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Leila Mousavifar, Gerard Vergoten, Guillaume Charron, René Roy. Comparative Study of Aryl O-, C-, and S-Mannopyranosides as Potential Adhesion Inhibitors toward Uropathogenic *E. coli* FimH. *Molecules*, 2019, 24 (19), pp.3566. 10.3390/molecules24193566 . hal-03095722

HAL Id: hal-03095722

<https://hal.univ-lille.fr/hal-03095722v1>

Submitted on 8 Feb 2021

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Article

Comparative Study of Aryl *O*-, *C*-, and *S*-Mannopyranosides as Potential Adhesion Inhibitors toward Uropathogenic *E. coli* FimH

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Academic Editors: Diego Muñoz-Torrero, F. Javier Luque and Marçal Pastor-Anglada

Received: 9 September 2019; Accepted: 1 October 2019; Published: 2 October 2019



Abstract: A set of three mannopyranoside possessing identical 1,1'-biphenyl glycosidic pharmacophore but different aglyconic atoms were synthesized using either a palladium-catalyzed Heck cross coupling reaction or a metathesis reaction between their corresponding allylic glycoside derivatives. Their X-ray structures, together with their calculated 3D structures, showed strong indicators to explain the observed relative binding abilities against *E. coli* FimH as measured by a improved surface plasmon resonance (SPR) method. Amongst the *O*-, *C*-, and *S*-linked analogs, the *C*-linked analog showed the best ability to become a lead candidate as antagonist against uropathogenic *E. coli* with a K_d of 11.45 nM.

Keywords: carbohydrate; D-mannosides; uropathogenic infections; *E. coli*; lectin; FimH; Heck reaction; metathesis; SPR; X-ray

1. Introduction

Multidrug resistant bacteria constitutes a serious public health burden with approximately 700,000 people dying each year from infections caused by antibiotic resistant organisms. This number is expected to reach 10 million people annually by 2050 with related cost ranging \$100 trillion US dollars [1]. Although medicinal chemistry has allowed the relatively fast discoveries of diverse families of potent antibiotics working under a wide range of bactericidal or bacteriostatic mechanisms [2], bacteria have similarly developed a variety of ingenious defense mechanisms [3]. Therefore, there is an urgent need to fight bacterial infections by mechanisms that alleviate multidrug resistances. One of these approaches, showing increasing successes, is directed at preventing the early step of bacterial adhesion and colonization, and biofilm formation pathways [4]. Since several infection mechanisms are initiated by molecular recognition between the host cell surface receptors and virulence factors on the bacterial surfaces [5], such strategies are likely to be unavoidable without impeding the bacteria's own capacity to recognize and adhere to the targeted tissues. Urinary tract infections (UTIs) represent one of the most common bacterial infections, affecting one out of two women during their lifetimes. UTIs are serious health problems as they reach close to \$2.5 billion in annual healthcare costs in the US alone [6] which are caused by Gram-negative uropathogenic *Escherichia coli* (UPEC) strains [7].

UPEC are the major cause of urinary tract infections and intestinal bowel diseases (IBD) including Crohn's disease (CD) [1,6]. These strains enter urinary tracts to form colonies. The adhesions of

various pathogenic *E. coli* strains to host cells are primarily mediated through carbohydrate-protein interactions involving bacterial fimbriae which can recognize specific glycoconjugate receptors on host cells. Of particular interests are the FimH and the PapG fimbriae that bind to mannosylated glycoproteins or glycolipids, respectively [8].

The FimH bacterial lectins adhere to the highly mannosylated glycoprotein uroplakin 1a [9,10] on the bladder cells and on glycoprotein 2 (GP2) exclusively expressed in the small intestine [7,11]. Once the infection process is initiated, it is usually followed by thick biofilm formation [12,13]. Biofilms have a pronounced impact in clinical settings because they render bacterial infections resilient to antibiotics [14]. Given that bacterial adhesion to host cells is the preliminary step toward the release of toxic proteins, the design of potent *E. coli* FimH antagonists has been the target of several efforts [15,16]. Several glycomimetic analogs of the naturally occurring complex oligomannosides have been synthesized [8,10] that provided potent α -D-mannopyranoside-based antiadhesins. Since small molecule antagonists represent the foremost choice of the pharmaceutical industry, these glycomimetics constitute most promising candidates for the replacement of the natural lead oligomannosides onto which *E. coli* adhere to [10,17].

To circumscribe the effect of anomeric linkages upon the design of uropathogenic *E. coli* FimH antagonists based on mannopyranosides, we compare herein three candidates possessing aromatic aglycones capable to fit within the well-established hydrophobic tyrosine gate present in the mannopyranoside binding pocket. Hence, the tridimensional structures, relative binding affinities of *O*-, *C*-, and *S*-linked mannopyranosides harboring a common 1,1'-biphenyl pharmacophore are described.

2. Results and Discussion

2.1. Glycomimetic Synthesis

Since *O*-substituted mannopyranosides are susceptible to hydrolysis by mannosidases and low pHs of the gut, there is a need to identify alternative candidates suitable under in vivo conditions. *N*- [13], *C*- [10,18,19], or *S*-linked [20] glycomimetics have the potential to possess the required stabilities and appropriate pharmacokinetic and pharmacodynamic parameters.

Analogues of *O*- and *C*-linked α -D-mannopyranosides with hydrophobic and aryl substituents have already been identified as potent *E. coli* FimH antagonists having low nanomolar Kds or IC₅₀s [5,10,18,21,22]. *Para*-substituted biphenyl derivatives were shown to be particularly appealing owing to their numerous favorable binding interactions within the identified tyrosine gate formed between Tyr48 and Tyr137 [18,21,23–25]. In turn, some derivatives were shown to bind better when the Tyr-gate is either open, half-open, or close, further pointing toward rationally design glycol therapeutics [20,22,26–28]. In fact, using the centroid positions of each of the Tyr48 and Tyr137 phenyl groups in crystal data sets, we calculated that the Tyr-gates are 8.77 Å, 7.79 Å, and 6.99 Å apart in the open (PDB 4AV0) [29], half-open (PDB 4BUQ) [29], and close states (PDB 4AUU) [30], respectively. Since this family of α -D-mannopyranoside derivatives showed high promises as antagonists, we decided to further evaluate their comparative 3-D structures, relative binding capacities, and biophysical characteristics, by exploring the roles played by substituting the anomeric linkages with heteroatoms, while keeping constant the *para*-biphenyl moieties. Toward this goal, we first accessed a series of α -D-mannopyranosides 1–6 (Figure 1) known to serve as comparative standards. Compounds 1 (Me α Man) and 2 (PNP α Man) are commercial products, while 3 (Heptyl α Man), the golden standard with a reported Kd of 5 nM (SPR), together with compounds 4 (see SI) were prepared by literature procedures. The 1, 1'-biphenyl derivatives with an anomeric *O*-linkage (5) and the corresponding *C*-linked analog (6), albeit with one atom shorter, were also prepared by published procedures [18]. Note that we successfully crystallized compounds 5 and 6 for the first time (see discussion below).

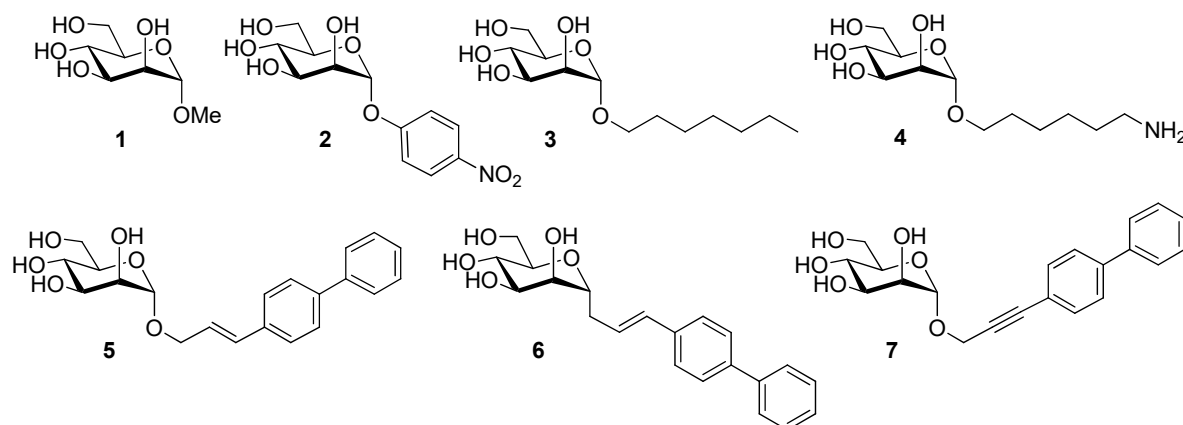
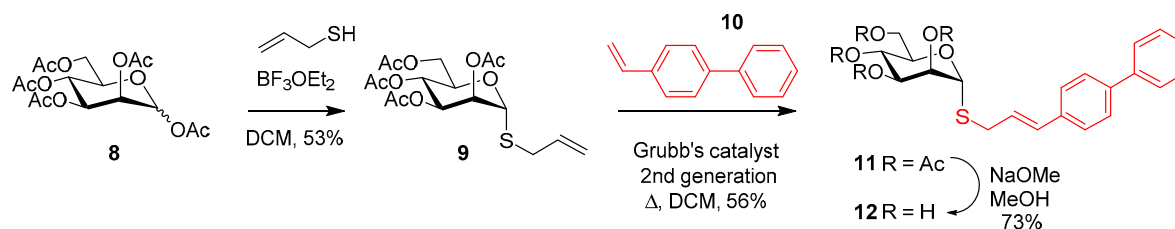


Figure 1. Structures of the known α -D-mannopyranoside *E. coli* FimH antagonists.

We describe herein the synthesis of the related *S*-linked α -D-mannopyranoside (**12**) (Scheme 1). Similarly to its *O*- and *C*-linked congeners, we chose the *S*-allyl α -D-mannopyranoside **9** as key starting materials. However, rather than using a palladium catalyzed Heck reaction and a series of aryl iodides for the cross-coupling as before [18,21], we decided to extend its alkenyl functionality by a metathesis reaction using 4-vinyl-1,1'-biphenyl (**10**) and Grubbs 2nd generation catalyst. This alternative strategy was considered equally appealing owing to its potential to access a diversified library of analogs. In this way, we can demonstrate the versatility of these two complementary chemical strategies leading to these important families of *O*-, *C*-, and *S*-linked derivatives.



Scheme 1. Synthetic scheme leading to the *S*-linked mannopyranoside analog **12** using a metathesis reaction.

Eventhough the necessary intermediate **9** was claimed to be prepared in a single step from D-mannose [31], the procedure failed in our hands to provide reproducible and high yields. Hence, we relied on the more classical Ferrier's conditions [32–34] from peracetylated D-mannose **8** (Scheme 1) which, when treated with prop-2-ene-1-thiol (allyl mercaptan) and BF₃·Et₂O in DCM, provided the α -anomer **9** in 53% yield after purification. All the physical data agreed with the published one [34], thus providing unambiguous evidence for an α -isomer. When **9** was further treated with 4-vinyl-1,1'-biphenyl (**10**) and Grubbs 2nd generation catalyst under refluxing DCM [24,35,36], the desired *trans* analog **11** was obtained in 56% yield after silica gel purification. Classical Zemplén deprotection (NaOMe, MeOH) afforded **12** quantitatively.

2.2. Structural and Conformational Analyses

Having secured entry to our three heteroatom-linked α -D-mannopyranosides **5**, **6**, and **12**, we evaluated their detailed 3D structures as it was anticipated that the relative orientation of their *p*-biphenyl pharmacophores with the *E. coli* FimH binding site would be determinant for their relative binding abilities. This is particularly true given the difference in the width of the Tyr gate in the open, half-open, and close states, as mentioned above and that they may individually bind to one form or another unpredictably. Fortunately, we accessed to two crystalline structures, namely, those of the *O*- and *C*-linked analogues **5** and **6**, respectively. In addition, we previously resolved fourteen X-ray

co-crystal data of several *O*-linked analogues bound to the *E. coli* FimH, thus giving us access to several docking opportunities [10,18,21,29,30]. One of them in particular (7) is of interest (PDB 4AV5) [29] as it also contains a 1,1'-biphenyl ring, albeit with a more rigid alkynyl linkage (see Figure S37 in SI for superimposed figure containing 5 and 7). Interestingly, while 7 showed a K_d of 405 nM, 5 was 135 fold better with a K_d of 3 nM (SPR) [18], again demonstrating the importance of pharmacophore orientation within the tyrosine gate. Figure S37 (SI) clearly shows compound 5 forming a closer contact with Tyr137 in the open form. Its docking within the FimH active site was done by superimposition of the mannoside residues with that of known 7.

The crystalline structure of 5 alone is shown in Figure 2. Also of interest is the fact that the two biphenyl rings are not tilted with respect to one another. This is in strike contrast with the situation observed from other *p*-biphenyl candidates including our own above (7, PDB 4AV5) [28,29,37]. In addition; it is also noteworthy that the conformation around the C5-C6 bond in the mannose ring is in the *gg* orientation, as it is the case of most crystal structures of mannosides within the binding pocket of FimH.

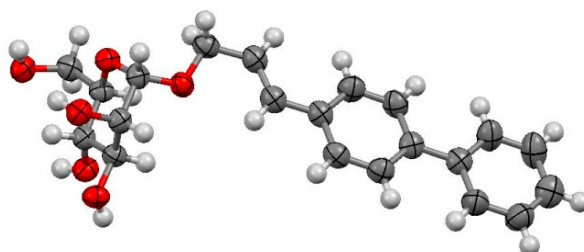


Figure 2. ORTEP diagram for the X-Ray structure of *O*-linked mannoside 5.

The *C*-linked analogue 6, having a C-O-C bond shorter, deserves some specific comments on its own because of the controversial arguments about the preferred conformational states of this family of glycomimetics. Indeed, several literature cases argued for the fact that *C*-linked α -D-mannopyranosides do not exist preferentially in the “usual” 4C_1 chair conformations, at least for simple aglycones [10,38–41]. As already shown by us with *O*-linked allyl α -D-mannopyranoside [10], the 4C_1 conformation was the most stable form with the inverted 1C_4 chair being next in line with a difference of 8.10 kcal/mole difference. To see if this observation still holds for compound 6, we undertook a conformational search using the Monte Carlo method [42]. The retained structure was then optimized using molecular mechanics [43–45] and the density functional theory (DFT) with the hybrid functional B3LYP, base 6-31G* (see SI info). The result clearly showed, once again that this additional *C*-linked α -D-mannopyranoside 6 also predominates in the desired 4C_1 conformation.

Fortuitously again, we were able to crystallize compound 6 alone as they appear in Figure 3 as 6A and 6B. However, the situation was complicated by the fact that the crystal lattice showed two sugar residues stacking to one another (Figure 3, 6A and 6B). Apparently, both biphenyl moieties were forming strong CH- π interactions [10,46,47], and likely reinforced in that state since the C3-OH group (6A) was hydrogen bonded to the C6-O (6B, 3.914 Å). In addition, one sugar (6A) had the C5-C6 bond in the *tg* conformation while 6B was in the *gg* state. Moreover, both of the *p*-substituted biphenyl rings were flat, as opposed to most observed cases discussed above [23,24] when the sugars are bound to the FimH. This is also seen in the calculated conformer 4C_1 (Figure 3, 6C) where the two rings are tilted by -56.0° in close agreement to those observed for PDB 3MCY [23] (-36.21°) and PDB 5F2F [23] (-37.47°) [24]. Additionally, the C5-C6 bond was in the *tg* conformation.

Clearly, for the situation seen when 6 crystallized alone, the self-assembly into a dimeric form becomes a dominant factor which could not be seen in either the calculated values nor with the FimH binding site. Other significant differences were observed between the three forms denoted 6A, 6B, and 6C for the crystalline states and calculated one, respectively: $\Phi A = -35.84^\circ$, $\Phi B = -54.84^\circ$, $\Phi C = +54.48^\circ$; $\Psi A = -54.50^\circ$; $\Psi B = +65.63^\circ$, $\Psi C = -91.69^\circ$, wherein the Φ torsion angle is define between H1-C1-C1'-C2' and the Ψ torsion angle is define as C1-C1'-C2'-C2'H.

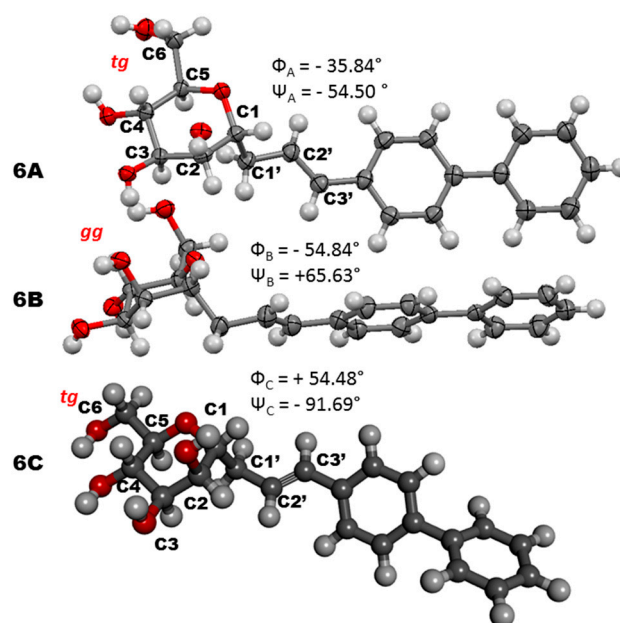


Figure 3. ORTEP diagram for the X-Ray structure of C-linked mannoside **6** as it crystallized in self-assembled stacked dimeric forms (**6A** and **6B**) and as obtained (**6C**) through molecular modelling (see above).

The fact that FimH is known to adopt three different conformations at the tyrosine gate (Tyr48-Tyr137; *open*, *close*, *half open*) depending to which ligands it is bound to, constitutes an additional example of molecular adaptation that has been previously observed with C-linked oligomannoside analogues [47,48], a process of conformational selection suggesting that the FimH likely binds these glycomimetics similarly to the way that it binds the natural analogues [24,49].

Contrary to the above situation for the *O*- and *C*-linked analogues **5** and **6**, we could not obtain a crystalline structure for the *S*-linked analogue **12**. However, it is well known that thioglycosides also obey to the *endo*- as well as the *exo*-anomeric effects [50]. Therefore; we undertook to perform the above Monte Carlo and DFT molecular dynamic calculation on **12** to glance at its most likely 3D structure. As anticipated, the C1-S and the S-C1' bonds length were 1.862 and 1.853 Å, respectively, longer than the equivalent C1-O and O-C1' bonds lengths in **5** (1.399 Å, 1.496 Å). Moreover, the Φ was calculated to be -45.93° in good agreement with the *exo*-anomeric effect for **5** ($\Phi = -57.69^\circ$) (Figure 4).

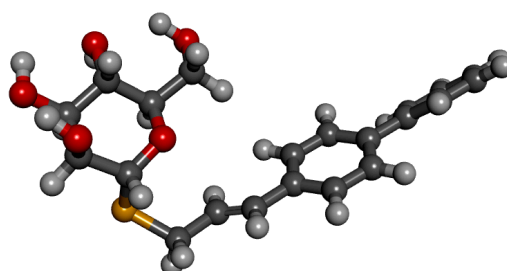


Figure 4. 3D-Structure of the minimum energy level of the 4C_1 chair conformation of **12** as calculated by Monte Carlo molecular mechanics followed by DFT determination with the hybrid functional B3LYP, base 6-31G*.

2.3. Relative FimH Binding Affinity

It has been previously observed that the analogous *O*-, *C*- and *S*-linked heptyl α -D-mannopyranosides had IC_{50} s in the low nanomolar range when measured by surface plasmon resonance (SPR) against the *E. coli* FimH [20]. However, the order was *O*- ($IC_{50} = 160$ nM), *C*- ($IC_{50} = 424$

nM), and the S-linked ($IC_{50} = 547$ nM), thus showing that the O-linked analogue was still unsurpassed and that the nature of the aglyconic heteroatoms had a critical effect on the binding affinities.

To evaluate the relative binding affinities of our own series of compounds **5**, **6**, and **12**, we undertook surface plasmon resonance (SPR) measurements using an improved competition method. To this end, we first immobilized 6-aminohexyl α -D-mannopyranoside (**4**) on the CM5 gold chip using the conventional carbodiimide method to which we allowed FimH to bind with and without our competitive inhibitors (see sensorgram of **4** in the SI). The equilibrium dissociation constant (Kd) of the interaction was then normalized, the data sets are illustrated in Figure 5 and values given in Table 1 (see experimental details). This modification from the previous protocol [18] using 8-aminooctyl α -D-mannoside was found much more practical and reliable given its very high affinity to FimH which made the gold chip regeneration from sample to sample too complicated (very slow k_{off}).

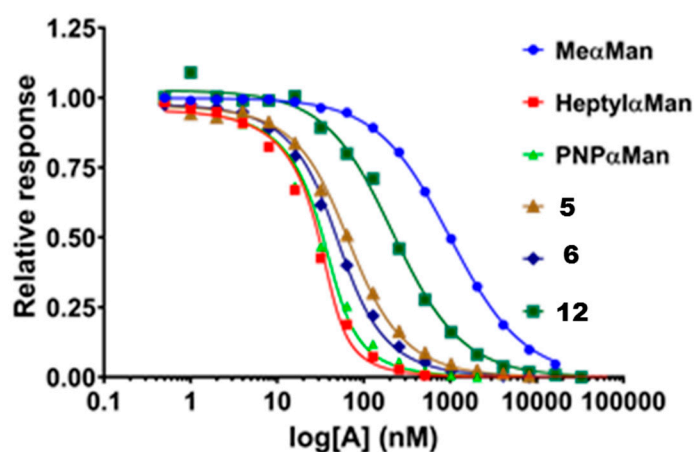


Figure 5. Normalized SPR data sets for competitive inhibition of binding of standards (1–3) and compounds **5**, **6**, and **12** on immobilized 6-aminohexyl α -D-mannopyranoside (**4**) on the CM5 gold chip.

Table 1. Relative binding affinities as measured by SPR.

Compound	Kd (nM)	Relative Potency	ΔG° (kcal mol ⁻¹)
MeαMan (1)	475.4 \pm 5.7	1	−8.62
PNPαMan (2)	4.21 \pm 0.72	113	−11.42
HeptylαMan (3)	2.37 \pm 0.50	200	−11.76
4	38.58 \pm 0.11	12	−10. 11
5	19.90 \pm 1.22	24	−10. 50
6	11.45 \pm 0.63	42	−10. 83
12	94.4 \pm 6.2	5	−9. 58

The binding data of methyl- (**Me α Man**, **1**), *p*-nitrophenyl α -D-mannopyranoside (**PNP α Man**, **2**), and heptyl- (**heptyl α Man**, **3**), were also included for comparison purpose. Their relative binding affinities are in the same order as those usually measured by other SPR methods [46], thus justifying the validity of our new procedure. As can be seen, the C-linked analog **6** was a better ligand (Kd = 11.45 nM) than the corresponding O-linked (**5**) (Kd = 19.90) and the S-linked (**12**) (Kd = 94.4 nM). Given the subtle changes in the 3D structures between **5** and **12**, the difference in their respective Kds is somewhat surprising (about 5 fold). However, the situation is similar for the related series of O- and S-linked heptyl α -D-mannopyranosides (see above, 3.4 fold). The major difference here is that the C-linked analog **6** was better than the other two and close to some of the best known *E. coli* FimH antagonists [10,18]. This could be attributed to a better π - π stacking of the biphenyl substituent of **6** within the tyrosine gate.

3. Materials and Methods

Reactions were carried out under Nitrogen using commercially available ACS grade solvents which were stored over 4 Å molecular sieves. Solutions in organic solvents were dried over anhydrous Na₂SO₄, filtered, and concentrated under reduced pressure. Reagents were obtained from Sigma Aldrich (Toronto, ON, Canada). The FimH residues 1-158 were isolated from the *E. coli* strain UTI89. The source was obtained from a synthetic gene expressed into pET24a [51–53]. Methyl α -D-mannopyranoside (**Me Man**, **1**) and *para*-nitrophenyl α -D-mannopyranoside (**PNP Man**, **2**) are commercially available from Alfa Aesar (Tewksbury, MA, USA) and Sigma Aldrich, respectively. Heptyl α -D-mannopyranoside (**Heptyl Man**, **3**) was prepared according to published procedure [18,53]. Melting points were measured on a Fisher Jones apparatus and are uncorrected. Reactions were monitored by thin-layer chromatography using silica gel 60 F254 coated plates (E. Merck). NMR spectra (Merck, Darmstadt, Germany). Were recorded on Varian Inova AS600 and Bruker Avance III HD 600 MHz spectrometer. Proton and carbon chemical shifts (δ) are reported in ppm relative to the chemical shift of residual CHCl₃, which was set at 7.28 ppm (¹H) and 77.16 ppm (¹³C{¹H}). Coupling constants (*J*) are reported in Hertz (Hz) and the following abbreviations are used for peak multiplicities: Singlet (s), doublet (d), doublet of doublets (dd), doublet of doublet with equal coupling constants (*t*_{ap}), triplet (t), multiplet (m). Assignments were made using COSY (Correlated Spectroscopy) and HSQC (Heteronuclear Single Quantum Coherence) experiments. High-resolution mass spectra (HRMS) were measured with a LC-MS-TOF (Liquid Chromatography Mass Spectrometry Time of Flight) instrument (Agilent Technologies Santa Clara, CA, USA). In positive electrospray mode by the analytical platform of UQAM. SPR were performed with a Biacore T200 on a CM5 sensor chip (GE Healthcare Life Sciences, Mississauga, ON, Canada).

SPR Solution Affinity of Ligands for the E. coli FimH Adhesin

First, the equilibrium dissociation constant (*K*_D) of FimH against immobilized 6-aminohexyl α -D-mannopyranoside (**4**) was determined. The ligand 6-aminohexyl α -D-mannopyranoside (10 mM in H₂O) was immobilized via standard amine carbodiimide coupling chemistry (and blocking with ethanolamine as per the manufacturer's procedure) to obtain 364 response units (RU) (see sensorgram in SI). The reference channel was treated similarly without ligand. A concentration series of the analyte FimH (1024 to 8 nM in running buffer, 2-fold dilutions) was then injected over the surface (700 s association time; 1200 s dissociation time; 30 μ L/min flow rate; 25 °C temperature; running buffer: PBS pH 7.45, 3 mM EDTA supplemented with 0.005% Tween-20; regeneration: 30 s of 10 mM Glycine pH 1.5) and the response was double-referenced by subtracting the reference channel and the response with no FimH (running buffer only). Since equilibrium could not be reached during injection, *K*_D was obtained from the extracted kinetic rate constants (*K*_D = *k*_{off} / *k*_{on}) from a non-linear globally fitted 1:1 Langmuir binding isotherm containing a mass transfer constant (*t*_c) and locally fitted refractive index constant (RI) (Biacore T200 evaluation software v1.0, GE Healthcare Life Sciences) of the obtained sensorgrams.

Second, the equilibrium dissociation constants (*K*_D) of FimH against the compounds of interest were determined by measuring the amount of free FimH in a series of concentrations of the compounds of interest at a constant concentration of FimH (close to the *K*_D of FimH: 6-aminohexyl α -D-mannopyranoside) after equilibrium in solution. A concentration series of the ligand of interest (65536 to 0.5 nM in running buffer, 2-fold dilutions) was equilibrated overnight with analyte FimH (40 nM) and then injected over the previous ligand 6-aminohexyl α -D-mannopyranoside-immobilized surface (300 s association time; 300 s dissociation time; 30 μ L/min flow rate; 25 °C temperature; running buffer: 5% DMSO, PBS pH 7.45, 3 mM EDTA supplemented with 0.05% Tween-20; regeneration: 30 s of 10 mM Glycine pH 1.5) and the response was single-referenced by subtracting the reference channel and solvent corrected with a DMSO calibration curve. Since equilibrium could not be reached during injection, the responses (free [FimH], Y-axis) at the end of the injections were normalized with the response from FimH only ([compound of interest] = 0) and then plotted over the compounds of interest

concentrations (X-axis). The obtained curves were then non-linearly fitted according to the following equation [53] using GraphPad Prism (v6.07, GraphPad Software, San Diego, CA, USA) to extract R_{max} and K_S :

$$Y = \left(\frac{[FimH] - A}{K_C + [FimH] - A} \right) R_{max}$$

$$A = \frac{[FimH] + X + K_S - \sqrt{([FimH] + X + K_S)^2 - 4[FimH]X}}{2}$$

where: Y is the concentration of free FimH (RU); R_{max} is the binding capacity of the SPR surface at saturation of all binding sites; K_C and K_S are the equilibrium dissociation constants (K_D) for binding of FimH to immobilized 6-aminohexyl α -D-mannopyranoside ligand on the sensor surface and with the compound of interest in solution, respectively; [FimH] is the constant concentration of FimH (40 nM); X is the concentration of the compound of interest (nM).

4. Conclusions

The work presented herein which describes the syntheses, tridimensional structures, binding parameters using a modified SPR procedure of a series of *O*-, *C*-, and *S*-linked mannopyranoside derivatives incorporating the identical 1,1'-biphenyl pharmacophore but diverse aglyconic atoms illustrates the necessity to explore this aspect when exploring novel *E. coli* FimH antagonists. The fact that our series of biphenyl analogues differs from those previously obtained from the corresponding heptyl mannopyranosides, clearly justify the deepen analysis undertaken in this report. Clearly, this study also confirms the fact that our *C*-linked α -D-mannopyranoside unambiguously exists in its preferred 4C_1 chair conformation as opposed to several claims in the literature. It also further confirms that *S*-glycoside to possess an *exo*-anomeric effect. Even though, none of the compounds exhibited higher affinities than the golden *O*-linked heptyl α -D-mannopyranoside standard, this work expanded our knowledge on the *E. coli* FimH tyrosine gate which showed that further binding contacts are worth exploring. This fact on its own strongly argues for revisiting some of the recent best antagonists identified thus far and that they can be further optimized. Further studies in vitro and PK/PD evaluation will warrant deepen our approach toward truly therapeutic candidates.

Allyl 2,3,4,6-tetra-O-acetyl-1-thio- α -D-mannopyranoside (9). To a solution of peracetylated mannose (8), (405 mg, 1.12 mmol) in 2 mL of dried methylene chloride was added, under N_2 , allyl mercaptan (374 μ L, 4 equiv.). The mixture was cooled at 0 °C and BF_3Et_2O (1.4 mL, 10 equiv.) was added. The reaction mixture was stirred at room temperature for 20 h. The reaction was quenched with 2 mL of water under stirring, then diluted with methylene chloride and was washed with 30 mL of 5% aqueous solution of $NaHCO_3$. The organic layer then dried over Na_2SO_4 , and concentrated under reduced pressure. The residue was purified by silica gel column chromatography using Hexane/ EtOAc (7:3) to afford compound 9 as a colorless oil with $R_f = 0.35$ (Hexane/ EtOAc (7:3)), (240 mg, 0.593 mmol, 53%). ($c = 0.3$, $CHCl_3$); $[\alpha]^{20}_D = 66.1$, m.p. 66.5–69 °C. 1H NMR (300 MHz, $CDCl_3$): 5.84–5.76 (m, 1 H, $CH=CH_2$), 5.36–5.30 (m, 3H, H-4,H-3,H-2), 5.29–5.19 (m, 3H, H-1, $CH=CH_2$), 4.42–4.35, (m, 2H, H-5,H-6a), 4.11 (dd, 1H, $J_{6a,6b} = 14.8$ Hz, $J_{5,6a} = 8.4$ Hz, H-6b), 3.29–3.15 (m, 2H, H1'a,1'b), 2.18, 2.1, 2.06, 2.00 (4s, 12H, 4 \times COCH₃),. ${}^{13}C\{{}^1H\}$ NMR ($CDCl_3$): δ 170.6, 169.8, 169.8, 169.7 (4 \times CO), 132.6 (C-2'), 118.8 (C-3'), 80.9 (C-1), 71.2 (C-4), 69.8 (C-3) 69.0 (C-5), 65.9 (C-2), 62.7 (C-6), 32.8 (C-1'). ESI+-HRMS: $[M + Na]^+$ calcd for $C_{17}H_{24}NaO_9S$: 427.1039. Found 427.0994. Physical data matched those published using a related procedure [34].

(2E)-3-(1,1'-biphenyl-2-propen-1-yl) 2,3,4,6-tetra-O-acetyl-1-thio- α -D-mannopyranoside (11). To a solution of 9 (53 mg, 0.131 mmol) in dry DCM (2 mL) were added to 4-vinyl-1,1'-biphenyl (10) (28 mg, 1.2 equiv), and 10% tricyclohexylphosphine[1,3-bis(2,4,6-trimethylphenyl)imidazol-2-ylidene] [2-thienylmethylene]ruthenium(II) dichloride (Grubbs' catalyst) under N_2 . The solution was refluxed

overnight. The solution was evaporated under reduced pressure. The organic layer was separated, dried over Na₂SO₄, filtered, concentrated and the crude residue purified by silica gel column chromatography using Toluene/EtOAc (9.6:0.4) to afford compound **11** as a colorless oil with R_f = 0.24 (Toluene/EtOAc (9.6:0.4)) (41 mg, 0.073 mmol, 56%). [α]²⁰_D = +70 (c = 0.1, CHCl₃). ¹H NMR (300 MHz, CDCl₃) δ : 7.62 (dd, 4H, ³J_{H-H} = 14.8 Hz, ⁴J_{H-H} = 6.3 Hz, H-arom), 7.46 (dd, 4H, ³J_{H-H} = 15.5 Hz, ⁴J_{H-H} = 8.3 Hz, H-arom), 7.38 (dd, 1H, ³J_{H,H} = 8.6, ⁴J_{H-H} = 5.8 Hz, H-arom), 6.56 (d, 1H, J_{2',3'} = 15.6 Hz, CH=CH-biPh), 6.27–6.15 (m, 1 H, CH=CH-biPh), 5.4–5.2 (m, 4H, H-3, H-4, H2, H1), 4.41 (m, 2H, H-5, H-6a), 4.15 (dd, 1H, J_{6a,6b} = 12.2 Hz, J_{5,6b} = 2.4 Hz, H-6b), 3.58–3.29 (m, 2H, H-1'a, H-1'b), 2.16, 2.15, 2.08, 2.01 (4s, 12H, 4×COCH₃). ¹³C{¹H} NMR (CDCl₃): δ 170.6, 169.8, 169.8, 169.7, (4×CO), 1405, 135.3, 133.4 (Carom-q), 133.4(C-3'), 129.0 (Carom), 128.2, 127.2, 127.0, 126.9 (Carom), 123.7 (C-2'), 80.7 (C-1), 70.8 (C-4), 69.7 (C-3), 68.9 (C-5), 66.3 (C-2), 62.9 (C-6), 32.6 (C-1'), 20.9, 20.8, 20.7, 20.6 (4×Ac). ESI+-HRMS: [M + Na]⁺ calcd for C₂₉H₃₂NaO₉S: 579.1659. Found, 579.1669.

(2E)-3-(1,1'-biphenyl-2-propen-1-yl) 1-thio- α -D-mannopyranoside (**12**). The acetylated mannoside **11** was dissolved in dry MeOH (3 mL), a solution of sodium methoxide (1M in MeOH, 0.5 equiv) was added and the reaction mixture was stirred at room temperature until disappearance of the starting material after 5 hr. The solution was neutralized by addition of H⁺ ion-exchange resin (Amberlite IR 120), filtered, washed with MeOH and the solvent was removed in vacuum. The residue was lyophilized to yield the fully deprotected mannoside **12** (20 mg, 0.07 mmol, 73%) after purification by RP-HPLC (see SI). [α]²⁰_D = +133 (c = 0.2, MeOH). ¹H NMR (300 MHz, CDCl₃): δ 7.51 (d, 2H, ³J_{H-H} = 7.5 Hz, H-arom), 7.48 (d, 2H, ³J_{H-H} = 8.1 Hz, H-arom), 7.39 (d, 2H, ³J_{H-H} = 8.1, H-arom), 7.32 (t, 2H, ³J_{H-H} = 7.6, H-arom), 7.22 (t, 1H, ³J_{H-H} = 7.3, H-arom), 6.50 (d, 1H, J_{2',3'} = 15.7 Hz, CH=CH-biPh), 6.26–6.17 (m, 1 H, CH=CH-biPh), 5.14 (s, 1H, H-1), 3.88–3.84 (m, 1H, H-5), 3.82–3.76 (m, 2H, H-4, H-2), 3.66 (dd, 1H, J_{2,3} = 6.0 Hz, J_{3,4} = 11.9 Hz, H-3), 3.60–3.54 (m, 2H, H-6a, H-6b), 3.41 (dd, J_{1'b,1'a} = 13.5 Hz, J_{1'a,2'} = 9.2 Hz, H-1'a), 3.24 (dd, J_{1'a,1'b} = 13.5 Hz, J_{1'b,2'} = 5.6 Hz, H-1'b). ¹³C{¹H} NMR (CDCl₃): δ 141.8, 141.4, 137.1 (Carom-q), 133.3 (C-3'), 129.7 (Carom), 128.2, 127.9, 127.7, 127.6 (Carom), 126.1 (C-2'), 84.6 (C-1), 74.9 (C-5), 73.4 (C-2), 73.3 (C-4), 68.8 (C-3), 62.5 (C-6), 33.1 (C-1'). ESI+-HRMS: [M + NH₄]⁺ calcd for C₂₁H₂₈NO₅S: 406.1688. Found 406.1699.

X-Ray crystal-structure of 5 and refinement data: formula, (C₂₁H₂₄O₆), orthorhombic, space group P212121, a 6.2463 (3) Å, b 7.5145 (3) Å, c 39.1736 (18) Å, α 90°, β 90°, γ 90°, V 1838.72(14) Å³, D_{calcd} 1.345 g/cm³. Crystallographic data for the structure reported in this paper has been deposited at the Cambridge Crystallographic Data Centre (CCDC) with deposition no: 1840503 for C₂₁H₂₄O₆. Supplementary data can be obtained free of charge from CCDC, 12 Union Road, Cambridge CB2 1EZ, UK (fax: (+44)1223-336-033; e-mail: deposit@ccdc.cam.ac.uk).

X-Ray crystal-structure of 6 and refinement data: formula, (C₂₁H₂₄O₅), orthorhombic, space group P1 21 1, a 9.7448 (4) Å, b 8.1887 (4) Å, c 21.6060 (9) Å, α 90°, β 92.097°, γ 90°, V 1722.94 (13) Å³, D_{calcd} 1.374 g/cm³. Crystallographic data for the structure reported in this paper has been deposited at the Cambridge Crystallographic Data Centre (CCDC) with deposition no: 1871374 for C₂₁H₂₄O₅. Supplementary data can be obtained free of charge from CCDC, 12 Union Road, Cambridge CB2 1EZ, UK (fax: (+44)1223-336-033; e-mail: deposit@ccdc.cam.ac.uk).

Supplementary Materials: Supplementary materials are available online, detailed synthesis of 6-aminoethyl α -D-mannopyranoside (**4**); ¹H- and ¹³C-NMR spectra, COSY, HSQC, and MS of all compounds; HPLC profile of compounds **7** and **11**; FT-IR of compound **3**; X-Ray data for compounds **3** and **7**.

Author Contributions: L.M. performed the experimental parts include designing, and synthesis, characterization and crystallization, writing- original draft. G.V. did the molecular modeling and calculated the energy levels by Monte Carlo molecular mechanics and DFT. G.S. performed SPR experiment and data curation. R.R. conceived the paper, designed the thematic, wrote and finalized the paper, supervised the project, and got funding.

Funding: This work was supported by grants from the Natural Science and Engineering Research Council of Canada (NSERC) to R. Roy including a Canadian Research Chair and the Fonds du Québec – Nature et Technologies to R.R, grant No. 2018-PR-206987.

Conflicts of Interest: The authors declare no conflicts of interest.

References

1. Adeyi, O.O.; Baris, E.; Jonas, O.B.; Irwin, A.; Berthe, F.C.J.; Le Gall, F.G.; Marquez, P.V.; Nikolic, I.A.; Plante, C.A.; Schneidman, M.; et al. Drug-resistant infections: A Threat to Our Economic Future. *World Bank Rep.* **2017**, *2*, 1–132.
2. Mousavifar, L.; Roy, R. Alternative therapeutic strategies to fight bacterial infections. *Frontiers in Drug, Chem. Clin. Res.* **2018**, *1*, 1–4.
3. Ritter, T.K.; Wong, C.-H. Carbohydrate-Based Antibiotics: A New Approach to Tackling the Problem of Resistance. *Angew. Chem. Int. Ed.* **2001**, *40*, 3508. [[CrossRef](#)]
4. Spaulding, C.N.; Klein, R.D.; Schreiber, H.L.; Janetka, J.W.; Hultgren, S.J. Precision antimicrobial therapeutics: The path of least resistance? *NPJ Biofilms Microbiomes* **2018**, *4*, 1–7. [[CrossRef](#)] [[PubMed](#)]
5. Klein, T.; Abgottspon, D.; Wittwer, M.; Rabbani, S.; Herold, J.; Jiang, X.; Kleeb, S.; Luthi, C.; Scharenberg, M.; Bezencon, J. FimH Antagonists for the Oral Treatment of Urinary Tract Infections: From Design and Synthesis to in Vitro and in Vivo Evaluation. *J. Med. Chem.* **2010**, *53*, 8627–8641. [[CrossRef](#)] [[PubMed](#)]
6. Griebing, T.L. Urologic diseases in America project: Trends in resource use for urinary tract infections in men. *J. Urol.* **2005**, *173*, 1288–1294. [[CrossRef](#)] [[PubMed](#)]
7. Flores-Mireles, A.L.; Walker, J.N.; Caparon, M.; Hultgren, S.J. Urinary tract infections: Epidemiology, mechanisms of infection and treatment options. *Nat. Rev. Microbiol.* **2015**, *13*, 269–284. [[CrossRef](#)]
8. Terlizzi, M.E.; Gribaudo, G.; Maffei, M.E. UroPathogenic Escherichia coli (UPEC) infections: Virulence factors, bladder responses, antibiotic, and non-antibiotic antimicrobial strategies. *Front. Microbiol.* **2017**, *8*, 1–23. [[CrossRef](#)]
9. Zhou, G.; Mo, W.-J.; Sebbel, P.; Min, G.; Neubert, T.A.; Glockshuber, R.; Wu, X.-R.; Sun, T.-T.; Kong, X.-P. Uroplakin Ia is the urothelial receptor for uropathogenic Escherichia coli: Evidence from in vitro FimH binding. *J. Cell Sci.* **2001**, *114*, 4095–4103.
10. Mousavifar, L.; Touaibia, M.; Roy, R. Development of Mannopyranoside Therapeutics against Adherent-Invasive Escherichia coli Infections. *Acc. Chem. Res.* **2018**, *51*, 2937–2948. [[CrossRef](#)]
11. Ohno, H.; Hase, K. Glycoprotein 2 (GP2) grabbing the fimH+ bacteria into m cells for mucosal immunity. *Gut. Microbes* **2010**, *1*, 407–410. [[CrossRef](#)] [[PubMed](#)]
12. Spaulding, C.; Hultgren, S. Adhesive Pili in UTI Pathogenesis and Drug Development. *Pathogens* **2016**, *5*, 30. [[CrossRef](#)] [[PubMed](#)]
13. Chalopin, T.; Brissonnet, Y.; Sivignon, A.; Deniaud, D.; Cremet, L.; Barnich, N.; Bouckaert, J.; Gouin, S.G. Inhibition profiles of mono- and polyvalent FimH antagonists against 10 different Escherichia coli strains. *Org. Biomol. Chem.* **2015**, *13*, 11369–11375. [[CrossRef](#)] [[PubMed](#)]
14. Hung, C.S.; Bouckaert, J.; Hung, D.; Pinkner, J.; Widberg, C.; DeFusco, A.; Auguste, C.G.; Strouse, R.; Langermann, S.; Waksman, G.; et al. Structure basis of tropism of Escherichia coli to the bladder during urinary tract infection. *Mol. Microbiol.* **2002**, *44*, 903–915. [[CrossRef](#)] [[PubMed](#)]
15. Wurlpel, D.J.; Beatson, S.A.; Totsika, M.; Petty, N.K.; Schembri, M.A. Chaperone-Usher Fimbriae of Escherichia coli. *PLoS ONE* **2013**, *8*, 1–11. [[CrossRef](#)] [[PubMed](#)]
16. Abgottspon, D.; Ernst, B. In vivo Evaluation of FimH Antagonists – A Novel Class of Antimicrobials for the Treatment of Urinary Tract Infection. *Chimia* **2012**, *66*, 166–169. [[CrossRef](#)] [[PubMed](#)]
17. Mydock-McGrane, L.K.; Cusumano, Z.T.; Janetka, J.W. Mannose-derived FimH antagonists: A promising anti-virulence therapeutic strategy for urinary tract infections and Crohn’s disease. *Expert Opin. Ther. Pat.* **2016**, *26*, 175–197. [[CrossRef](#)]
18. Touaibia, M.; Krammer, E.-M.; Shiao, T.C.; Yamakawa, N.; Wang, Q.; Glinschert, A.; Papadopoulos, A.; Mousavifar, L.; Maes, E.; Oscarson, S.; et al. Sites for Dynamic Protein-Carbohydrate Interactions of O- and C-Linked Mannosides on the E. coli FimH Adhesin. *Molecules* **2017**, *22*, 1101. [[CrossRef](#)] [[PubMed](#)]
19. Hartmann, M.; Lindhorst, T.K. The bacterial lectin FimH, a target for drug discovery—Carbohydrate inhibitors of type 1 fimbriae-mediated bacterial adhesion. *Eur. J. Org. Chem.* **2011**, 3583–3609. [[CrossRef](#)]

20. Dorta, D.A.; Sivignon, A.; Chalopin, T.; Dumych, T.I.; Roos, G.; Bilyy, R.O.; Deniaud, D.; Krammer, E.M.; de Ruyck, J.; Lensink, M.F.; et al. The Antiadhesive Strategy in Crohn's Disease: Orally Active Mannosides to Decolonize Pathogenic Escherichia coli from the Gut. *ChemBioChem*. **2016**, *17*, 936–952. [[CrossRef](#)]
21. Mousavifar, L.; Vergoten, G.; Roy, R. Deciphering the conformation of C-linked α -D-mannopyranosides and their application toward the synthesis of low nanomolar *E. Coli* FimH ligands. *Arkivoc*. **2018**, *7*, 384–397. [[CrossRef](#)]
22. Han, Z.; Pinkner, J.S.; Ford, B.; Obermann, R.; Nolan, W.; Wildman, S.A.; Hobbs, D.; Ellenberger, T.; Cusumano, C.K.; Hultgren, S.J.; et al. Structure-based drug design and optimization of mannoside bacterial fimH antagonists. *J. Med. Chem.* **2010**, *53*, 4779–4792. [[CrossRef](#)] [[PubMed](#)]
23. Mydock-McGrane, L.; Cusumano, Z.; Han, Z.; Binkley, J.; Kostakioti, M.; Hannan, T.; Pinkner, J.S.; Klein, R.; Kalas, V.; Crowley, J.; et al. Antivirulence C-Mannosides as Antibiotic-Sparing, Oral Therapeutics for Urinary Tract Infections. *J. Med. Chem.* **2016**, *59*, 9390–9408. [[CrossRef](#)] [[PubMed](#)]
24. Fiege, B.; Rabbani, S.; Preston, R.C.; Jakob, R.P.; Zihlmann, P.; Schwardt, O.; Jiang, X.; Maier, T.; Ernst, B. The tyrosine gate of the bacterial lectin FimH: A conformational analysis by NMR spectroscopy and x-ray crystallography. *ChemBioChem*. **2015**, *16*, 1235–1246. [[CrossRef](#)] [[PubMed](#)]
25. Rabbani, S.; Krammer, E.M.; Roos, G.; Zalewski, A.; Preston, R.; Eid, S.; Zihlmann, P.; Prévost, M.; Lensink, M.F.; Thompson, A.; et al. Mutation of Tyr137 of the universal Escherichia coli fimbrial adhesin FimH relaxes the tyrosine gate prior to mannose binding. *IUCr* **2017**, *4*, 7–23. [[CrossRef](#)] [[PubMed](#)]
26. Sivignon, A.; Yan, X.; Dorta, D.A.; Bonnet, R.; Bouckaert, J.; Fleury, E.; Bernard, J.; Gouin, S.G.; Darfeuille-Michaud, A.; Barnich, N. Development of heptyl mannoside-based glycoconjugate antiadhesive compounds against adherent-invasive escherichia coli bacteria associated with crohn's disease. *MBio* **2015**, *6*, 1–9. [[CrossRef](#)] [[PubMed](#)]
27. Spaulding, C.N.; Klein, R.D.; Ruer, S.; Kau, A.L.; Schreiber, H.L.; Cusumano, Z.T.; Dodson, K.W.; Pinkner, J.S.; Fremont, D.H.; Janetka, J.W.; et al. Selective depletion of uropathogenic *E. coli* from the gut by a FimH antagonist. *Nature* **2017**, *546*, 528–532. [[CrossRef](#)] [[PubMed](#)]
28. Tomašič, T.; Rabbani, S.; Gobec, M.; Raščan, I.M.; Podlipnik, Č.; Ernst, B.; Anderluh, M. Branched α -D-mannopyranosides: A new class of potent FimH antagonists. *Med. Chem. Commun.* **2014**, *5*, 1247–1253. [[CrossRef](#)]
29. Wellens, A.; Lahmann, M.; Touaibia, M.; Vaucher, J.; Oscarson, S.; Roy, R.; Remaut, H.; Bouckaert, J. The tyrosine gate as a potential entropic lever in the receptor-binding site of the bacterial adhesin FimH. *Biochemistry* **2012**, *51*, 4790–4799. [[CrossRef](#)]
30. Roos, G.; Wellens, A.; Touaibia, M.; Yamakawa, N.; Geerlings, P.; Roy, R.; Wyns, L.; Bouckaert, J. Validation of reactivity descriptors to assess the aromatic stacking within the tyrosine gate of FimH. *ACS Med. Chem. Lett.* **2013**, *4*, 1085–1090. [[CrossRef](#)]
31. Tiwari, P.; Agnihotri, G.; Misra, A.K. Modified One-Pot Protocol for the Preparation of Thioglycosides from Unprotected Aldoses via S-Glycosyl Isothiouonium Salts. *J. Carbohydr. Chem* **2005**, *24*, 723–732. [[CrossRef](#)]
32. Pachamuthu, K.; Schmidt, R.R. Synthetic routes to thiooligosaccharides and thioglycopeptides. *Chem. Rev.* **2006**, *106*, 160–187. [[CrossRef](#)] [[PubMed](#)]
33. Ferrier, R.J.; Furneaux, R.H. Synthesis of 1,2-trans-Related 1-Thioglycoside Esters. *Carbohydr. Res.* **1976**, *52*, 63–68. [[CrossRef](#)]
34. Crich, D.; Mataka, J.; Zakharov, L.N.; Rheingold, A.L.; Wink, D.J. Stereoselective Formation of Glycosyl Sulfoxides and Their Subsequent Equilibration: Ring Inversion of an r-Xylopyranosyl Sulfoxide Dependent on the Configuration at Sulfur. *J. Am. Chem. Soc.* **2002**, *124*, 6028–6036. [[CrossRef](#)]
35. Dominique, R.; Das, S.K.; Roy, R. Alkenyl O- and C-glycopyranoside homodimerization by olefin metathesis reaction. *Chem. Commun.* **1998**, 2437–2438. [[CrossRef](#)]
36. Roy, R.; Das, S.K. Recent applications of olefin metathesis and related reactions in carbohydrate chemistry. *Chem. Commun.* **2000**, 519–529. [[CrossRef](#)]
37. De Ruyck, J.; Lensink, M.F.; Bouckaert, J. Structures of C-mannosylated anti-adhesives bound to the type 1 fimbrial FimH adhesion. *IUCr* **2016**, *3*, 163–167. [[CrossRef](#)]
38. Choumane, M.; Banchet, A.; Probst, N.; Gérard, S.; Plé, K.; Haudrechy, A. The synthesis of d-C-mannopyranosides. *Comptes Rendus Chim.* **2011**, *14*, 235–273. [[CrossRef](#)]

39. George, T.G.; Szolcsányi, P.; Koenig, S.G.; Paterson, D.E.; Isshiki, Y.; Vasella, A. Preparation of an advanced intermediate for the synthesis of stable analogues of guanofosfocin. *Helv. Chim. Acta* **2004**, *87*, 1287–1298. [[CrossRef](#)]
40. Schwardt, O.; Rabbani, S.; Hartmann, M.; Abgottsporn, D.; Wittwer, M.; Kleeb, S.; Zalewski, A.; Smieško, M.; Cutting, B.; Ernst, B. Design, synthesis and biological evaluation of mannosyl triazoles as FimH antagonists. *Bioorg. Med. Chem.* **2011**, *19*, 6454–6473. [[CrossRef](#)]
41. Malapelle, A.; Coslovi, A.; Doisneau, G.; Beau, J.M. An expeditious synthesis of N-acetylneuraminic acid α -C-glycosyl derivatives (“ α -C-glycosides”) from the anomeric acetates. *Eur. J. Org. Chem.* **2007**, *19*, 3145–3157. [[CrossRef](#)]
42. Jorgensen W., L.; Tirado-Rives, J. Molecular modeling of organic and biomolecular systems using BOSS and MCPRO. *J. Comput. Chem.* **2005**, *26*, 1689–1700. [[CrossRef](#)] [[PubMed](#)]
43. Derreumaux, P.; Vergoten, G. A new spectroscopic molecular mechanics force field. Parameters for proteins. *J. Chem. Phys.* **1995**, *102*, 8586–8605. [[CrossRef](#)]
44. Vergoten, G.; Mazur, I.; Lagant, P.; Michalski, J.C.; Zanetta, J.P. The SPASIBA force field as an essential tool for studying the structure and dynamics of saccharides. *Biochimie* **2003**, *85*, 65–73. [[CrossRef](#)]
45. Lagant, P.; Nolde, D.; Stote, R.; Vergoten, G.; Karplus, M. Increasing normal modes analysis accuracy: The SPASIBA spectroscopic force field introduced into the CHARMM program. *J. Phys. Chem. A* **2004**, *108*, 4019–4029. [[CrossRef](#)]
46. Bouckaert, J.; Berglund, J.; Schembri, M.; De Genst, E.; Cools, L.; Wuhler, M.; Hung, C.S.; Pinkner, J.; Slättegård, R.; Zavialov, A. Receptor binding studies disclose a novel class of high-affinity inhibitors of the Escherichia coli FimH adhesion. *Mol. Microbiol.* **2005**, *55*, 441–455. [[CrossRef](#)]
47. Mikkelsen, L.M.; Hernáiz, M.J.; Martín-Pastor, M.; Skrydstrup, T.; Jiménez-Barbero, J. Conformation of glycomimetics in the free and protein-bound state: Structural and binding features of the C-glycosyl analogue of the core trisaccharide α -D-Man-(1 \rightarrow 3)-[α -D-Man-(1 \rightarrow 6)]-D-Man. *J. Am. Chem. Soc.* **2002**, *124*, 14940–14951. [[CrossRef](#)] [[PubMed](#)]
48. Fernández-Alonso, C.M.; Díaz, D.; Berbis, M.Á.; Marcelo, F.; Cañada, J.; Jiménez-Barbero, J. Protein-carbohydrate interactions studied by NMR: From molecular recognition to drug design. *Curr. Protein Pept. Sci.* **2012**, *13*, 816–830. [[CrossRef](#)]
49. Eris, D.; Preston, R.C.; Scharenberg, M.; Hulliger, F.; Abgottsporn, D.; Pang, L.; Jiang, X.; Schwardt, O.; Ernst, B. The Conformational Variability of FimH: Which Conformation Represents the Therapeutic Target? *ChemBioChem* **2016**, *17*, 1012–1020. [[CrossRef](#)]
50. Crich, D.; Mataka, J.; Sun, S.; Lam, K.C.; Rheingold, A.L.; Win, D.J. Stereoselective sulfoxidation of α -mannopyranosyl thioglycosides: The exo-anomeric effect in action. *Chem. Commun.* **1998**, *24*, 2763–2764. [[CrossRef](#)]
51. Chen, S.L.; Hung, C.S.; Pinkner, J.S.; Walker, J.N.; Cusumano, C.K.; Li, Z.; Bouckaert, J.; Gordon, J.I.; Hultgren, S.J. Positive selection identifies an in vivo role for FimH during urinary tract infection in addition to mannose binding. *PNAS* **2009**, *106*, 22439–22444. [[CrossRef](#)] [[PubMed](#)]
52. André, S.; Cañada, F.J.; Shiao, T.C.; Largartera, L.; Diercks, T.; Bergeron-Brlek, M.; Papadopoulos, A.; Ribeiro, J.P.; Touaibia, M.; Solís Menéndez, D.M.; et al. Fluorinated Carbohydrates as Lectin Ligands: Biorelevant Sensors with Capacity to Monitor Anomer Affinity in 19F-NMR-Based Inhibitor Screening. *Eur. J. Org. Chem.* **2012**, 4354–4364.
53. de Mol, N.J. Affinity Constants for Small Molecules from SPR Competition Experiments. In *Surface Plasmon Resonance*; de Mol, N.J., Fischer, M.J.E., Eds.; Humana Press of Springer Science+Business Media: New York, NY, USA, 2010; pp. 101–113. ISBN 9781607616696.

Sample Availability: Samples of the compounds are not available from the authors.



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