

# Automated Synthesis of Algal Fucoidan Oligosaccharides

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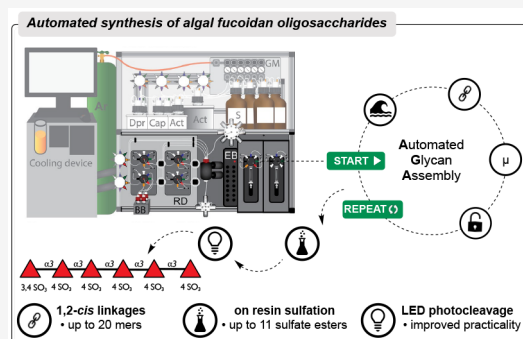


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**ABSTRACT:** Fucoidan, a sulfated polysaccharide found in algae, plays a central role in marine carbon sequestration and exhibits a wide array of bioactivities. However, the molecular diversity and structural complexity of fucoidan hinder precise structure–function studies. To address this, we present an automated method for generating well-defined linear and branched  $\alpha$ -fucan oligosaccharides. Our syntheses include oligosaccharides with up to 20 *cis*-glycosidic linkages, diverse branching patterns, and 11 sulfate monoesters. In this study, we demonstrate the utility of these oligosaccharides by (i) characterizing two *endo*-acting fucoidan glycoside hydrolases (GH107), (ii) utilizing them as standards for NMR studies to confirm suggested structures of algal fucoidans, and (iii) developing a fucoidan microarray. This microarray enabled the screening of the molecular specificity of four monoclonal antibodies (mAb) targeting fucoidan. It was found that mAb BAM4 has cross-reactivity to  $\beta$ -glucans, while mAb BAM2 has reactivity to fucoidans with 4-*O*-sulfate esters. Knowledge of the mAb BAM2 epitope specificity provided evidence that a globally abundant marine diatom, *Thalassiosira weissflogii*, synthesizes a fucoidan with structural homology to those found in brown algae. Automated glycan assembly provides access to fucoidan oligosaccharides. These oligosaccharides provide the basis for molecular level investigations into fucoidan's roles in medicine and carbon sequestration.



## INTRODUCTION

Polysaccharides are the central metabolic fuel of the marine carbon cycle. Annually, algae sequester petagrams of carbon dioxide into a rich diversity of glycans.<sup>1</sup> The unique structure of each glycan dictates its residence time and flow within marine ecosystems.<sup>2</sup> Macroalgae and diatoms synthesize and secrete fucose-containing sulfated polysaccharides, termed fucoidan, into the environment.<sup>3,4</sup> The structural diversity of fucoidan poses challenges to marine bacteria, necessitating evolution of complex enzymatic cascades for its degradation.<sup>5</sup> Fucoidan that escapes microbial degradation can self-assemble into particles,<sup>6</sup> sink to the deep ocean, and store carbon for centuries.<sup>7,8</sup> Moreover, fucoidan also displays a range of biological activities that are under investigation in drug development and cosmetics.<sup>9,10</sup> A current limitation is that poor knowledge exists regarding the exact molecular determinants of fucoidan that mediate a specific bioactivity or if there are precise structures in fucoidan that best mediate carbon sequestration.

To uncover the molecular mechanisms governing fucoidan carbon sequestration and its bioactivities, well-defined standards are imperative. Extraction from biological systems does not lead to homogeneous samples due to the nontemplate-encoded nature of glycans. Consequently, chemical synthesis stands out as a distinct method to obtain precisely defined organic matter.<sup>11</sup> An automated process would significantly

enhance the accessibility of defined fucoidan oligosaccharides. These defined standards would form the basis for a variety of investigations including: creating microarrays,<sup>12–16</sup> delineating the activities of carbohydrate-active enzymes (CAZymes),<sup>17–20</sup> and serving as standards for NMR experiments.<sup>21–26</sup>

The chemical synthesis of complex glycans is challenging,<sup>11</sup> however, advances in automated approaches have enabled high-throughput assembly of both oligosaccharides and polysaccharides.<sup>27,28</sup> The automated chemical synthesis of fucoidan oligosaccharides faces three primary challenges: (i) the stereocontrolled formation of 1,2 *cis*-glycosidic bonds,<sup>29–33</sup> (ii) the high degree of sulfation,<sup>34–37</sup> and (iii) the high reactivity of fucose glycosyl donors.<sup>38–40</sup>

Here, we present an automated glycan assembly (AGA) process for synthesizing well-defined fucoidan oligosaccharides, encompassing linear  $\alpha$ -fucans up to 20-mers, branched fucoidan oligosaccharides, and glycans that contain up to 11 sulfate esters. These glycans served as standards for NMR spectroscopy, aided in delineation of CAZymes activities, and

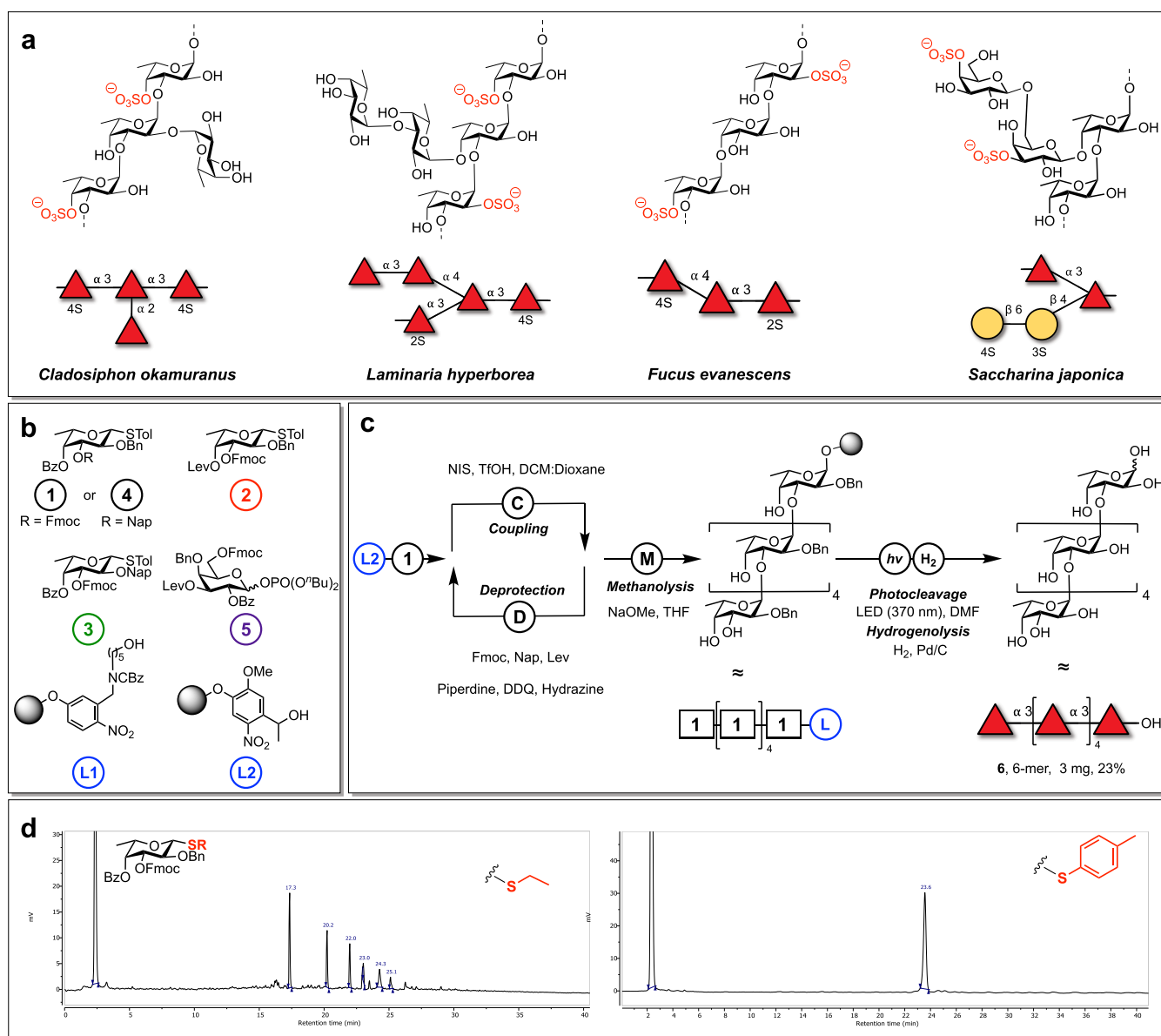
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**Figure 1.** Structural diversity of fucoidan. (a) Four examples of fucoidan found in brown algae. From left: *Cladosiphon okamuranus* contains an  $\alpha$ -1,3-backbone and is known to possess a degree of 4-*O*-sulfation.<sup>41,42</sup> *Laminaria hyperborea* is an  $\alpha$ -1,3 linked fucan with a defined number of motifs including those with  $\alpha$ -1,4 and  $\alpha$ -1,2 branches with sulfate esters primarily on C-2 and C-4.<sup>43</sup> *Fucus evanescens* is an  $\alpha$ -1,3/1,4 linked fucoidan.<sup>44</sup> *Saccharina japonica* is an  $\alpha$ -1,3 linked fucoidan known to contain  $\beta$ -1,4 galactopyranosyl branches.<sup>45</sup> (b) Building blocks used in this study. (c) Automated assembly of fucoidan hexasaccharide. (d) Representative example of HPLC traces from AGA of fucoidan hexasaccharide **6**. Left trace with a thioethyl thioglycoside, right trace with a 4-methylphenyl thioglycoside.

enabled the creation of a fucoidan microarray. Utilizing this microarray, we elucidated the specificity of fucoidan-directed antibodies, which, in turn, suggests brown algae-type fucoidan motifs can be found in marine diatoms.

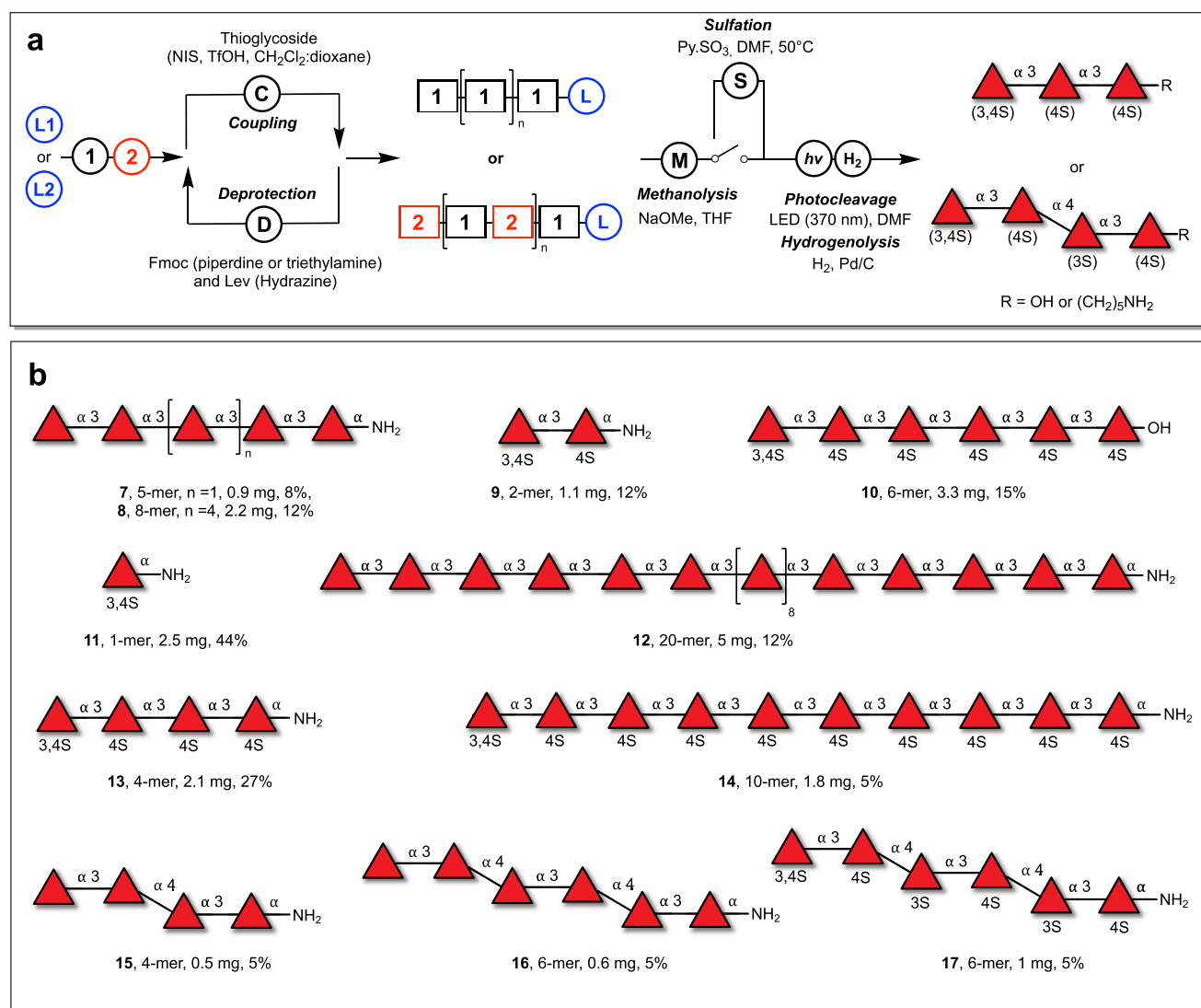
## RESULTS AND DISCUSSION

### Retrosynthetic Analyses and Building Block Design.

Homo- and heterofucans represent two classes of fucoidan. Homofucans consist of either  $\alpha$ -(1  $\rightarrow$  3)-linked structures or alternating  $\alpha$ -(1  $\rightarrow$  3)- $\alpha$ -(1  $\rightarrow$  4)-linked L-fucose linkages. On the other hand, heterofucans do not have a defined glycan backbone and can consist of galactose,<sup>46</sup> mannose, or glucuronic acid with fucose branches.<sup>47</sup> The structural diversity of homofucans is increased by the presence of sulfate esters, acetylation, and saccharide modifications like galactose,

glucuronic acid, or xylose.<sup>41,48</sup> Despite the diverse structures within homofucans across brown algae, each species contains distinct motifs. For instance, fucoidan from *Laminaria hyperborea* primarily features  $\alpha$ -(1  $\rightarrow$  3)-linkages, accompanied by smaller quantities of  $\alpha$ -(1  $\rightarrow$  2) and  $\alpha$ -(1  $\rightarrow$  4)-linkages.<sup>43</sup> Presently, defining precise sulfation patterns in fucoidan polysaccharides is technically challenging. However, *Cladosiphon okamuranus* possess a high degree of 4-*O*-sulfation (Figure 1a).<sup>42</sup>

Retrosynthetic analyses identified thioglycoside building blocks 1–5 as suitable candidates for assembling different fucoidan oligosaccharides (Figure 1b). A constraint on the L-fucose building block design is the requirement of non-participating protecting groups at the 2-hydroxyl, to allow for  $\alpha$ -selectivity in glycosylations. Building block 1 is equipped



**Figure 2.** Automated glycan assembly of two major types of fucoidan backbone. (a) Automated assembly of two types of fucoidan backbone,  $\alpha$ -1,3-linked structures and those with alternating  $\alpha$ -1,3/ $\alpha$ -1,4-linked *L*-fucose linkages. (b) Collection of fucoidan oligosaccharides synthesized.

with a nonparticipating benzyl ether on the 2-hydroxyl, a temporary fluorenylmethoxycarbonyl (Fmoc) group on the 3-*O*-position, and the 4-*O*-position was protected with a benzoate ester. The 4-*O*-benzoate ester serves as a long-range participating group (LRP) to support 1,2-*cis*-glycoside formation.<sup>37,49–51</sup> Additionally, it can be cleaved on the resin to enable 4-*O*-sulfation. Building block 2 carries a 4-*O*-levulinate ester (Lev) for the formation of 1  $\rightarrow$  4 linkages and could be utilized for precise 4-*O*-sulfation. Building block 3 bears a nonparticipating 2-naphthylmethyl ether (Nap) protecting group that can be selectively removed to install 1  $\rightarrow$  2 linkages or 2-*O*-sulfate esters. Building block 4 used a Nap ether at the 3-OH, permitting Lev and Fmoc related protecting group manipulations elsewhere in the oligosaccharide. This Nap ether could later be removed to proceed with the backbone synthesis. Finally, building block 5 allowed for the synthesis of galactopyranoside branches.

**Altering the Thioglycoside Leaving Group Improves Automated Glycan Assembly of Fucoidan Oligosaccharides.** Initially, thioethyl glycosides were used for AGA of  $\alpha$ -(1 $\rightarrow$ 3)-homofucans, resulting in significant quantities of deletion sequences and reproducibility problems (SI Table 1,

Figure 1c and d, SI Figure 1). Efforts to enhance the efficiency of the glycosylation by trialling different Lewis acids, varying temperatures, and employing double coupling cycles still produced inconsistent results (SI Table 1, entries 1–4). Considering that fucosyl donors are often highly reactive,<sup>38,52</sup> and coupling temperatures below  $-40$  °C are difficult to adopt at current automated synthesizers,<sup>53</sup> dibutyl glycosyl phosphate donors were tested but did not improve yields (SI Table 1, entries 5 and 6).

Modifying thioglycoside reactivity by changing the protecting groups was impractical due to the structural complexity of the oligosaccharide targets that include branches and sulfate esters. Instead, the thioglycoside aglycon leaving group was modified to regulate glycosyl donor reactivity,<sup>54</sup> which can adjust the activation temperature of thioglycosides by as much as  $+10$  °C.<sup>38</sup> Initially, 4-methylthiophenol thioglycosides were chosen for their availability and low cost, and this was found to enhance both the quality and reproducibility of the glycosylation modules during AGA. Under these conditions, no detectable deletion sequences by HPLC or MALDI-MS were identified (Figure 1d and SI Figure 1, SI Table 1, entries 7 and 8). The optimized glycosylation modules involved *N*-

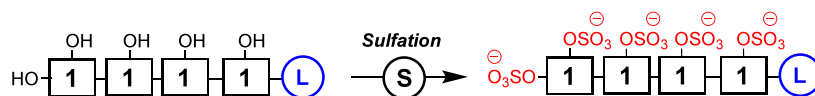


Figure 3. Optimization of on-resin sulfation using tetrasaccharide 13.

Table 1. On Resin Sulfation Optimization

| entry | solvent                      | reagent                           | temperature | time              | comment        | reference |
|-------|------------------------------|-----------------------------------|-------------|-------------------|----------------|-----------|
| 1     | DMF                          | NMe <sub>3</sub> :SO <sub>3</sub> | 90 °C       | 30 min (2 cycles) | incomplete     | 53        |
| 2     | DMF                          | NMe <sub>3</sub> :SO <sub>3</sub> | 90 °C       | 90 min (6 cycles) | incomplete     | 53        |
| 3     | DMF:Py (1:1)                 | Py:SO <sub>3</sub>                | 40 °C       | 12 h              | incomplete     | 58        |
| 4     | DMF                          | Py:SO <sub>3</sub>                | 50 °C       | 16 h              | irreproducible | this work |
| 5     | DMF                          | NEt <sub>3</sub> :SO <sub>3</sub> | 50 °C       | 16 h              | irreproducible | this work |
| 6     | DMF:Py (80:20)               | Py:SO <sub>3</sub>                | 50 °C       | 16 h              | complete       | this work |
| 7     | DMF:NEt <sub>3</sub> (80:20) | NEt <sub>3</sub> :SO <sub>3</sub> | 50 °C       | 16 h              | complete       | this work |

iodosuccinimide (NIS) and triflic acid (TfOH) with a reaction sequence of  $-20\text{ }^{\circ}\text{C}$  for 15 min, followed by  $0\text{ }^{\circ}\text{C}$  for 35 min using five equivalents of the donor (CH<sub>2</sub>Cl<sub>2</sub>/dioxane, 2:1).

Subsequent investigations focused on assessing the stereoselectivity of glycosylations, on-resin methanolysis, photocleavage, sulfation, and hydrogenolysis.

**Automated Synthesis of Fucoidan Oligo- and Polysaccharides.** The chemical synthesis of 1,2-*cis*-glycosides in a stereocontrolled fashion is not a solved problem.<sup>11,29–33</sup> However, long-range remote assistance is useful for the synthesis of 1,2-*cis* glucosides and fucosides.<sup>37,49,50</sup> Leveraging the optimized AGA conditions, a series of  $\alpha$ -(1 $\rightarrow$ 3)-linked fucoidan oligosaccharides, pentamer 7, hexamer 10, octamer 8, and a 20-mer 12, were prepared (Figure 2).

Polystyrene resins were either equipped with a 5-aminopentanol to release glycans with a terminal amine for coupling to microarray surfaces or a “traceless” photocleavable linker,<sup>55</sup> which permits the synthesis of free-reducing end glycosides for enzyme assays.<sup>56</sup> Each coupling cycle consisted of an acidic wash with trimethylsilyl trifluoromethanesulfonate (TMSOTf), followed by NIS-TfOH promoted glycosylation. Subsequently, the resin underwent incubation with a solution of acetic anhydride (Ac<sub>2</sub>O) and methanesulfonic acid (MsOH) to “cap” any unreacted nucleophile (Figure 2a).<sup>57</sup> For  $\alpha$ -(1 $\rightarrow$ 3)-linked fucans, the temporary Fmoc protective group was removed using a piperidine solution (20% in dimethylformamide) to expose the nucleophile for the subsequent coupling cycle. The coupling cycles were reiterated five times for 7, six times for 10, and eight times for 8.

For the synthesis of 20-mer fucan 12, an initial assembly of a 10-mer was undertaken, monitoring the process by cleaving and analyzing a small resin sample via MALDI-MS and analytical HPLC (SI Figure 2a). Subsequently, the synthesis continued to reach the targeted 1,2-*cis*-linked 20-mer (SI Figure 2b and c).

The second major backbone of fucoidan comprises alternating  $\alpha$ -fucopyranosyl-(1 $\rightarrow$ 3)- $\alpha$ -fucopyranosyl-(1 $\rightarrow$ 4) linkages. AGA of these oligosaccharides relied on the iterative use of building blocks 1 and 2, leading to the production of tetramer 15 and hexamer 16. During the synthesis of these mixed linkage oligosaccharides, the Fmoc protecting group removal module was completed by using a 20% triethylamine solution in dimethylformamide. This method was chosen as Lev esters can be sensitive to the treatment with piperidine.<sup>28</sup> Smooth coupling between the axial 4-OH acceptor and

thioglycoside 1 was observed without any reactivity issues (SI Figure 3).

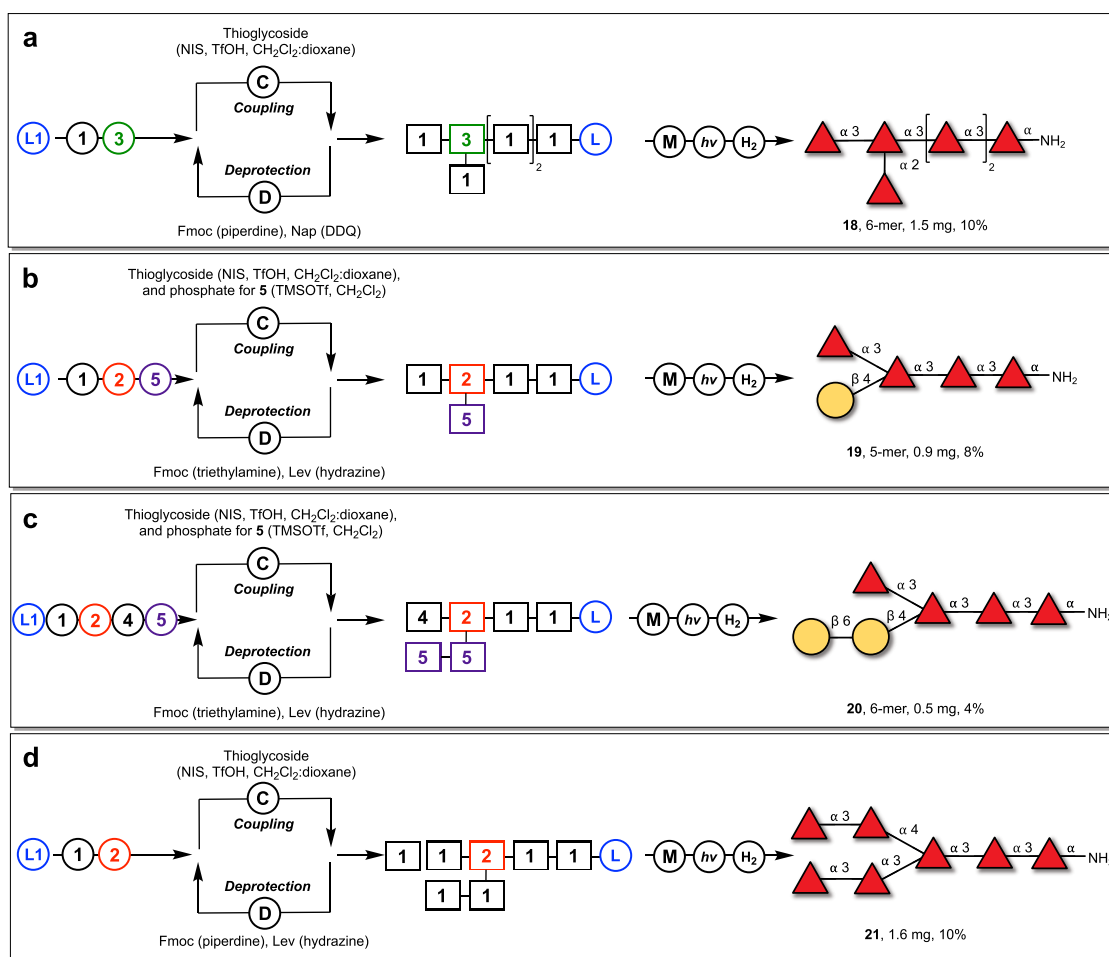
The methanolysis module, employed for removing base-labile protecting groups, was not effective under “standard conditions”, even after extended incubation periods of 168 h (10% 0.5 M NaOMe in anhydrous THF, 5 mL, v/v).<sup>58,59</sup> Utilizing a reduced volume (<5%) of sodium methoxide was necessary for efficiently removing the benzoate esters in under 16 h. Oligosaccharides larger than 10-mers necessitated prolonged incubation periods of 70 h.

Oligosaccharides 6 and 16 were released from the solid support using a flow-based photoreactor,<sup>60</sup> followed by hydrogenolysis using 5% Pd/C in THF/*t*-BuOH/H<sub>2</sub>O (60:10:30, v/v/v).<sup>61,62</sup> The two distinct fucoidan backbones were individually purified via reverse-phase HPLC (Hypercarb, gradient 0 to 80%, acetonitrile/water) to yield hexasaccharides 6 and 16.

NMR analysis confirmed that both fucoidan backbones were prepared with complete  $\alpha$ -selectivity, where the only observed  $\beta$ -linkage was associated with the free-reducing end at 4.57 ppm (d,  $J = 7.9\text{ Hz}$ ) in 6. Comparison of the synthetic fucoidan oligosaccharide NMR spectra to polysaccharides extracted from brown algae showed agreement.<sup>43</sup> Fucoidan oligosaccharides with  $\alpha$ -1,3-backbone shifts occurred at  $\sim 5.06\text{ ppm}$  (<sup>1</sup>H NMR) and 95.5 ppm (<sup>13</sup>C NMR), while the  $\alpha$ -1,4-backbone occurred at  $\sim 4.96\text{ ppm}$  (<sup>1</sup>H NMR) and 100.2 ppm (<sup>13</sup>C NMR).<sup>43</sup> The use of the nucleophilic 5-aminopentanol linker can lead to stereoselectivity issues.<sup>63</sup> However, NMR analysis of compounds 7, 8, 9, and 11–17 contain exclusively 1,2-*cis* linkages in NMR, with an  $\alpha$ -linked anomeric proton at  $\sim 4.88\text{ ppm}$  (<sup>1</sup>H NMR) and  $>98.2\text{ ppm}$  (<sup>13</sup>C NMR).

**Automated Assembly of Sulfated Fucoidan Oligosaccharides.** The precise pattern and degree of fucoidan sulfation depend on a range of factors, including environmental conditions, growth stages, and extraction methods. Moreover, our understanding of how different sulfation patterns impact fucoidan’s biological functions remains limited. This is in contrast to glycosaminoglycans (GAGs), where well-defined oligosaccharides have played a pivotal role, allowing for a detailed molecular-level understanding of the roles that individual sulfate groups play in modulating bioactivity.<sup>64–68</sup>

Sulfated oligosaccharides prepared via AGA to date contain up to four sulfate esters.<sup>58,69</sup> However, to decode glycan–protein interactions with high fidelity, sulfated tetra- to dodecasaccharides are ideal. Therefore, we initially targeted the synthesis of tetrasaccharide 13 with five sulfate esters.



**Figure 4.** AGA of branched fucoidan oligosaccharides. (a) Synthesis of oligosaccharide **18** with a  $\alpha$ -(1 $\rightarrow$ 2)-branch found in *Fucus vesiculosus*. (b) Synthesis of an oligosaccharide **19** with a  $\beta$ -(1 $\rightarrow$ 4)-galactopyranoside branch found in *Saccharina japonica*. (c) Synthesis of oligosaccharide **20** with a  $\beta$ -(1 $\rightarrow$ 6)-gal- $\beta$ -(1 $\rightarrow$ 4)-galactopyranoside found in *Saccharina japonica*. (d) Synthesis of oligosaccharide **21** with a  $\alpha$ -(1 $\rightarrow$ 4)-branch found in *Laminaria hyperborea*.

AGA, followed by methanolysis, provided the tetrasaccharide that was attached to the solid support. Two methods reported for on-resin sulfation did not yield the desired penta-*O*-sulfated compound (Figure 3, Table 1, entries 1–4, SI Figure 4).<sup>53,58</sup> These results may be due to the low nucleophilicity of the axial C4 hydroxyl group vs primary hydroxyl groups,<sup>53,70</sup> or the steric demands involved in placing several sulfate esters in close proximity on the solid support.<sup>58</sup>

Solution-phase syntheses of sulfated glycans, such as GAGs, can necessitate prolonged reaction times,<sup>50,64</sup> that in turn renders sulfation in an automated fashion not always practical (Table 1, entries 1 and 2).<sup>53</sup> While, a recently published on-resin approach, conducted in plastic syringes could not provide the precise atmospheric and temperature control necessary for extended reaction times (Table 1, entry 3).<sup>58</sup> Sealable silanized microwave vials proved ideal for carrying out long sulfation reactions in an aluminum heating block (SI Figure 16, Table 1, entries 4–7).

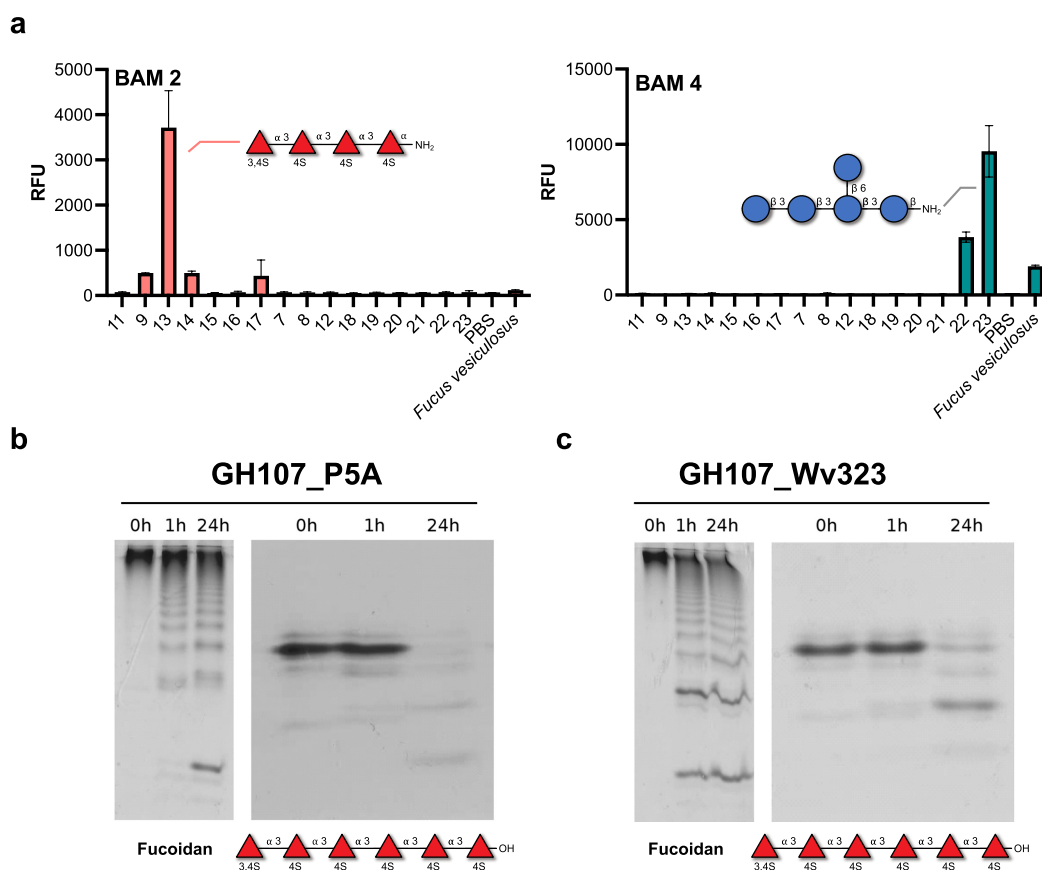
In sealed microwave vials, both sulfur trioxide pyridine (Py-SO<sub>3</sub>) and triethylamine (NEt<sub>3</sub>-SO<sub>3</sub>) complexes yielded comparable results (Table 1, entries 4 and 5). Prebuffering the sulfation solution with an appropriate base,<sup>71</sup> such as pyridine for sulfation reactions using Py-SO<sub>3</sub>, helped reduce the batch-to-batch variability in the quality of sulfur trioxide reagents (Table 1, entries 6 and 7).<sup>72</sup> Even then sometimes,

multiple cycles of the sulfation module were required to achieve full sulfation, for example, hexa- (**10**) and deca-saccharides (**14**).

To monitor solid-phase sulfation reactions, microcleavage must be performed, which releases minute quantities of oligosaccharides for HPLC and MS analysis. Here, we employed dimethylformamide (DMF) as a photocleavage solvent, for its good swelling of polystyrene resins and its ability to solubilize the released sulfated glycans.<sup>73,74</sup> Released glycans were analyzed using quadrupole time-of-flight mass spectrometry (Q-TOF MS, SI Figure 5) and reverse-phase HPLC (CS Luna, 5% ACN to 100), with the HPLC analysis only effective for oligosaccharides with fewer than six sulfate esters (SI Figure 4).

Using these conditions (see SI, modules h1 and h2), solid-phase sulfation from a mono- to a deca-saccharide was completed, with oligosaccharide **14** containing 11 sulfate esters (**9–11**, **13**, **14**, and **17**). <sup>1</sup>H–<sup>13</sup>C HSQC NMR analysis confirmed the sulfation of the 4-OH, with the carbon atoms shifting characteristically downfield compared to nonsulfated fucoidan oligosaccharides from 68 to 79 ppm.

**Photocleavage Using a LED Lamp Allows for Parallel Cleavage of Multiple Resins.** Reported approaches for cleaving sulfated oligosaccharides from the solid support involve a mercury lamp flow-reactor setup, utilizing a



**Figure 5.** Synthetic fucoidans as tools for marine glycobiology. (a) Mapping the reactivity of BAM monoclonal antibodies with a fucoidan microarray. Shown is the 1:50 antibody dilution. (b) Left gel shows the activity of GH107\_P5A on fucoidan from *L. hyperborea* by CPAGE. Right gel demonstrates the activity of GH107\_P5A on synthetic  $\alpha$ -1,3 fucan oligosaccharide **10**. (c) Left gel is the activity of GH107\_P5A on fucoidan from *L. hyperborea* by CPAGE. Right gel shows the activity of GH107\_Wv323 on synthetic  $\alpha$ -1,3 fucan oligosaccharide **10**. Enzyme incubations were complete at 1  $\mu$ M for 0, 1, and 24 h with each lane containing  $\sim$ 4  $\mu$ g of the initial substrate. The products resulting from enzymatic degradation are separated according to the size and degree of sulfation and visualized with Stains-All.

DCM–methanol mixture.<sup>58,60</sup> This approach requires multiple passages through the flow cell, which are required to achieve good photocleavage results;<sup>60</sup> this may be due to poor resin-swelling properties of methanol.<sup>58,60</sup> Similar results were obtained when sulfated oligosaccharide **10** was cleaved from the solid support using photolysis. Therefore, we alternatively employed an LED lamp (370 nm) with DMF as a photocleavage solvent.<sup>75,76</sup> DMF was chosen as it can solubilize the released amphiphilic sulfated glycans due to its good resin-swelling properties.<sup>73,74</sup> The LED lamp can facilitate the parallel cleavage of multiple resins (SI Figure 6).

Following the release of the oligosaccharides from the solid support, the crude material was subjected to hydrogenolysis (5% Pd/C, THF/*t*-BuOH/ $H_2O$ , 50:20:30, v/v/v), and the sulfated glycans were purified using RP-HPLC, size-exclusion chromatography, or a combination of both methods. The choice of purification method depended on the number of sulfate esters on the oligosaccharide. Glycans containing more than five sulfate esters were best purified using a combination of size-exclusion chromatography and HPLC. Using this approach, a series of sulfated fucans (**9–11**, **13**, **14**, and **17**) with different backbones, lengths, and sulfation patterns was prepared (Figure 2b).

**Automated Assembly of Branched Fucoidan Oligosaccharides.** Brown algae synthesize a diverse range of fucoidans with distinct branching and sulfation patterns. The

utility of the AGA platform to synthesize such branched fucoidans was demonstrated by four examples. These glycans contain  $\alpha$  and  $\beta$  branching residues and cover the branching patterns found in fucoidan (2-OH, 3-OH, and 4-OH).

$\alpha$ -(1 $\rightarrow$ 2)-Fucopyranoside branches occur in various brown algae species, including *Cladosiphon okamuranus* and *Laminaria hyperborea* (Figure 1a). Therefore, we prepared hexasaccharide **18** that contains this motif (Figure 4a).<sup>48</sup> Two cycles of building block 3 were required to fully convert the acceptor trisaccharide to the desired tetrasaccharide (SI Figure 11). Selective oxidative cleavage of the Nap group facilitated regioselective glycosylation of the 2-OH acceptor (SI Figure 12). Subsequently, the Fmoc removal and glycosylation produced the desired protected oligosaccharide intermediate, with HPLC displaying a major peak at 20 min (SI Figure 13). Following methanolysis, the presence of the semideprotected hexasaccharide was confirmed by MALDI-TOF (SI Figure 14). Photocleavage released the oligosaccharide from the resin; following hydrogenolysis and HPLC purification, 1.5 mg (10%) of  $\alpha$ -fucan **18** was isolated. NMR analysis of **18** revealed a distinct upfield chemical shift at 5.33 ppm (d,  $J$  = 3.8 Hz, 1H), previously annotated for  $\alpha$ -(1 $\rightarrow$ 2)-linkages in *Laminaria hyperborea*,<sup>43</sup> therefore, the synthetic oligosaccharide supported the assigned structure.

Subsequently, two *Saccharina japonica* motifs were synthesized, one featuring a  $\beta$ -(1 $\rightarrow$ 4)-galactopyranoside branch, and

the other presenting a more extended gal- $\beta$ -(1 $\rightarrow$ 6)-gal- $\beta$ -(1 $\rightarrow$ 4) branching pattern.<sup>77</sup> AGA of pentasaccharide **19** relied on building blocks **1** and **2** to construct the tetrasaccharide backbone (Figure 4b). Following the removal of the 4-*O*-Lev ester, galactosylation was completed using the dibutyl phosphate donor **5** (TMSOTf,  $-35$  °C for 5 min  $\rightarrow$   $-20$  °C for 30 min, 5.5 equiv).<sup>78</sup> HPLC analysis revealed a major peak at 32 min for the pentasaccharide (SI Figure 9). To assemble glycan **20** with the gal- $\beta$ -(1 $\rightarrow$ 6)-gal- $\beta$ -(1 $\rightarrow$ 4)-branch, AGA utilized building blocks **1** and **2**, alongside **4**, containing a 3-Nap ether, permitting future extension of the fucan backbone following the assembly of the galactose chain (Figure 4c). HPLC analysis of the hexasaccharide displayed a major peak at 30.52 min (SI Figure 10). Following methanolysis, photocleavage, hydrogenolysis, and HPLC purification yielded glycans **19** (0.9 mg, 8%) and **20** (0.5 mg, 4%). NMR analysis of **19** showed a  $\beta$ -(1 $\rightarrow$ 4)-linkage at 4.52 ppm (d,  $J$  = 7.3 Hz, 1H), while **20** contained an additional  $\beta$ -linkage at 4.32 ppm (SI Table 2).

Heptasaccharide **21** contains an  $\alpha$ -(1 $\rightarrow$ 4)-fucopyranosyl branch found in *Laminaria hyperborea* (Figure 1a, Figure 4d).<sup>43</sup> This oligosaccharide was assembled using building blocks **1** and **2**, with the Lev ester on **2** allowing the synthesis of the  $\alpha$ -(1 $\rightarrow$ 4)-branch. Microcleavage analysis at the pentasaccharide stage revealed a single major peak at 22.2 min in the HPLC (SI Figure 7). Subsequently, the two Fmoc protecting groups were removed, and two coupling cycles of **1** produced the protected branched oligosaccharide (SI Figure 8). Methanolysis prior to photolytic release from the solid-phase was followed by hydrogenolysis. Reverse-phase HPLC (Hypercarb, 0 to 80 H<sub>2</sub>O) yielded 1.6 mg (10%) of the  $\alpha$ -(1 $\rightarrow$ 4)-containing fucan **21**. NMR analysis of **21** revealed a 1,2-*cis*-linked oligosaccharide, with the anomeric carbon of the  $\alpha$ -(1 $\rightarrow$ 4)-linkage occurring at 5.15 ppm (d,  $J$  = 4.0 Hz, 1H), downfield of the  $\alpha$ -(1 $\rightarrow$ 3)-backbone linkages (SI Table 2).<sup>43</sup>

**Synthetic Glycan Defines the Activity of Two *endo*-Fucoidan Hydrolases.** The microbial degradation of fucoidan involves hundreds of enzymes.<sup>5,79</sup> Enhancing our understanding of fucoidan-active CAZymes and their substrate tolerances will advance our mechanistic understanding of how certain fucoidan structures resist degradation, leading to carbon sequestration. Furthermore, characterized enzymes can serve as biocatalytic assays to assist in the detection and quantification of fucoidan in the environment.<sup>80,81</sup>

CAZymes of the glycoside hydrolase family 107 (GH107) cleave in midchain glycosidic bonds of algal fucoidans.<sup>82</sup> All current GH107 members are *endo*-fucoidanases targeting either  $\alpha$ -1,3 or  $\alpha$ -1,4 fucosyl linkages. Several  $\alpha$ -1,4-*endo*-fucoidanases have been functionally validated, e.g., MfFcnA from *Mariniflexile fucanivorans* and MefI from *Allomuricauda eckloniae*, for which also protein structures were obtained.<sup>83,84</sup> On the other hand, only one  $\alpha$ -1,3-*endo*-fucoidanase has been characterized.<sup>85</sup>

GH107\_PSAFcnA from *Psychromonas* sp. SWSA displays activity against fucoidan from *Laminaria hyperborea*—a fucoidan consisting predominantly of sulfated  $\alpha$ -1,3-linked fucan—while it is inactive on fucoidans with alternating  $\alpha$ -1,3/ $\alpha$ -1,4-linked fucan backbone.<sup>43,83</sup> This suggests that GH107\_PSAFcnA is an  $\alpha$ -1,3-*endo*fucoidanase but requires functional validation. Therefore, we used the synthetic  $\alpha$ -(1 $\rightarrow$ 3) fucan oligosaccharide **10** to test the activity of GH107\_PSAFcnA. Recombinant, purified GH107\_PSAFcnA was obtained as previously described,<sup>83</sup> and the enzyme was

incubated with fucoidan and oligo **10**. Enzyme activity and product formation were assayed over time by CPAGE (Figure 5b). The results show that GH107\_PSAFcnA degrades both fucoidan from *L. hyperborea* as well as oligo **10** and confirms that the enzyme cleaves  $\alpha$ -1,3-linked sulfated fucan. Next, we used the protein sequence of GH107\_PSAFcnA to search for homologue enzymes at National Center for Biotechnology Information (NCBI). A putative GH107 (WP\_179351272) from the marine flavobacterium *Winogradskyella vidalii* showed 59% identity (>90% coverage) with GH107\_PSAFcnA. Genomic studies have linked *Winogradskyella* spp. to fucoidan utilization, but so far this has not been biochemically verified.<sup>86</sup> We found that pure recombinant GH107 from *W. vidalii* displayed similar activity as GH107\_PSAFcnA against fucoidan and  $\alpha$ -1,3-linked sulfated fucan oligosaccharide (Figure 5c). As such, both enzymes are  $\alpha$ -1,3-*endo*-fucoidanases that are able to initiate the degradation of fucoidan derived from *L. hyperborea*.

According to the CPAGE results, both GH107s show activity against fucoidan after 1 h, whereas longer incubation time is needed to degrade the oligosaccharide. This suggests that the enzymes prefer substrates with longer glycan chains or different sulfation patterns than **10**—structures that seem to occur in the native fucoidan. For example, fucoidan from *L. hyperborea* can be C2 sulfated in addition to C4.<sup>43</sup> Nevertheless, we demonstrate here the  $\alpha$ -(1 $\rightarrow$ 3)-fucan specificity of two GH107 fucoidanases derived from marine bacteria. These results demonstrate the utility of synthetic oligosaccharides in the discovery and characterization of fucoidan-degrading enzymes.

**Glycan Microarrays Map Specificity of Fucoidan-Directed Antibodies.** Understanding the relationship between the structure of fucoidans and their functional properties is challenging due to the heterogeneity of polysaccharides extracted from algae. Therefore, we constructed a glycan microarray to investigate the binding of protein to our synthetic fucoidan library. Amine-functionalized fucoidan oligosaccharides along with two control  $\beta$ -(1 $\rightarrow$ 3)-glucans, **22** and **23**, were covalently attached to *N*-hydroxylsuccinimide (NHS)-functionalized glass slides. Each glycan was printed in quadruplicate at a concentration of 100  $\mu$ M using a robotic printer.

The binding specificity of four monoclonal antibodies (mAbs) targeting fucoidan, BAM1–BAM4, was investigated. These antibodies allow the visualization of algae and diatom cell walls.<sup>3,4,12</sup> Furthermore, they aid in the environmental detection and quantification of fucoidan in seawater and sediments.<sup>3,4,7</sup> A limitation of these antibodies is that their epitopes are not precisely defined.<sup>12</sup>

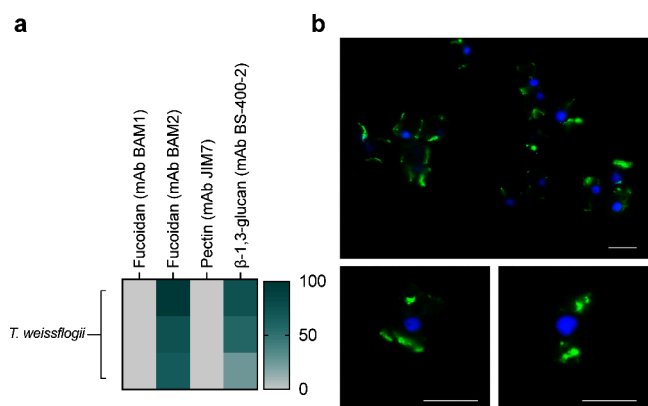
Microarray analysis of BAM1 and BAM3 revealed no binding to any fucoidan oligosaccharides on the array, suggesting interactions with structural epitopes not present in the current library (Figure 5a). mAb BAM4 demonstrated no binding to any fucoidan structures on the array but did display reproducible binding to a  $\beta$ -(1 $\rightarrow$ 3)-glucan tetrasaccharide **22** and **23**  $\beta$ -(1 $\rightarrow$ 3)-glucan tetrasaccharide with a central  $\beta$ -(1 $\rightarrow$ 6)-branch (Figure 5a; SI Figure 15).

In the case of mAb BAM2, binding was observed to sulfated fucoidan oligosaccharides **11**, **9**, **13**, **14**, and **17** (Figure 5a). Low level binding to glycan **11**, a monosaccharide with a di-3,4-*O*-sulfation pattern, was only observed at a higher concentration of mAb BAM2, suggesting that this di-*O*-sulfated monosaccharide poorly represents the BAM2 epitope.

While binding to  $\alpha$ -(1 $\rightarrow$ 3)-fucoidan oligosaccharides (9, 13, and 14) with 4-*O*-sulfate esters and a terminal di-3,4-*O*-sulfation pattern provided more robust binding across a range of concentrations (SI Figure 15). mAb BAM2 also bound weakly to sulfated fucoidan oligosaccharide 17, with an  $\alpha$ -(1 $\rightarrow$ 3)- $\alpha$ -(1 $\rightarrow$ 4)-linked backbone. Collectively, this suggests that these 4-*O*-sulfated oligosaccharides contain the BAM2 epitope.<sup>14,86</sup>

**Binding of BAM2 to *Thalassiosira weissflogii* Suggests the Diatom Synthesizes a Fucoidan Structurally Similar to Those in Brown Algae.** The formation of sinking particles in the ocean promotes carbon sequestration, and microalgal polysaccharides are involved in this process. Recent findings, utilizing the fucoidan-specific monoclonal antibodies BAM1 and BAM2, have revealed that diatoms *Chaetoceros spp.* and *Thalassiosira weissflogii* produce fucose-containing sulfated polysaccharides (FCSP). These polysaccharides form particles that promote aggregation, sinking, and, consequently, carbon sequestration.<sup>4,87</sup> FCSP is a broad term used to imply the presence of fucoidan-like structures but does not refer to a particular structure.

Analysis of polysaccharide extracts from the diatom *Thalassiosira weissflogii* using microarrays suggested the presence of distinct fucoidan that was reactive to BAM2 but not BAM1 (Figure 6a),<sup>87</sup> implying that diatom species



**Figure 6.** Binding of mAb BAM2 detects that *Thalassiosira weissflogii* synthesizes fucoidan. (a) *T. weissflogii* water extracts were printed onto microarrays, and those were probed with a panel of monoclonal antibodies (mAbs), each row represents a replicate. In addition to the antifucoidan BAM1 and BAM2, JIM7 (antipectin) and BS-400-2 (anti- $\beta$ -1,3-glucan) were included as negative and positive controls, respectively. The heatmap shows normalized mAb binding intensity in the extraction triplicates. (b) Representative images showing that mAb BAM2 binds to *T. weissflogii* cells. Fluorescence microscopy images demonstrate that the BAM2 epitope is present on the diatom cell surfaces.  $\alpha$ -1,3-Linked fucoidan (green), DAPI (blue). Scale bars: 10  $\mu$ m. Experiments were performed three times with similar results.

synthesize diverse fucoidan structures. In this study, we mapped the specificity of the mAb BAM2, and hypothesized that this means *T. weissflogii* produces a fucoidan with either  $\alpha$ -(1 $\rightarrow$ 3)-linked fucose or alternating  $\alpha$ -(1 $\rightarrow$ 3)- $\alpha$ -(1 $\rightarrow$ 4)-linked fucose and 4-*O*-sulfation. Microscopy analysis of diatom cells post-roller tank experiments, which were employed to induce aggregation, revealed the presence of the BAM2 fucoidan epitope surrounding the diatom cell aggregates.<sup>87</sup> Furthermore, here we demonstrate that individual diatom cells produce this

fucoidan epitope and present it on their cell surfaces (Figure 6b).

The presence of structures known for their carbon sequestration capacity, within a globally distributed diatom suggests that the synthesis of molecules is more common than previously assumed. Detailed analysis of the glycans present in marine ecosystems will allow for a deeper understanding of the marine carbon cycle. In this process, structurally defined synthetic oligosaccharides served as a missing link in various existing tools, including enzymatic, immunological, and spectroscopic methods.

## CONCLUSIONS

Automated glycan assembly provides access to fucoidan oligosaccharides, reaching lengths of up to 20-mers, with diverse branching patterns, and up to 11 sulfate esters. Modulating the reactivity of building blocks, by altering the thioglycoside aglycon from an alkyl to a less reactive aryl group, enabled the synthesis of the oligosaccharides. NMR experiments confirmed that these synthetic fucoidan oligosaccharides contain structural features that are found in brown algae. A synthetic oligosaccharide also enabled the characterization of two GH107 *endo*-fucoidanases from marine bacteria, with both enzymes capable of degrading  $\alpha$ -(1 $\rightarrow$ 3)-linked fucoidan sulfated oligosaccharides. A fucoidan microarray was used to map the specificity of four monoclonal antibodies (mAbs) directed toward fucoidan. The mAb BAM4 was discovered to have cross-reactivity toward  $\beta$ -(1 $\rightarrow$ 3)-glucans, both with and without  $\beta$ -(1 $\rightarrow$ 6)-glucose branches. The mAb BAM2 bound to fucoidan oligosaccharides with  $\alpha$ -(1 $\rightarrow$ 3)- and  $\alpha$ -(1 $\rightarrow$ 3)- $\alpha$ -(1 $\rightarrow$ 4)-linkages bearing 4-*O*-sulfate esters. This molecular understanding of mAb BAM2 specificity provides evidence that *Thalassiosira weissflogii*, a globally abundant diatom, synthesizes fucoidan structurally similar to that found in brown algae. Synthetic fucoidan oligosaccharides are tools to investigate fucoidan's roles in carbon sequestration and medicine.

## ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/jacs.4c02348>.

Supporting figures, tables, experimental procedures, and spectral data for all reactions and compounds (PDF)

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