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Mukul Raj Gupta, Vijayasimha Molakaseema and René Roy

Glycodendrimers: endlessly evolving architecture design, synthetic strategies, and biological applications

Review article

Glycodendrimers: endlessly evolving architecture design, synthetic strategies, and biological applications

Mukul Raj Gupta [a], **Vijayasimha Molakaseema** [a] **and René Roy** [ORCID],*,[a]

[a] Glycosciences and Nanomaterial Laboratory, Université du Québec à Montréal, P.O. Box 8888, Succ. Centre-Ville, Montréal, Québec H3C 3P8, Canada

E-mail: roy.rene@uqam.ca (R. Roy)

Abstract. The synthesis of dendritic architectures has steadily advanced to overcome challenges associated with biological applications. Key issues—including reduced toxicity, enhanced target specificity, improved stability, and efficient delivery of drugs, vaccines, and genes—have posed significant hurdles, particularly when scalability for therapeutic translation is required. Among these, optimizing dendrimer systems to engage in carbohydrate-related biological functions has remained a central focus since the introduction of glycodendrimers. The emergence of glycodendrimers has proven especially effective in contexts demanding multivalent binding interactions. This review highlights advancements in the design of multivalent carbohydrate-based nanomaterials, tracing their evolution from the early development of glycopolymers to the emergence of glycodendrimers. It presents key developments in the use of scaffolds featuring an expanded array of surface functional groups alongside variations in the chemical building blocks traditionally employed in classical dendrimer synthesis—culminating in the so-called "onion-peel strategy". These innovations are further complemented by the integration of orthogonal ligation techniques. Additionally, efforts to streamline the synthesis of multivalent glycoarchitectures have led to the emergence of simplified methodologies, including transition-metal-templated assembly and the self-organization of glycodendrimersomes.

1. Introduction

Cell surface glycoconjugates (glycoproteins, glycolipids, and glycoRNA) are fundamental in cellular communications through carbohydrate–protein interactions [1–3]. In addition, glycans can recognize each other, thus allowing multivalent intercellular events. These numerously and variably exposed saccharides comprising glycoproteins, glycolipids, and polysaccharides, such as glycosaminoglycans, are responsible for a plethora of biochemical processes, including signaling, immune function, cell adhesion, and pathogen recognition (Figure 1). Not surprisingly, they are intrinsically involved in cancer vaccines for viral and bacterial infections, and they constitute highly efficient targeting elements for gene and drug delivery, particularly as nanoparticles.

Conspicuously, carbohydrate–protein interactions are multivalent by nature and result in precise and robust binding interactions (avidity, nM range) as opposed to mM affinity when taken individually
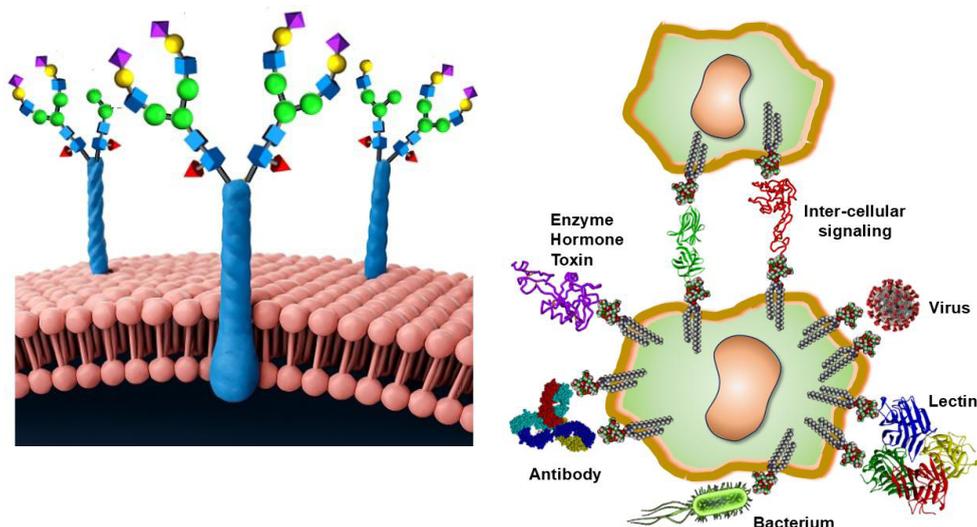
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Figure 1. Left panel: typical cell surface carbohydrates exposing multiple copies of saccharides [1–6]. Right panel: glycoproteins, glycolipids, and glycoRNA are involved in a wide range of biochemical functions.

(affinity) [7,8]. Therefore, designing artificial carbohydrate ligands capable of imitating cell surface carbohydrates necessitates three major parameters encompassing the following: (1) the exact saccharides (or oligosaccharides) involved, (2) their multivalent presentation, and importantly (3) their relative architectural topologies [9–11].

For these multiple reasons, we embarked several years ago on the design of *glycodendrimers* [12–18], following initial activities in *glycopolymers* [19–22]. As opposed to ill-defined polymers, glycodendrimers are a fascinating class of synthetic monodispersed macromolecules that combine the structural precision of dendrimers with the biological functionality of carbohydrates, usually exposed at the dendritic surfaces. The first version of glycopolymers was based on polyacrylamide and resulted in very high molecular weight materials having large polydispersity (Figure 2). These problems were only partly fixed with more modern methods such as the RAFT polymerization method and others [23–25]. Even though these glycopolymers were tremendously powerful in binding to their cognate receptors and lectins [26,27], they still lacked the ability to optimize geometrical constraints while minimizing the required number of specific glycosidic residues [28–30]. Importantly, in our opinion, obtaining precise thermodynamic binding parameters on a per saccharide basis is still an unanswered question. An additional improvement in

the field of glycopolymers was achieved with the syntheses of *dendronized glycopolymers* [31–35].

Originally, dendrimers have been designed through rather simplified synthetic strategies involving a multifunctional scaffold (typically two) such as in poly(amidoamine) (PAMAM) dendrimers followed by incremental branching with identical subunits [36]. Their construction initially followed two major paths: divergent and convergent methods. Fortunately, the strategies have rapidly evolved into more sophisticated approaches in order to expedite the synthesis of dendrimers while increasing monodispersity. This was accomplished by accelerated synthetic strategies [31,37,38].

As mentioned, these typical convergent, divergent, and accelerated methodologies are commonly created on repetitive synthetic sequences built with identical chemical motifs. Yet, to address the complexity, multivalency, and structural variety of carbohydrate receptors, we embarked on an alternative strategy through which glycodendrimers were built with varied chemical building blocks that offer the required flexibilities and specific targeting abilities [39]. The strategy, designated as "onion peel", allows the synthesis of "heterolayered" glycodendrimers encompassing a wide range of chemical scaffolds varying in chemical, physical, and biochemical properties [32,33,40–47]. This short review highlights our own work in this area.
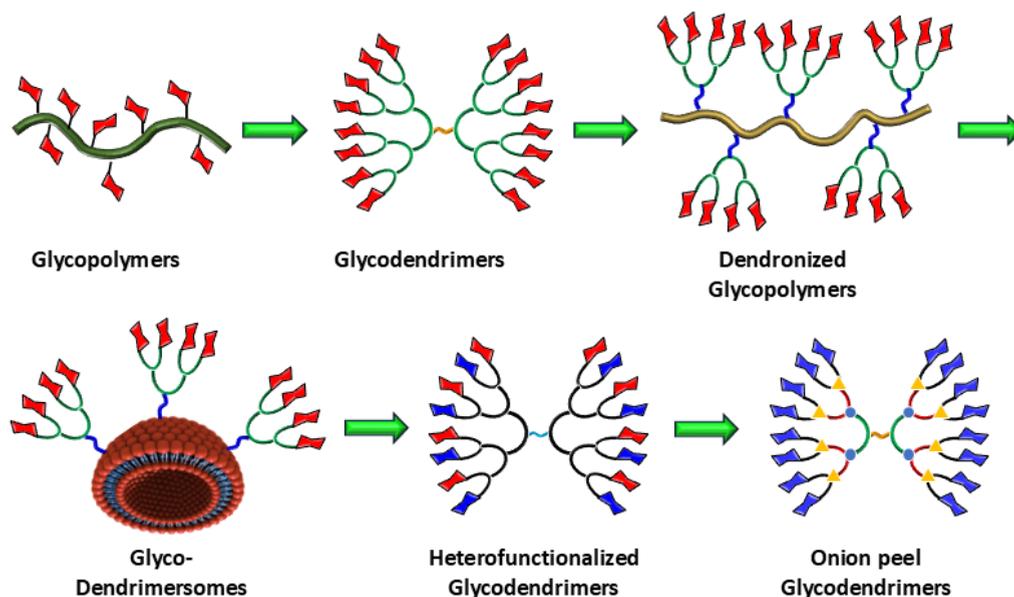
Figure 2. The course of evolution of multivalent glyconanoparticles.

2. Synthetic methodologies toward glycodendrimers

Glycodendrimers [48–50] have been constructed by chemical [38,51–57], chemoenzymatic [58,59], and self-assembling approaches [60]. In one of the later cases, transition metal complexes were first synthesized and appropriately functionalized saccharide dendrons, possessing metal ligands, were allowed to self-assemble through strong coordination/chelation chemistry. In an alternative method, glycolipids, including dendritic glycolipids, are allowed to self-assemble into glycoliposomes or glycodendrimersomes [61,62].

To further expedite the synthetic developments, highly functionalized scaffolds have also been used to accelerate multivalency [63–65]. Figure 3 illustrates some of these high valency scaffolds that have been used in the construction of glycodendrimers. For instance, Majoral–Caminade cyclotriphosphazene has been successfully used as the core molecule for the expedient synthesis of highly branched α-D-mannopyranosides [65–69]. Similarly, suitably functionalized hexaphenylbenzenes were also employed in the rapid synthesis of glycodendrimers having six exposed saccharide residues in the first generation. Analogously, aromatic scaffolds such

as calix[n]arenes [70], corannulenes [71], pyrenes, fullerenes, and single-walled carbon nanotubes also constituted high valency building blocks [72]. Interestingly, carbohydrates themselves have also been utilized as multivalent scaffolds. For example, mono-, di- and oligosaccharides represented excellent cases for the accelerated building of chiral glycodendrimers.

2.1. *Choosing high valency scaffolds*

As a typical illustration of the high potential and efficiency in the use of multifunctional scaffolds [73], our team has used β-cyclodextrin [74–76]. Cyclodextrins constitute a family of cyclic oligosaccharides derived from starch by enzymatic treatment. There are three major structures of cyclodextrins (α, β, and γ) forming 6-, 7-, and 8-membered rings through (1-4)-α-D-glucopyranosidic linkages, respectively. For instance, using β-cyclodextrin as a scaffold (**1**), the hydroxy groups were transformed into peracetylated heptakis iodo β-cyclodextrin (**2**), which upon treatment with per-*O*-acetylated 1-thio-α-D-mannose (**3**) (Cs_2CO_3, DMF) followed by the removal of *O*-acetyl protecting groups produced glycodendrimer **4** in high yield (Scheme 1). The expedient synthesis of heptakis α-D-mannopyranosides provided potent
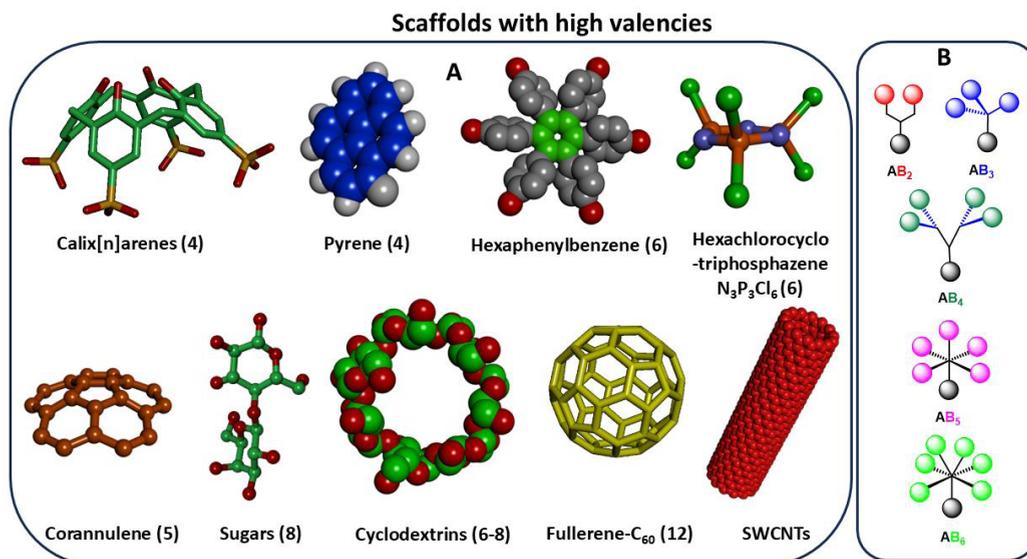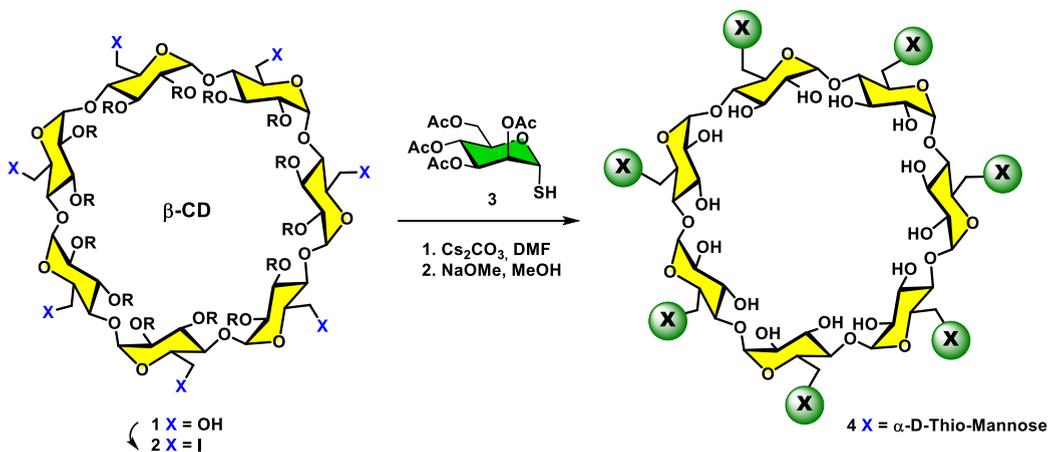
Figure 3. Multifunctional building blocks used as core molecules toward accelerated syntheses of glycodendrimers: (A) number in parentheses refers to the potential valency (exposed surface groups) on the chosen scaffolds; (B) interlayer building blocks.

Scheme 1. Using a highly functionalized scaffold such as β-cyclodextrin harboring seven primary hydroxyl groups, glycodendrimers with several copies of carbohydrate ligands (here α-D-mannopyranoside) can be rapidly constructed.

glycoclusters capable of rapidly interacting with the homotetrameric leguminous lectin isolated from *Canavalia ensiformis* (Concanavalin A [ConA]) as demonstrated by their capacity to form stable insoluble cross-linked lattices. In this work, several related families of mannosylated β-cyclodextrin were similarly prepared, some of which had aromatic linkers while others had extended linking arms. The results further demonstrated the general usefulness of carbohydrate-containing aromatic aglycones in the binding of lectins. The use of cyclodextrins as

multivalent scaffolds toward the construction of hyperbranched glycodendrimers has been comprehensively detailed in a seminal review by a Spanish group [77,78].

Several additional examples using high valency scaffolds for the construction of glycodendrimers have been reported since their early discovery in 1993 [12] (Figure 3). As mentioned, examples included calix[n]arenes [16,70,79], corannulenes [71,80,81], fullerenes [82], and hexaphenylbenzene [83]. The scope of using vastly functionalized
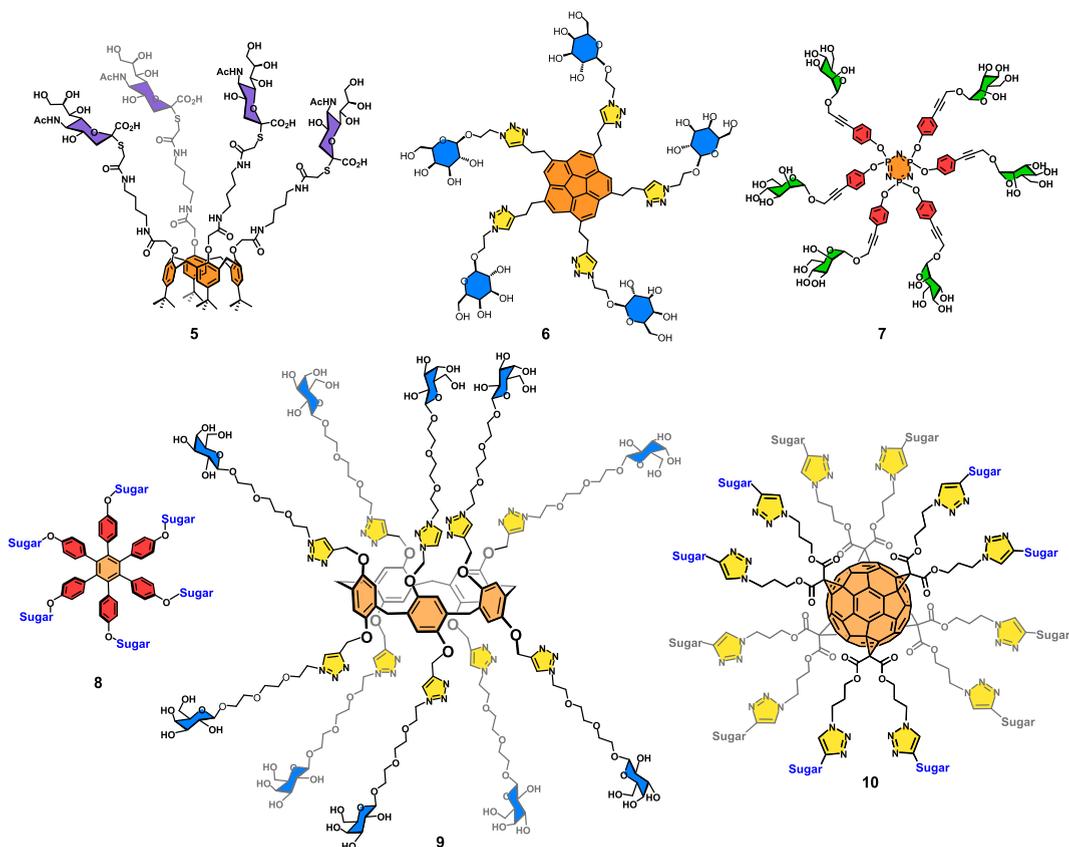
Figure 4. Typical examples of glycodendrimers built on high-valency-containing scaffolds.

aromatic scaffolds has been included in a review article by authors [71]. Figure 3 illustrates some of the most significant samples. Our group has been the first to use calix[4]arene as a tetrameric scaffold for the build-up of glycodendrimers. For instance, compound **5** harboring four sialic acid residues, an important carbohydrate in viral infections, particularly involving flu viruses, was synthesized using a neuraminidase resistant α-thiosialic acid precursor [26,84] (Figure 4). An additional noteworthy application of glyco-calix[4]arene, exposing lactoside residues (not shown), was identified, and it demonstrated anti-trypanosomal activity comparable to the drug benznidazole in in vitro anti-parasite assays [79]. Notably, analogous constructs encompassing the tumor-associated carbohydrate antigen, Thomsen nouveau (T$_N$; α-D-GalNAc) [85], were found to adsorb on polystyrene microtiter plates through lipophilic *tert*-butyl tails. This glycodendrimer was found to be ten times more

potent than its monomeric counterparts toward the binding of a model leguminous lectin *Vicia villosa* agglutinin in a competitive enzyme-linked lectin assay against asialoglycoprotein [16,70].

The Siegel group synthesized an interesting aromatic glycodendrimer based on corannulene, which harbored five β-D-galactopyranoside moieties (**6**) [80] (Figure 4). The pentakis bioconjugate was prepared using the increasingly successful copper catalyzed azide–alkyne cycloaddition reaction (CuAAC; click chemistry). Our own group was deeply involved in the design of a uropathogenic inhibitor based on *Escherichia coli* FimH infection [86]. Toward this goal, a wide range of α-D-mannopyranoside dendrimers were successfully produced, among which is hexakis mannoside **7** built on a cyclotriphosphazene core [44,45,66]. It is to be noted that the cyclotriphosphazene scaffold exposes three of its six appendages on top of the molecules while the other three are in the opposite
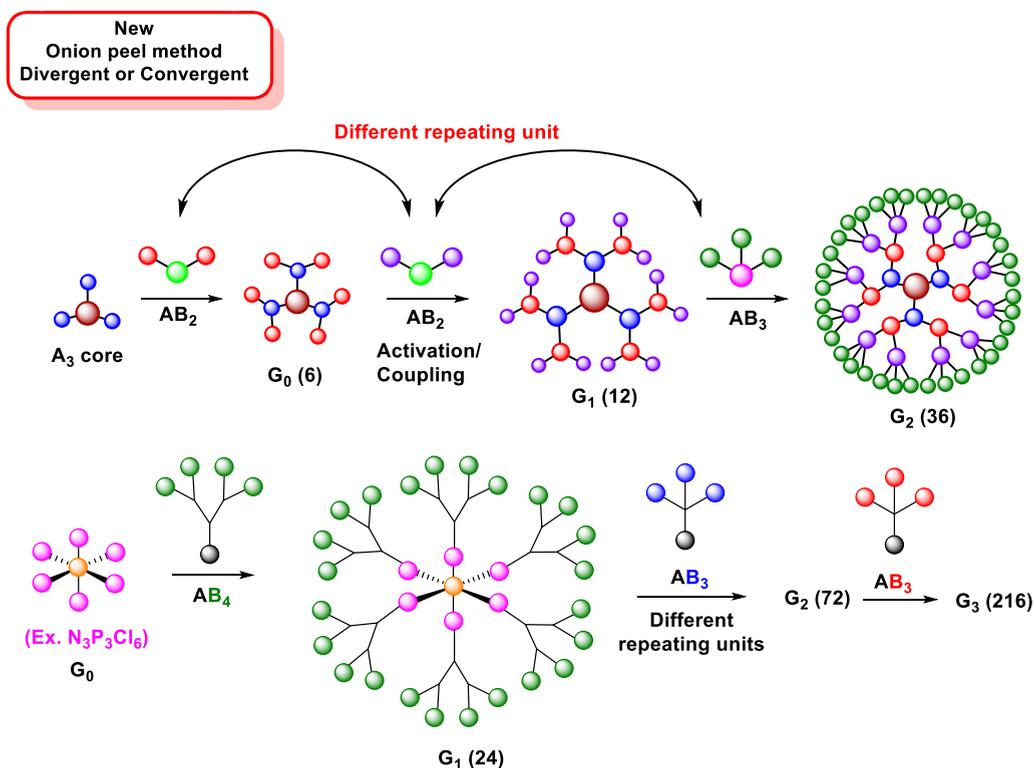
Figure 5. Onion-peel strategy for the construction of high-density surface groups possessing different building blocks at each layer.

downward directions [32,44]. This geometry is in sharp contrast when hexaphenylbenzenes are used as the central building block. A glycodendrimer such as **8** constitutes a case where the carbohydrate residues are planarly exposed [83].

Steadily increasing numbers of biologically valuable carbohydrates presented as surface groups have been aptly exemplified with the use of pillar[5]arenes (**9**) having ten exposed galactoside residues [87]. The same group pioneered the practice of fullerenes such as **10** and having twelve saccharides on the surface (Figure 3) [82]. Both cases further illustrate the usefulness of click chemistry in the expedient synthesis of glycoclusters [88–90]. Remarkably, both compounds **9** and **10** were applied as potent ligands in the fight against bacterial infections (Figures 3 and 4).

2.2. Onion-peel strategy

As opposed to conventional dendrimer syntheses that use low valency scaffolds and identical building blocks in repetitive sequences requiring a large number of steps to achieve high surface group densities, the onion-peel strategy starts with highly functionalized scaffolds such as that shown in Figure 2 and including dipentaerythritol 19 (Scheme 3). By using different building blocks at each layer, a much wider chemical space can be reached, thus enabling improved control over chemical, physical, and biochemical properties in addition to providing high surface group density in much fewer steps. Figure 5 presents the onion-peel strategy. Clearly, not only the methodology offers a wide range of chemical possibilities but also it offers a rapid build-up and reduced number of steps to create a higher number of surface groups.

As a typical example, taking trivalent scaffolds (trimesic acid, tris(2-aminoethyl)amine, cyanuric acid) and AB_2 type interlayer residues (3,4-, 3,5-dihydroxybenzoic acids), glycodendrimers with 6, 12, and 36 residues can be obtained in generations G0, G1, and G2, respectively [41,91]. This is in sharp contrast when PAMAM dendrimers are employed,

which provide only 4, 8, and 16 surface bearing moieties in identical generations.

Analogously, if a hexachlorocyclotriphosphazene ($N_3P_3Cl_6$) core is used in combination with alternating AB_4 and AB_3 interlayer building blocks, glycodendrimers having 24, 72, and 216 residues can be obtained as early as generations G1, G2, and G3, respectively [32,43,44,92].

As an illustration of this approach, the synthesis of a glycodendrimer having 36 LacNAc disaccharides as surface groups was published (Scheme 2). For this purpose, a hexachlorocyclotriphosphazene ($N_3P_3Cl_6$) core was initially chosen, which was treated with monopropargylated benzene-1,4-diol (hydroquinone) **11** (Cs_2CO_3, THF, 66 °C, 18 h) to provide the known hexapropargylated cyclotriphosphazene **12** as a crystalline adduct in 83% yield [32,44,93]. The next step involved a thiol–yne photocatalyzed addition of 3-mercaptopropanoic acid (**13**) to form dendrimer **14** having twelve carboxylic acids as surface groups in 76% yield. Trispropargylated pentaerythritol (**15**) serving as an AB_3 wedge was subsequently anchored via carbodiimide-mediated esterification to achieve the construction of hypercore **16** in good yield (72%). This sequence of reactions corresponds to $6 \times 2 \times 3$ (for 36 groups) illustrating an A_6 core, followed by an equivalent AB_2 interlayer, and finally an AB_3 building block. The final step necessitated the use of a LacNAc azide (**17**), which upon CuAAC coupling produced targeted glycodendrimer **18** in 76% yield. The strategy allowed the use of an unprotected glycoside derivative. Spectroscopic analyses (NMR, MS) were all consistent with the anticipated structure. A low-resolution mass spectral value was obtained by the MALDI-TOF technique ([M+Na]$^+$ adducts) for which compound **18** exhibited a molecular weight of m/z 19774.8 for a calculated value of m/z 19779.9.

Surface plasmon resonance (SPR) studies were conducted to assess the protein-binding abilities of **18** relative to glycodendrimer congeners possessing fewer LacNAc surface groups. For this purpose, the LacNAc-specific leguminous lectin (ECA) from *Erythrina cristagalli* was used. In these studies, the lectin was immobilized onto a CM5 gold sensor surface. Solutions with various concentrations of LacNAc-functionalized dendrimers were passed over the surface-bound lectin, and significant interactions were determined for each glycodendrimer

and compared to monovalent standard LacNAc (**17**). Determination of the kinetic parameters relative to the glycodendrimer–lectin interactions were fitted by using a 1:1 Langmuir model. In general, multivalent compounds exhibited high nanomolar affinities with the dimeric ECA. All glycodendrimers exhibited high relative potencies when compared to monomeric **17**. In one case, a maximum of 216-fold enhancement in global affinity was observed for the best candidate harboring only 24 LacNAc groups. This work demonstrated once more that highly dense glycodendrimers do not necessarily outperform those with lower carbohydrate residues. Our overall experience in this field indicated that not only the *number* of carbohydrate residues exposed as surface groups represents beneficial key factors but also their inter-residue *accessibility* (steric hindrance), the *linker*'s length, the structure of their *aglycones*, and obviously the *multivalency* of the required protein receptors.

In this work, the relative binding affinities of closely related glycodendrimers were evaluated by competitive SPR using the β-D-galactoside specific bacterial lectin from the gram-negative bacteria *Pseudomonas aeruginosa*. This protein constitutes a virulence factor and is involved in the pathogenesis of the bacteria in cystic fibrosis patients. The SPR experiments clearly demonstrated that all glycodendrimers exhibited much higher binding affinity compared to their corresponding monovalent derivatives due to the "multivalent or glycoside cluster effect". The incorporation of a tetraethylene glycol linker to the galactoside aglyconic portion allowed a twofold enhancement of the affinity for the lectin by providing better accessibility to the deep binding pocket of the lectin. Galactodendrimer **25** provided one of the best ligands known to date with an IC$_{50}$ value of 37 nM. This result unambiguously highlights the key role of linkers in interactions with lectin, with a counterbalanced entropic cost due to their flexibility. Additionally, tridimensional distribution of terminal and optimized galactosides crucially contributed to high potencies since a substantial improvement (32-fold) was observed for each ligand in **25** when compared to a monomeric reference.

## 2.3. *Chemical ligation and orthogonal chemistry*

This objective may be accomplished through the design and synthesis of heterofunctionalized
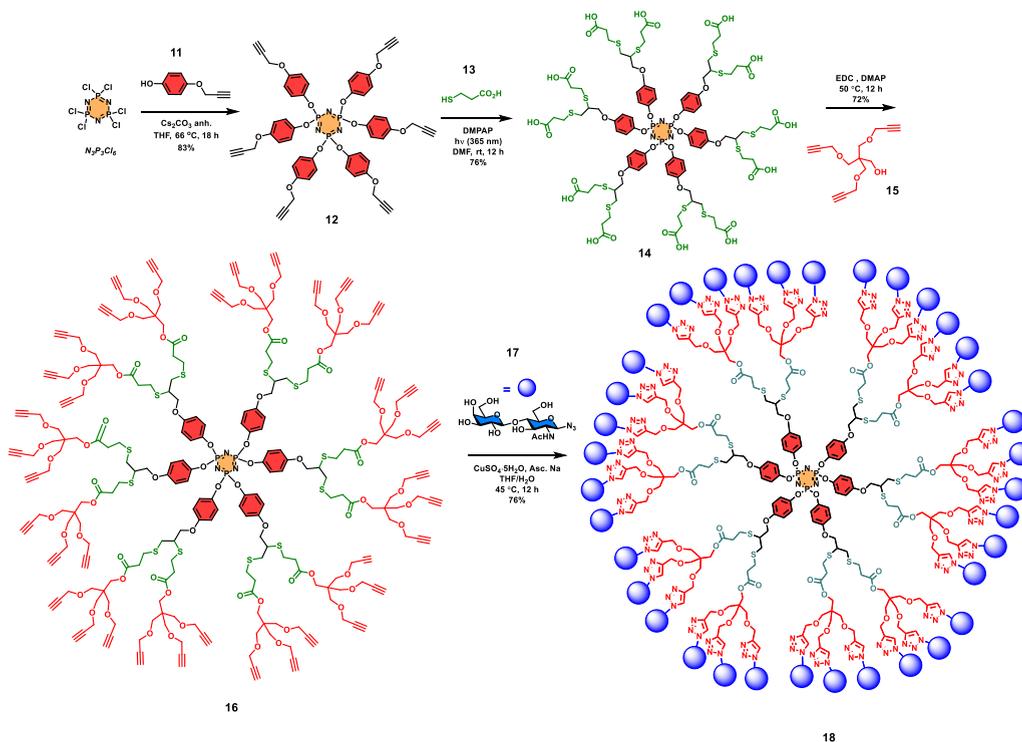
Scheme 2. Typical example of a fast growing onion-peel glycodendrimer using high-density scaffold (A_6) together with different interlayer building blocks (AB_2 and AB_3) that resulted in a dendrimer having 36 N-acetyllactosamine residues as surface groups [32].
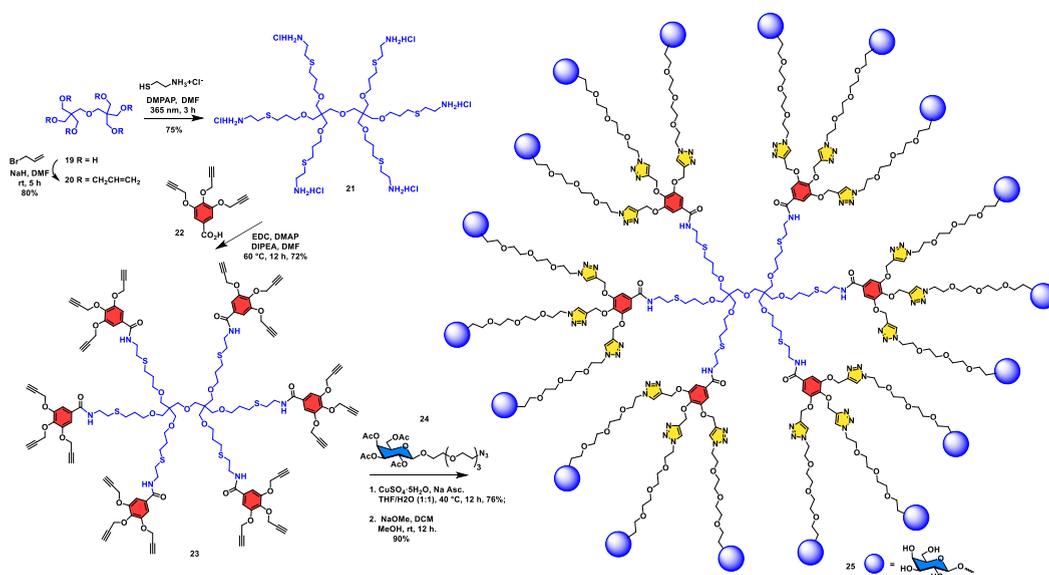
Scheme 3. Dipentaerythritol has been underused as a high-density scaffold with its six primary hydroxyl functionalities (A_6 core), which when combined with an AB_3 interlayer moiety formed a glycodendrimer having 36 surface residues in five consecutive steps. This example further illustrates the possibility of adopting both divergent and convergent approaches using the onion-peel concept [40].
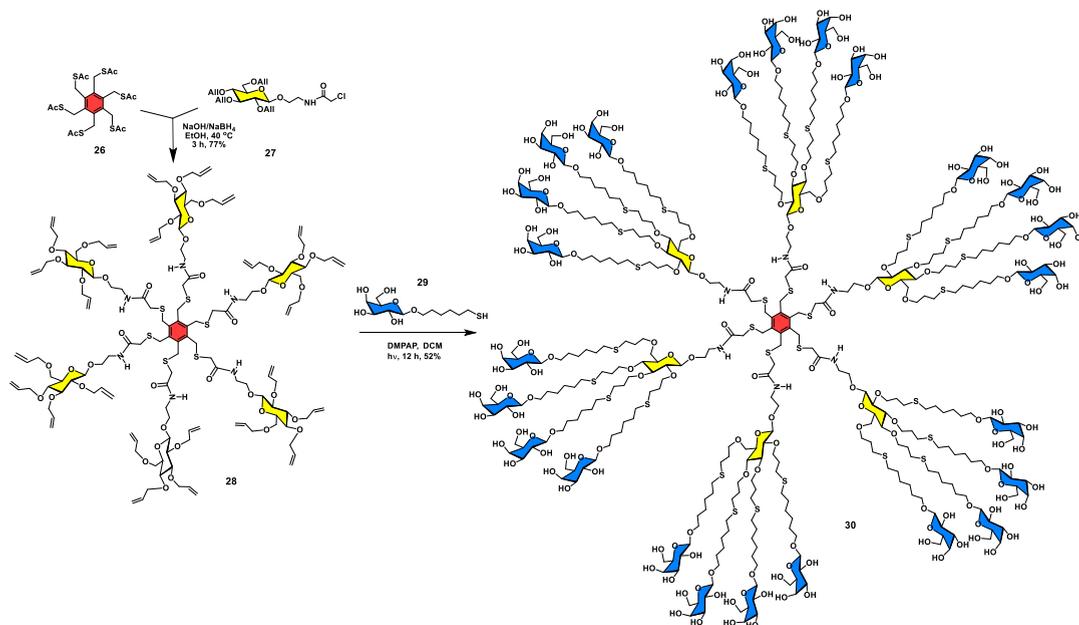
Scheme 4. Accelerated glycodendrimer synthesis through: highly functional core, glucoside as chiral interlayer building blocks, and orthogonal ligation.

macromolecules with precise molecular architecture and the integration of multiple bioactive moieties. In our view, recent progress in orthogonal chemoselective ligation methodologies positions these strategies among the most effective for constructing such biomacromolecular systems in a controlled and reproducible manner—a prerequisite for ensuring consistent biological outcomes [94–96].

Due to their exceptional reactivity and stability, suitably functionalized carbohydrates represent key molecular building blocks in this context. Alongside, glycans and other dendrimer scaffolds can be modified with compatible functional groups such as alkenes, alkynes, thiols, and azides. These approaches, for instance, may enable the development of synthetic multiantigenic vaccines incorporating diverse tumor-associated carbohydrate and peptide antigens [3]. Moreover, the high modularity and broad compatibility of orthogonal ligation techniques with carbohydrates, peptides, and nucleic acids could facilitate the creation of novel biomolecular assemblies with unprecedented biological functions and therapeutic potential.

The use of glycosides as both a high-density core and as interlayer residues has also been a valuable concept in the construction of dendrimers in general and of glycodendrimers in particular [97]. The example that follows combines several principles presented in this review. By including a carbohydrate moiety in the design, a key concept has been added through which chirality can be included, hence providing an additional architectural parameter (Scheme 4) [40].

In the present example, an orthogonal coupling strategy was developed by combining thiol–ene and S_N2 reactions, which was subsequently applied for the accelerated synthesis of multifunctional dendrimers using carbohydrate as both building blocks and surface groups. In this work, we first established the orthogonality of a thiol–ene against an S_N2 reaction. Toward this goal, a glucoside-based AB$_4$ monomer (**27**), possessing four allyl appendages and a N-(2-(glucosyloxy)ethyl)-2-chloroacetamide group as the aglycone, was used as an interlayer building block (Scheme 4). The ensuing nucleophilic substitution was executed with hexakis(thioacetoxymethyl)benzene (**26**) (A$_6$ core) under reducing conditions to avoid disulfide formation to provide intermediate **28** in 77% yield [40]. Notably, dendrimer **28** is a chiral entity encompassing 24 surface groups that was next successfully treated with thiogalactoside **29** under photocatalyzed thiol–ene
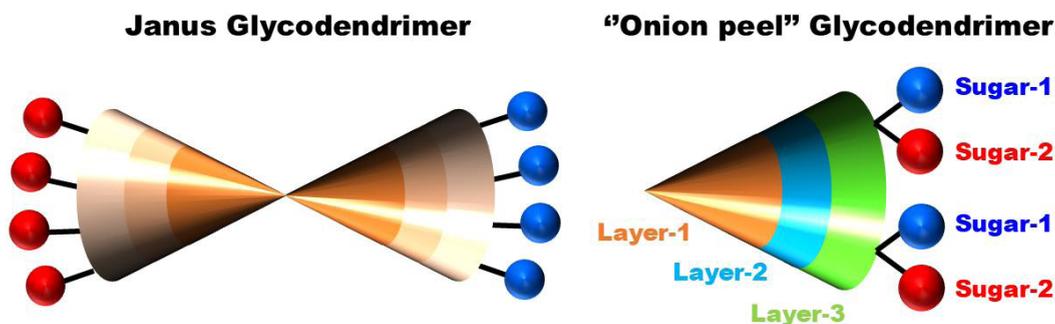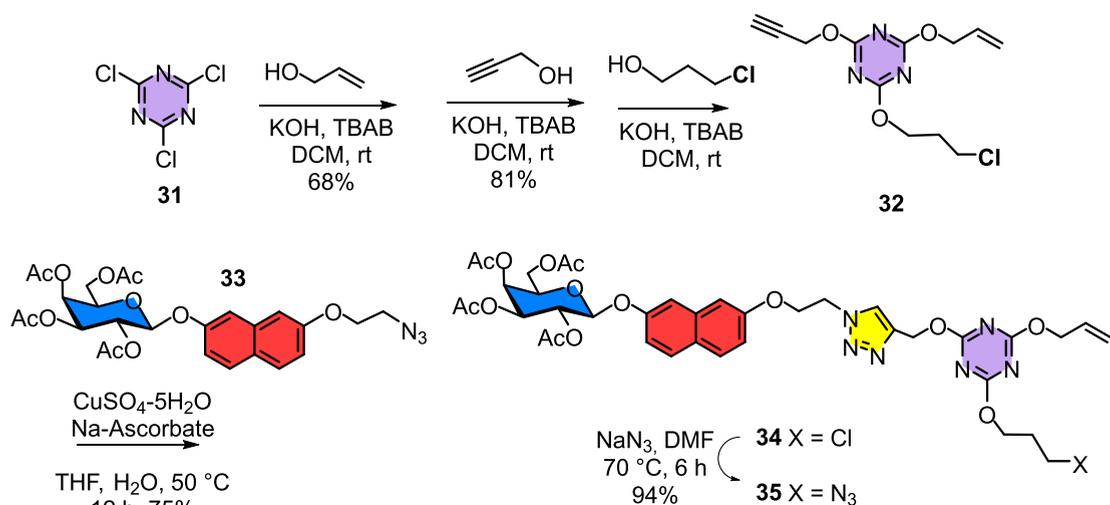
Figure 6. Janus glycodendrimer with two different saccharide moieties on each side of the "faces" (left) and "onion-peel" glycodendrimer having alternating and different glycoside residues (right).


Scheme 5. Building an orthogonally functionalized s-triazine triol scaffold to be used for the divergent synthesis of an heterolayered onion-peel glycodendrimer.

reaction to form glycodendrimer **30** (52%). Using SPR studies, the β-D-galactopyranoside-coated dendrimer exhibited nM binding affinity against the bacterial LecA lectin from *P. aeruginosa*.

In our endeavors to create glycodendrimers with high biological values, we initially focused our efforts on constructing multivalent architectures with just a single carbohydrate moiety at the surface. This was deemed necessary in order to target only one protein receptor at a time with the highest possible specificity and affinity. However, there are situations for which different carbohydrate residues are necessary to block a precise biological function. This is particularly the case with bacterial lectins expressing two or more carbohydrate-binding proteins such as those found in *P. aeruginosa* [98]. For this sce-

nario, we synthesized Janus-type glycodendrimers containing for instance both α-L-fucopyranoside on one end and β-D-galactopyranoside at the opposite face (Figure 6). With the next glycodendrimer generation, as mentioned, an onion-peel strategy was employed for a subsequent modulation of biophysical properties.

An additional variant was further included through which both the Janus dendrimer and the onion-peel concept were combined together with optimized saccharide aglycones to provide specificities toward each targeted lectin (Figure 6) [42,43]. Hence, the implementation of the orthogonal ligation strategy was also incorporated into our design. For this purpose, a photolytic thiol–ene reaction was combined with click chemistry (Schemes 5 and 6).
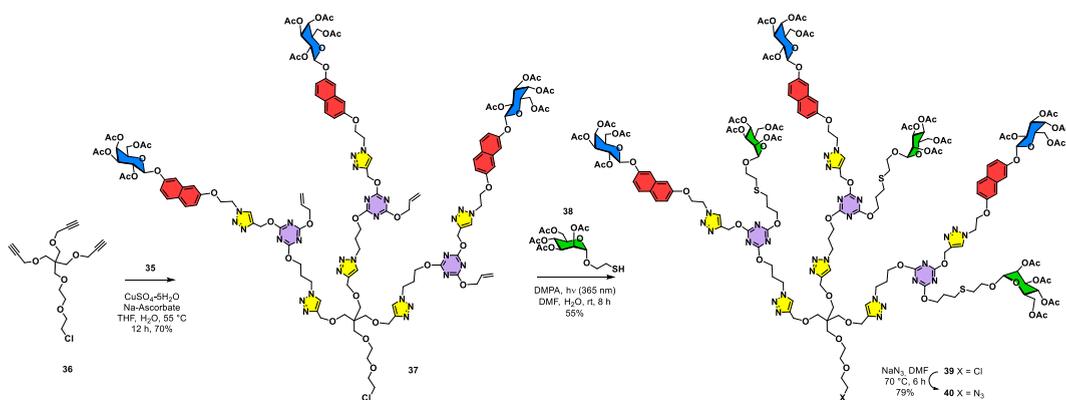
Scheme 6. Synthesis of a heterobifunctional glycodendron harboring both galactose and mannose residues together with orthogonal azide at the focal point.
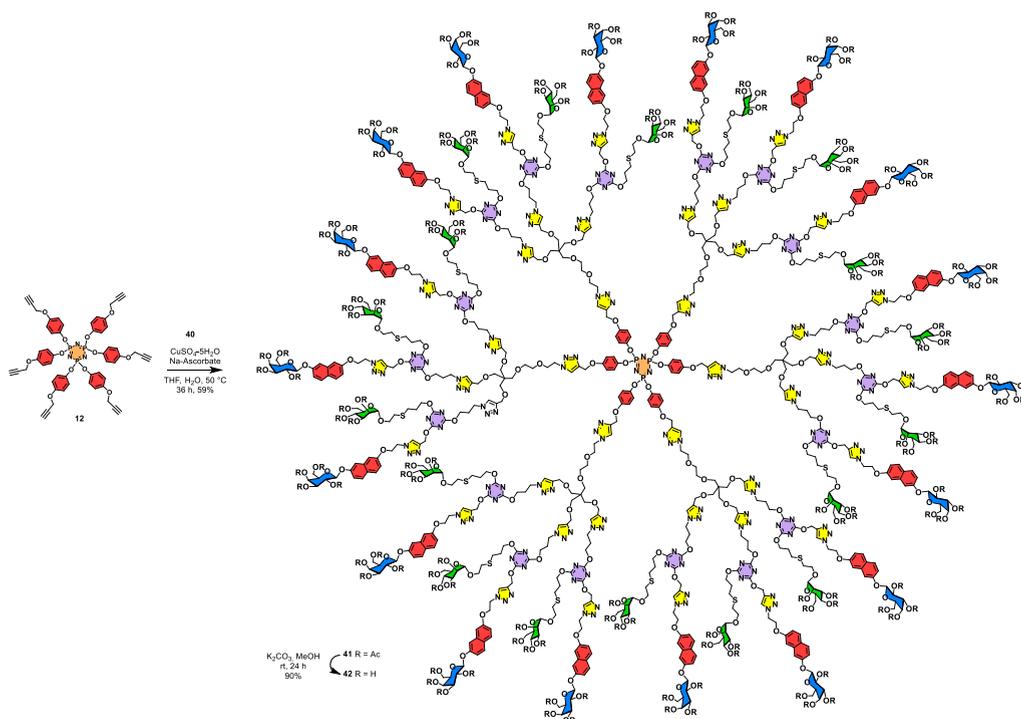
Scheme 7. Convergent building of an alternating heterobifunctional glycodendrimer harboring a total of 36 galactoside and mannoside residues using a cyclotriphosphazene core.

Finally, the target glycodendrimer was cleanly built on the hexakispropargylated cyclotriphosphazene core **12** using powerful click chemistry to yield **41** (59%) (Scheme 7). It is to be noted that these complex peracetylated compounds could be readily purified by standard silica gel column chromatography. Conventional transesterification under Zemplén conditions (K$_2$CO$_3$, MeOH) provided glycodendrimer **42** (90%) containing 36 alternat-

ing carbohydrate surface groups, of which 18 were β-D-galactopyranosides and a further 18 were α-D-mannopyranosides. This case constitutes an ideal example corresponding to a 6 × 3 × 2 methodology (A$_6$ core, AB$_3$ interlayer, and AB$_2$ interlayer).

The biochemical functionality of the aforementioned hybrid glycodendrimer (**42**) was evaluated using kinetic studies through DLS. Consequently, **42** (~3–5 nm) readily formed large

cross-linked aggregates in the presence of the ho-
motetrameric mannose-specific leguminous lectin
ConA (~8 nm) from *C. ensiformis* (jack bean), result-
ing in glycodendrimer–protein cross-linked aggre-
gates of ~3500 nm within 2 min [9]. At the same time,
the galactose-specific homotetrameric lectin LecA
from *P. aeruginosa* also formed large cross-linked
lattices (~5000 nm) within 4 min, thus demonstrat-
ing the accessibility of either glycoside with larger
protein receptors.

Given the high potential of glycodendrimers in
biological settings and the complexity of carbohy-
drate entities with their multiple and varied func-
tional groups together with the necessity for reli-
able and selective ligation chemistry, we illustrate the
most common of these glycoconjugation partners in
Table 1. As can be seen, the carbohydrate moieties
usually employed in glycoconjugate synthesis such
as amines, acids, azides, and alkynes represent the
most frequently encountered examples that are tra-
ditionally utilized.

3. Self-assembling glycodendrimers

Self-assembling is an important phenomenon in
biological systems; for example, the formation of
cell membranes and metal-based chelation (met-
alloproteins) represent truly inspiring examples of
supramolecular chemistry [134]. Therefore, its im-
plementation in the field of glycodendrimers can
greatly reduce the number of steps necessary for
the construction of well-defined multivalent archi-
tectures having nanometer size. In this section, we
illustrate the evolution of our thought process from
the initial description of a randomly disposed gly-
copolymer strategy to describing case studies with
transition-metal-templated self-assembly and glyco-
dendrimersomes.

3.1. *Transition-metal-templated self-assembly: glyco-metallo-dendrimers*

The concept of self-assembly—where molecules
spontaneously organize into structured
arrangements—has revolutionized the way we de-
sign functional materials. Among the most versatile
tools in this domain are transition metals, which
serve as dynamic templates guiding the formation of
complex architectures. Transition-metal-templated

self-assembly leverages the coordination chemistry
of metals like iron, copper, palladium, and ruthe-
nium to direct the spatial arrangement of ligands
and molecular building blocks [135–138]. In essence,
transition-metal-templated self-assembly is not
just a synthetic strategy—it is a design philosophy
that blends chemistry, physics, and engineering to
build order from chaos. The metal-templated self-
assembly strategy has also been effectively extended
into the field of glycodendrimers by combining dy-
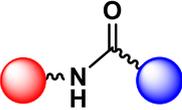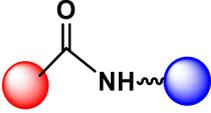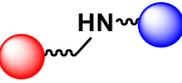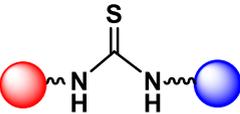namic combinatorial chemistry [139,140].

In the following example, a 2,2′-bipyridine-4,4′-
dicarboxylic acid core was selected as a Cu-based
assembly. A bivalent glycodendron composed of
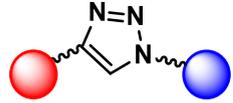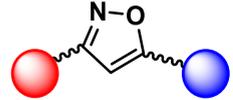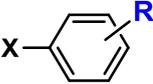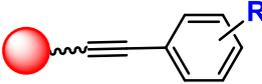*N*-acetyl-D-galactosamine (GalNAc) (**43**) was assem-
bled using classic amide bonding to form a tetrava-
lent subunit for the coordination with copper sulfate
(Scheme 8). The self-assembly was simply achieved
with only two equivalents of **43**—one equivalent
of $CuSO_4 \cdot 5H_2O$ in water at 23 °C for two days—to
deliver the metal-based symmetrical self-assembled
glycodendrimer (**44**) as a blue-purple colored powder
after lyophilization. The readily formed octavalent
GalNAc cluster **44** showed high inhibitory properties
in the binding of asialoglycophorin to the legumi-
nous lectin *V. villosa* with a 259-fold higher potency
in comparison to its monomeric counterpart [141].

3.2. *Lipid-templated self-assembly: glycoden-drimersomes*

Glycolipids [142], amphiphilic molecules composed
of a hydrophilic carbohydrate head group and a
hydrophobic lipid tail, exhibit remarkable self-
assembly behavior in aqueous environments due
to their dual chemical nature. This intrinsic am-
phiphilicity drives the spontaneous formation of or-
ganized supramolecular structures such as micelles,
vesicles, and fibrous networks depending on molec-
ular geometry, concentration, and environmental
conditions. The self-assembly process is governed by
noncovalent interactions, including hydrogen bond-
ing, van der Waals forces, and hydrophobic effects,
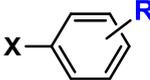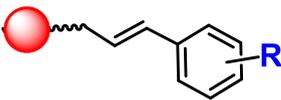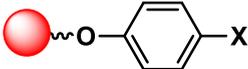which collectively stabilize the resulting nanostruc-
tures.

Microbial glycolipids are particularly notable for
their ability to form reversible, stimulus-responsive
assemblies. These systems can transition between
morphologies—for example, from spherical micelles

Table 1. Most common ligation chemistry used in the synthesis of glycodendrimers

Entry	Carbohydrate functionality	Scaffold functionality	Ligation	Refs.
1	ᴧNH$_2$	HO$_2$C ᴧ	ᴧN(H)–C(O)–	[99,100]
2	ᴧCO$_2$H	H$_2$N ᴧ	–C(O)–NH ᴧ	[99,101]
3	ᴧCHO	H$_2$N ᴧ	HN ᴧ	[102–104]
4	ᴧX X = OAc, SR, Cl, Br I, CO(NH)CCl$_3$ (Glycosyl donors)	HO ᴧ	ᴧO ᴧ	[105]
5	ᴧNCS	H$_2$N ᴧ	ᴧN(H)–C(S)–N(H) ᴧ	[106–108]
6	ᴧOCOOR	H$_2$N ᴧ	ᴧOCONH ᴧ	[109–111]
7	ᴧSH	(allyl) ᴧ	–S–	[72,97,112]
8	ᴧONH$_2$	OHC–C(O) ᴧ	ᴧO–N=	[3,113,114]
9	ᴧN$_3$	(alkyne) ᴧ	(triazole)	[32,115,116]
10	(alkyne)	N$_3$ ᴧ	(triazole)	[32,115,116]
11	ᴧ≡N$^{\oplus}$–O$^{\ominus}$	(alkyne) ᴧ	(isoxazole)	[117]
12	(alkyne)	X–(aryl)–R	ᴧ≡–(aryl)–R	[118]
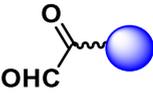
Table 1. (continued)

Entry	Carbohydrate functionality	Scaffold functionality	Ligation	Refs.
13				[119]
14				[120,121]
15				[122,123]
16				[124–126]
17				[127,128]
18				[129]
19				[130]
20				[91,131]
21				[32,132]
22				[72,133]
23				[115]
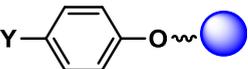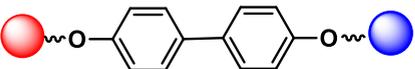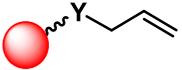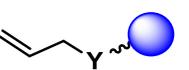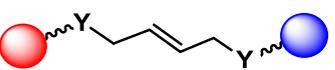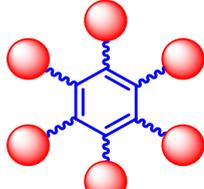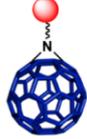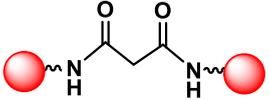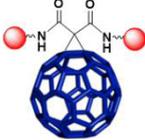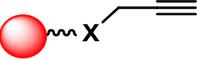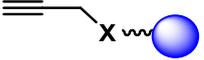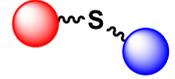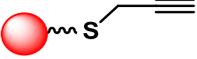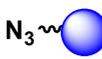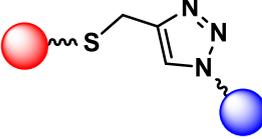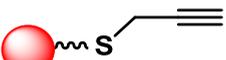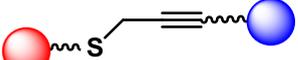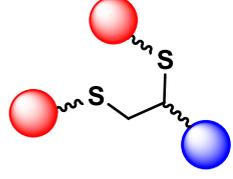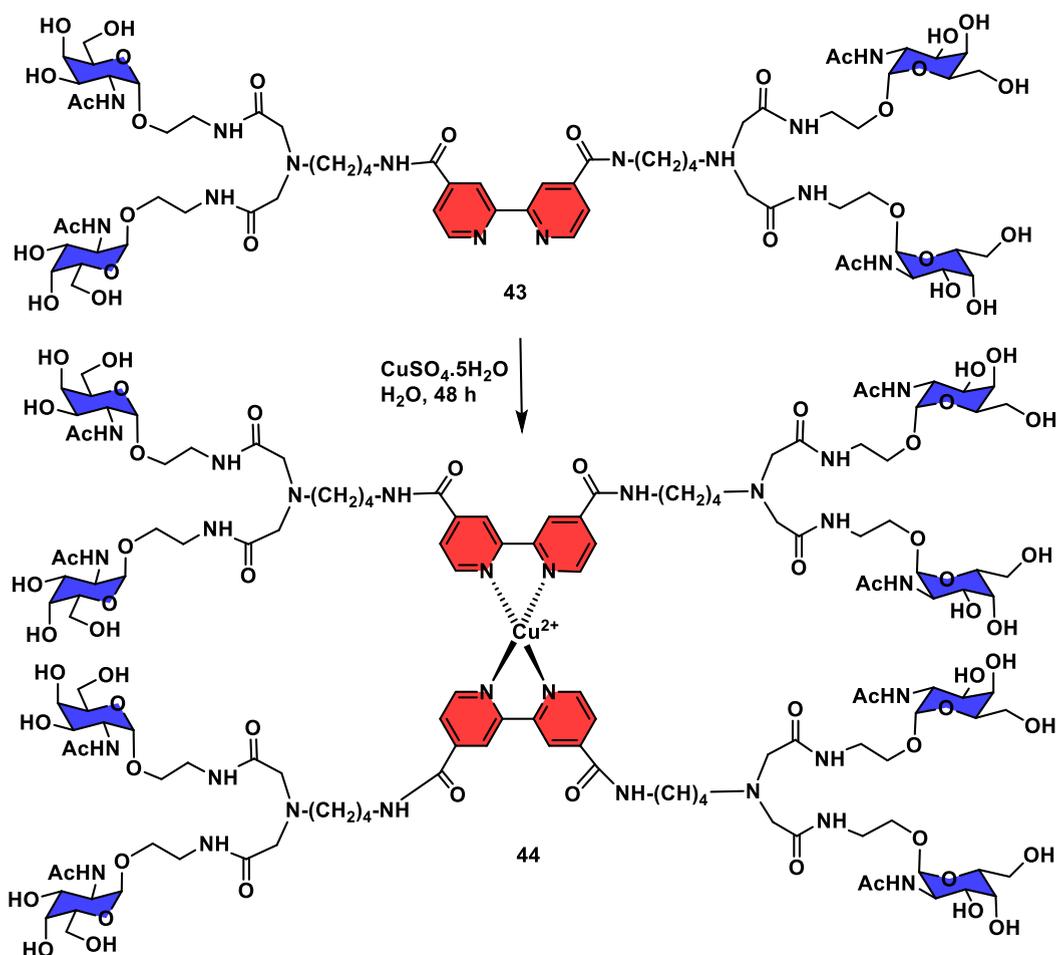
Scheme 8. Transition metal used in the self-assembly process toward the construction of glycodendrimers [141].

to elongated fibers—based on external cues, making them highly versatile for biomedical and industrial applications. Additionally, amphiphilic glycolipid glycoconjugates have been employed to encapsulate hydrophobic drugs and dyes, enhancing solubility, stability, and targeted delivery.

The biocompatibility, biodegradability, and sustainable synthesis of glycolipids via enzymatic or chemoenzymatic routes from renewable feedstocks further underscore their potential in green chemistry and nanomedicine. Their ability to self-organize into functional architectures without the need for harsh processing conditions positions glycolipids as promising candidates for next-generation soft materials and bioactive platforms.

In collaboration with the Percec group, we synthesized seven libraries containing 51 self-assembling amphiphilic Janus glycodendrimers composed of monosaccharides D-mannose and D-galactose and the disaccharide D-lactose denoted "glycodendrimersomes" using different hydrophobic aromatic cores and saccharide-based hydrophilic head groups having different linkers (Figure 7) [143–145]. Scheme 9 illustrates a typical example for the construction of these biologically relevant amphiphilic nanosized (~100 nm) materials using click chemistry.

These self-assembled glycodendrimersomes were stable over time in water and in buffer, exhibited narrow molecular-weight distribution, and displayed dimensions that are programmable by the concentration of the solution from which they are injected. This study expanded the molecular principles leading to single-type soft glycodendrimersomes assembled from amphiphilic Janus glycodendrimers.
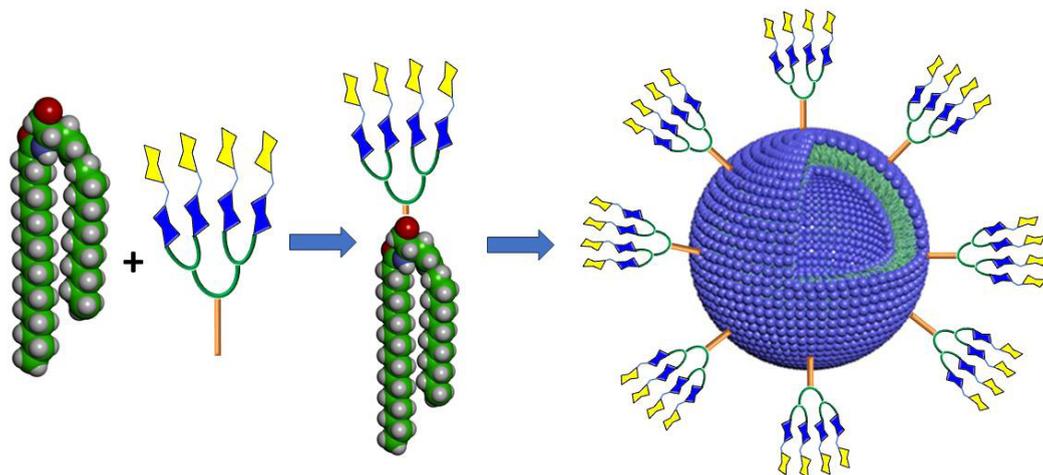
Figure 7. Principle used in the construction of self-assembling glycodendrimersomes from lipophilic tails and glycodendrons [143–145].
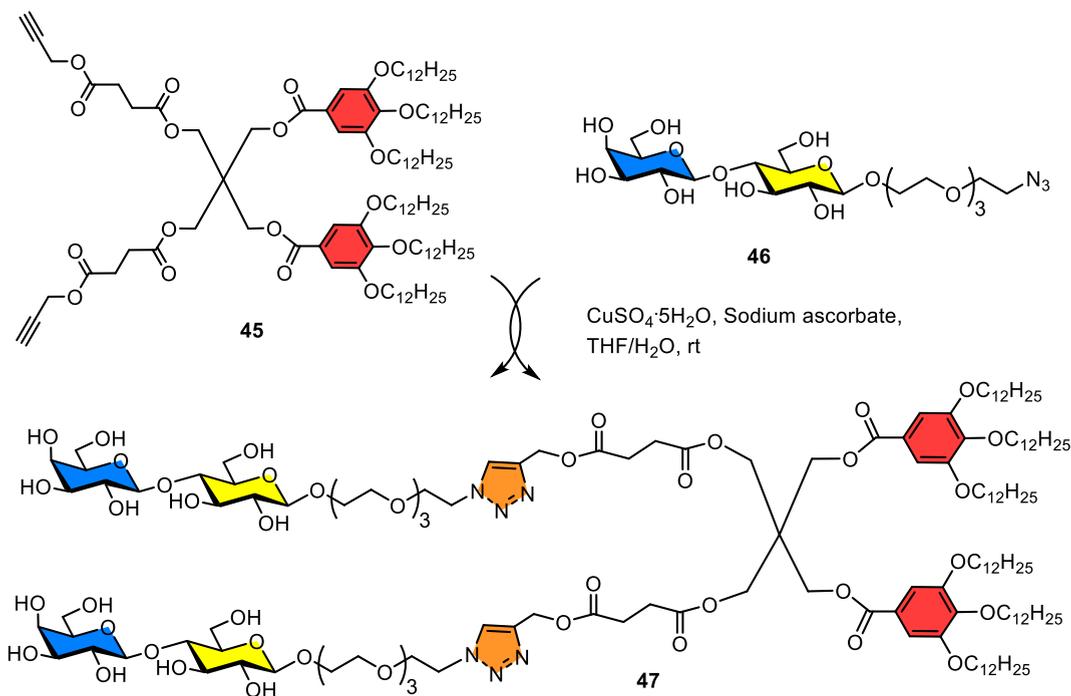
Scheme 9. Typical example used in the syntheses of the first glycodendrimersomes [143–145].

The multivalency of glycodendrimersomes with different sizes and their ligand bioactivity were demonstrated by selective agglutination assays with a diversity of glycoside-binding protein receptors such as the plant lectins ConA (D-mannose/D-glucose), the highly toxic mistletoe *Viscum album* agglutinin (D-Gal/D-GalNAc), and the bacterial lectin PA-IL from *P. aeruginosa* (D-Gal). The galactoside-containing glycodendrimersomes are of significant biomedical relevance since they were shown capable of binding to human adhesion/growth-regulatory galectin-3 and galectin-4. These results demonstrated the

candidacy of glycodendrimersomes as new mimics of biological membranes with programmable glycan ligand presentations, as supramolecular lectin blockers, vaccines, and targeted delivery devices.

4. Conclusion

In this review, recent advances in the design of chemical architectures incorporating carbohydrate moieties are discussed. The multivalency of the resulting ligands significantly enhances lectin-binding avidity, thereby modulating the ability of carbohydrates to engage in improved supramolecular recognition processes. These interactions facilitate the transfer of biological information and enable targeted engagement with complementary lectin receptors.

However, current models often fail to account for the heterogeneity and dynamic nature of densely glycosylated cell membranes. They also overlook the potential influence of high-density, nanoscale arrangements on interactions with the broader spectrum of carbohydrate-binding proteins. For years, it has been tacitly assumed the following: (i) effective lectin recognition generally necessitates high densities of the primary ligand and (ii) the mechanisms underlying carbohydrate motif recognition by lectins versus glycosidases are fundamentally distinct. We proposed a more generalized multivalent construction strategy to better address issues related to the topographical chemistry of carbohydrate and glycomimetic structures in biological systems. Within this context, (hetero)multivalency functions as a multimodal switch, channeling encoded information through diverse molecular pathways. We firmly believe that further development would be essential to translate these insights into practical applications, particularly in the promising field of glycodendrimers.

Acknowledgments

Declaration of interests

The authors do not work for, advise, own shares in, or receive funds from any organization that could benefit from this article, and have declared no affiliations other than their research organizations.

References

[1] J. I. Quintana, U. Atxabal, L. Unione, A. Ardá and J. Jiménez-Barbero, "Exploring multivalent carbohydrate–protein interactions by NMR", *Chem. Soc. Rev.* **52** (2023), pp. 1591–1613.

[2] N. Sharon, "Protein–carbohydrate interactions: At the heart of biochemistry", *Biochemistry* **28** (2006), pp. 13–17.

[3] R. Roy and L. Mousavifar, "Carrier diversity and chemical ligations in the toolbox for designing tumor-associated carbohydrate antigens (TACAs) as synthetic vaccine candidates", *Chem. Soc. Rev.* **52** (2023), pp. 3353–3396.

[4] A. Varki, "Biological roles of glycans", *Glycobiology* **27** (2017), pp. 3–49.

[5] R. A. Flynn, K. Pedram, S. A. Malaker, et al., "Small RNAs are modified with N-glycans and displayed on the surface of living cells", *Cell* **184** (2021), pp. 3109–3124.

[6] R. A. Dwek, "Glycobiology: Toward understanding the function of sugars", *Chem. Rev.* **96** (1996), pp. 683–720.

[7] R. Roy and F. D. Tropper, "Carbohydrate protein interactions. Syntheses of agglutination inhibitors of wheat germ agglutinin by phase transfer catalysis", *Can. J. Chem.* **69** (1991), pp. 817–821.

[8] Y. C. Lee and R. T. Lee, "Carbohydrate-protein interactions: Basis of glycobiology", *Acc. Chem. Res.* **28** (1995), pp. 321–327.

[9] L. Mousavifar, M. R. Gupta, M. Rivat, et al., "Self-assembling PEGylated mannolipids for liposomal drug encapsulation of natural products", *Mater. Adv.* **6** (2025), pp. 365–377.

[10] H.-J. Gabius, M. Cudic, T. Diercks, et al., "What is the sugar code?", *ChemBioChem* **23** (2022), article no. e202100327.

[11] M. Touaibia and R. Roy, "Glycodendrimers as anti-adhesion drugs against type 1 fimbriated *E. coli* uropathogenic infections", *Mini Rev. Med. Chem.* **7** (2007), pp. 1270–1283.

[12] R. Roy, D. Zanini, S. J. Meunier and A. Romanowska, "Solid-phase synthesis of dendritic sialoside inhibitors of influenza A virus haemagglutinin", *J. Chem. Soc., Chem. Commun.* **24** (1993), pp. 1869–1872.

[13] R. Roy, D. Zanini, S. J. Meunier and A. Romanowska, "Synthesis and antigenic properties of sialic acid based dendrimers. In synthetic oligosaccharides", *ACS Sympos. Ser.* **560** (1994), pp. 104–119.

[14] R. Roy, W. K. C. Park, Q. Wu and S.-N. Wang, "Synthesis of hyper-branched dendritic lactosides", *Tetrahedron Lett.* **36** (1995), pp. 4377–4380.

[15] D. Zanini, W. K. C. Park and R. Roy, "Synthesis of novel dendritic glycosides", *Tetrahedron Lett.* **36** (1995), pp. 7383–7386.

[16] S. J. Meunier and R. Roy, "Polysialosides scaffolded on p-Tert-butylcalix[4]arene", *Tetrahedron Lett.* **37** (1996), pp. 5469–5472.

[17] T. C. Shiao and R. Roy, "Glycodendrimers as functional antigens and antitumor vaccines", *New J. Chem.* **36** (2012), pp. 324–339.

[18] R. Roy, T. C. Shiao and K. Rittenhouse-Olson, "Glyco-dendrimers: versatile tools for nanotechnology", *Braz. J. Pharm. Sci.* **49** (2013), pp. 85–108.

[19] R. Roy, "Blue-prints, synthesis and applications of gly-copolymers", *Trends Glycosci. Glycotechnol.* **8** (1996), pp. 79–99.

[20] R. Roy, F. D. Tropper and A. Romanoska, "New strategy in glycopolymer syntheses. Preparation of antigenic water-soluble poly (acrylamide-co-p-acrylamidophenyl. beta.-lactoside)", *Bioconjugate Chem.* **3** (1992), pp. 256–261.

[21] R. Roy and C. A. Laferrière, "Synthesis of protein conju-gates and analogues of N-acetylneuraminic acid", *Can. J. Chem.* **68** (1990), pp. 2045–2454.

[22] N. Saha and K. K. Banerjee, "Carbohydrate-dependent binding of the cell-free hemagglutinin of Vibrio cholerae to glycoprotein and glycolipid", *J. Bacteriol.* **177** (1995), pp. 758–764.

[23] J. Zhu, J. Lin, R. Wang, Z. Ma, W. Zuo and M. Zhu, "Enhanced photosensitizer-embedded glycopoly-mers through self-catalytic PET-RAFT polymerization for targeted PDT", *Biomacromolecules* **26** (2025), pp. 2654–2664.

[24] S. Roy, S. Paul, S. Mukherjee, P. De and A. Mukher-jee, "Unraveling mechanism and enhancing selectiv-ity of a RuII-bis-bipyridyl-morphocumin complex with RAFT-generated glycopolymer exploiting Warburg effect in cancer", *Chemistry* **31** (2025), article no. e202403695.

[25] R. J. Mancini, J. Lee and H. D. Maynard, "Trehalose gly-copolymers for stabilization of protein conjugates to en-vironmental stressors", *J. Am. Chem. Soc.* **134** (2012), pp. 8474–8479.

[26] D. Zanini and R. Roy, "Synthesis of new α-thiosialodendrimers and their binding properties to the sialic acid specific lectin from Limax flavus", *J. Am. Chem. Soc.* **119** (1997), pp. 2088–2095.

[27] R. Roy, W. Park, D. Zanini, C. Foxall and O. Srivastava, "Dendritic 3′-Sulfo-Lewisensuremath^ x-(Glc) As potent L-and E-selectin antagonists", *Carbohydr. Lett.* **2** (1997), pp. 259–266.

[28] S. Mignani, J. Rodrigues, R. Roy, X. Shi, V. Ceña, S. El Kaz-zouli and J.-P. Majoral, "Exploration of biomedical den-drimer space based on in-vitro physicochemical param-eters: key factor analysis (Part 1)", *Drug Discov. Today* **24** (2019), pp. 1176–1183.

[29] S. Mignani, J. Rodrigues, R. Roy, X. Shi, V. Ceña, S. El Kaz-zouli and J.-P. Majoral, "Exploration of biomedical den-drimer space based on in-vivo physicochemical parame-ters: Key factor analysis (Part 2)", *Drug Discov. Today* **24** (2019), pp. 1184–1192.

[30] S. Mignani, X. Shi, J. Rodrigues, R. Roy, Á. Muñoz-Fernández, V. Ceña and J.-P. Majoral, "Dendrimers to-ward translational nanotherapeutics: Concise key step analysis", *Bioconjugate Chem.* **31** (2020), pp. 2060–2071.

[31] H. Sashiwa, Y. Shigemasa and R. Roy, "Highly convergent synthesis of dendrimerized chitosan–sialic acid hybrid", *Macromolecules* **34** (2001), pp. 3211–3214.

[32] R. Sharma, K. Naresh, Y. M. Chabre, R. Rej, N. K. Saadeh and R. Roy, ""Onion peel" dendrimers: A straightforward synthetic approach towards highly diversified architec-tures", *Polym. Chem.* **5** (2014), pp. 4321–4331.

[33] R. Sharma, I. Zhang, L. Abbassi, R. Rej, D. Maysinger and R. Roy, "A fast track strategy toward highly functionalized dendrimers with different structural layers: An "onion peel approach"", *Polym. Chem.* **6** (2015), pp. 1436–1444.

[34] V. Aseyev, E. Kontturi, H. Rosilo, M. Österberg, N. Houbenov and O. Ikkala, "Supracolloidal multivalent in-teractions and wrapping of dendronized glycopolymers on native cellulose nanocrystals", *J. Am. Chem. Soc.* **136** (2014), pp. 866–869.

[35] X. Wang, M. Wang, C. Wang, W. Deng and M. Liu, "Carbohydrate–lectin recognition of well-defined hetero-geneous dendronized glycopolymers: systematic studies on the heterogeneity in glycopolymer–lectin binding", *Polym. Chem.* **12** (2021), pp. 4722–4735.

[36] D. A. Tomalia, H. Baker, J. Dewald, et al., "A new class of polymers: Starburst-dendritic macromolecules", *Polym. J.* **17** (1985), pp. 117–132.

[37] Y. M. Chabre and R. Roy, "Recent trends in glyco-dendrimer syntheses and applications", *Curr. Top. Med. Chem.* **8** (2008), pp. 1237–1285.

[38] H. Sashiwa, Y. Makimura, Y. Shigemasa and R. Roy, "Chemical modification of chitosan: preparation of chitosan–sialic acid branched polysaccharide hybrids", *Chem. Commun.* **11** (2000), pp. 909–910.

[39] J. D. Reuter, A. Myc, M. M. Hayes, et al., "Inhibition of vi-ral adhesion and infection by sialic-acid-conjugated den-dritic polymers", *Bioconjugate Chem.* **10** (1999), pp. 271–278.

[40] R. Sharma, N. Kottari, Y. M. Chabre, L. Abbassi, T. C. Shiao and R. Roy, "A highly versatile convergent/divergent "onion peel" synthetic strategy toward potent multi-valent glycodendrimers", *Chem. Commun.* **50** (2014), pp. 13300–13303.

[41] C. Sehad, T. C. Shiao, L. M. Sallam, A. Azzouz and R. Roy, "Effect of dendrimer generation and aglyconic linkers on the binding properties of mannosylated dendrimers prepared by a combined convergent and onion peel ap-proach", *Molecules* **23** (2018), article no. 8.

[42] R. S. Bagul, M. M. Hosseini, T. C. Shiao and R. Roy, ""Onion peel" glycodendrimer syntheses using mixed tri-azine and cyclotriphosphazene scaffolds", *Can. J. Chem.* **95** (2017), pp. 975–983.

[43] R. S. Bagul, M. Hosseini, T. C. Shiao, N. K. Saadeh and R. Roy, "Heterolayered hybrid dendrimers with optimized sugar head groups for enhancing carbohydrate–protein interactions", *Polym. Chem.* **8** (2017), pp. 5354–5366.

[44] L. Abbassi, Y. M. Chabre, N. Kottari, A. A. Arnold, S. André, J. Josserand, H.-J. Gabius and R. Roy, "Multifaceted glycodendrimers with programmable bioactivity through convergent, divergent, and accelerated approaches using polyfunctional cyclotriphosphazenes", *Polym. Chem.* **6** (2015), pp. 7666–7683.

[45] L. M. Sallam, T. C. Shiao, C. Sehad, A. Azzouz and R. Roy, "Accelerated synthesis of surface functionalized mannosylated dendrimers built on cyclotriphosphazene core", *MRS Adv.* **4** (2019), pp. 3187–3198.

[46] N. Katir, N. El Brahmi, A. El Kadib, S. Mignani, A.-M. Caminade, M. Bousmina and J. P. Majoral, "Synthesis of onion-peel nanodendritic structures with sequential functional phosphorus diversity", *Chem. Eur. J.* **21** (2015), pp. 6400–6408.

[47] R. Roy, D. Pagé, S. Figueroa Perez and V. Verez Bencomo, "Effect of shape, size, and valency of multivalent mannosides on their binding properties to phytohemagglutinins", *Glycoconj. J.* **15** (1998), pp. 251–263.

[48] R. Roy, "Glycodendrimers: a new class of biopolymers", *Polymer News* **21** (1996), pp. 226–232.

[49] S. André, P. J. Cejas Ortega, M. A. Perez, R. Roy and H.-J. Gabius, "Lactose-containing starburst dendrimers: influence of dendrimer generation and binding-site orientation of receptors (plant/animal lectins and immunoglobulins) on binding properties", *Glycobiology* **9** (1999), pp. 1253–1261.

[50] L. Mousavifar and R. Roy, "synthetic strategies, and therapeutic applications of heterofunctional glycodendrimers", *Molecules* **26** (2021), article no. 2428.

[51] R. Dominique, B. Liu, S. K. Das and R. Roy, "Synthesis of molecular Asterisks' via sequential cross-metathesis, Sonogashira and Cyclotrimerization reactions", *Synthesis* **6** (2000), pp. 862–868.

[52] M. Llinares and R. Roy, "Multivalent neoglycoconjugates: solid-phase synthesis of N-linked α-sialodendrimers", *Chem. Commun.* **21** (1997), pp. 2119–2120.

[53] H. Sashiwa, Y. Shigemasa and R. Roy, "Chemical modifications of chitosan. Part 2. Novel N-alkylation of chitosan via Michael type reaction", *Chem. Lett.* **8** (2000), pp. 862–863.

[54] H. Sashiwa, J. M. Thompson, S. K. Das, Y. Shigemasa, S. Tripathy and R. Roy, "Chemical modification of chitosan: Preparation and lectin binding properties of α-galactosyl-chitosan conjugates. Potential inhibitors in acute rejection following xenotransplantation", *Biomacromolecules* **1** (2000), pp. 303–305.

[55] H. Sashiwa, Y. Shigemasa and R. Roy, "Chemical modification of chitosan. 10. Synthesis of dendronized chitosan–sialic acid hybrid using convergent grafting of preassembled dendrons built on gallic acid and Tri (ethylene glycol) backbone", *Macromolecules* **34** (2001), pp. 3905–3909.

[56] Y. M. Chabre, C. Contino-Pépin, V. Placide, T. C. Shiao and R. Roy, "Expeditive synthesis of glycodendrimer scaffolds based on versatile TRIS and mannoside derivatives", *J. Org. Chem.* **73** (2008), pp. 5602–5605.

[57] A. Papadopoulos, T. C. Shiao and R. Roy, "Diazo transfer and click chemistry in the solid phase syntheses of lysine-based glycodendrimers as antagonists against Escherichia coli FimH", *Mol. Pharm.* **9** (2012), pp. 394–403.

[58] D. Zanini and R. Roy, "Chemoenzymatic synthesis and lectin binding properties of dendritic N-acetyllactosamine", *Bioconjugate Chem.* **8** (1997), pp. 187–192.

[59] M. M. Palcic, H. Li, D. Zanini, R. S. Bhella and R. Roy, "Chemoenzymatic synthesis of dendritic sialyl Lewis x", *Carbohydr. Res.* **305** (1997), pp. 433–442.

[60] D. Pagé and R. Roy, "Glycodendrimers as novel biochromatography adsorbents", *Int. J. Biochromatogr.* **3** (1997), pp. 231–243.

[61] R. Roy, "Recent developments in the rational design of multivalent glycoconjugates", in *Glycoscience Synthesis of Substrate Analogs and Mimetics* (H. Driguez and J. Thiem, eds.), Springer: Berlin, Heidelberg, 1998, pp. 241–274.

[62] R. Roy, "A decade of glycodendrimer chemistry", *Trends Glycosci. Glycotechnol.* **15** (2003), pp. 291–310.

[63] Y. M. Chabre and R. Roy, "Design and creativity in synthesis of multivalent neoglycoconjugates", *Adv. Carbohydr. Chem. Biochem.* **63** (2010), pp. 165–393.

[64] R. Roy, P. V. Murphy and H.-J. Gabius, "Multivalent carbohydrate-lectin interactions: How synthetic chemistry enables insights into nanometric recognition", *Molecules* **21** (2016), article no. 629.

[65] D. Kushwaha and V. K. Tiwari, "Click chemistry inspired synthesis of glycoporphyrin dendrimers", *J. Org. Chem.* **78** (2013), pp. 8184–8190.

[66] M. Touaibia and R. Roy, "First synthesis of "majoral-type" glycodendrimers bearing covalently bound α-D-mannopyranoside residues onto a hexachlocyclotriphosphazene core", *J. Org. Chem.* **73** (2008), pp. 9292–9302.

[67] D. Pagé, D. Zanini and R. Roy, "Macromolecular recognition: effect of multivalency in the inhibition of binding of yeast mannan to concanavalin A and pea lectins by mannosylated dendrimers", *Bioorg. Med. Chem.* **4** (1996), pp. 1949–1961.

[68] D. Pagé, S. Aravind and R. Roy, "Synthesis and lectin binding properties of dendritic mannopyranoside", *Chem. Commun.* **16** (1996), pp. 1913–1914.

[69] D. Pagé and R. Roy, "Synthesis of divalent α-D-mannopyranosylated clusters having enhanced binding affinities towards concanavalin A and pea lectins", *Bioorg. Med. Chem. Lett.* **6** (1996), pp. 1765–1770.

[70] R. Roy and J. M. Kim, "Amphiphilic p-tert-Butylcalix[4]arene scaffolds containing exposed carbohydrate dendrons", *Angew. Chem. Int. Ed.* **38** (1999), pp. 369–372.

[71] Y. M. Chabre and R. Roy, "Multivalent glycoconjugate syntheses and applications using aromatic scaffolds", *Chem. Soc. Rev.* **42** (2013), pp. 4657–4708.

[72] M. Gingras, Y. M. Chabre, M. Roy and R. Roy, "How do multivalent glycodendrimers benefit from sulfur chemistry?", *Chem. Soc. Rev.* **42** (2013), pp. 4823–4841.

[73] O. Renaudet and R. Roy, "Multivalent scaffolds in glycoscience: an overview", *Chem. Soc. Rev.* **42** (2013), pp. 4515–4517.

[74] J. J. García-López, F. Hernández-Mateo, J. Isac-García, J. M. Kim, R. Roy, F. Santoyo-González and

A. Vargas-Berenguel, "Synthesis of per-glycosylated β-cyclodextrins having enhanced lectin binding affinity", *J. Org. Chem.* **64** (1999), pp. 522–531.

[75] M. N. Tahir, Y. Cao, A. Azzouz and R. Roy, "Host-guest chemistry of the sulfasalazine-β-cyclodextrin inclusion complex", *Tetrahedron* **85** (2021), article no. 132052.

[76] R. Roy, F. Hernández-Mateo and F. Santoyo-González, "Synthesis of persialylated β-cyclodextrins", *J. Org. Chem.* **65** (2000), pp. 8743–8746.

[77] Á. Martínez, C. Ortiz Mellet and J. M. García Fernández, "Cyclodextrin-based multivalent glycodisplays: covalent and supramolecular conjugates to assess carbohydrate–protein interactions", *Chem. Soc. Rev.* **42** (2013), pp. 4746–4773.

[78] G. Rivero-Barbarroja, J. M. Benito, C. Ortiz Mellet and J. M. García Fernández, "Cyclodextrin-based functional glyconanomaterials", *Nanomaterials* **10** (2020), article no. 2517.

[79] E. Galante, C. Geraci, S. Sciuto, et al., "Glycoclusters presenting lactose on calix [4] arene cores display trypanocidal activity", *Tetrahedron* **67** (2011), pp. 5902–5912.

[80] M. Mattarella and J. S. Siegel, "Sym-(CH 2 X) 5-corannulenes: molecular pentapods displaying functional group and bioconjugate appendages", *Org. Biomol. Chem.* **10** (2012), pp. 5799–5802.

[81] M. Mattarella, L. Berstis, K. K. Baldridge and J. S. Siegel, "Synthesis of Bioconjugated sym-Pentasubstituted Corannulenes: Experimental and Theoretical Investigations of Supramolecular Architectures", *Bioconjugate Chem.* **25** (2014), pp. 115–128.

[82] I. Nierengarten and J.-F. Nierengarten, "Fullerene sugar balls: a new class of biologically active fullerene derivatives", *Chem. Asian J.* **9** (2014), pp. 1436–1444.

[83] Y. M. Chabre, P. P. Brisebois, L. Abbassi, S. C. Kerr, J. V. Fahy, I. Marcotte and R. Roy, "Hexaphenylbenzene as a rigid template for the straightforward syntheses of "star-shaped" glycodendrimers", *J. Org. Chem.* **76** (2011), pp. 724–727.

[84] R. Roy, "Designing novel multivalent glycotools for biochemical investigations related to sialic acid", *J. Carbohydr. Chem.* **21** (2007), pp. 769–798.

[85] M.-G. Baek, K. Rittenhouse-Olson and R. Roy, "Synthesis and antibody binding properties of glycodendrimers bearing the tumor related T-antigen", *Chem. Commun.* **3** (2001), pp. 257–258.

[86] L. Mousavifar and R. Roy, "Recent development in the design of small 'drug-like' and nanoscale glycomimetics against *Escherichia coli* infections", *Drug Discov. Today* **26** (2021), pp. 2124–2137.

[87] K. Buffet, I. Nierengarten, N. Galanos, et al., "Pillar [5] arene-based glycoclusters: Synthesis and multivalent binding to pathogenic bacterial lectins", *Chem. Eur. J.* **22** (2016), pp. 2955–2963.

[88] A. K. Agrahari, M. K. Jaiswal, M. S. Yadav and V. K. Tiwari, "CuAAC mediated synthesis of cyclen cored glycodendrimers of high sugar tethers at low generation", *Carbohydr. Res.* **508** (2021), article no. 108403.

[89] A. K. Agrahari, A. S. Singh, R. Mukherjee and V. K. Tiwari, "An expeditious click approach towards the synthesis of galactose coated novel glyco-dendrimers and dentromers utilizing a double stage convergent method", *RSC Adv.* **10** (2020), pp. 31553–31562.

[90] S. V. Pawar, P. K. Upadhyay, S. Burade, N. Kumbhar, R. Patil and D. D. Dhavale, "Synthesis and anti-leishmanial activity of TRIS-glycine-β-alanine dipeptidic triazole dendron coated with nonameric mannoside glycocluster", *Carbohydr. Res.* **485** (2019), article no. 107815.

[91] S. J. Meunier, Q. Wu, S.-N. Wang and R. Roy, "Synthesis of hyperbranched glycodendrimers incorporating α-thiosialosides based on a gallic acid core", *Can. J. Chem.* **75** (1997), pp. 1472–1482.

[92] R. Sharma, I. Zhang, T. C. Shiao, G. M. Pavan, D. Maysinger and R. Roy, "Low generation polyamine dendrimers bearing flexible tetraethylene glycol as nanocarriers for plasmids and siRNA", *Nanoscale* **8** (2016), pp. 5106–5119.

[93] E. Cavero, M. Zablocka, A.-M. Caminade and J. P. Majoral, "Design of bisphosphonate-terminated dendrimers", *Eur. J. Org. Chem.* **14** (2010), pp. 2759–2767.

[94] R. E. Bird, S. A. Lemmel, X. Yu and Q. A. Zhou, "Bioorthogonal chemistry and its applications", *Bioconjugate Chem.* **32** (2021), pp. 2457–2479.

[95] S. L. Scinto, D. A. Bilodeau, R. Hincapie, et al., "Bioorthogonal chemistry", *Nat. Rev. Meth. Primers* **1** (2021), article no. 30.

[96] T. K. Tiefenbrunn and P. E. Dawson, "Chemoselective ligation techniques: Modern applications of time-honored chemistry", *Pept. Sci.* **94** (2010), pp. 95–106.

[97] R. Roy and T. C. Shiao, "Glyconanosynthons as powerful scaffolds and building blocks for the rapid construction of multifaceted, dense and chiral dendrimers", *Chem. Soc. Rev.* **44** (2015), pp. 3924–3941.

[98] I. Deguise, D. Lagnoux and R. Roy, "Synthesis of glycodendrimers containing both fucoside and galactoside residues and their binding properties to Pa-IL and PA-IIL lectins from Pseudomonas aeruginosa", *New J. Chem.* **31** (2007), pp. 1321–1331.

[99] M.-G. Baek and R. Roy, "Synthesis and protein binding properties of T-antigen containing GlycoPAMAM dendrimers", *Bioorg. Med. Chem.* **10** (2002), pp. 11–17.

[100] D. Zanini and R. Roy, "Novel dendritic α-sialosides: Synthesis of glycodendrimers based on a 3,3′-Iminobis (propylamine) core", *J. Org. Chem.* **61** (1996), pp. 7348–7354.

[101] R. Clayton, J. Hardman, C. C. LaBranche and K. D. McReynolds, "Evaluation of the synthesis of sialic acid-PAMAM glycodendrimers without the use of sugar protecting groups, and the anti-HIV-1 properties of these compounds", *Bioconjugate Chem.* **22** (2011), pp. 2186–2197.

[102] A. Nelson and J. F. Stoddart, "Synthesis of lactoside glycodendrons using photoaddition and reductive amination methodologies", *Carbohydr. Res.* **339** (2004), pp. 2069–2075.

[103] W. B. Turnbull, S. A. Kalovidouris and J. F. Stoddart, "Large oligosaccharide-based glycodendrimers", *Chem. Eur. J.* **8** (2002), pp. 2988–3000.

[104] S. A. Kalovidouris, W. B. Turnbull and J. F. Stoddart, "Glycodendrimers based on cellobiosyl-derived monomers", *Can. J. Chem.* **80** (2002), pp. 983–991.

[105] A. Marra, M.-C. Scherrmann, A. Dondoni, R. Ungaro, A. Casnati and P. Minari, "Sugar calixarenes: Preparation of calix [4] arenes substituted at the lower and upper rims with O-glycosyl groups", *Angew. Chem. Int. Ed. Engl.* **33** (1995), pp. 2479–2481.

[106] D. Zanini and R. Roy, "Practical synthesis of starburst PAMAM α-thiosialodendrimers for probing multivalent carbohydrate–lectin binding properties", *J. Org. Chem.* **63** (1998), pp. 3486–3491.

[107] M. L. Wolfenden and M. J. Cloninger, "Mannose/glucose-functionalized dendrimers to investigate the predictable tunability of multivalent interactions", *J. Am. Chem. Soc.* **127** (2005), pp. 12168–12169.

[108] D. Pagé and R. Roy, "Synthesis and biological properties of mannosylated starburst poly (amidoamine) dendrimers", *Bioconjugate Chem.* **8** (1997), pp. 714–723.

[109] P. I. Kitov, J. M. Sadowska, G. Mulvey, G. D. Armstrong, H. Ling, N. S. Pannu, R. J. Read and D. R. Bundle, "Shiga-like toxins are neutralized by tailored multivalent carbohydrate ligands", *Nature* **403** (2000), pp. 669–672.

[110] V. Wittmann and S. Seeberger, "Combinatorial solid-phase synthesis of multivalent cyclic neoglycopeptides", *Angew. Chem., Int. Ed.* **39** (2000), pp. 4348–4352.

[111] R. Roy, "Syntheses and some applications of chemically defined multivalent glycoconjugates", *Curr. Opin. Struct. Biol.* **6** (1996), pp. 692–702.

[112] N. Kottari, Y. M. Chabre, T. C. Shiao, R. Rej and R. Roy, "Efficient and accelerated growth of multifunctional dendrimers using orthogonal thiol–ene and S$_N$2 reactions", *Chem. Commun.* **50** (2014), pp. 1983–1985.

[113] C. Pifferi, B. Thomas, D. Goyard, N. Berthet and O. Renaudet, "Heterovalent glycodendrimers as epitope carriers for antitumor synthetic vaccines", *Chem. Eur. J.* **23** (2017), pp. 16283–16296.

[114] C. Pifferi, G. C. Daskhan, M. Fiore, T. C. Shiao, R. Roy and O. Renaudet, "Aminooxylated carbohydrates: Synthesis and applications", *Chem. Rev.* **117** (2017), pp. 9839–9873.

[115] V. K. Tiwari, B. B. Mishra, K. B. Mishra, N. Mishra, A. S. Singh and X. Chen, "Cu-catalyzed click reaction in carbohydrate chemistry", *Chem. Rev.* **116** (2016), pp. 3086–3240.

[116] T. C. Shiao, R. Rej, M. Rose, G. M. Pavan and R. Roy, "Synthesis of dense and chiral dendritic polyols using glyconanosynthon scaffolds", *Molecules* **21** (2016), article no. 448.

[117] R. Roy, S. K. Das, R. Dominique, M. C. Trono, F. Hernández-Mateo and F. Santoyo-Gonzalez, "Transition metal catalyzed neoglycoconjugate syntheses", *Pure Appl. Chem.* **71** (1999), pp. 565–571.

[118] B. Liu and R. Roy, "Olefin self-metathesis as a new entry into xenotransplantation antagonists bearing the Galili antigen", *Chem. Commun.* **6** (2002), pp. 594–595.

[119] D. Giguère, M. A. Bonin, P. Cloutier, R. Patnam, C. St-Pierre, S. Sato and R. Roy, "Synthesis of stable and selective inhibitors of human galectins-1 and-3", *Bioorg. Med. Chem.* **16** (2008), pp. 7811–7823.

[120] E. Smits, J. B. F. N. Engberts, R. M. Kellogg and H. A. van Doren, "Reliable method for the synthesis of aryl β-D-glucopyranosides, using boron trifluoride–diethyl ether as catalyst", *J. Chem. Soc., Perkin Trans. 1* **24** (1996), pp. 2873–2877.

[121] M. Bergeron-Brlek, D. Giguère, T. C. Shiao, C. Saucier and R. Roy, "Palladium-catalyzed Ullmann-type reductive homocoupling of iodoaryl glycosides", *J. Org. Chem.* **77** (2012), pp. 2971–2977.

[122] R. Roy, R. Dominique and S. K. Das, "Extended alkenyl glycosides by ruthenium-catalyzed cross-metathesis reaction and application toward novel C-linked pseudodisaccharides", *J. Org. Chem.* **64** (1999), pp. 5408–5412.

[123] R. Roy and S. K. Das, "Recent applications of olefin metathesis and related reactions in carbohydrate chemistry", *Chem. Commun.* **7** (2000), pp. 519–529.

[124] R. Dominique, S. K. Das and R. Roy, "Alkenyl O- and C-glycopyranoside homodimerization by olefin metathesis reaction", *Chem. Commun.* **22** (1998), pp. 2437–2438.

[125] K. P. C. Vollhardt, "Cobalt-mediated [2 + 2 + 2]-cycloadditions: a maturing synthetic strategy [new synthetic methods (43)]", *Angew. Chem. Int. Ed. Engl.* **23** (1984), pp. 539–556.

[126] R. J. Kaufman and R. S. Sidhu, "Synthesis of aryl cluster glycosides by cyclotrimerization of 2-propynyl carbohydrate derivatives", *J. Org. Chem.* **47** (1982), pp. 4941–4947.

[127] S. Yoon, S. H. Hwang and W. B. Ko, "Ultrasound-assisted cycloadditions of [70]fullerene with various 2-azidoethyl per-O-acetyl glycosides", *Full., Nanotubes, Carb. Nanostruct.* **17** (2009), pp. 496–506.

[128] S. Yoon, S. H. Hwang and W. B. Ko, "Synthesis of glycosyls fullerene [C70] under ultrasonic irradiation", *Colloids Surf. A: Physicochem. Eng. Asp.* **313–314** (2008), pp. 304–307.

[129] P. R. Ashton, S. E. Boyd, C. L. Brown, N. Jayaraman, S. A. Nepogodiev and J. F. Stoddart, "A convergent synthesis of carbohydrate-containing dendrimers", *Chem. Eur. J.* **2** (1996), pp. 1115–1128.

[130] M. Ghirardello, K. Öberg, S. Staderini, et al., "Thiol-ene and thiol-yne-based synthesis of glycodendrimers as nanomolar inhibitors of wheat germ agglutinin", *J. Polym. Sci., Part A: Polym. Chem.* **52** (2014), pp. 2422–2433.

[131] D. Heyl, E. Rikowski, R. C. Hoffmann, J. J. Schneider and W. D. Fessner, "A "clickable" hybrid nanocluster of cubic symmetry", *Chem. Eur. J.* **16** (2010), pp. 5544–5548.

[132] B. Trastoy, M. E. Pérez-Ojeda, R. Sastre and J. L. Chiara, "Octakis (3-azidopropyl) octasilsesquioxane: A versatile nanobuilding block for the efficient preparation of highly functionalized cube-octameric polyhedral oligosilsesquioxane frameworks through click assembly", *Chem. Eur. J.* **16** (2010), pp. 3833–3841.

[133] C. Boyère, G. Broze, C. Blecker, C. Jérôme and A. Debuigne, "Monocatenary, branched, double-headed, and bolaform surface active carbohydrate esters via photochemical thiol-ene/-yne reactions", *Carbohydr. Res.* **380** (2013), pp. 29–36.

[134] S. Moffett, T. C. Shiao, L. Mousavifar, S. Mignani and R. Roy, "Aberrant glycosylation patterns on can-

cer cells: Therapeutic opportunities for glycodendrimers/metallodendrimers oncology", *Wiley Interdiscip. Rev. Nanomed. Nanobiotechnol.* **13** (2021), article no. e1659.

[135] V. Balzani, S. Campagna, G. Denti, A. Juris, S. Serroni and M. Venturi, "Designing dendrimers based on transition-metal complexes. Light-harvesting properties and pre-determined redox patterns", *Acc. Chem. Res.* **31** (1998), pp. 26–34.

[136] G. R. Newkome, E. He and C. N. Moorefield, "Suprasuper-molecules with novel properties: Metallodendrimers", *Chem. Rev.* **99** (1999), pp. 1689–1746.

[137] F. Zeng and S. C. Zimmerman, "Dendrimers in supramolecular chemistry: from molecular recognition to self-assembly", *Chem. Rev.* **97** (1997), pp. 1681–1712.

[138] D. Wetzel and U. Lüning, "Second generation self-assembling dendrimer combining supramolecular and dynamic covalent chemistry", *Eur. J. Org. Chem.* **43** (2022), article no. e202200983.

[139] P. Reeh and J. de Mendoza, "Dynamic multivalency for carbohydrate-protein recognition through dynamic combinatorial libraries based on FeII-bipyridine complexes", *Chem. Eur. J.* **19** (2013), pp. 5259–5262.

[140] O. Ramström, S. Lohmann, T. Bunyapaiboonsri and J.-M. Lehn, "Dynamic combinatorial carbohydrate libraries: probing the binding site of the concanavalin A lectin", *Chem. Eur. J.* **10** (2004), pp. 1711–1715.

[141] R. Roy and J. M. Kim, "Cu (II)-self-assembling bipyridyl-glycoclusters and dendrimers bearing the Tn-antigen cancer marker: syntheses and lectin binding properties", *Tetrahedron* **59** (2003), pp. 3881–3893.

[142] L. Mousavifar, S. Abdullayev and R. Roy, "Recent development in the design of neoglycoliposomes bearing arborescent architectures", *Molecules* **26** (2021), article no. 14.

[143] V. Percec, P. Leowanawat, H.-J. Sun, et al., "Modular synthesis of amphiphilic Janus glycodendrimers and their self-assembly into glycodendrimersomes and other complex architectures with bioactivity to biomedically relevant lectins", *J. Am. Chem. Soc.* **135** (2013), pp. 9055–9077.

[144] S. Zhang, R.-O. Moussodia, H.-J. Sun, et al., "Mimicking biological membranes with programmable glycan ligands self-assembled from amphiphilic janus glycodendrimers", *Angew. Chem. Int. Ed.* **126** (2014), pp. 11079–11083.

[145] S. Zhang, R.-O. Moussodia, C. Murzeau, et al., "Dissecting molecular aspects of cell interactions using glycodendrimersomes with programmable glycan presentation and engineered human lectins", *Angew. Chem. Int. Ed.* **127** (2015), pp. 4108–4112.