predominantly through S1N and S2 (Fig. 5g and S8b†), the same as in the native HS tetrasaccharide and similar to the crystal structure (Table S1†). In addition, **K** adopts a partly folded conformation (Fig. S8b†), enabled by the flexibility of the triazole spacer. This conformation facilitates partial



interactions of S3 with Lys27 and binding of S4 to both Lys27 and Arg121 (Fig. 5g and S8b†). These lesser variations in S3/S4 interactions are the main point of difference between the bound structures of **K** and native HS tetrasaccharide. Overall, these results show that the HS mimetic is capable of reproducing the sulfate-dependent binding behavior of its native counterpart.

The simulation results also provide further insights on the sulfate-mediated binding of HS-mimetic oligomers. From the protein-binding studies by microarray and SPR, we find that oligomers **G–K** show strong binding affinity to FGF2 (Fig. 4a and c). With the exception of **G**, these mimetic oligomers all possess *N*- and 2-*O*-sulfation on adjacent residues. This is consistent with our simulations, which indicate that the predominant interactions of the glycomimetic oligomers with the positively charged domain of FGF2 occurs through adjacent *N*- and 2-*O*-sulfates (*i.e.*, S1*N* and S2 in Fig. 5g). The importance of these two sulfation positions is also observed in native HS (Fig. 5e).<sup>39,40,45</sup> In addition, the strong binding oligomers (**G–K**) have the highest sulfation density, with 4–5 sulfate groups across four monosaccharide units. The simulated binding conformation of **K** illustrates how up to four of these sulfates can simultaneously participate in interactions with FGF2 (Fig. 5g and S8b†), which explains the contribution of the additional sulfate groups in **G–K**. Binding by these additional sulfate groups is also seen in native HS tetrasaccharides (Fig. 5e).<sup>45</sup>

Overall, our simulations suggest that the greater flexibility and weaker ion interactions of the HS-mimetic oligomers can compensate for the effects of the triazole spacer. This can lead to a high degree of similarity in protein binding between HS-mimetic oligomers and native HS, as shown by their sulfate-dependent interactions with FGF2.

## Conclusions

In conclusion, we developed a new approach to HS glycomimetics based on iterative assembly of clickable disaccharide building blocks. Mimicking the disaccharide repeating units of native HS, these clickable disaccharides can undergo solution-phase iterative assembly *via* click chemistry to generate sequence-defined glycooligomers with variable sulfation patterns. To demonstrate the versatility of this general approach, we constructed a library of HS-mimetic oligomers with defined sulfation patterns that can be readily sequenced by tandem mass spectrometry. Microarray and SPR experiments confirmed that, despite incorporating non-natural triazole spacers, these HS-mimetic oligomers bind FGF2 in a sulfation-dependent manner consistent with that of the native HS. This structure–function relationship was further confirmed by MD simulations of the HS-mimetic tetramer and the corresponding native HS tetrasaccharide. As a future direction, we envision that the combinatorial assembly of modular HS disaccharide structures *via* flexible non-natural linkers will allow access to sulfation patterns and conformations both within native HS and beyond. Such an ability makes the method presented herein readily applicable to generating glycomimetic oligomers capable of interacting with other HS-binding proteins. In

parallel to our work, Huang and coworkers demonstrated that the amidation chemistry widely used in peptide synthesis can also be used to construct HS-mimetic oligomers with diverse sulfation patterns.<sup>52</sup> Collectively, these results established general approaches to HS glycomimetics that can serve as promising alternatives to native HS in functional studies in both fundamental research and disease models.

## Data availability

MD simulation files are included in the electronic attachment (HS\_simulation\_files.zip); all trajectory data are available on request. All other data, including the experimental data and characterizations of the new compounds, are available in the manuscript or the ESI.†

## Author contributions

CY, YD and JN conceived and designed the project and wrote the manuscript. JN oversaw the project. CY, YD, PB, CL, SH, JF, MC, AS, CL, and ACZ synthesized the clickable disaccharides and the HS glycomimetics. YW and UM performed the computational studies. AYM, KJB, and RDC performed the microarray experiment. CX and CL conducted the MS/MS analysis of the glycomimetic oligomers. All authors discussed the results and commented on the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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