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## Controlled density glycodendron microarrays for studying carbohydrate–lectin interactions†

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Glycodendron microarrays with defined valency have been constructed by on-chip synthesis on hydrophobic indium tin oxide (ITO) coated glass slides and employed in lectin–carbohydrate binding studies with several plant and human lectins. Glycodendrons presenting sugar epitopes at different valencies were prepared by spotwise strain-promoted azide–alkyne cyclo-addition (SPAAC) between immobilised cyclooctyne dendrons and azide functionalised glycans. The non-covalent immobilisation of dendrons on the ITO surface by hydrophobic interaction allowed us to study dendron surface density and SPAAC conversion rate by *in situ* MALDI-TOF MS analysis. By diluting the dendron surface density we could study how the carbohydrate–lectin interactions became exclusively dependant on the valency of the immobilised glycodendron.

Glycan microarrays are now well established high-throughput research tools for studying the specificity of novel lectins and carbohydrate processing enzymes.<sup>1–4</sup>

Monovalent carbohydrate–glycan interactions are usually weaker than most specific protein–protein interactions but the lectin binding affinity of glycans can be enhanced by over 3 orders of magnitude by the dense and multivalent presentation of glycans immobilised on a microarray surface.<sup>5</sup> The surface display of glycans can favour the rapid rebinding of ligands to receptor domains and the chelation of multiple ligands with multivalent receptors leading to an overall increase in avidity. In nature, glycolipid clustering, *N*-glycan

branching and repetition of recognition motifs on extended biopolymers like mucins or polysaccharides are mechanisms to achieve a similar increase in glycan–protein avidity.

For the preparation of microarrays, glycan ligands are either isolated from natural sources or prepared by chemical and enzymatic synthesis, and conveniently derivatized for printing onto microarray surfaces by a variety of chemical or physical methods.<sup>4</sup>

Glycans and their conjugates are usually synthesised in solution before immobilisation or isolated from natural sources but on-chip chemo-enzymatic assembly of glycan ligands as pioneered by our group and others can rapidly provide high density glycan microarrays with potentially very high savings in time and resources.<sup>9–11</sup> Monitoring the conversion of surface-based reactions is important for assessing ligand homogeneity but at the same time challenging due to the limited number of compatible available analytical methods.

Fluorescently labelled lectins that bind specifically to starting materials and products can be used to qualitatively monitor the progress of enzymatic glycosylations using recombinant glycosyltransferases.<sup>11</sup>

With the development of surfaces that are compatible with analysis by mass spectrometry and microarray technology, the analysis of surface based reactions including the construction of glycan arrays, was finally possible.<sup>12–17</sup> As a variable surface density can affect the strength of interaction between glycans and lectins, several groups have developed strategies to homogenize and control glycan surface density.<sup>18,19</sup>

A number of approaches employ macromolecules like polymers, cyclic peptides, peptide nucleic acids, proteins or dendrons that have been functionalized with glycans at a discrete inter-glycan spacing.<sup>20–22</sup> Immobilisation of these well-defined glyco-macromolecules however, results in variable inter-glycan distances due to the non-homogenous functionalisation of the surface with the glycoconjugates.<sup>23</sup>

Glycan surface densities have been assessed using a fluorescent linker for attachment<sup>24</sup> for direct quantification of immobilised species, by wash off experiments with fluorescent

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ligands<sup>5</sup> or by autoradiography after enzymatic extension of ligands with a tritium labelled probe.<sup>6</sup> *Via* tritium imaging we have previously visualized how printing of different glycans at the same concentration lead to a variable surface density presumably due to differences in size, hydrophobicity and structure dependant kinetics for the immobilisation reaction.<sup>6–8</sup>

With the exception of the tritium imaging method, the mentioned methods only provide a rough estimation of ligand density as a function of the ligand printing concentration but no data on individual ligand densities. Being able to control and assess glycan ligand density however is key for the development of well-defined and reproducible glycan microarrays for quantitative applications in clinical diagnostics and glycomics research. In addition, it is helpful for evaluating multivalent display systems for solution phase applications by ensuring that the multivalent presentation on the surface matches those used in solution phase, eliminating the congestion surface effect.<sup>19,25</sup> Here we present a strategy for preparing glycan arrays with a homogenous and constant ligand density by a process comprising (1) the non-covalent coating of a slide with a layer of a NHS-functionalized lipid, (2) attachment of a dendron spacer carrying a bioorthogonal cyclooctyne group and (3) complete functionalisation of the spacer with different glycans by SPAAC.

Recently Mende *et al.* published a similar strategy based on the on-chip synthesis of multivalent alkyne-functionalized peptides which were coupled with azido glycans by copper(I)-catalyzed azide–alkyne cycloaddition (CuAAC).<sup>26</sup>

The authors prepared a homogenous surface layer of alkyne-functionalized tetrapeptides by an on-chip laser assisted printing process termed cLIFT. While the described process is likely to produce more homogenous neoglycopptide arrays than by printing of pre-synthesized multivalent

ligands, the surface homogeneity or density was not directly assessed.

Our approach towards glycan arrays with constant glycan density made use of a previously developed microarray platform where glycan ligands are attached *via* hydrophobic interactions to a lipophilic indium tin oxide (ITO) glass slide.<sup>12</sup> Microarrays prepared in this manner are fully compatible with readout both by fluorescence imaging and MALDI-TOF MS which allowed us to monitor all reactions carried out on the slide. For the glycan microarray employed in this study, we prepared a novel class of cyclooctyne activated mono-bi- and tetravalent dendrons (1–3) based on a bis-MPA (2,2-bismethyl-propionic acid) synthetic strategy.<sup>12</sup> These dendrons are functionalised with an amine group at the focal point allowing the immobilisation on the surface, a bis-MPA skeleton and cyclooctyne groups at the terminal points, ready to be involved for the SPAAC conjugation with a selection of simple azido ethylglycosides (4–7)<sup>27,28</sup> (Fig. 1).

For the assembly of the glycodendron arrays we functionalised commercial indium tin oxide slides first with a layer of octadecylphosphonate onto a NHS-activated carbamate linker (Fig. 2a) by spray-coating as reported previously.<sup>13</sup>

This hydrophobic bilayer was then functionalised with cyclooctyne-dendrons (1–3), at different concentrations and in separate wells (Step I, Fig. 1). To eliminate any congestion surface effects we explored surface functionalisation at low dendron concentrations between 5–30  $\mu$ M (Fig. 2). At higher dendron dilution we reasoned that the glycan surface density would be easily adjustable by employing dendron spacers with one, two or four terminal cyclooctyne groups (Fig. 1).

The coating density with cyclooctyne dendrons 1–3 was determined by mass spectrometry as the ratio of dendron-conjugate and unreacted free NHS activated linker ( $m/z$  1040)

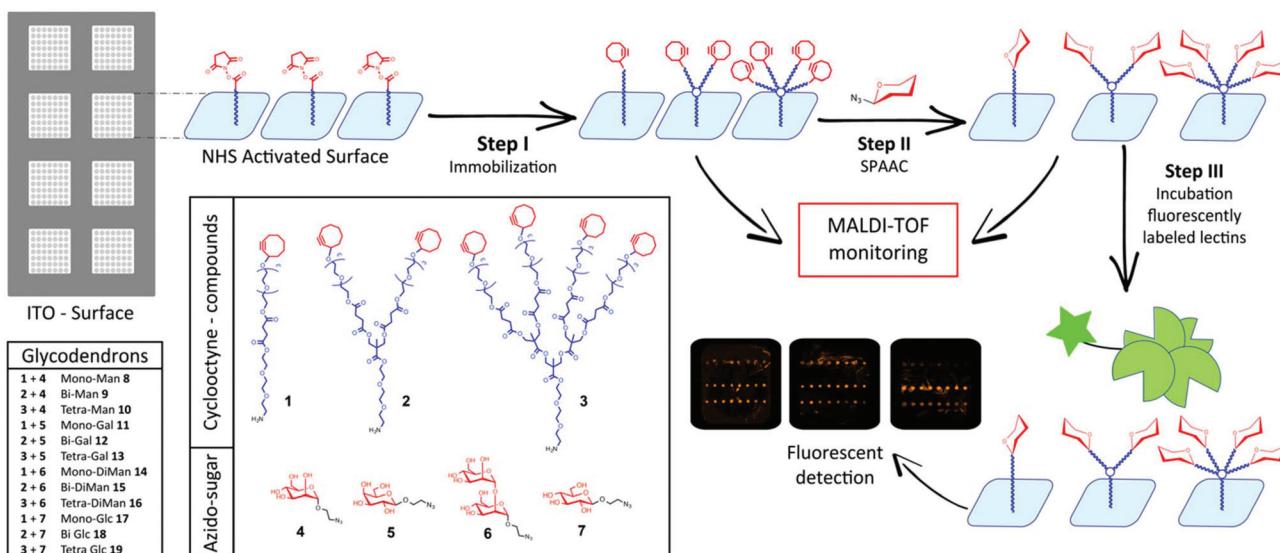
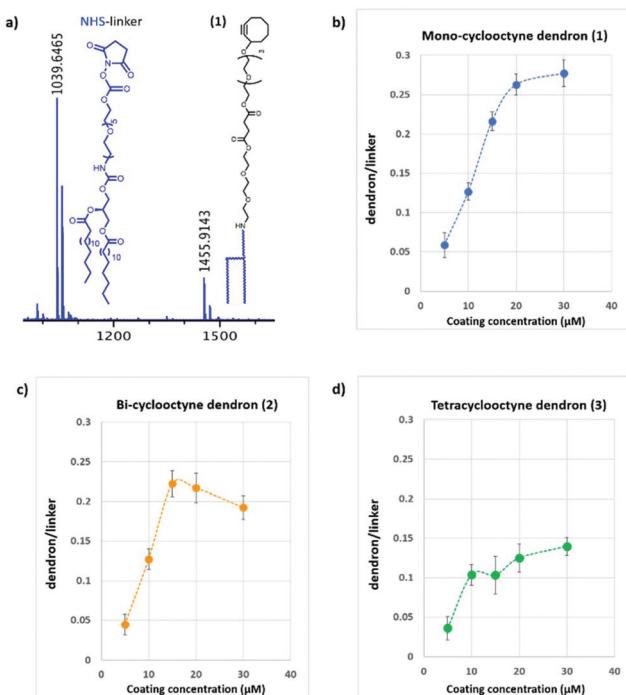


Fig. 1 General outline of the glycodendron array strategy.





**Fig. 2** (a) MALDI-TOF spectrum showing immobilized Mono-cyclooctyne (**1**) mass peak ( $m/z$  1456) vs unreacted NHS-linker mass peak ( $m/z$  1040). (b-d) Graphs of the dendron/linker ratio against coating concentration.

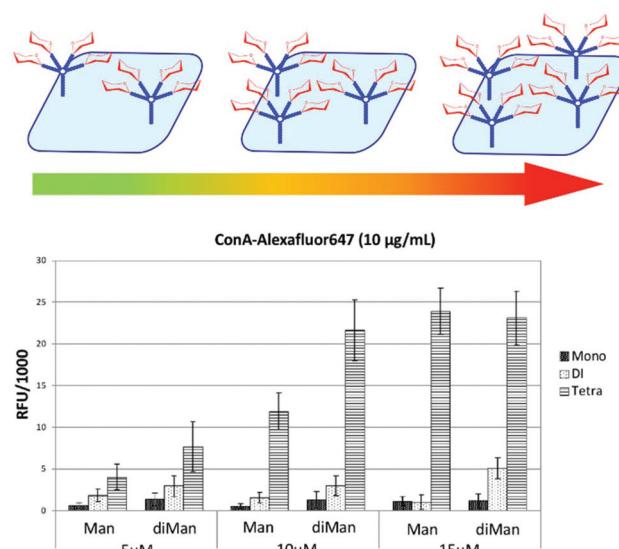
which was used as internal standard. The effect of linker concentration used for coating on the surface density measured as dendron/linker ratio is depicted in Fig. 2 for mono-, bi- and tetravalent cyclooctyne dendrons **1–3**. As a general trend, we observed a linear increase of the relative surface density with increasing dendron concentration up to a maximum where saturation in surface density occurred. We found a compound-dependant surface saturation at 20  $\mu$ M for the monovalent ligand **1** while the bulkier bi- and tetra-valent dendrons **2** and **3** saturated at lower coating concentrations around 15 and 10  $\mu$ M respectively (Fig. 2b-d).

The variation observed in dendron/linker ratios, for the different dendrons, is likely to reflect both the structure-dependent ionization and the spatial requirements during immobilisation.

It is of note that at lower concentration (5–10  $\mu$ M) the three dendrons showed a similar density while at higher coating concentrations the dendron structures strongly affect the immobilization ratio.

These results suggest that a comparable ligand density for the three multivalent scaffolds (**1–3**) can be achieved for a coating concentration between 5–10  $\mu$ M. Moreover, surfaces functionalized with dendrons at high dilution lead to low surface densities where the binding event is likely determined by dendron valency and architecture alone and with minimal “bridging” of lectins with 2 more immobilised ligand molecules.

To study the effect of the ligand density on multivalent binding, dendron surfaces (**1–3**) were prepared at three coating



**Fig. 3** Fluorescence quantification of Concanavalin A binding to Man (**8–10**) and  $\alpha$ Man1,2Man-based (**14–16**) glycodendrons at different dendrons coating concentration (5, 10 and 15  $\mu$ M).

concentrations (5, 10, and 15  $\mu$ M), coupled to azido linked mannose (**4**) and  $\alpha$ 1,2-dimannose (**6**) via spot-wise printing and screened against fluorescently labelled Concanavalin A (Fig. 3).

At lower density (5–10  $\mu$ M) tetravalent glycodendrons functionalized either with  $\alpha$ 1,2-dimannose (**6**) or mannose (**4**) bound Con A stronger than the mono- and bivalent scaffolds. The dimannoside ligand bound stronger to ConA than the mannose monosaccharide at all dendron valencies studied (Fig. 3). At higher glycodendron densities (15  $\mu$ M), tetravalent mannose (**10**) and  $\alpha$ 1,2-dimannose (**16**) showed similar binding strength to ConA. Moreover, no differences in binding between the mono- (**8**) and the bivalent (**9**) mannose scaffolds were observed.

Based on these initial results, we chose to functionalize the microarray surface with dendrons at a low printing concentration of 5  $\mu$ M. Arrays of mono, bi- and tetravalent glycodendrons were then prepared via SPAAC by spotting azido-ethyl glycosides (**4–7**) solutions onto cyclooctyne surfaces (**1–3**) (Step II, Fig. 1). On-chip SPAAC reactions were evaluated by MALDI TOF MS to ensure complete conversion for all printed glycosides. The surface coating at low dendron densities, together with the complete conversion in SPAAC conjugation (ESI $\dagger$ ), provided homogeneous glycodendron presentations for the study of valency dependant carbohydrate lectin binding, excluding the multivalent effect from the array. For our binding study (Step III, Fig. 1) we chose the two commercially available plant lectins *Pisum sativum* (PSA) and *Wisteria floribunda* (WFA) and the extracellular domains (ECD) of the human lectins receptors *DC-SIGN*, *DC-SIGNR*, *Langerin* and *Dectin-2* (Fig. 4).

Most glycan arrays, currently employed in glycomics research, are printed at ligand concentrations between

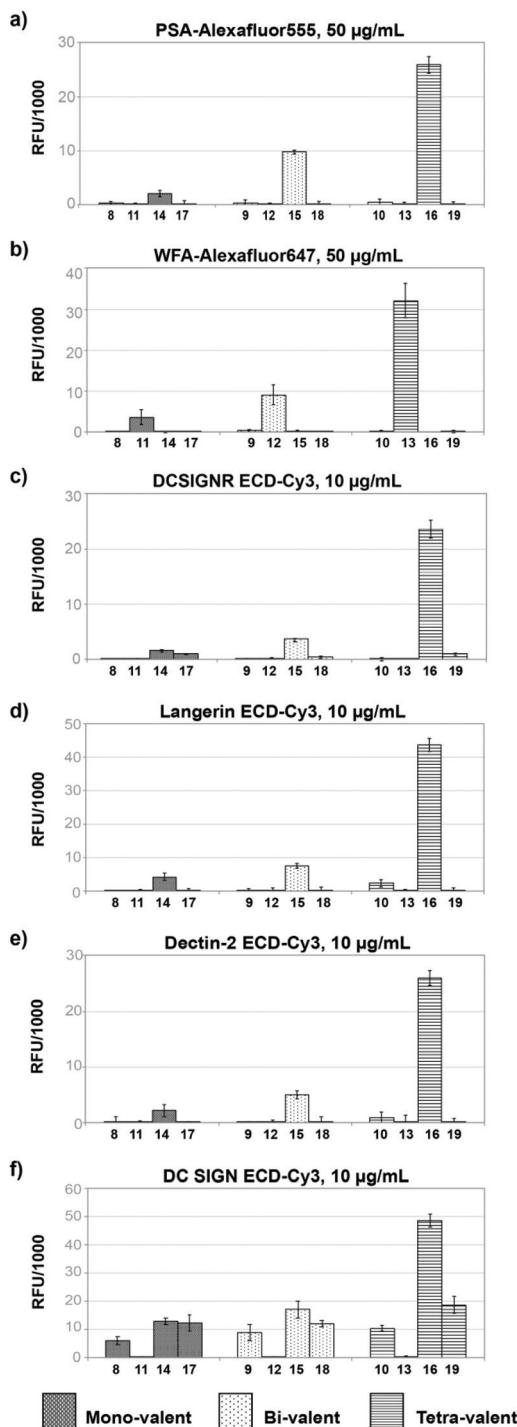


Fig. 4 Glycodendron binding profiles to (a) PSA, (b) WFA, (c) DC-SIGNR ECD, (d) Langerin ECD, (e) Dectin-2 ECD, (f) DC-SIGN ECD.

50–100  $\mu$ M for which saturated ligand immobilisation was observed for the majority of glycans. Here, we chose to functionalize surfaces with the dendrons at a highly dilute concentration of 5  $\mu$ M to deliberately achieve low surface densities where the binding event would be largely determined by the dendron valency and architecture thus avoiding crosslinking of lectins to two or more immobilised ligand molecules.

## Plant lectins

In our assay dimeric PSA<sup>29</sup> with preference for binding Fuc $\alpha$ 1,6GlcNAc and  $\alpha$ -Man residues showed no binding to the mannose (8–10), galactose (11–13) or glucose derivatives (17–19) independent of their valency. Only structures presenting disaccharide Man $\alpha$ 1,2Man (14–16) were bound by the lectin with a strong increase in binding when moving from the mono to the tetravalent species (Fig. 4a). Dendrons functionalised with mannose monosaccharide were not bound at this low surface concentration.

In the case of monomeric WFA,<sup>30</sup> a lectin with preference for terminal GalNAc structures, the lectin bound the galactoside derivatives (11–13) with a strong effect of ligand valency in the binding (Fig. 4b).

## Human lectins

The human tetravalent C-type lectin DC-SIGNR ECD,<sup>31,32</sup> trivalent Langerin ECD<sup>33,34</sup> and the monomeric Dectin-2 ECD<sup>35</sup> showed highly selective binding only to constructs presenting the Man $\alpha$ 1,2Man epitope, (14–16) with increasing binding intensity for higher valency. As with PSA lectin, no binding was observed towards dendrons functionalised with mannose monosaccharide (8, 9, 10).

For DC-SIGN ECD,<sup>13</sup> we observed similar binding strength towards mono- and divalent dendrons displaying mannose monosaccharide (8,9), Man $\alpha$ 1,2Man (14,15) and glucose (17,18) but strongest binding to the Man $\alpha$ 1,2Man disaccharide (16) in the tetravalent form (Fig. 4f). In line with the known binding specificity of DC-SIGN, no binding was observed for the dendrons presenting galactose residues (11–13).

This nearly 5-fold increase in affinity for the Man $\alpha$ 1,2Man in a tetravalent presentation shows the gain in selectivity through a particular multivalent presentation in an otherwise more promiscuous lectin. Brewer *et al.* have explained this behaviour with a threshold epitope density required to produce a productive interaction between lectins and glycans that is capable to trigger a biological signalling event *e.g.* in cellular immunity.<sup>23</sup>

As the lectins show a different degree of labelling (DOL) the fluorescence values (RFU) obtained should not be compared directly between each other without prior normalisation (see ESI†).

In conclusion, we have developed a rapid screening platform to study multivalent carbohydrate lectin interactions as a function of dendron valency and carbohydrate epitope. The glycodendron on-chip synthesis was assisted by MALDI-TOF MS analysis to monitor the conversion of surface based SPAAC reactions.

The dilute ligand presentation on the array surface implied that ligand valency was mainly determined by the dendron architecture, and in consequence allowed us to identify matched glycan epitope/dendron combinations with high avidity and much-improved selectivity over other structurally

similar compounds. We anticipate this type of glycan array, which includes defined ligand presentation as an additional parameter, to be a valuable tool for the discovery of multi-valent glycoconjugates as antagonists to interfere in clinically relevant carbohydrate–lectin interactions such as pathogen infection or glycoimmunity therapy.

## Conflicts of interest

No conflicts of interest are declared.

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