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Stereoselective synthesis of (+)-5-thiosucrose and (+)-5-thioisosucrose†

Atsushi Ueda, *[ab] Jinhong Pi,[a] Yui Makura,[b] Masakazu Tanaka[b] and Jun'ichi Uenishi*[ac]

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(+)-5-Thiosucrose **1**, a novel isosteric sulfur analog of sucrose, was synthesized stereoselectively for the first time *via* indirect β-D-fructofuranosidation involving selective β-D-psicofuranosidation, followed by stereoinversion of the secondary hydroxy group at the C-3 position on the furanose ring. Glycosidation of protected 5-thio-D-glucose with a D-psicofuranosyl donor provided β-D-psicofuranosyl 5-thio-α-D-glucopyranoside and that with D-fructofuranosyl donor gave α-D-fructofuranosyl 5-thio-α-D-glucopyranoside. Two anomeric stereocenters of the glycosyl donor and acceptor were controlled correctly to provide a single disaccharide among four possible anomeric isomers in the glycosylation. Conversion of the resulting disaccharides afforded (+)-5-thiosucrose **1** and (+)-5-thioisosucrose **2** in excellent yields, respectively. Inhibitory activities of **1** and **2** against α-glucosidase *in vitro* were also examined.

Introduction

Sucrose (Fig. 1) is one of the common sugars in nature and is an important carbohydrate for energy source in human health. At the same time, it is also a favorite sweetener in our daily life. So far, a variety of synthetic analogs of sucrose have been synthesized as low- or noncalorie alternatives for a healthy dietary purpose. For example, sucralose (Splenda®) is a widely known analog of sucrose in which three hydroxy groups are replaced by chlorine atoms, leading to a hundredfold increase in sweetness compared to that of sucrose.[1] Many other derivatives of sucrose with different substituents have also been reported.[2] However sucrose analogs in which an atom in the sucrose ring system is changed are far less common. As rare examples, C-sucrose, in which the glycosidic oxygen is replaced by a carbon atom, was synthesized by Kishi *et al.*, and hemicarbasucrose, a carba-analog of sucrose, was reported by Jiménez-Barbero and Sollogoub *et al.*[3] Although sucrose is an actual substrate for α-glucosidase, these modified sucrose analogs are not substrates of α-glycosidase and non-notable activities in inhibition against α-glycosidase were reported.

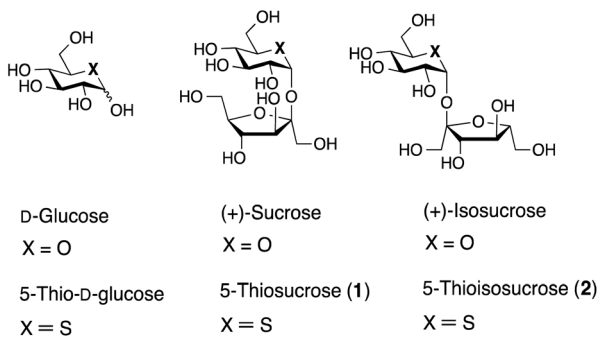

Fig. 1 Structures of glucose, sucrose, isosucrose, and their sulfur analogs.

Thiosugars replace a ring oxygen atom with a sulfur atom in carbohydrate, and are extremely rare in nature with the exception of 5-thio-D-mannose[4] and salacinols.[5] It should be noted that salacinol has potent enzymatic inhibitory activity against α-glucosidase. In fact, it has been already approved and used commercially in the context of dietary drinks or supplement of foods. Hetero monosaccharides, including thiosugars,[6] azasugars,[7] and carbasugars,[8] and their disaccharide analogs[9] have been synthesized as sugar mimics and their biological behaviors and functions involving α-glucosidase inhibitory activity were examined.[10] However, there has been nothing potent beyond salacinol concerning α-glucosidase inhibition.[11,12]

α-Glucosidase hydrolyzes sucrose to fructose and glucose, and interestingly, 5-thio-D-glucose inhibits this process.[11a,f] A pyranose ring oxygen is essentially required for the substrates of α-glucosidase.[13] As an isosteric analog of sucrose, 5-thiosucrose **1** and 5-thioisosucrose **2** possesses a 5-thioglucose moiety and

[a]*Kyoto Pharmaceutical University, Misasagi, Yamashina, Kyoto 607-8412, Japan. E-mail: aueda@nagasaki-u.ac.jp; juenishi@phs.osaka-u.ac.jp*

[b]*Graduate School of Biomedical Sciences, Nagasaki University, 1-14 Bunkyo-machi, Nagasaki 852-8521, Japan*

[c]*Graduate School of Pharmaceutical Sciences, Osaka University, Suita, Osaka 565-0871, Japan*

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would resemble the conformation of sucrose. We thought that **1** would be of interest as a sweetener as well as an inhibitor superior than 5-thio-D-glucose. This research is focused on the stereoselective synthesis of 5-thiosucrose **1** and 5-thioisosucrose **2**, and herein we report their synthesis and some biological properties.

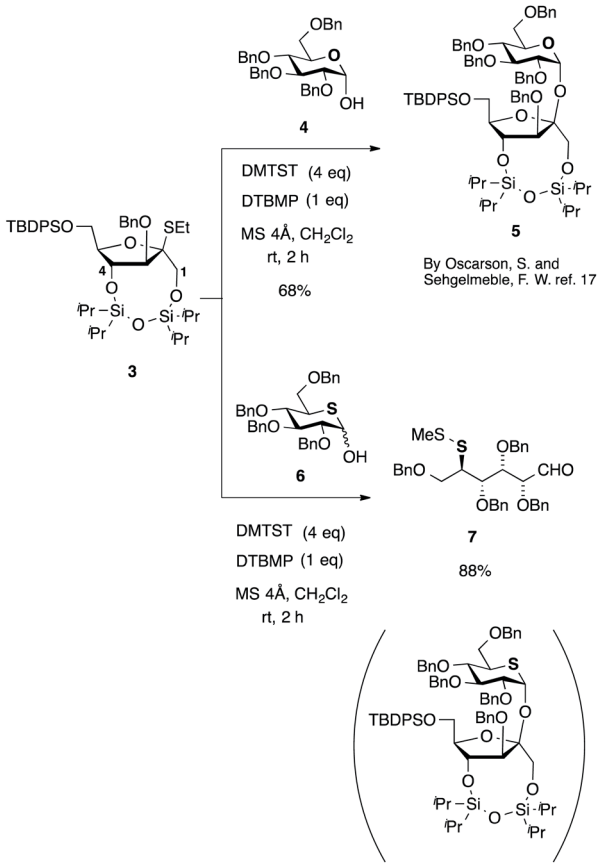
Results and discussion

Although many methods for the stereoselective glycosylation have been reported and utilized,[14] early attempts at the synthesis of sucrose[15] revealed the difficulties involved with the stereocontrol of its two anomeric centers. For the synthesis of **1** and **2**, two anomeric carbon centers connected with β,α- and α,α-O-linkage [**1**; β-D-Fru*f*-(2↔1)-α-D-Glc*p* and **2**; α-D-Fru*f*-(2↔1)-α-D-Glc*p*] need to be controlled in a glycosylation step. An α-selectivity rather than β-selectivity was reported for glycosidation of D-fructose as an either glycosyl acceptor or donor.[14a] In fact, glycosidation of D-fructofuranose affords an α-anomer or α-predominant mixtures in most cases.[16] On the other hand, 5-thio-D-glucose has been used as a glycosyl donor to form an α-glycosidic bond by the anomeric effect of the sulfur ring.[6] However, it has never been used as an acceptor in glycosidation reaction to our best knowledge.

Despite these failures in β-D-fructofuranosidation, the natural occurrence of β-D-furanoside can be found in sucrose and inulin. Therefore, β-directing D-fructofuranosylation has been a challenging task and this has encouraged carbohydrate chemists to develop selective β-D-fructofuranosylation. There is only one elegant example in β-D-fructofuranosylation, reported by Oscarson *et al.*[17] As shown in Scheme 1, they used ethyl thioglycoside **3** as a D-fructofuranosyl donor, in which the C-1 and C-4 hydroxy groups are fixed with a connection of cyclic disiloxyether to block an attack from the α-face of the furanose ring. Stereoselective glycosylation of the acceptor **4** with **3** promoted by dimethyl(methylthio)sulfonium triflate (DMTST) in the presence of 2,6-di-*tert*-butyl-4-methylpyridine (DTBMP) and 4 Å molecular sieves occurred to give β-furanosyl α-pyranoside **5** with 68% yield, exclusively.[17a]

Accordingly, we initially attempted the Oscarson's method for the synthesis of **1**. However, glycosidation of 2,3,4,6-tetra-O-benzyl-5-thio-D-glucopyranose (**6**)[18] with **3** gave no desired disaccharide. Instead, disulfide **7** was isolated in 88% yield. Activation of sulfide by DMTST occurred on the *endo*-sulfide of the acceptor **6** instead of the *exo*-sulfide of the donor **3**. Therefore, this method could be useful for general oxygen sugars but not for thiosugars.

Since Oscarson's β-D-fructofuranosylation method was found to be incompatible with the synthesis of **1**, we turned our attention to indirect synthesis through β-D-psicofuranosylation, which was employed in the stereoselective synthesis of sucrose previously.[19] We have reported that glycosidation of D-glucopyranose with D-psicofuranosyl donor[20] protecting 3,4-diols with acetonide gave β-D-psicofuranosyl α-D-glucopyranoside, which afforded sucrose after several steps. D-Psicofuranosyl donor was regarded as β-D-fructofuranosyl donor in the disaccharide syntheses.[19,21] On the other hand, glycosylation of D-glucopyranose with D-fructofuranosyl donor occurred to give α-D-fructofuranosyl α-D-glucopyranoside predominantly which was used for the synthesis of isosucrose. The course of α-D-fructofuranosylation or β-D-psicofuranosylation could be governed by the stereochemistry of the C-3 hydroxy substituent. Glycosidation of acceptor with D-fructofuranosyl donor is not sufficient for the syntheses of β-D-fructofuranosides, but is suitable for α-D-fructofuranoside. Based on the above results, synthetic plan for **1** and **2** is depicted in Fig. 2.



Scheme 1 Reaction of Oscarson's glycosyl donor **3** with acceptors **4** and **6**.

D-Fructofuranosyl donor with 5-thioglucose acceptor would give α-D-fructofuranosyl 5-thio-α-D-glucopyranoside, of which anomeric centers would matched with the stereochemistry of **2**. Glycosylation of 5-thioglucose acceptor with D-psicofuranosyl donor would give β-D-psicofuranosyl 5-thio-α-D-glucopyranoside of which anomeric centers would matched with the stereochemistry of **1**. This disaccharide will lead to **1** after a stereoinversion at the C-3 hydroxy group. In both cases, α-glycoside on the anomeric center of 5-thioglucopyranose would be formed by the strong anomeric effect of the thiane ring.

Synthesis of 5-thioisosucrose (2)

Several D-fructofuranosyl donors are available. Fructofuranosyl halide,[16a] fructofuranosyl phosphite,[16c] 2-O-acetylfructofuranose,[16e] 2-thiofructofuranoside,[16d] and fructofuranosyl *N*-phenyltrifluoroacetimidate[16f] have been reported. Nevertheless,

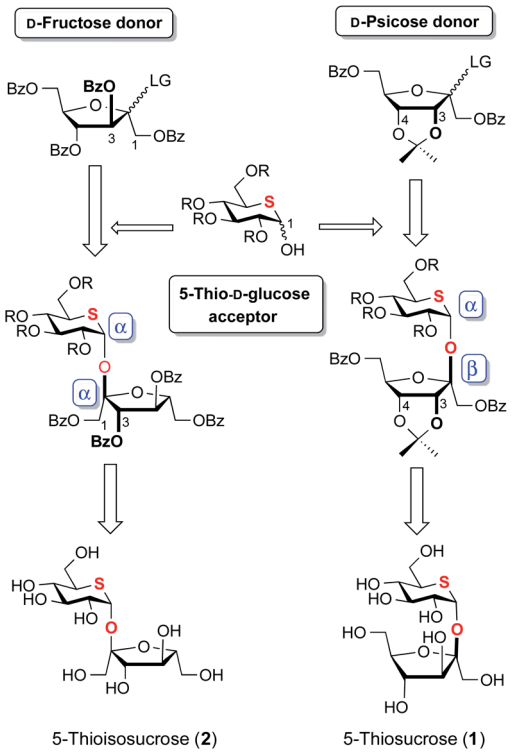


Fig. 2 Synthetic plan for 5-thiosucrose **1** and 5-thioisosucrose **2**.

most of these glycosidations entail synthetic difficulties since D-fructose possesses a sterically congested anomeric hydroxy group due to the structure of ketohexofuranose. In addition, the choice of leaving groups is restricted in this case because a sensitive cyclic sulfide unit exists in the glycosyl acceptor for the synthesis of **2**.

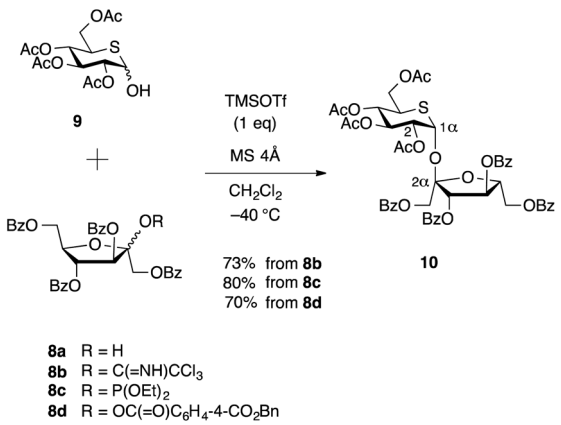
As shown in Scheme 2, we examined three different D-fructofuranosyl donors, thus imidate donor[22] **8b**, phosphite donor **8c**,[16c] and benzyl phthalate donor[23] **8d**. These donors were prepared from 1,3,4,6-tetra-O-benzoyl-D-fructofuranose (**8a**).[16b] Trimethylsilyl trifluoromethanesulfonate (TMSOTf) promoted glycosylation of 2,3,4,6-tetra-O-acetyl-5-thio-D-glucose **9** (ref. 18) with each glycosyl donor **8b**, **8c**, or **8d**[24] at −40 °C in dichloromethane in the presence of 4 Å molecular sieves resulted in the formation of 2α,1α-disaccharide **10** [α-D-Fru*f*-(2↔1)-α-D-Glc*p*] as a single isomer with 73%, 80%, and 70% yields, respectively. It is noteworthy that a single isomer (2α,1α) was formed exclusively among the four possible anomeric isomers (2β,1α, 2α,1α, 2α,1β, and 2β,1β) in this glycosidation, while D-glucopyranosyl acceptor reported in the synthesis of isosucrose produced two anomeric isomers (2α,1α, 2α,1β) in moderate selectivities (47 : 53,[16a] 4 : 1,[16c] and 84 : 16 (ref. 16*e*)). Although the three fructofuranosyl donors used in this study showed similar reactivity and selectivity, phthalate donor **8d** is regarded as the most convenient donor because it is readily prepared and stable under storage, in comparison with other donors **8b** and **8c**.

Stereochemistry at the anomeric positions in **10** was determined by the coupling constant in the ^{1}H NMR spectrum and the chemical shifts in the $^{13}C\{^1H\}$ NMR spectrum. The $J_{1,2}$ value of the pyranose ring proton is 3.1 Hz, which is in the typical value of α-glucopyranoside, and it is identical to that of 1,2-*cis*-α-D-glucopyranoside. The $^{13}C\{^1H\}$ NMR chemical shift of the anomeric position of the furanose ring exhibits at 109.0 ppm, which is identified to that of α-D-fructofuranoside.[17a,25] These data supported the structure of the disaccharide **10**.

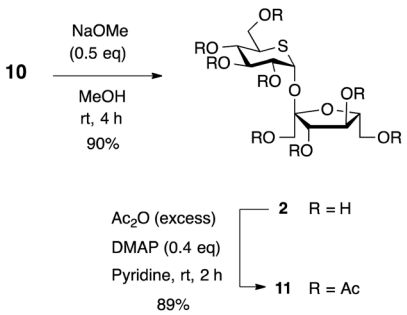
Finally, treatment of **10** with NaOMe in MeOH furnished the synthesis of **2** in 90% yield (Scheme 3). Compound **2** was synthesized in two steps from D-fructose donor with strict stereocontrol of two anomeric centers. The corresponding octaacetate **11** was prepared in 89% yield in order to compare the analytical data with the related disaccharides shown in Table 1.

Synthesis of 5-thiosucrose (1)

According to the synthetic plan, we started the synthesis of **1** through β-D-psicofuranosidation. The initial step of this synthesis involved β-selective glycosidation of **6** with the O-protected D-psicofuranosyl donor **12**,[18,19,24] which was prepared from D-psicose in 49% overall yield in five steps.[20b] Reaction of **6** with **12** in the presence of TMSOTf in dichloromethane at −40 to −20 °C afforded the desired glycoside **13** in 76% yield as a single stereoisomer (Scheme 4). The configuration of the anomeric center on the pyranoside ring in **13** was identified as that of α-D-glucopyranoside by a $J_{1,2}$ value of 2.9 Hz. The anomeric center in D-psicofuranoside was identical to β-



Scheme 2 α-D-Fructofuranosylation of 5-thio-D-glucopyranose **9**.

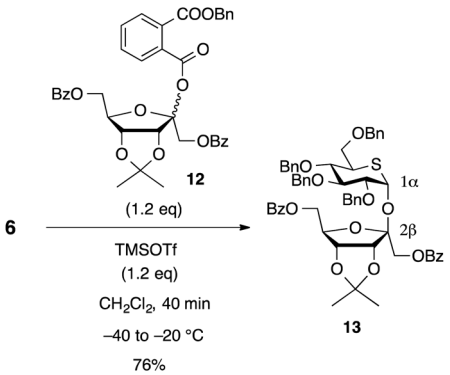


Scheme 3 Synthesis of 5-thioisosucrose **2** and its acetate **11**.

Table 1 Comparison of octaacetyl disaccharides **18**, **11**, **19**, and **20**

Compound	Specific rotation[a]	Chemical shifts of furanose protons[b] (δ)							Coupling constants of furanose protons[b] (Hz)			
	$[\alpha]_D$	H-1a	H-1b	H-3	H-4	H-5	H-6a	H-6b	$J_{3,4}$	$J_{4,5}$	$J_{5,6a}$	$J_{5,6b}$
18	+67.6	4.38	4.28	5.71	5.62	4.22	4.52	4.52	5.9	5.6	5.5	5.5
11	+147.8	4.52	4.29	5.69	4.87	4.55	4.54	4.30	0.5	3.3	4.1	7.7
19[c]	+60.0	4.37	4.29	5.71	5.55	4.20	4.41	4.41	5.4	5.7	5.4	5.4
20[c]	+83.5	4.79	3.98	5.69	4.84	4.45	4.39	3.96	1.0	3.0	4.0	2.0

[a] $CHCl_3$ was used as a solvent. [b] Benzene-d_6 was used as a solvent. [c] These values were obtained from the literature.[30]



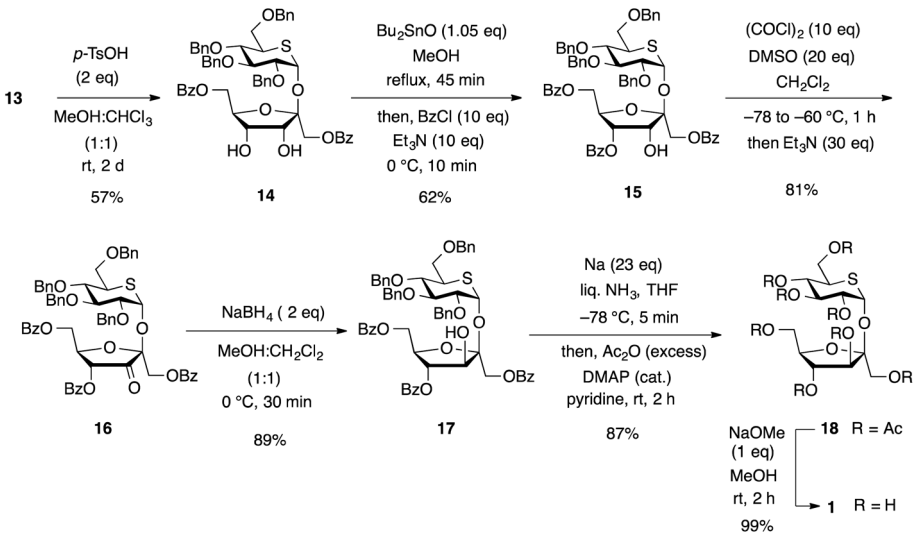
Scheme 4 Psicofuranosylation with 5-thio-α-D-glucopyranose **6**.

glycoside. In general, the C2-carbon of psicofuranoside appears between 107–109 ppm for β anomers and between 103–105 ppm for α anomers,[19,20] and the $^{13}C\{^1H\}$ NMR chemical shift of the anomeric carbon existed at 109.9 ppm in **13**.

In this reaction, protected 5-thioglucose **6** performed an α-directing acceptor as same as **9** in the synthesis of 5-thio-isosucrose. The glycosidation occurred on the β-face of the furanose donor **12** because of the steric influence of the acetonide group existing on the α-side of the ring. Considering that D-psicofuranosylation of the corresponding α-D-glucose gave a mixture in the ratio of 2 : 1 (α-glucopyranoside *vs.* β-glucopyranoside),[19] it should be noted that the predominant selectivity for 5-thio-D-glucopyranose **6** *vs.* D-glucopyranose is quite interesting in psicofuranosylation of 5-thiopyranose and pyranose donors.[26]

Conversion of β-D-psicofuranoside **13** to β-D-fructofuranoside **17** was carried out in four steps (Scheme 5):

Deprotection of acetonide group in **13** with *p*-toluenesulfonic acid in MeOH gave diol **14** in 57% yield. Benzoate **15** was obtained from **14** *via* a stannylene intermediate. Treatment of **14** with Bu_2SnO in MeOH at reflux temperature followed by benzoylation on the C-4 hydroxy group with benzoyl chloride gave **15** selectively.[27] The Swern conditions will be a choice for oxidation of the secondary hydroxy group in the presence of cyclic sulfide. The secondary alcohol of **15** was oxidized smoothly to give ketone **16** in 81% yield. Then, reduction of the ketone **16** with $NaBH_4$ occurred from the bottom of the furanose ring selectively to convert to β-D-fructofuranoside **17** in 89% yield. The direction of hydride attack in the reduction of 3-ketone is controlled by the adjacent 2-β-glycosidic bond to give 2,3-*syn*-product.[28] Removal of both O-benzoyl and O-benzyl groups in disaccharide **17** under the Birch conditions and



Scheme 5 Synthesis of **1**.

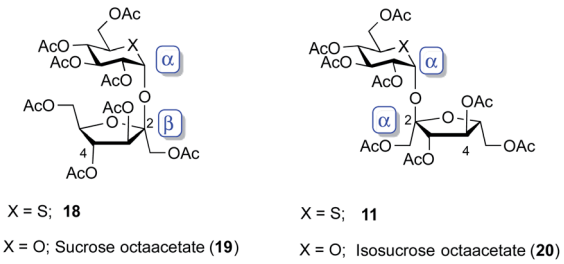


Fig. 3 Octaacetate of 5-thiosucrose **18**, 5-thioisososucrose **11**, sucrose **19**, and isosucrose **20**.

successive acetylation gave octaacetate **18** in 87% yield. After deprotection of all acetyl groups under Zemplén's conditions,[29] the synthesis of **1** was accomplished in 99% yield.

Stereochemistry

The specific rotations, and chemical shifts and coupling constant of ^{1}H NMR for octaacetates of 5-thiosucrose and 5-thioisosucrose (**18** and **11**) are summarized with the corresponding data reported for sucrose and isosucrose (**19** and **20**)[30] in Table 1. Their structures are shown in Fig. 3. Specific rotation values of 5-thioisosucrose and isosucrose (**11** and **20**) are relatively larger than those of 5-thiosucrose and sucrose (**18** and **19**). Chemical shifts of each α,α-anomers (**11** and **20**) and those of β,α-anomers (**18** and **19**) are comparable in ^{1}H NMR. The chemical shifts of the H-4 protons in furanose ring are characteristic. Their difference (*ca.* 0.7 ppm) can be observed between **19** and **20**, and thioanalogs **18** and **11**. Coupling constants of $J_{3,4}$ and $J_{4,5}$ in sucrose **19** and thiosucorse **18** are larger than those of isosucrose **20** and thioisosucorse **11**. Thus, all these results supported the structures of 5-thiodisaccharides **11** and **18**.

Biological study

Inhibitory activities against α-glycosidase for compounds **1**, **2**, and 5-thio-D-glucose were examined *in vitro* using rat intestinal α-glucosidase. In literature, 5-thio-D-glucose is reported to be a weak to moderate inhibitor for α-glucosidase.[11f,12] In the present study, 5-thio-D-glucose showed 48% inhibition at 8 mM. It was regrettable that neither **1** nor **2** exhibited any inhibition at 8 mM, while commonly used α-glucosidase inhibitor such as acarbose and voglibose work at nM levels. Although 5-thio-D-glucose has a sweet taste, **1** was found to be a little bitter rather than sweet, in rough-and-ready taste analyses.

Conclusions

The first stereoselective syntheses of **1** and **2** were achieved by stereoselective glycosidation. The key steps involved D-fructofuranosylation and D-psicofuranosylation of protected 5-thio-D-glucose acceptors **9** and **6** to afford α-D-fructofuranosyl 5-thio-α-D-glucopyranoside **10** and β-D-psicofuranosyl 5-thio-α-D-glucopyranoside **13** with high stereoselectivity in excellent yields, respectively. The configurations of their two-anomeric centers were strictly controlled in a single glycosidation step, in which the strong anomeric effect of 5-thio-D-glucopyranose was observed. We have demonstrated that 5-thio-D-glucopyranose works as an α-directing glycosyl acceptor for the first time. Although neither **1** nor **2** exhibit α-glucosidase inhibitory activity or sweetness, current results will aid in the design of new α-glucosidase inhibitors and the synthesis of other disaccharide of thiosugar derivatives.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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