Synthesis and spectroscopic investigation of a novel sensitive and selective fluorescent chemosensor for Ag⁺ based on a BINOL–glucose derivative†

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Based on a versatile 2,2'-binaphthol (BINOL) backbone, a novel BINOL–glucose derivative fluorescent sensor was synthesized using a click reaction. The fluorescence responses of the BINOL–glucose derivative (S,β-d)-1 conclude that it can be used as a specific fluorescent chemical sensor for Ag⁺ in the presence of a large number of competing metal ions without any obvious interference from other metal ions. Mass spectrometric and NMR spectroscopic data were used to study the mechanism, and implied the formation of a 1 + 1 complex between BINOL–glucose 1 and Ag⁺. Both the oxygen atoms of S-BINOL and two nitrogen atoms of triazole were involved in coordinating the silver ion.

Introduction

It is very important to discover and design new highly selective fluorescent sensors that can detect heavy metal ions, which may cause serious harm to human health and the living environment. Among the important heavy metals, silver has been paid more attention because of its adverse effects on biological harmony and the environment, due to its ability to easily bind to mercapto groups in proteins and inactivate respiratory enzymes.1,2 It is very difficult to identify silver ions from other heavy metal ions because of the moderate coordination ability of Ag⁺. However, almost all fluorescent sensors containing triazole experience interference from other transition metals, especially from mercury ions when attempting to identify silver ions,3–8 which leads to the limitation of the use of specific fluorescent sensors for silver ions. Only a few fluorescent sensors9–11 that can detect Ag⁺ have been reported in recent years. So, it is still a challenge to design an effective chemical method to target only Ag⁺.

The highly sensitive and rapid identification of isomers and metal ions is the highlight of using optically active BINOL derivatives as a fluorescent chemical sensor. With a versatile BINOL backbone, it can be easily modified through steric stabilization and electronic effects, and due to this, BINOL and its derivatives14–21 have attracted great attention in the field of fluorescent chemical sensors24–30 and synthetic catalysis.

Moreover, sugar is a hot research topic due to its natural existence, good biocompatibility and structural diversity without being toxic. Furthermore, its good water solubility is considered to be a very ideal feature for using it as a component in fluorescent chemical sensors. In recent years, triazole modified functional sugar derivatives have attracted continuous interest in research31–34 and various sugar-based fluorescent chemosensors have been synthesized using click reactions. In this way, a novel BINOL–glucose derivative fluorescent sensor was synthesized through the Cu-catalyzed 1,3-dipolar cycloaddition35–40 of an alkyne and azole. The synthesis was based on a versatile (S)-BINOL backbone, and the (S)-BINOL and two triazole units were used to represent the fluorophore and recognition group, respectively. The synthesized sensor could then be used as a specific fluorescent chemical sensor for Ag⁺ in the presence of a large number of competing cations, as anticipated.

Experimental

Materials and methods

All the solvents were of analytical purity, and the materials were obtained from commercial suppliers or prepared by our laboratory, with no further purification of the commercially supplied chemicals before use. If not otherwise specified, the solvents used in the optical spectroscopic studies were of spectroscopic purity. Various metal ion solutions (0.1 M) were prepared from their respective nitrates in distilled-deionized water, except for K⁺, Hg²⁺, Mn²⁺, and Ba²⁺, which were made from their chloride salts. AgNO₃ was used as the Ag⁺ source unless otherwise stated.

1H nuclear magnetic resonance (NMR) and 13C NMR were measured on a Bruker AM-400WB spectrometer using
tetramethylsilane (TMS) as an internal standard and CDCl₃ or CD₂OD as solvents. All UV-Vis absorptions were recorded on an Agilent 8453 UV-Vis spectrometer. Fluorescence emission spectra were obtained using a Hitachi F-4500 fluorescence spectrometer at 298 K, unless otherwise stated. Electrospray ionization mass spectrometric (ESI-MS) data were recorded using a Thermo Fisher LCQ. Melting points were measured on a WRS-1B melting point apparatus. Optical rotation was carried out using a Rudolph AUTOPOL IV automatic polarimeter.

**Synthesis of (S,β-ν)-2**

S-2′,2′-Bis(prop-2-yn-1-yloxy)-1,1′-binaphthalene (0.63 g, 1.75 mmol) and 2,3,4,6-tetra-O-acetyl-beta-D-glucopyranosyl azide (1.4 g, 3.67 mmol) were added to 50 mL of tetrahydrofuran (THF) with stirring at 273 K under an argon atmosphere and the mixture was stirred for five minutes. Sodium ascorbate (0.69 g, 3.48 mmol) and CuSO₄·5H₂O (0.44 g, 1.76 mmol) were added to the mixture and the temperature slowly rose to room temperature, after which it was stirred for 12 h under Ar₂. After the reaction was completed, the mixture was poured into ice water. The mixture was extracted three times with EtOAc and then the organic layer was washed with brine and dried over anhydrous MgSO₄. After evaporation of the organic solvent, the crude product was purified directly by column chromatography on silica (petroleum ether : EtOAc = 1 : 1, v/v) to give the desired product (S,β-ν)-2 (1.7 g, 41.9%) as a white solid: [α]D25 ¼ 41.2 (c 0.1, CH3OH). ¹H NMR (400 MHz, CDCl₃) δ 7.99 (d, J = 9.0 Hz, 1H), 7.93 (d, J = 8.1 Hz, 1H), 7.49 (d, J = 9.0 Hz, 1H), 7.39 (t, J = 7.4 Hz, 1H), 7.25 (d, J = 7.0 Hz, 1H), 7.19–7.12 (m, 2H), 5.79 (d, J = 9.4 Hz, 1H), 5.42–5.30 (m, 2H), 5.25 (t, J = 9.2 Hz, 1H), 5.19 (s, 2H), 4.30 (dd, J = 12.6, 5.1 Hz, 1H), 4.13 (d, J = 12.7 Hz, 1H), 3.93 (dd, J = 9.5, 3.1 Hz, 1H), 2.11–2.02 (m, 9H), 1.73 (d, J = 20.0 Hz, 1H), ¹C NMR (400 MHz, CDCl₃) δ 170.32, 169.57, 168.55, 153.68, 154.53, 133.93, 131.53–130.59, 130.04, 128.87, 127.29, 126.91–124.70, 124.70–124.15, 123.23, 122.24, 120.43, 116.66, 115.07, 86.28, 84.69, 74.72, 77.10, 76.78, 75.63, 74.22, 73.88, 72.06, 71.00, 69.46, 68.38, 66.85, 65.10, 63.41, 61.94, 60.21, 22.60, 21.95, 21.30, 20.3, 19.97–19.70, 19.70–17.92 ppm. HRMS (ESI⁻): calcd for [C₅₋H₄₋N₆₋O₁₂₋+ Cl⁻] 844.3; found 844.3.

**Synthesis of (S,β-ν)-1**

A mixture of (S,β-ν)-2 (1.3 g, 1.17 mmol), NaOH (0.47 g, 11.7 mmol), and methanol (100 mL) was stirred at room temperature for 12 h. After the reaction was completed, the solvent was removed under reduced pressure to obtain the crude product. The product was purified directly by flash column chromatography on silica gel using dichloromethane and methanol in a ratio of 2 : 1 (v/v) as the eluent to give the desired product 1 (0.75 g, 82.8%) as a white solid: [α]D25 ¼ 54 (c 0.07, CH3OH). ¹H NMR (400 MHz, MeOD) δ 8.02 (d, J = 6.3 Hz, 1H), 7.92 (d, J = 8.1 Hz, 1H), 7.69–7.54 (m, 1H), 7.34 (d, J = 9.8 Hz, 2H), 7.23 (s, 1H), 7.06 (d, J = 8.5 Hz, 1H), 5.45 (d, J = 9.1 Hz, 1H), 5.27–5.04 (m, 2H), 3.87 (dd, d = 10.9 Hz, 1H), 3.68 (dd, J = 10.8, 6.9 Hz, 2H), 3.62–3.41 (m, 3H), 3.31 (s, 4H) ppm. ¹³C NMR (101 MHz, MeOD) δ 153.77, 144.31, 133.93, 129.77, 129.49, 127.90, 125.73, 125.16, 123.86, 122.80, 120.48, 115.77, 87.99, 79.65, 77.19, 72.65, 69.82, 63.15–62.26, 61.12 ppm. HRMS (ESI²⁺): calcd for [C₃₀H₄₄N₆O₁₂₋+ Cl⁺] 807.2; found 807.2.

**Fluorescence and UV-visible measurements**

A stock solution of (S,β-ν)-1 (2.0 × 10⁻⁵ mol L⁻¹) prepared in methanol and stock solutions of the metal ions (0.1 mol L⁻¹ M in H₂O) were freshly prepared before testing each performance. For each fluorescence quenching measurement, varying equivalents of Ag⁺ stock solution were added to the sensor solutions in a 5 mL volumetric flask at 298 K. The competition experiments involved mixing of each metal ion solution with a stock solution of Ag⁺ of the same concentration (2.0 × 10⁻⁵ mol L⁻¹). The Job plot for the complexation of (S,β-ν)-1 with Ag⁺ was obtained by recording the fluorescence response of (S,β-ν)-1 with different ratios of Ag⁺. A 2.0 mM stock solution of (S,β-ν)-1 dissolved in methanol and a 2.0 mM AgNO₃ in H₂O were freshly prepared for each measurement.

**Preparation of the samples for NMR tests**

AgNO₃ (0.1 M in D₂O) was gradually added to an NMR tube containing (S,β-ν)-1 (0.4 mL, 25 mM) to obtain 1 : 0, 1 : 0.1, 1 : 0.3, 1 : 0.5, 1 : 0.7, and 1 : 1 ratios of the sensor to Ag⁺. The resulting solutions were allowed to equilibrate at 298 K for 4 h before testing.

**Results and discussion**

The BINOL–glucose derivative fluorescent sensor was synthesized using “click chemistry”, as shown in Scheme 1. According to the previous literature, the dipropargyl 3 (ref. 41) derivative of (S)-BINOL was prepared (Scheme 1). The reaction of dipropargyl 3 and azide-functionalized glucose 4 was carried out in THF at room temperature in the presence of sodium ascorbate and copper(II) sulfate to afford BINOL–glucose derivative 2 in moderate yield after a simple purification process. The new fluorescent sensor 1 was obtained when 2 was hydrolyzed in methanol, in 83% yield. The structures of the desired compounds were determined from ¹H NMR, ¹³C NMR, and ESI-MS measurements.

**Fluorescence study**

The fluorescence responses of (S,β-ν)-1 were studied in the presence of Ag⁺, Ba²⁺, Cd²⁺, Mg²⁺, Ca²⁺, Cr³⁺, Al³⁺, Ca²⁺, Co²⁺, Cu²⁺, K⁺, Ni²⁺, Mn²⁺, Zn²⁺, Hg²⁺, Sn²⁺, Pb²⁺, and Sr²⁺ ions in

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**Scheme 1** Synthetic route for compound (S,β-ν)-1
different solutions using fluorescence spectroscopy. First, the fluorescence measurements of \((S,\beta-v)-1\) were carried out in \(\text{CH}_3\text{OH}\) \([([S,\beta-v]-1) = 20 \text{ mM}\). As shown in Fig. 1, only the addition of \(\text{Ag}^+\) ions to a solution of \((S,\beta-v)-1\) could result in an almost complete quenching of the fluorescence, however, other metal ions induced no obvious change in the fluorescence response. When the fluorescence selectivity experiments of \((S,\beta-v)-1\) were tested in THF solution (as shown in Fig. S8†), no significant difference in the fluorescence response was found. The discrimination between the different metal ions showed a sensitive dependence on the solvent. So, all of the fluorescence measurements of \((S,\beta-v)-1\) were investigated in methanol solution, in which \((S,\beta-v)-1\) was found to be highly selective and sensitive towards \(\text{Ag}^+\). An unexpected quenching of the fluorescence of the excimer emission upon addition of \(\text{Ag}^+\) ions to a solution of 1 may be due to PET (photoinduced electron transfer). In other words, the metal ions combined with electron acceptor triazole units and the glucose units behaved as a PET donor. This means that the two triazole units of \((S,\beta-v)-1\) form a selective and effective metal ion binding site.

**Metal ion competition studies**

For an effective cation probe, the key factor is the ability to detect a specific metal ion in the presence of different metal ions. Fig. 2 shows the results of competition experiments we conducted in which we tested the ability of the probe to selectively detect one metal ion over other metal ions. The addition of 5.0 equiv. of \(\text{Ag}^+\) combined with 5.0 equiv. of the other metal ions in methanol solutions of \((S,\beta-v)-1\), respectively, were used in the competitive experiments. The quenching ratio of the \(I_F/I_0\) value at 375 nm (where \(I_0\) represents the fluorescence intensity of only \((S,\beta-v)-1\) and \(I_F\) represents the fluorescence intensity upon the addition of a mixture of competitive metal ions and \(\text{Ag}^+\)) for the majority of the competitive metal ions was almost 0.17. No significant interference was observed in the presence of the various competitive metal ions. This indicated that the BINOL–glucose sensor could be used to detect \(\text{Ag}^+\) in the presence of various competitive ions without any obvious interference from the other metal ions. Based on the fluorescence results, we found that the BINOL–glucose derivative has high selectivity and sensitivity to \(\text{Ag}^+\) and therefore can be used as an \(\text{Ag}^+\) sensor.

The binding tests of \((S,\beta-v)-1\) towards \(\text{Ag}^+\) were also studied upon the addition of different amounts of \(\text{Ag}^+\) to \((S,\beta-v)-1\) in \(\text{CH}_3\text{OH}\), using UV-Vis absorption spectroscopy. As shown in Fig. 3, the maximum absorption wavelength of \((S,\beta-v)-1\) was found to be around 229 nm upon the addition of 0 to 3 equivalent of \(\text{Ag}^+\). However, there was found to be a slight change in the UV-Vis absorption spectrum of \((S,\beta-v)-1\) in the presence of various concentrations of \(\text{Ag}^+\), where the absorbance at 210 nm was observed to begin to increase, which may indicate the formation of a \((S,\beta-v)-1–\text{Ag}^+\) complex. It is a pity that the slight absorbance intensity shift did not induce any obvious color change. The interaction between the host and guest was evaluated using fluorescence spectroscopy. From Fig. 3, it can be seen that \((S,\beta-v)-1\) has three absorption peaks at 229 nm, 290 nm and 335 nm. The maximum absorption wavelength \(\lambda_{\text{max}}\) of \((S,\beta-v)-1\) was around 229 nm with weak fluorescence emission. However, the absorption at 290 nm was observed to have a stronger fluorescence intensity (see Fig. S9†), and so, \(\lambda_{\text{ex}} = 290\text{ nm}\).
As shown in Fig. 4, the fluorescence spectra of (S,β-D)-1 upon adding different equivalents of Ag⁺ were used to determine the nature of the complex formed between (S,β-D)-1 and Ag⁺. When the amount of Ag⁺ used was above 3 equiv., the intensity of the fluorescence emission underwent a strong change. We assumed that the stoichiometry ratio of (S,β-D)-1–Ag⁺ complex is 1 : 1, so the association constant of (S,β-D)-1 with Ag⁺ was found to be 2.4 × 10⁶ M⁻¹ (R = 0.997) from the Lineweaver–Burk plot of 1/(F₀ – F) versus 1/[Ag⁺]. As shown in Fig. 5, the maximum fluorescence quenching of (S,β-D)-1 provided by Ag⁺ takes place at a ratio of 1 : 1. Based on the fluorescence analyses, the BINOL–glucose compound and Ag⁺ were proposed to form a 1 : 1 complex. The Job plot for the complex provided further direct evidence that the (S,β-D)-1–Ag⁺ complex stoichiometry ratio was 1 : 1. Respectively, the detection limit (LOD) of (S,β-D)-1 to Ag⁺ was assessed to be 1.57 × 10⁻⁹ mol L⁻¹ using the following equation: LOD = 3σ/s, where σ represents the standard deviation of the (S,β-D)-1 solution and s represents the slope between the fluorescence intensity versus the Ag⁺ concentration (Fig. S11†). Further evidence for the 1 : 1 complex stoichiometry ratio was obtained from ESI-MS spectra data (see Fig. S7, ESI data†). The peak at m/z = 881.1 (calcd 881) was found to correspond to [1–Ag⁺ + H⁺], which supports our assumption that there is strong binding between (S,β-D)-1 and Ag⁺.

The 1H NMR experiments were carried out in CD₃OD to find out further detailed information on the binding of Ag⁺ with (S,β-D)-1. As shown in Fig. 6, different equiv. (from 0 to 1 equiv.) of Ag⁺ were added to a solution of (S,β-D)-1 and resulted in an obvious downshift in the chemical shifts, however, the shifts tended to saturation when the amount of Ag⁺ was beyond 1 equiv. The H₅ nuclear magnetic peaks of the –OCH₂⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻吸入一枚の例文を示しています。
complex between BINOL–glucose 1 and Ag⁺ without any interference from various different metal ions. The fluorescence quenching was ascribed to both the triazole nitrogen atoms and the oxygen atoms of (S,b-f)-1 involved in the coordination with Ag⁺. These results show the possibility of preparing highly selective fluorescent sensors based on the versatile BINOL backbone for the detection of various metal ions.

Conflicts of interest

There are no conflicts to declare.

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