

Cite this: *RSC Adv.*, 2019, 9, 28241Received 26th July 2019
Accepted 3rd September 2019

DOI: 10.1039/c9ra05797h

rsc.li/rsc-advances

Synthesis of a 1,2-*cis*-indoxyl galactoside as a chromogenic glycosidase substrate†

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A synthetically challenging 1,2-*cis*-indoxyl galactoside, X- α -galactoside, was first prepared in this study using a cyclic ketone indoxyl acceptor and a glycosyl trichloroacetimidate donor to produce an enol glycoside and a 4,6-*O*-di-*tert*-butylsilylene-protected galactosyl donor to complete the synthesis. The target compound shows enzyme activity in the presence of α -galactosidase.

Introduction

Indoxyl glycosides are chromogenic substrates that are useful in detecting glycosidase activity. These substrates have been widely used for this purpose in histochemical, biochemical, and bacteriological studies.¹ In this process, the hydrolysis of an indoxyl substrate by glycosidase releases a sugar moiety and a free indoxyl, the liberated molecule of which then undergoes oxidation in the air followed by dimerization to generate an indigoid dye (Fig. 1). Indigoid dyes are insoluble in water, making the diffusion of the slightest amount of dye from enzymatic hydrolysis sites possible to determine the precise location of glycosidases.² The color and physical properties (*e.g.*, solubility and affinity for protein) of indigoid dyes depend on the nature of the halogen substituents located on the indole ring of the compound. For example, 5-bromo-4-chloro (greenish blue)-, 5-bromo-6-chloro (magenta)-, 5-chloro (red)-, and 5-iodo (purple)-substituted substrates are well known, and these different-colored compounds are used for tailored applications.¹

Many indoxyl glycosides are now commercially available, but halogen-substituted indoxyl glycosides are expensive because they are difficult to prepare. Their synthesis is difficult because of the low nucleophilicity of the indole hydroxy group, which suppresses the formation of the glycoside, in addition to side reactions that occur because of the indoxyl structure, leading to a decrease in the coupling yield. Furthermore, another synthetic challenge that exists for indoxyl glycosides is that they exist in the enol form and can be readily cleaved under normal

glycosylation (acidic) conditions to release an aglycone in the keto form. Therefore, to date, reports on the synthesis of enol glycosides under acidic conditions have been extremely limited.³ S_N2-like reaction between glycosyl halides and *N*-acetyl-indoxyl derivatives in the presence of NaOH in acetone or Na in MeOH is a commonly used synthetic approach used to produce indoxyl glycosides, which generally results in very low reaction yields of less than 50% (Fig. 2A-i).⁴ These basic conditions are required to generate the enolate form from the cyclic ketone version of the free indoxyl, which exists in keto-enol equilibrium in the reaction medium and almost none of the corresponding enol forms in nonpolar solvents.⁵ Unfortunately, these conditions result in low yields, in addition to the formation of various side products, including indigogenic compounds and acetyl migration and/or elimination products generated from the donor. In 2013, Thiem and coworkers synthesized indoxylic acid allyl ester **1** to improve the reactivity and instability of the indoxyl acceptor by blocking the reactive 2-position.⁶ In this study, they found that phase transfer glycosylation using **1** suppressed the formation of indigogenic side products, resulting in a drastic increase in the yields by up to 86% (Fig. 2A-ii).⁷ Recently, Wei *et al.* found that solid-liquid phase transfer catalysis (S-L PTC) glycosylation between acetobromo sugars and an indoxyl acceptor could be used to synthesize indoxyl glycosides, resulting in moderate

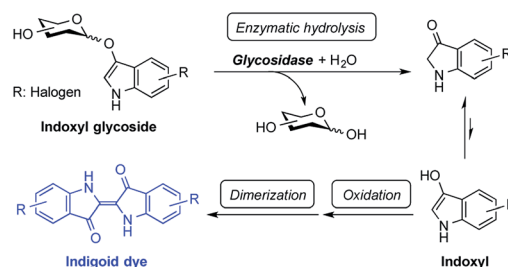


Fig. 1 Using indoxyl glycosides to detect glycosidase activity.

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† Electronic supplementary information (ESI) available: NMR data for all new compounds. See DOI: 10.1039/c9ra05797h



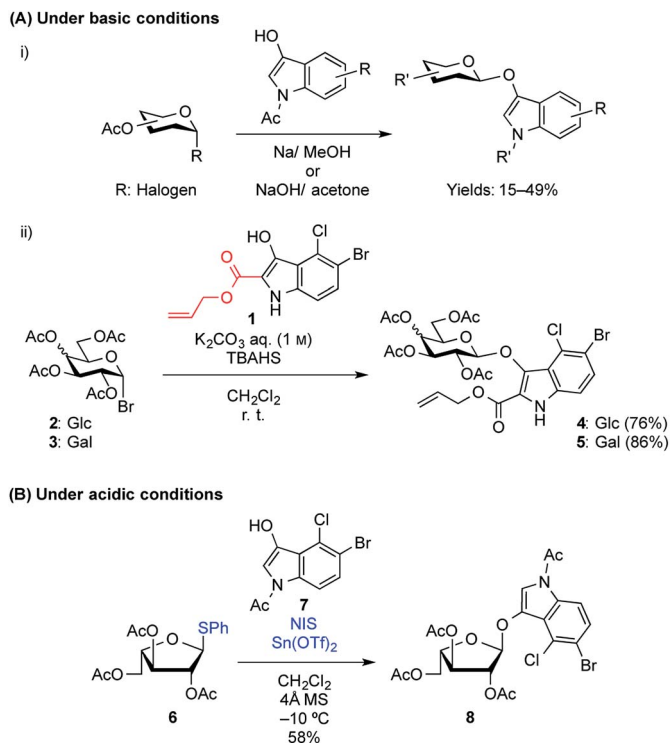


Fig. 2 Examples of the successful formation of 1,2-*trans*-indoxyl glycosides *via* (A) under basic conditions and (B) under acidic conditions. TBAHS: tetrabutylammonium hydrogen sulfate; NIS: *N*-iodosuccinimide.

glycosylation yields of 46–65%.⁸ However, the formation of indoxyl glycosides *via* a standard glycosylation reaction under acidic conditions has scarcely been reported, although there are limited reports on the synthesis of *X*- α -L-arabinofuranosides using the Koenigs–Knorr method⁹ or thioglycoside activation.¹⁰ In the latter, the authors used thiophenyl L-arabinofuranosyl donor **6** and indoxyl **7** in the presence of NIS-Sn(OTf)₂ in CH₂Cl₂ to give **8** in a 58% yield (Fig. 2B). To date, various indoxyl glycosides, including β -Glc,¹¹ β -Gal,^{11,12} β -GlcNAc,^{11a} β -GlcA,¹³ α -L-Araf,^{9,10} β -LacNAc,^{7b} and β -xylobioside,¹⁴ have been chemically synthesized, but they are all 1,2-*trans*-oriented glycosides. To the best of our knowledge, there have been no reports on the synthesis of 1,2-*cis*-indoxyl glycosides, which is due not only to the difficulty in preparing these compounds, but also to the lack of reliable methods available to control the anomeric stereoselectivity in 1,2-*cis*-glycoside formation. However, a large number of 1,2-*cis*-glycosides (*e.g.*, α -Gal, α -Glc, and α -Fuc) and the corresponding glycosidases can be found in nature, where they play pivotal roles in various biological processes.¹⁵ Thus, developing a synthetic route for 1,2-*cis*-indoxyl glycosides is, therefore, of great importance.

Results and discussion

In this study, for the first time, the synthesis of *X*- α -(1,2-*cis*-oriented)-galactoside (**9**, shown in Fig. 3, where X is the common abbreviation for 5-bromo-4-chloro-3-indoxyl) is

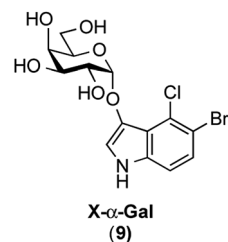
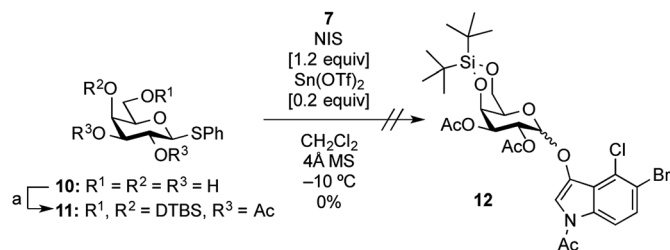


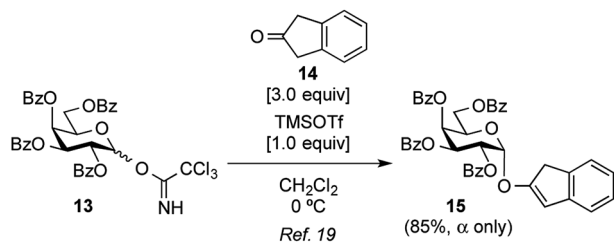
Fig. 3 *X*- α -galactoside, the target molecule in this study. X = 5-bromo-4-chloro-3-indoxyl.

reported. The first attempt to synthesize the target *X*- α -galactoside was made using thiophenyl 4,6-*O*-di-*tert*-butylsilylene- (DTBS-) protected galactosyl donor **11**, readily prepared from known thiophenyl galactoside **10**,¹⁶ and known indoxyl acceptor **7** (ref. 11b) to give **12**. Based on our previous success in the synthesis of a large number of α -galactosides using DTBS-protected Gal donors, it was thought that this route would lead to a promising result.¹⁷ Furthermore, the abovementioned report on the synthesis of *X*-Araf using a thiophenyl arabinofuranosyl donor mediated by an NIS-Sn(OTf)₂ system warranted further examination. However, the glycosylation reaction performed under the reported conditions did not proceed at all, and only the oxidative dimerization of **7** was observed, leading to a 91% recovery of the donor (Scheme 1). In this attempt, although 1.2 equiv. of the acceptor was initially used according to the preparation described in the literature,¹⁰ all of the acceptor was consumed in 1 h, resulting in the precipitation of a blue dye. The use of 3.0 equiv. of the acceptor under the same conditions did not change the outcome of the reaction. These results suggest that the use of a thiophenyl pyranosyl donor was not feasible for the formation of *X*-glycosides, which might be due to the pyranoside donor being less reactive than its furanoside counterpart. In addition, in order to prevent the oxidative dimerization of the indoxyl acceptors, the use of thioglycosides, which usually have to be oxidatively activated, might be unsuitable in this study.

Attention was then turned to the acid activation of a trichloroacetimidate donor.¹⁸ Ding and coworkers reported the efficient direct α -glycosylation of a cyclic ketone using a glycosyl trichloroacetimidate donor, in which the coupling of 2,3,4,6-tetra-*O*-benzoyl-galactosyl trichloroacetimidate donor **13** (ref.



Scheme 1 Glycosylation of indoxyl **7** using DTBS-protected Gal thioglycoside donor **11**. Reagents and conditions: (a) DTBS(OTf)₂, Py, r.t.; Ac₂O, r.t., 97% (2 steps in one-pot fashion).



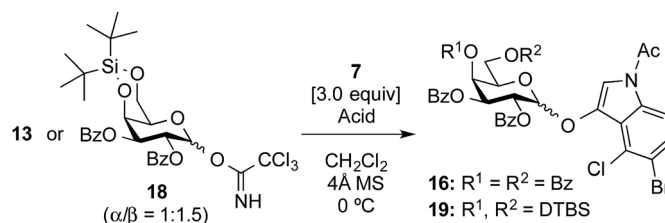
Scheme 2 The successful conditions for the stereoselective glycosylation of cyclic ketone derivative **14** using Gal trichloroacetimidate donor **13** reported by Liu *et al.*

19) and cyclic ketone **14** in the presence of TMSOTf (1.0 equiv.) in CH_2Cl_2 at 0°C provided the corresponding α -enol glycoside **15** as the sole product in an 85% yield (Scheme 2).²⁰ The high α -stereo-selectivity observed was a result of the acid-catalyzed *in situ* anomerization of the kinetically formed β -isomer to the thermodynamic α -isomer. These glycosylation conditions were, thus, used in the present study (Table 1, entry 1). As a result, the reaction between **13** and **7** quantitatively proceeded to give the desired X-galactoside **16** contaminated with the dimerized dye product of **7**, which was hard to remove by column chromatography. This result led to the first preparation of X-pyranosides under acidic conditions. However, the stereoselectivity in this reaction was very poor, with an α/β ratio of 1 : 1.5, favoring the unwanted β -isomer. The conversion of the β -isomer into the α -isomer was then attempted *via in situ* anomerization, in which the isolated β -isomer and **7** were stirred together under the above glycosylation conditions for 20 h. However, the formation of the α -isomer was not observed, with the reaction only resulting in the degradation of the β -isomer to the corresponding lactol and **7**. Despite its poor stereoselectivity, the successful formation of an X-glycoside using a trichloroacetimidate donor prompted us to use DTBS-protected galactosyl donors again to improve the α -selectivity.

Using 4,6-O-DTBS-protected galactosyl trichloroacetimidate donor **18**, readily prepared from **10** *via* the corresponding thiophenyl glycoside **17**, the glycosylation of **7** in the presence of TMSOTf (1.0 equiv.) was investigated (Table 1, entry 2). As anticipated, the reaction afforded the desired X-galactoside **19** in an almost quantitative yield with excellent α -selectivity (α/β ratio = >20 : 1). The installation of the DTBS group to endow the product with low polarity made the removal of the dye from the reaction mixture easy, but separation of the stereoisomers was not possible at this stage. However, the structure of **19 α** as the major product was confirmed by ^1H NMR spectrum of the crude mixture, where the $J_{1,2}$ value of 3.5 Hz for H-1 of the Gal residue (δ 5.94, doublet) and the presence of H-2 of the indole ring (δ 7.16, singlet) indicated the α -configuration of X-galactoside. In entry 3 of Table 1, it can be seen that reducing the amount of TMSOTf to 0.1 equiv. led to a sluggish reaction as well as a decrease in the coupling yield to 84%. Next, the acid catalyst was changed to triflic acid (Table 1, entry 4), but the outcome of the glycosylation was almost the same as that using TMSOTf. Subsequently, tin(II) triflate, which was used in the abovementioned thioglycoside activation, gave a result similar to that of TMSOTf (Table 1, entry 5). The results of both experiments, shown in entries 4 and 5 in Table 1, suggest that the nature of the acid species is not important in this reaction. Consequently, the use of a trichloroacetimidate donor was found to be of great importance in the synthesis of indoxyl glycosides.

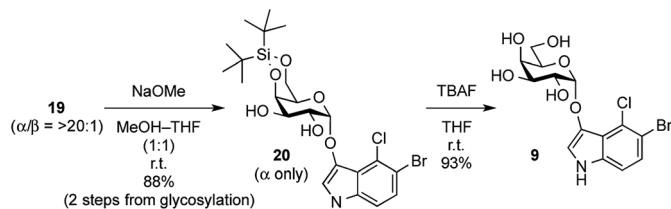
In order to complete the synthesis of X- α -galactoside, its protecting groups, including acyl groups and the DTBS group, were removed (Scheme 3). Before acyl group deprotection, it was thought that the 1,2-*cis*-enol glycosides would be alkali-sensitive because the 1,2-*cis*-configuration could lead to β -elimination at the anomeric carbon, producing the enolate form of the indoxyl and the 1,2-ene derivative of the sugar. Fortunately, these fears proved to be unfounded and the acyl groups in **19** (a mixture of α - and β -isomers) were cleaved under Zemplén conditions,

Table 1 Reaction scheme and conditions used for the glycosylation of **7** using Gal trichloroacetimidate donors **13** or **18**



Entry	Donor ^a	Acid (equiv.)	Time [h]	Prod.	% Yield ^b	α/β ratio ^c
1	13	TMSOTf (1.0)	0.5	16	>100 ^d	1 : 1.5
2	18 ^e	TMSOTf (1.0)	0.5	19	96	>20 : 1
3	18	TMSOTf (0.1 + 0.1 + 0.1)	2	19	84	>20 : 1
4	18	TfOH (1.0)	0.5	19	91	>20 : 1
5	18	Sn(OTf) ₂ (1.0)	0.5	19	90	>20 : 1

^a One equivalent of the donor was used. ^b Isolated yield. ^c Determined based on ^1H NMR spectra of the isomer mixtures. ^d Contaminated with the dimerization product of **7** (dye). ^e The trichloroacetimidate donor **18** was prepared from **10** over four steps *via* the corresponding thiophenyl glycoside **17**.



Scheme 3 Global deprotection of fully protected X- α -Gal.

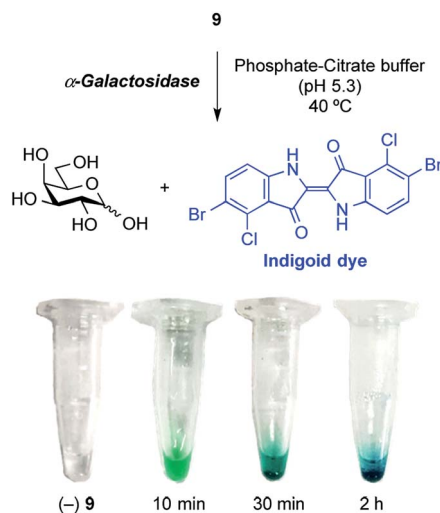


Fig. 4 Monitoring of α -galactosidase activity using the synthesized X- α -Gal compound.

affording **20** as a single isomer in good yield without any degradation of the enol glycoside. Subsequently, clean removal of the DTBS group by TBAF in THF furnished the target X- α -Gal **9** in excellent yield. During the deprotection process, it should be noted that the protected X-glycoside derivatives **19** and **20** were not very stable during even short-term storage in the freezer and underwent degradation to the corresponding lactol and indigoid dye. Therefore, after the glycosylation of the indoxyl, global deprotection should be performed as soon as possible.

Finally, enzymatic hydrolysis of the synthesized X- α -Gal **9** using an α -galactosidase (from green coffee bean) was carried out (Fig. 4). The glycosidase activity in a phosphate-citrate buffer (pH 5.3) at 40 °C was monitored in Eppendorf tubes for 10 min, 30 min, and 2 h. A greenish-blue color evolved over 10 min, after which the precipitation of dye particles was observed after 30 min and 2 h. In the tube, the substrate was hydrolyzed by glycosidase, and the liberated indoxyl was gradually oxidized and then dimerized to form a water-insoluble indigoid dye, providing a clear indication of enzyme activity. This result demonstrated that the synthesized X- α -Gal was applicable to the detection of α -galactosidase activity.

Conclusions

This was the first successful preparation of a synthetically challenging 1,2-*cis*-oriented indoxyl galactoside, the reaction

steps of which proceeded in a facile and high-yielding manner. The formation of the α -galactosidic linkage was achieved using a 4,6-*O*-DTBS-protected galactosyl donor with excellent stereo-selectivity. Importantly, the use of a trichloroacetimidoyl leaving group was found to be effective in the formation of an enol glycoside. Upon exposure to α -galactosidase, the solution of the synthesized X- α -Gal in a buffer underwent a color change as the substrate was hydrolyzed by the enzyme, oxidized by the air, and then dimerized, showing that the target X- α -Gal exhibits enzyme activity. The successful synthesis of a 1,2-*cis*-indoxyl galactoside in this study can be expanded upon to allow access to a variety of indoxyl glycosides. We are currently underway with the efficient synthesis of other 1,2-*cis*-indoxyl glycosides including X- α -D-glucoside and X- α -L-fucoside. The results will be reported in the near future.

Experimental

Synthetic procedures

General methods. All reactions were performed under a positive argon pressure, unless otherwise stated. All chemicals were obtained from commercial suppliers and used without further purification, unless otherwise stated. Molecular sieves were purchased from Wako Chemicals Inc. and dried at 300 °C for 2 h in a muffle furnace before use. TLC analysis including high performance TLC (HPTLC) was conducted on Merck TLC (silica gel 60F₂₅₄ on glass plate). Compounds were detected either by exposure to UV light (2536 Å) or by soaking in a solution of 10% H₂SO₄ in ethanol followed by heating. Silica gel (80 mesh and 300 mesh) manufactured by Fuji Silysia Chemical Ltd. was employed for flash column chromatography. The quantity of silica gel was usually estimated as 100- to 150-fold the weight of the sample to be charged. Solvent systems in chromatography are specified in v/v. Evaporation and concentration were performed *in vacuo*. High-resolution mass spectrometry (ESI-TOF MS) was performed on a Bruker micrOTOF instrument. Optical rotations were measured using a “Horiba SEPA-300” high-sensitivity polarimeter. The ¹H NMR and ¹³C NMR spectra were recorded using Bruker Biospin AVANCE III 500 spectrometer. Chemical shifts in ¹H NMR spectra are expressed in ppm (δ) relative to the Me₄Si signal, adjusted to δ 0.00 ppm. Data are presented as follows: chemical shift, multiplicity (s = singlet, d = doublet, t = triplet, dd = doublet of doublet, td = triplet of doublet, m = multiplet and/or multiple resonance), integration, coupling constant in Hertz (Hz), and position of the corresponding proton.

Enzyme assay. α -Galactosidase from green coffee bean (pH optimum 5.3 and no data for temperature optimum) was purchased from Sigma-Aldrich. To a suspension of the enzyme (*ca.* 25 U) in 3.2 M ammonium sulfate (pH 6.0) (100 μ L) was added 1.0 mL of phosphate-citrate buffer prepared at pH 5.3, giving the enzyme solution. The synthetic X- α -Gal (1.0 mg) was added to the enzyme solution (275 μ L) and the enzyme activity at 40 °C was monitored in 10 min, 30 min, and 2 h.

1-Acetyl-5-bromo-4-chloroindol-3-ol (7). Known indoxyl acceptor **7** was prepared from commercially available 1-acetyl-5-bromo-4-chloro-3-indolyl acetate according to the literature.^{11b}

To 95% sulfuric acid (4.65 mL) with vigorously stirring was added portionwise 1-acetyl-5-bromo-4-chloro-3-indolyl acetate (1.00 g, 3.04 mmol) at 0 °C. After stirring for 2 min at 0 °C, the completion of the reaction was confirmed by TLC (1 : 1 *n*-hexane/EtOAc). The reaction mixture was poured on crushed ice in a beaker and stirred quickly by glass rod. The precipitate formed was filtered off and washed with generous quantities of 1% solution of sodium acetate followed by water. The resulting residue was exposed to high vacuum, giving **7** (856 mg, 98%) as yellow solid. Prepared **7** should be stored in a temperature of less than –60 °C until use. ¹H NMR (500 MHz, CDCl₃): δ 8.42 (d, 1H, ³J_{ortho} = 8.5 Hz, Ar), 7.82 (d, 1H, Ar), 4.36 (s, 2H, CH₂), 2.33 (s, 3H, Ac); ¹³C NMR (125 MHz, CDCl₃): δ 190.7, 168.1, 154.1, 140.9, 131.8, 122.8, 119.0, 117.8, 56.4, 24.3. HRMS (ESI-TOF): *m/z* calcd for C₈H₅BrClNO: 267.9135 [M + Na]⁺; found: 267.9135.

Phenyl 2,3-di-O-acetyl-4,6-O-di-*tert*-butylsilylene-1-thio-β-D-galactopyranoside (11). To a solution of known phenyl 1-thio-β-D-galactopyranoside **10** (1.00 g, 3.67 mmol) in pyridine (36.7 mL) was added di-*tert*-butylsilyl bis(trifluoromethanesulfonate) (1.43 mL, 4.41 mmol) at 0 °C. After stirring for 45 min at rt, the completion of the reaction was confirmed by TLC (5 : 1 *n*-hexane/EtOAc). Subsequently, acetic anhydride (3.45 mL, 36.7 mmol) was added to the reaction mixture at 0 °C. After stirring for 1 h at rt, the completion of the reaction was confirmed by TLC (5 : 1 *n*-hexane/EtOAc). The reaction was quenched by the addition of dry methanol at 0 °C. Solvents were removed by co-evaporation with toluene, and then the residue was diluted with CHCl₃, washed with 2 M HCl, H₂O, satd aq NaHCO₃, and brine. The organic layer was subsequently dried over Na₂SO₄, filtered, and concentrated. The resulting residue was purified by silica gel column chromatography (5 : 1 *n*-hexane/EtOAc) to give **11** (1.76 g, 97%) as white foamy material. [α]_D +33.8 (*c* 1.0 in CHCl₃); ¹H NMR (500 MHz, CDCl₃): δ 7.50–7.25 (m, 5H, Ph), 5.47 (t, 1H, *J*_{1,2} = *J*_{2,3} = 10.0 Hz, H-2), 4.80 (dd, 1H, *J*_{3,4} = 3.0 Hz, H-3), 4.73 (d, 1H, H-1), 4.69 (d, 1H, H-4), 4.25 (dd, 1H, *J*_{5,6a} = 1.5 Hz, *J*_{gem} = 13.0 Hz, H-6a), 4.22 (dd, 1H, *J*_{5,6b} = 2.0 Hz, H-6a), 3.49 (s, 1H, H-5), 2.10 (s, 3H, Ac), 2.08 (s, 3H, Ac), 1.11 (s, 9H, ^tBu), 1.01 (s, 9H, ^tBu); ¹³C NMR (125 MHz, CDCl₃): δ 170.6, 169.5, 134.0, 132.0, 128.9, 127.7, 87.2, 77.3, 74.9, 74.8, 70.1, 67.4, 67.0, 27.5, 27.5, 23.2, 20.9, 20.8, 20.6. HRMS (ESI-TOF): *m/z* calcd for C₂₄H₃₆O₇SSi: 519.1843 [M + Na]⁺; found: 519.1844.

***N*-Acetyl-5-bromo-4-chloroindoxyl 2,3,4,6-tetra-O-benzoyl-D-galactopyranoside (16).** A solution of **13** (200 mg, 0.271 mmol) in CH₂Cl₂ (1.0 mL) was transferred into a two-neck round bottom flask, in which pre-activated 4 Å molecular sieves (200 mg) were placed, using a cannula at 0 °C. To the suspension was transferred a solution of **7** (233 mg, 0.813 mmol) in CH₂Cl₂ (3.0 mL) using a cannula at 0 °C. Subsequently, TMSOTf (49.0 μL, 0.271 mmol) was added to the mixture at 0 °C. After stirring for 10 h at the same temperature, the completion of the reaction was confirmed by HPTLC (1 : 1 *n*-hexane/EtOAc). The reaction was quenched by the addition of satd aq NaHCO₃ at 0 °C. The precipitate was filtered through Celite pad and the pad was washed with EtOAc. The combined filtrate and washings were washed with satd aq NaHCO₃, 2 M HCl, and brine. The organic layer was subsequently dried over Na₂SO₄, filtered, and concentrated. The resulting residue was purified by silica gel

column chromatography (7 : 1 *n*-hexane/EtOAc) to give **16** (a mixture of the stereoisomers, α/β = 1/1.5), **7**, and the corresponding dye (426 mg as the mixture). **16β** (contaminated with some compounds): ¹H NMR (500 MHz, CDCl₃): δ 8.23–7.27 (m, 22H, Ar), 7.25 (s, 1H, =CH–N), 6.20 (dd, 1H, *J*_{1,2} = 7.5 Hz, *J*_{2,3} = 10.0 Hz, H-2), 6.08 (d, 1H, *J*_{3,4} = 3.5 Hz, H-4), 5.72 (dd, 1H, H-3), 5.36 (d, 1H, H-1), 4.76 (dd, 1H, *J*_{5,6a} = 7.0 Hz, *J*_{gem} = 12.0 Hz, H-6a), 4.58 (dd, 1H, *J*_{5,6b} = 4.5 Hz, H-6b), 4.54 (m, 1H, H-5), 2.24 (s, 3H, Ac); ¹³C NMR (125 MHz, CDCl₃): δ 168.3, 166.2, 165.6, 165.6, 165.2, 139.7, 133.8, 133.6, 133.5, 133.4, 133.4, 130.5, 130.1, 129.8, 129.8, 129.8, 129.7, 129.3, 129.1, 128.8, 128.6, 128.6, 128.5, 128.4, 125.3, 122.5, 118.4, 116.1, 112.8, 101.2, 72.4, 71.4, 69.1, 68.2, 63.0, 23.4. HRMS (ESI-TOF): *m/z* calcd for C₄₄H₃₃BrClNO₁₁: 888.0818 [M + Na]⁺; found: 888.0814.

Phenyl 2,3-di-O-benzoyl-4,6-O-di-*tert*-butylsilylene-1-thio-β-D-galactopyranoside (17). To a solution of **10** (1.00 g, 3.67 mmol) in pyridine (18.4 mL) was added di-*tert*-butylsilyl bis(trifluoromethanesulfonate) (1.43 mL, 4.41 mmol) at 0 °C. After stirring for 1 h at rt, the completion of the reaction was confirmed by TLC (2 : 1 *n*-hexane/EtOAc). Subsequently, benzoyl chloride (3.41 mL, 29.4 mmol) and a catalytic amount of *N,N*-dimethyl-4-aminopyridine were added to the reaction mixture at 0 °C. After stirring for 1 h at rt, the completion of the reaction was confirmed by TLC (5 : 1 *n*-hexane/EtOAc). The reaction was quenched by the addition of dry methanol at 0 °C. Solvents were removed by co-evaporation with toluene, and then the residue was diluted with CHCl₃, washed with 2 M HCl, H₂O, satd aq NaHCO₃, and brine. The organic layer was subsequently dried over Na₂SO₄, filtered, and concentrated. The resulting residue was purified by silica gel column chromatography (8 : 1 *n*-hexane/EtOAc) to give **17** (2.00 g, 88%) as white foamy material. [α]_D +96.5 (*c* 1.0 in CHCl₃); ¹H NMR (500 MHz, CDCl₃): δ 8.14–7.21 (m, 15H, Ph), 5.93 (t, 1H, *J*_{1,2} = *J*_{2,3} = 10.0 Hz, H-2), 5.22 (dd, 1H, *J*_{3,4} = 3.0 Hz, H-3), 4.95 (d, 1H, H-1), 4.89 (d, 1H, H-4), 4.34 (d, 1H, *J*_{gem} = 12.0 Hz, H-6a), 4.30 (dd, 1H, *J*_{5,6b} = 2.0 Hz, H-6b), 3.66 (s, 1H, H-5), 1.16 (s, 9H, ^tBu), 0.96 (s, 9H, ^tBu); ¹³C NMR (125 MHz, CDCl₃): δ 166.1, 165.5, 133.8, 133.3, 133.2, 132.5, 129.8, 129.8, 129.6, 129.4, 128.9, 128.4, 128.4, 127.8, 87.5, 75.5, 75.0, 70.5, 68.1, 67.1, 27.5, 27.5, 23.3, 20.7. HRMS (ESI-TOF): *m/z* calcd for C₃₄H₄₀O₇SSi: 643.2156 [M + Na]⁺; found: 643.2155.

2,3-Di-O-benzoyl-4,6-O-di-*tert*-butylsilylene-D-galactopyranosyl trichloroacetimidate (18). To a solution of **17** (1.00 g, 1.61 mmol) in the mixed solvent (16.1 mL, 5 : 1 acetone–H₂O) was added *N*-bromosuccinimide (1.43 g, 8.05 mmol) at 0 °C. After stirring for 15 min at rt, the completion of the reaction was confirmed by TLC (2 : 1 *n*-hexane/EtOAc). The reaction mixture was diluted with EtOAc, washed with satd aq Na₂SO₃ and brine. The organic layer was subsequently dried over Na₂SO₄, filtered, and concentrated. The resulting residue was purified by silica gel column chromatography (7 : 1 *n*-hexane/EtOAc) to give the lactol compound (746 mg, 88%). The lactol (600 mg, 1.14 mmol) was then dissolved in CH₂Cl₂ (11.4 mL). To the solution were added CCl₃CN (1.14 mL, 11.4 mmol) and 1,8-diazabicyclo[5.4.0]undec-7-ene (136 μL, 0.912 mmol) at 0 °C. After stirring for 6 h at 0 °C as the reaction was monitored by TLC (1 : 1 *n*-hexane/EtOAc), the reaction mixture was concentrated. The resulting

residue was purified by silica gel column chromatography (15 : 1 → 2 : 1 *n*-hexane/EtOAc) to give **18** (711 mg, 93%, $\alpha/\beta = 1/1.5$) as white foamy material. **18** α : ^1H NMR (500 MHz, CDCl_3): δ 8.56 (s, 1H, NH), 8.03–7.35 (m, 10H, Ph), 6.79 (d, 1H, $J_{1,2} = 3.5$ Hz, H-1), 6.04 (dd, 1H, $J_{2,3} = 10.5$ Hz, H-2), 5.69 (dd, 1H, $J_{3,4} = 3.0$ Hz, H-3), 5.01 (d, 1H, H-4), 4.34 (dd, 1H, $J_{5,6a} = 2.0$ Hz, $J_{\text{gem}} = 12.5$ Hz, H-6a), 4.30 (dd, 1H, $J_{5,6b} = 1.5$ Hz, H-6b), 4.17 (s, 1H, H-5), 1.15 (s, 9H, $t\text{Bu}$), 0.98 (s, 9H, $t\text{Bu}$); ^{13}C NMR (125 MHz, CDCl_3): δ 166.2, 165.7, 160.8, 133.4, 133.3, 129.9, 129.7, 129.6, 129.0, 128.4, 124.8, 94.6, 91.0, 70.9, 70.7, 69.8, 67.3, 66.6, 27.5, 27.2, 23.3, 20.8. HRMS (ESI-TOF): m/z calcd for $\text{C}_{30}\text{H}_{36}\text{Cl}_3\text{NO}_7\text{Si}$: 694.1168 [M + Na] $^+$; found: 694.1172.

N-Acetyl-5-bromo-4-chloroindoxyl 2,3-di-O-benzoyl-4,6-O-di-tert-butylsilylene-D-galactopyranoside (19). A solution of **18** (200 mg, 0.298 mmol) in CH_2Cl_2 (1.0 mL) was transferred into a two-neck round bottom flask, in which pre-activated 4 Å molecular sieves (222 mg) were placed, using a cannula at 0 °C. To the suspension was transferred a solution of **7** (258 mg, 0.894 mmol) in CH_2Cl_2 (3.0 mL) using a cannula at 0 °C. Subsequently, TMSOTf (54.0 μL , 0.298 mmol) was added to the mixture at 0 °C. After stirring for 0.5 h at the same temperature, the completion of the reaction was confirmed by HPTLC (1 : 1 *n*-hexane/EtOAc). The reaction was quenched by the addition of satd aq NaHCO_3 at 0 °C. The precipitate was filtered through Celite pad and the pad was washed with EtOAc. The combined filtrate and washings were washed with satd aq NaHCO_3 , 2 M HCl, and brine. The organic layer was subsequently dried over Na_2SO_4 , filtered, and concentrated. The resulting residue was purified by silica gel column chromatography (9 : 1 *n*-hexane/EtOAc) to give **19** (229 mg, 96%, $\alpha/\beta = 20/1$) as purple powder. **19** α : ^1H NMR (500 MHz, CDCl_3): δ 8.23 (d, 1H, $^3J_{\text{ortho}} = 8.5$ Hz, Ar), 8.05–7.36 (m, 11H, Ar), 7.16 (s, 1H, =CH-N), 5.96 (dd, 1H, $J_{1,2} = 3.5$ Hz, $J_{2,3} = 10.5$ Hz, H-2), 5.94 (d, 1H, H-1), 5.90 (dd, 1H, $J_{3,4} = 3.0$ Hz, H-3), 5.03 (d, 1H, H-4), 4.35 (dd, 1H, $J_{5,6a} = 2.0$ Hz, $J_{\text{gem}} = 13.0$ Hz, H-6a), 4.22 (dd, 1H, $J_{5,6b} = 1.5$ Hz, H-6b), 4.10 (s, 1H, H-5), 2.56 (s, 3H, Ac), 1.19 (s, 9H, $t\text{Bu}$), 1.00 (s, 9H, $t\text{Bu}$); ^{13}C NMR (125 MHz, CDCl_3): δ 168.1, 166.2, 166.0, 140.2, 133.6, 133.5, 133.2, 130.6, 129.9, 129.7, 129.0, 128.5, 128.5, 128.4, 125.4, 122.3, 118.4, 116.2, 107.9, 97.5, 77.2, 70.9, 70.7, 68.3, 68.1, 66.8, 27.6, 27.4, 24.0, 23.3, 20.8. HRMS (ESI-TOF): m/z calcd for $\text{C}_{38}\text{H}_{41}\text{BrClNO}_9\text{Si}$: 820.1315 [M + Na] $^+$; found: 820.1315.

5-Bromo-4-chloroindoxyl 4,6-O-di-tert-butylsilylene- α -D-galactopyranoside (20). To a solution of **19** (210 mg, 0.256 mmol, $\alpha/\beta = 20/1$) in the mixed solvent (2.6 mL, 1 : 1 MeOH-THF) was added sodium methoxide (1.0 M solution in MeOH, 77 μL , 76.8 μmol) at 0 °C. After stirring for 1.5 h at rt as the reaction was monitored by TLC (1 : 1 *n*-hexane/EtOAc), the reaction was neutralized with Muromac (H^+) resin. The resin was filtered off and washed with MeOH. The filtrate and washings were concentrated. The resulting residue was purified by silica gel column chromatography (1 : 1 *n*-hexane/EtOAc) to give **20** (225 mg, 88%) as white powder. $[\alpha]_{\text{D}} +106.7$ (*c* 1.0 in CHCl_3); ^1H NMR (500 MHz, CDCl_3): δ 7.87 (br s, 1H, NH), 7.37 (d, 1H, $^3J_{\text{ortho}} = 8.5$ Hz, Ar), 7.10 (d, 1H, $J_{\text{C}=\text{CH},\text{NH}} = 3.0$ Hz, C=CH), 7.09 (d, 1H, Ar), 5.46 (d, 1H, $J_{1,2} = 3.0$ Hz, H-1), 4.56 (d, 1H, $J_{3,4} = 1.5$ Hz, H-4), 4.30 (dd, 1H, $J_{5,6a} = 2.0$ Hz, $J_{\text{gem}} = 12.5$ Hz, H-6a), 4.17 (dd, 1H, $J_{5,6b} = 1.5$ Hz, H-6b), 4.04–3.97 (m, 2H, H-2, H-3), 3.95 (s,

1H, H-5), 2.67 (m, 1H, OH), 2.45 (m, 1H, OH), 1.08 (s, 9H, $t\text{Bu}$), 1.06 (s, 9H, $t\text{Bu}$); ^{13}C NMR (125 MHz, CDCl_3): δ 136.7, 133.4, 127.2, 124.3, 118.2, 113.5, 111.4, 110.3, 100.7, 77.2, 73.2, 71.5, 70.1, 68.2, 66.8, 27.6, 27.2, 23.4, 20.7. HRMS (ESI-TOF): m/z calcd for $\text{C}_{38}\text{H}_{41}\text{BrClNO}_9\text{Si}$: 570.0685 [M + Na] $^+$; found: 570.0688.

5-Bromo-4-chloroindoxyl α -D-galactopyranoside (9). To a solution of **20** (225 mg, 0.395 mmol) in THF (3.9 mL) were added acetic acid (54 μL , 0.948 mmol) and tetrabutylammonium fluoride (1.0 M solution in THF, 948 μL , 0.948 mmol) at 0 °C. After stirring for 0.5 h at rt, the completion of the reaction was confirmed by TLC (5 : 1 $\text{CHCl}_3/\text{MeOH}$). Solvents were removed by co-evaporation with toluene. The resulting residue was purified by silica gel column chromatography on Iatrobeds 6RS-8060 (10 : 1 → 5 : 1 $\text{CHCl}_3/\text{MeOH}$) to give **9** (150 mg, 93%) as light sky blue powder. $[\alpha]_{\text{D}} +117.2$ (*c* 1.0 in CHCl_3); ^1H NMR (500 MHz, CD_3OD): δ 7.28 (d, 1H, $^3J_{\text{ortho}} = 9.0$ Hz, Ar), 7.25 (s, 1H, C=CH), 7.14 (d, 1H, Ar), 5.38 (d, 1H, $J_{1,2} = 3.5$ Hz, H-1), 4.07 (t, 1H, $J_{5,6a} = J_{5,6b} = 6.0$ Hz, H-5), 4.05–4.00 (m, 2H, H-3, H-4), 3.97 (dd, 1H, $J_{2,3} = 10.0$ Hz, H-2), 3.73 (d, 2H, H-6a, H-6b); ^{13}C NMR (125 MHz, CD_3OD): δ 137.1, 135.1, 127.1, 125.0, 119.6, 114.1, 113.4, 112.7, 101.9, 73.4, 71.6, 71.0, 70.4, 62.7, 49.9. HRMS (ESI-TOF): m/z calcd for $\text{C}_{14}\text{H}_{15}\text{BrClNO}_6$: 429.9663 [M + Na] $^+$; found: 429.9668.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

This work was supported in part by JSPS KAKENHI Grant Number JP18K05455 (to A. I.). We thank Prof. Dr Thiem and Dr Böttcher (University of Hamburg) for technical support on the synthesis of indoxyl derivatives.

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