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Precision dendritic-supramolecular glycan assemblies for probing multivalent lectin interactions†

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Multivalent glycan–protein binding events participate in various physiological functions including cell signaling, immune response, and pathogen–host recognition, among others. The complexity of these processes has driven sustained interest in developing densely functionalized glycan nanoassemblies to probe and modulate these important interactions. While synthetic glyconanomaterials have demonstrated significant promise in nanomedicine and biotechnology applications, achieving precise control over their architecture remains challenging. To address this limitation, we present a new method of synthesizing molecular glycan nanoassemblies by integrating the dense functionalization of dendritic architectures with the preorganized scaffolding of metallosupramolecular frameworks. A family of Fe(II)-anchored superassemblies featuring 24, 36, and 72 peripheral mannosides was prepared and characterized by spectroscopic, microscopic, and light scattering techniques. These assemblies demonstrate strong binding with the lectins Concanavalin A (Con A) and Griffithsin (GRFT), as evaluated by isothermal titration calorimetry (ITC), with the 72-mannose derivative exhibiting low nanomolar binding affinity ($K_d = 28 \pm 4$ nM for Con A; 12 ± 1 nM for GRFT). The high binding strength of these assemblies highlights the potential of integrating dendritic architectures with rigid metallosupramolecular cores to enhance lectin recognition. Our findings present a new framework for probing glycan protein interactions and offer insights into the design of hybrid glycoassemblies as biomedically-relevant tools.

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Introduction

Protein–glycan interactions play key roles in many biological processes including molecular recognition events, cellular communication, and signal transduction, among others.^{1–3} The importance of these biomacromolecular interactions has driven sustained interest across multiple scientific disciplines toward uncovering and understanding the nature of these events, as demonstrated by approaches spanning chemical biology,^{4,5} materials science,^{6–8} and chemical synthesis.^{9–11} Probing these interactions has become increasingly important in biomedical and biotechnology applications,^{12–17} as these processes are central to many diseases and signalling pathways.^{18–20} At the molecular level, protein–glycan recognition occurs through multivalent binding, where multiple weak, non-covalent, and

reversible interactions work cooperatively to result in enhanced functional affinity that exceeds the binding strength of the single carbohydrate–protein interaction.²¹ This fundamental principle enables nature to achieve remarkably strong and selective molecular recognition, as evidenced by protein–glycan interactions that typically display micromolar to nanomolar binding affinities ($K_d \sim 10^{-6}$ to 10^{-9} M).^{12,22} A detailed understanding of the macromolecular mechanisms underlying carbohydrate recognition is therefore essential for driving the development of innovative tools, treatments, and biomedical agents.

Synthetic platforms that effectively mimic and probe these molecular recognition events hold potential in therapeutic and bioengineering applications, especially in physiological processes where protein–glycan interactions are critical mediators. As the synthesis of glycan assemblies continues to evolve, sustained progress is guided by the need for precise control over molecular architecture and the demand for access to complex structures reminiscent of nature's sophisticated biological ligands.^{11,23,24} A variety of platforms have been used to this end, including dendrimers,²⁵ small organic and inorganic molecules,^{26–31} polymers,^{15,32,33} peptides,³⁴ nanoparticles,³⁵ and templated systems that pre-organize glycans on protein or synthetic scaffolds.^{36–38} However, despite considerable advances

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Fig. 1 Superassembly framework showing the core structure, dendron generations, and terminal carbohydrate units.

in synthetic methodology, examples of structurally well-defined glycoassemblies remain scarce and many approaches lack the generalizable synthetic versatility needed to systematically investigate the relationships governing molecular architecture and binding efficiency.^{39–41} While many conventional platforms like polymers or nanoparticles achieve high affinity, these glycan arrays frequently exhibit intrinsic polydispersity in scaffold size and ligand valency, which can lead to unpredictable binding behaviours and reproducibility challenges.^{31,32,42,43} Monodisperse and structurally tunable ligand assemblies remain underexplored in glycomaterial development, however, despite their importance in the rational design of new multivalent glycomimetics. Motivated by this need, we sought to employ synthetic supramolecular systems that can probe these important recognition events while offering molecular precision and versatility.

Our approach leverages the complementary advantages of dendritic architectures and metallosupramolecular scaffolding to introduce well-defined hybrid glycan systems with high lectin binding affinity. Recent advances in dendrimer chemistry have provided powerful tools for achieving dense surface functionalization, which allows for the systematic incorporation of saccharide units at specifically defined positions.^{34,44–47} Unlike traditional polymer and nanoparticle platforms, dendrimers offer higher levels of structural precision through their step-wise synthesis. Although dendrimer scaffolds provide valuable architectural control, they often suffer from high conformational flexibility and limited structural preorganization, which is typically needed to maximize lectin binding efficiency.¹¹ To address these limitations, we incorporated coordination-driven subcomponent self-assembly in our approach. Coordination-driven subcomponent self-assembly is a powerful synthetic strategy that employs molecular synthons and non-covalent interactions to construct rigid and complex architectures while maintaining the same degree of precision and control

attainable in molecular synthesis.^{48,49} While metal-anchored supramolecular architectures provide well-defined frameworks with structured spatial arrangements of surface-grafted elements,^{50–53} these systems typically offer limited peripheral functionalization compared to dendritic platforms. By preserving the advantages of both approaches, this work integrates glycosylated dendritic fragments within rigid metal-organic polyhedral frameworks to introduce a family of densely grafted, architecturally precise superassemblies (Fig. 1) that demonstrate exceptionally strong lectin binding affinity.

Results and discussion

We recently reported work highlighting the versatility of coordination-driven subcomponent self-assembly⁵⁴ in the synthesis of structurally well-defined saccharide-grafted Fe(II)-anchored supramolecular assemblies.⁵⁵ This synthetic approach allowed for molecular-level control^{48,49} over assembly size, shape, topology, saccharide surface density, and charge.⁵⁶ Depending on the number of iron centres present in the structure, up to twelve saccharides were introduced on the assembly periphery in a straightforward manner, which enabled precise tuning of the structural valency. Building upon key design principles established from our work, the present study employs diethyleneglycol (DEG) linkages to tether saccharide units to supramolecular frameworks, as these linkers offered optimal carbohydrate spatial arrangements and flexibility that enhanced lectin binding affinity.⁵⁶

Here, we have expanded the synthetic landscape by introducing nanoassemblies with significantly higher valencies than previously reported. This report presents a new series of dendritic glycan picolinaldehyde ligands bearing two, three, and six mannose residues that were prepared through copper-catalysed azide-alkyne cycloaddition chemistry (CuAAC, Fig. 2). When paired with a tritopic amine subcomponent that enforces a tetrahedral, tetrametallic core, these combinations gave rise to imino-pyridine supported glycan superassemblies bearing 24, 36, and 72 mannosides through an iron(II)-templated synthesis (Scheme 1). Evaluation of these hybrid glycan assemblies revealed strong binding affinity toward the lectins Concanavalin A (Con A) and Griffithsin (GRFT), with dissociation constants reaching low nanomolar levels. Notably, binding to Con A surpassed that of our previously reported systems by over two orders of magnitude.⁵⁶

Dendron ligand synthesis

The first generation of di- and tri-substituted glycodendrimer picolinaldehyde ligands was synthesized starting from commercially available 3,5-dihydroxymethylbenzoate and methyl gallate reagents, respectively. These precursors were treated with propargyl bromide, followed by reduction with LiAlH₄. Bromination of the benzyl alcohol products with PBr₃ yielded the corresponding benzyl bromide analogues, which were treated with 5-hydroxypicolinaldehyde to afford the “clickable” terminal alkyne ligand precursors (a, b, Fig. 2). These products were subsequently treated with an excess of α -D-





Fig. 2 Synthetic schemes for the preparation of dendritic subcomponents **man-2**, **man-3**, and **man-6** from **a**, **b**, and **c** precursors, respectively, with corresponding DLS data (bottom right, measured in H_2O , 1 mM).

mannopyranosyl(diethyleneglycol) azide under CuAAC conditions with $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ as the catalyst and sodium ascorbate as the reducing agent. Flash liquid chromatography purification provided the acetate-protected di- (**man-2(OAc)**) and trimannoside (**man-3(OAc)**) picolinaldehyde derivatives in 73 and 79 percent yields, respectively. Characterization of these products by ^1H NMR spectroscopy demonstrated complete conversion of the terminal alkyne units of **a** and **b** into triazole moieties as evidenced by the disappearance of the alkyne resonances (δ 2.52 (**a**); 2.49, 2.46 (**b**) ppm in CDCl_3), and the appearance of triazolyl resonances located at δ 7.77 and 7.84 ppm for **man-2(OAc)** and **man-3(OAc)**, respectively. In the ^{13}C NMR spectrum, the signals located at δ 144.4 and 124.0 ppm for **man-2(OAc)**, and δ 144.4, 143.5, 124.6, and 124.3 ppm for **man-3(OAc)** additionally confirmed the presence of triazolyl carbons.^{9,57} The ^1H and ^{13}C NMR spectra reveal single species in solution that exhibit two-fold rotational symmetry, consistent with the proposed molecular structures.

The Fourier-transform infrared (FTIR) spectra of both subcomponents lack the characteristic signatures of alkyne or azide functional groups, which supports covalent attachment of the saccharide units to the picolinaldehyde anchor (ESI Section S2.1†), and clean isolation of the 1,2,3-triazole glycosides. Quantitative de-*O*-acetylation under standard Zemplén conditions (cat. NaOMe , MeOH , Fig. 2)⁵⁸ yielded the deprotected α -mannopyranoside derivatives, **man-2** and **man-3**, after neutralization with the acidic Amberlyst-15® ion-exchange resin. Full deprotection of the mannopyranose groups was confirmed by ^1H NMR spectroscopy based on the disappearance of acetate resonances located between δ 1.8–2.1 ppm (D_2O). Electrospray ionization mass spectrometry (ESI-MS(+)) analyses of the products in positive mode displayed signals corresponding to the protonated molecular ions $[\text{M} + n\text{H}]^{n+}$, which confirmed both the empirical formula and integrity of the species. Dynamic light scattering (DLS) measurements provided further

characterization and offered insights into the molecular dimensions of these compounds by determining their hydrodynamic particle size distributions in solution. Measurements conducted in aqueous media (1 mM) demonstrated well-defined size distributions for both **man-2** and **man-3**, with hydrodynamic diameters (D_h) of 1.6 ± 0.6 and 1.8 ± 0.1 nm respectively (Fig. 2). These dimensions align with the expected values for non-aggregated single particles, providing strong evidence for the structural integrity and monodispersity of these species in solution.

We next focused on expanding this platform by preparing a second-generation dendron ligand based on a branched hexaalkynylated picolinaldehyde core that would provide access to a hexa-mannoside subcomponent. The synthesis of hexa-functionalized dendritic architecture, **man-6**, mirrored the procedure described for the syntheses of **man-2** and **man-3** following the initial coupling of 3,4,5-trispropargyloxybenzyl bromide (two equiv.) with 3,5-dihydroxymethylbenzoate in the presence of K_2CO_3 and 18-crown-6.⁵⁹ Reduction of the methyl ester with LiAlH_4 followed by bromination provided the hexaalkynylated benzyl bromide dendrimer core, which enabled installation of the picolinaldehyde unit *via* $\text{S}_{\text{N}}2$ substitution chemistry. With this building block, the targeted sixfold-substituted glycan subcomponent, **man-6(OAc)**, was synthesized through a hexa-click reaction following the same CuAAC parameters outlined for the first-generation derivatives (*vide supra*). Characterization of **man-6(OAc)** by ^1H and ^{13}C NMR spectroscopy revealed signatures consistent with those observed for the **man-2(OAc)** and **man-3(OAc)** analogues, supporting successful synthesis and isolation of the desired two-fold symmetric glycoconjugate. FTIR spectroscopy was employed to confirm the absence of alkyne and azide functional groups, which further substantiated the identity of the desired compound. ESI-MS(+) analysis of the deprotected **man-6** glycan was consistent with the protonated $[\text{M} + 3\text{H}]^{3+}$ ion, thereby





Scheme 1 Fe(II)-directed subcomponent self-assembly reactions to prepare [1][BF₄]₈, [2][BF₄]₈, and [3][BF₄]₈. Only one of four face-capping ligands is shown for clarity.

confirming the identity and molecular composition of the ligand. The ¹H NMR spectrum of deprotected **man-6** displayed broadened signals when recorded at ambient temperature at concentrations suitable for spectroscopic analysis (10–30 mM), which is consistent with the tendency of densely functionalized glycan assemblies to form clustered aggregates in aqueous solution.⁹ Collection of ¹H NMR data in CD₃ OD, however, led to sharper signals and reduced aggregation, likely due to the disruption of carbohydrate–aromatic π interactions that occur in aqueous solution.⁶⁰ At significantly lower concentrations (1 mM) in water, however, a hydrodynamic diameter aligning with a single particle measuring $D_h = 2.4 \pm 0.2$ nm was observed (Fig. 2), further demonstrating the architectural integrity of the product.

Superassembly synthesis

With **man-2**, **man-3**, and **man-6** subcomponents in hand, we focused on preparing saccharide grafted supramolecular assemblies utilizing our established Fe(II)-driven subcomponent self-assembly methodology.^{55,56} The glycan ligands were paired with the isocyanurate-based tritopic amine subcomponent shown in Scheme 1. This subcomponent was selected due to its water-solubilizing polar groups that impart the aqueous

compatibility required for this work when paired with a water-soluble iron(II) salt. Furthermore, we demonstrated⁵⁶ that this tritopic amine subcomponent directs the formation of a face-capped Fe₄L₄ structural motif,^{61–63} where four Fe(II) centres occupy the vertices with pseudo-octahedral geometry, while trigonally-symmetric iminopyridine ligands span the faces.^{64,65} Since each iron centre coordinates three bidentate ligands based on single-crystal X-ray diffraction studies of structurally analogous M₄L₄ (M = Fe(II), Co(II), Zn(II)) assemblies,^{50,64–67} the resulting complexes supported by **man-2**, **man-3**, and **man-6** give rise to glycan superassemblies grafted with 24, 36, and 72 saccharides, respectively. This structural configuration provides access to architectures decorated with significantly higher saccharide loadings that are two-, three-, and six-fold greater than those of our previously reported systems.^{55,56} This enhanced functionalization offers potential for both stronger lectin binding affinity and broader applications for these constructs.

Accordingly, treatment of triamine isocyanurate (1 equiv.) with subcomponents **man-2**, **man-3**, and **man-6** (3 equiv.) in the presence of Fe(BF₄)₂·6H₂O (1 equiv.) in H₂O/MeCN (2 : 1) solvent mixtures at 25 °C afforded the 24- ([1][BF₄]₈), 36- ([2][BF₄]₈), and 72-fold ([3][BF₄]₈) mannose-substituted assemblies,





Fig. 3 Characterization data of $[3][\text{BF}_4]_8$: (A) ^1H NMR spectrum (D_2O , 45 °C); (B) UV-vis spectrum (H_2O , 50 μM); (C) ATR-IR spectrum; (D) AFM images and height profiles (tapping mode, air, 298 K) of a drop-cast H_2O solution (0.2 μM) on mica; (E) DLS measurement (50 μM , 85% v/v DMSO/ H_2O); (F) negative-contrast electron micrograph (uranyl acetate).

respectively (Scheme 1), which were subsequently purified *via* centrifugal ultrafiltration. To provide a thorough analysis of their structural and molecular properties, the three super-assemblies were characterized by spectroscopic, microscopic and light scattering techniques. The ^1H NMR spectra of $[1]^{8+}$, $[2]^{8+}$, and $[3]^{8+}$ collected in D_2O at 25 °C revealed broadened resonances with increased linewidths, which is attributed to slow dynamic motion of the cages in solution, and is typical of large molecular-weight molecules (MW $[1][\text{BF}_4]_8 = 13\,200$, $[2][\text{BF}_4]_8 = 17\,400$, $[3]^{8+} = 32\,400$ g mol^{-1}) that display short T_2 relaxation times.^{26,68–70} We therefore collected ^1H NMR spectra of these systems at slightly elevated temperatures to increase the overall molecular tumbling rate, which resulted in moderately sharpened and more well-defined signals for all three complexes (ESI Fig. S44, S47 and S50[†]). However, due to the limited thermal stability of these systems, we were unable to collect ^1H NMR data at temperatures exceeding 45 °C. Despite these limitations, the ^1H NMR spectra of $[1]^{8+}$, $[2]^{8+}$, and $[3]^{8+}$ display signals that are consistent with single, T -symmetric structures in solution, which enable confirmation of the structural integrity of the species. Imine-bond formation during the self-assembly reactions was confirmed by the disappearance of aldehyde resonances located at δ 9.55, 9.79, and 9.97 ppm

(D_2O), concomitant with the appearance of new, upfield-shifted signals attributed to the imine protons, located at δ 9.15, 8.85, and 8.77 ppm for complexes $[1][\text{BF}_4]_8$, $[2][\text{BF}_4]_8$, and $[3][\text{BF}_4]_8$, respectively. When measured at 45 °C, the ^1H NMR spectra of $[2][\text{BF}_4]_8$ and $[3][\text{BF}_4]_8$ (Fig. 3A) each displayed two distinct singlets (δ 7.99, 7.74 ppm for $[2][\text{BF}_4]_8$; δ 7.96, 7.77 ppm for $[3][\text{BF}_4]_8$), corresponding to the two chemically unique 1,2,3-triazole methine protons present in each complex. In contrast, the ^1H NMR spectrum of $[1][\text{BF}_4]_8$ contains a single resonance located at δ 8.21 ppm, which is consistent with one unique triazolyl methine proton.

Additional spectroscopic characterization was performed by UV-vis analysis (50 μM , H_2O) for all superassemblies, which displayed characteristic metal-to-ligand charge transfer excitations for low-spin iron(II) in an iminopyridine coordination environment ($\lambda_{\text{max}} = 505, 550$ nm, Fig. 3B), which align well with the data reported for our first series of Fe(II) glycan assemblies.^{55,56} The FTIR spectra of complexes $[1][\text{BF}_4]_8$, $[2][\text{BF}_4]_8$, and $[3][\text{BF}_4]_8$ provide further evidence supporting the formation of an imine bond during self-assembly reactions, as indicated by the absence of aldehyde bands and the presence of strong vibrations ascribed to the imino C=N stretching mode ($[3][\text{BF}_4]_8$ $\nu_{\text{C=N}} 1558$ cm^{-1} , Fig. 3C).⁷¹



site binding model based on monomeric Con A. The dissociation constants ($K_{d(\text{avg})}$) for the interactions of $[1]^{8+}$ and $[2]^{8+}$ with Con A were determined to be $1.1 \pm 0.1 \mu\text{M}$ and $290 \pm 4 \text{ nM}$ (Table 1), respectively, through three independent titrations of each complex. The K_d values illustrate significant affinity enhancements, with 17- and 67-fold increases compared to the binding of our previously reported twelve-mannose analogue with Con A ($[\text{Fe}_4\text{L}_4\text{-}12][\text{BF}_4]_8$; $K_d = 18.9 \pm 2.5 \mu\text{M}$).⁵⁶ We attribute the increase in binding affinity to the enhanced cluster glycoside effect^{83–85} resulting from the significantly higher density of peripheral saccharides.

Complex $[3]^{8+}$ demonstrated the strongest binding to Con A among the superassembly family, with a calculated dissociation constant of $28 \pm 4 \text{ nM}$ (Fig. 4). This binding strength surpasses α -D-mannose ($470 \mu\text{M}$),^{86,87} by approximately four orders of magnitude, representing the strongest interaction observed for any supramolecular glycoassembly in our current library of 17-mannosylated complexes.^{55,56} The binding affinity demonstrated by $[3]^{8+}$ approaches levels typically observed for densely-functionalized nanoparticle and polymer-based platforms with Con A, with only a few examples reaching such interaction strengths to our knowledge.^{86–89} Furthermore, when normalized on a per-saccharide basis, the immobilization of mannose units on the superassembly framework increases the binding strength of each mannoside by a factor of 231. This outcome is particularly noteworthy, as $[3]^{8+}$ exhibits a greater binding enhancement on a per-sugar basis than many existing polymeric, dendritic, and nanoparticle systems, which typically require significantly higher carbohydrate densities to achieve comparable affinity,^{86,87,90} which highlights the structural efficiency of this superassembly design.

Since the separation between dimeric Con A binding sites precludes chelation by the glycoassemblies, the enhanced affinity of these systems is more likely attributed to statistical rebinding and crosslinking multivalent mechanisms. These operating mechanisms are well-precedented for other multivalent platforms,⁹¹ and can substantially enhance apparent

affinity without requiring simultaneous engagement of multiple sites on a single lectin. This interpretation aligns with our prior work on the lower-valent, Fe_4L_4 dodeca-functionalized analogues,⁵⁶ where extensive DLS experiments provided evidence consistent with crosslinking and statistical rebinding.

We further attribute the enhanced binding capabilities of these systems to the preorganized supramolecular framework, which combines the structural rigidity of the inorganic cage core with the flexibility of the dendritic glycan ligand, enabling these systems to achieve a similar binding strength to systems with much higher valencies. Although the Fe_4 core is positioned away from the binding site (*ca.* 2 nm), its structural rigidity pre-organizes the terminal mannoside groups, likely reducing the configurational entropic penalty typically observed in flexible multivalent systems such as conventional glycodendrimers. It is well-documented that preorganized glycan architectures can enhance lectin binding while minimizing overall entropy cost.^{92,93} The combination of the rigid Fe_4L_4 core with the flexible dendron ligand fragments has the potential to simultaneously provide flexibility for the systems to access and adapt to the lectin CRD (carbohydrate recognition domain) while reasonably minimizing the conformational enthalpy penalty. Striking the right balance between rigidity and flexibility in multivalent glycosystems has become a central focus within the expanding field of glyconanotechnology, as architectural optimization has been shown to increase avidity by several orders of magnitude beyond what is possible through statistical rebinding events alone.^{11,94,95}

Building on these findings, we expanded this work to investigate the interactions between our superassembly platform and GRFT, a broad-spectrum antiviral lectin that specifically recognizes high-mannose glycans.⁹⁶ Consistent with the binding trend observed for the assemblies with Con A, glyco-cluster association with monomeric GRFT also displayed clear valency-dependent binding, with complex $[3]^{8+}$ exhibiting the highest affinity ($K_d = 12 \pm 1 \text{ nM}$). This binding strength represents a significant enhancement over that of monovalent α -D-mannose, which binds GRFT with a K_d value of approximately $100 \mu\text{M}$,⁹⁷ representing a nearly 8000-fold increase in overall binding strength, and a 110-fold enhancement per saccharide. These results demonstrate that the multivalent presentation of mannosides on the dendritic metallosupramolecular framework not only enhances affinity toward canonical targets like Con A but also extends to therapeutically relevant lectins such as GRFT. The ability of these systems to engage both lectins with nanomolar affinity highlights the potential utility of this approach as a broadly applicable glycan display platform for probing and modulating diverse carbohydrate-protein interactions.

We acknowledge that DLS, TEM, NMR, and AFM analyses collectively reveal some degree of aggregation of the supramolecular constructs in water, which likely stems from weak intermolecular carbohydrate-aromatic π -interactions rather than structural decomposition of the underlying assembly framework. This interpretation is supported by retained imine proton resonances in the ^1H NMR spectra in aqueous solution (δ 9.15–8.77 ppm) and unchanged UV-vis profiles (λ_{max} 505–550 nm) under various conditions, confirming the integrity of the



Fig. 4 Left: ITC thermogram (top) and fitted binding isotherm (bottom) for the binding of $[3][\text{BF}_4]_8$ to dimeric Con A. Right: Model of the binding interaction showing the Con A CRD (PDB code 1QDC).



metal–ligand bonds within the complexes. While these assemblies are architecturally well-defined and monodisperse under mixed solvent conditions, aggregation in aqueous environments may affect the apparent glycan valency during lectin binding and should be taken into consideration when interpreting affinity measurements. Since aggregation may lead to an underestimation of the effective valency, the K_d values must be interpreted with some caution, as the structural valency may not be entirely representative of monomeric binding. These assemblies, however, remain fully soluble in water, suggesting they remain dynamic and accessible for molecular interactions. Similar behaviour has been observed in other densely functionalized glycan architectures, where self-aggregation does not preclude their use in biomedical applications but rather represents an intrinsic structural feature that can influence binding interactions.^{9,72} Nevertheless, these supramolecular architectures represent unique and well-characterized examples of discrete glycoassemblies capable of high-affinity interactions, offering a versatile platform for further development in biomedical and therapeutic contexts.^{9,72}

Conclusions

The molecular recognition of carbohydrates in biological systems is directly tied to both the chemical composition and spatial presentation of surface glycans. Motivated by the need to understand and probe these complex interactions, we designed a new class of glycan superassemblies by introducing a synthetic strategy that merges dendritic architectures with coordination-driven self-assembly. This approach enabled the isolation of molecular glycoassemblies exhibiting exceptional lectin binding capabilities. Picolinaldehyde di-, tri-, and hexa-substituted mannoside subcomponents were synthesized employing CuAAC click–chemistry reactions and used to prepare molecular assemblies bearing 24, 36, and 72 saccharides. The three superassemblies were characterized by standard spectroscopic techniques (¹H NMR, FTIR, UV-vis), as well as DLS, TEM and AFM measurements. All three assemblies were evaluated as multivalent binders to Con A and GRFT, revealing K_d values in the low micromolar to nanomolar range. Notably, the 72-mannoside derivative demonstrated exceptionally strong binding, which likely stems from intentional architectural design that balances scaffold rigidity with localized flexibility, establishing important principles for developing increasingly sophisticated molecular systems capable of engaging in biomolecular recognition events. The versatile nature of this synthetic strategy allows for the systematic exploration and expansion of diverse scaffold architectures and functionalities as we aim to enhance our fundamental understanding of multivalent glycan recognition and provide synthetic approaches that leverage supramolecular chemistry as a biomedically-relevant tool.^{98,99}

Data availability

The data that support the findings of this study are available in the ESI† of this article.

Author contributions

J. M. S. designed the project and experiments. J. M. S. and T. M. B. carried out all synthetic work and the characterization of compounds. J. M. S. performed ITC binding studies, and G. J. M. performed TEM and DLS characterization. W. S. and G. J. M. performed AFM experiments, which were supervised by A. R. T. The GRFT protein was expressed by E. H. under the supervision of A. J. G. The manuscript was written by J. M. S., and all authors gave final approval.

Conflicts of interest

There are no conflicts to declare.

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