Chiral Cryptands Possessing Tetraazamacrocyclic and BINAM Moieties: Synthesis and Evaluation as Fluorescent Detectors

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Pd(0)-Catalyzed amination of N,N'-di(bromobenzyl) substituted tetraazamacrocycles (cyclen, cyclam) with (S)-2,2'-diamino-1,1'-binaphthalene (BINAM) was employed for the synthesis of a novel type of chiral cryptands. Better yields were obtained for cyclen derivatives. The compounds were modified with dansyl groups to enhance their fluorescent properties. Cryptands theirselves and their dansyl derivatives were evaluated for sensing enantiomers of 7 amino alcohols and 21 metal cations.

Keywords: Macrocycles, Pd catalysis, chirality, fluorescence, detection

Хиральные криптанды, содержащие фрагменты тетраазамакроциклов и БИНАМа: синтез и оценка в качестве флуоресцентных детекторов

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Pd(0)-Катализируемое аминирование N,N'-ди(бромбензил)замещенных тетраазамакроциклов (циклен, циклам) с (S)-2,2'-дiamoно-1,1'-бинафталеном (БИНАМом) использовано для синтеза хиральных криптандов нового типа. Более высокие выходы получены для производных циклена. Полученные соединения модифицировали даньсильными группами для улучшения флуоресцентных свойств соединений. Криптанды и их даньсильные производные были исследованы в детектировании энантиомеров 7 аминоспиртов и 21 катиона металлов.

Ключевые слова: Макроциклы, Pd каталаз, хиральность, флуоресценция, детектирование.
Introduction

Fluorescent analytical methods are highly demanded due to their sensitivity, selectivity and wide opportunities of adjusting receptor and signaling units to certain analytes. Beginning from 1992, 1,1′-binaphthol (BINOL) has been employed as a basic moiety for creating chemosensors for fluorescence detection of chiral organic compounds. This is due to unique properties of BINOL: it combines C2 chirality, fluorescent properties and possesses two oxygen atoms which can be employed in various structural modifications.[11] BINOL derivatives were used mainly for sensing amino acids and their derivatives. For instance, 3,3′-disubstituted BINOLs were found to be useful for detecting N-Boc-alanine and N-Boc-phenylalanine by fluorescence quenching,[2,3] emission enhancement was used for mandelic amino acids and their derivatives. For instance, 3,3′-disubstituted exocyclic chiral ethers containing macrocycles and their use in fluorescent enantiomers sensing with a BINOL-diamine derivative.[4,5]

The same analyte can be detected by a macrocyclic sensor combining two BINOL and two coplanar porphyrins was contrived for better optical response. Even a cryptand possessing two tert coordination.

This method was used for the synthesis of macrobicycles 6-9. A two-neck flask equipped with a magnetic stirrer and reflux condenser, flushed with dry argon, was charged with Pd(dbca) (9 mg, 0.016 mmol), BINAP (11 mg, 0.018 mmol), (S)-BINAM 5 (0.2 mmol, 57 mg), corresponding N,N′-di(bromomethyl) substituted cyclon or cyclam (0.2 mmol, 102 mg or 107 mg), absolute dioxane (8 ml) and sodium tert-butoxide (0.6 mmol, 58 mg). The reaction mixture was stirred under reflux for 24 h, cooled to ambient temperature, solution was filtered, residue was washed with 10 ml of dichloromethane, combined organic phases were evaporated in vacuo, and the residue was chromatographed on silica gel using a sequence of eluents: CH$_2$Cl$_2$, CH$_2$Cl$_2$-MeOH 10:1–3:1. Yield 29 mg (23 %), yellowish solid. (MALDI-TOF) found: 633.3671. C$_{31}$H$_{56}$N$_6$ requires 633.3706 [M+H]. 1H NMR (CDCl$_3$, 298 K) δ ppm: 2.41–2.47 m (2H), 2.58–2.75 m (6H), 2.83–2.93 m (6H, 2.94–3.02 m (2H), 3.57 d (2H, J = 12.8 Hz), 3.69 d (2H, J = 12.8 Hz), 5.76 s (2H), 6.74 d (2H, J = 7.6 Hz), 6.99 d (2H, J = 8.1 Hz), 7.07–7.10 m (4H), 7.14 t (2H, J = 7.8 Hz), 7.20–7.24 m (2H), 7.28–7.32 m (2H), 7.73 d (2H, J = 9.0 Hz), 7.84 d (2H, J = 7.8 Hz), 7.91 d (2H, J = 9.0 Hz), two NH protons were not assigned. 13C NMR (CDCl$_3$) (23 %), yellowish solid. (MALDI-TOF) found: 48.2 (2C), 48.5 (2C), 51.8 (2C), 52.6 (2C), 62.9 (2C), 116.3 (2C), 117.0 (2C), 118.8 (2C), 119.6 (2C), 123.2 (2C), 123.5 (2C), 124.5 (2C), 127.1 (2C), 128.1 (2C), 129.2 (2C), 129.3 (2C), 129.5 (2C), 134.0 (2C), 136.9 (2C), 140.0 (2C), 142.5 (2C).

Macrocyclit 10. Obtained as the second product in the synthesis of the cryptand 6. Eluent: CH$_2$Cl$_2$–MeOH–NH$_3$ (aq) 100:20:1. Yield 17 mg (13 %), yellowish solid. (MALDI-TOF) found: 1265.7626. C$_{31}$H$_{56}$N$_6$ requires 1265.7333 [M–H]. 1H NMR (CDCl$_3$, 298 K) δ ppm: 2.40–2.70 m (32H), 3.36 d (4H, J = 13.0 Hz), 3.46 d (4H, J = 13.0 Hz), 5.50 s (4H, 4.64 brd (4H, J$_{1,2}$= 7.0 Hz), 6.74–6.80 m (8H, 6.99 t (2H, J = 7.7 Hz), 7.12 d (4H, J = 8.2 Hz), 7.15–7.20 m (4H), 7.22–7.26 m (4H), 7.56 d (4H, J = 9.0 Hz), 7.80 d (4H, J = 8.5 Hz), 7.83 d (4H, J = 9.0 Hz), four NH protons were not assigned. 13C NMR (CDCl$_3$, 298 K) δ ppm: 55.8 (8C), 51.8 (8C), 60.0 (4C), 116.0 (4C), 117.8 (4C), 119.2 (4C), 121.2 (4C), 123.1 (4C), 123.4 (4C), 124.3 (4C), 127.0 (4C), 128.2 (4C), 129.1 (4C), 129.3 (4C), 129.4 (4C), 133.8 (4C), 140.1 (4C), 140.8 (4C), 142.5 (4C).

Crypand 7. Eluent: CH$_2$Cl$_2$–MeOH–NH$_3$ (aq) 100:20:1. Yield 44 mg (34 %), yellowish solid. (MALDI-TOF) found: 633.38. C$_{31}$H$_{56}$N$_6$ requires 633.37 [M–H]; found: 631.3593. C$_{42}$H$_{41}$N$_6$ requires 629.3393 [M–H]. 1H NMR (CDCl$_3$, 298 K) δ ppm: 2.35–2.50 m (2H), 2.51–2.65 m (6H), 2.65–2.80 m (8H), 3.40 d (2H, J = 14.0 Hz), 3.54 d (2H, J = 14.0 Hz), 4.66 brs (2H), 5.58 s (2H), 6.73 d (4H, J = 8.3 Hz), 6.96 d (4H, J = 8.3 Hz), 7.24 d (2H, J = 7.8 Hz), 7.29–7.33 m (2H), 7.35–7.39 m (2H), 7.48 d (2H, J = 8.9 Hz), 7.85–7.88 m (4H). 13C NMR (CDCl$_3$, 298 K) δ ppm: 46.8 (2C), 51.3 (2C), 52.4 (2C), 52.6 (2C), 61.1 (2C), 120.2 (2C), 120.9 (4C), 123.8 (2C), 124.6 (2C), 127.1 (2C), 128.3 (2C), 128.9 (3C), 129.7 (3C), 132.9 (2C), 134.0 (2C), 140.6 (2C), 142.4 (2C).

Crypand 8. Eluent: CH$_2$Cl$_2$–MeOH–NH$_3$ (aq) 100:20:3. Yield 22 mg (16 %), yellowish solid. (MALDI-TOF) found: 661.3967. C$_{31}$H$_{56}$N$_6$ requires 661.4019 [M–H]. 1H NMR (CDCl$_3$, 298 K) δ ppm: 1.58–1.66 m (1H), 1.66–1.78 m (2H), 1.78–1.86 m (1H), 2.23–2.74 m (12H), 2.81–2.96 m (9H), 3.02 d (1H, J = 14.0 Hz), 3.05–3.09 m (1H), 3.13 d (1H, J = 14.0 Hz), 3.57 d (1H, J = 14.0 Hz), 3.83 d (1H, J = 14.0 Hz), 5.75 s (1H), 5.81 s (1H), 6.75 d (1H, J = 7.5 Hz), 6.79 d (1H, J = 7.5 Hz), 6.92 d (1H, J = 7.8 Hz), 7.00 d (1H, J = 8.2 Hz), 7.06–7.14 m (4H), 7.19–7.21 m (6H), 7.51 d (1H, J = 9.0 Hz), 7.62 d (1H, J = 9.0 Hz), 7.79–7.84 m (3H), 7.85 d (1H, J = 9.0 Hz), two NH protons were not assigned.

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\(^{13}\)C NMR (CDCl\(_3\), 298 K) \(\delta\) ppm: 25.6, 25.9, 46.8, 49.0, 49.8, 50.1, 51.0, 52.5, 54.6 (2C), 57.3, 57.7, 116.4, 117.5, 118.0 (2C), 119.1, 119.4, 121.5, 122.9, 123.0, 123.1, 123.2, 123.4, 124.4, 124.6, 126.8 (2C), 128.0, 128.1, 128.4, 128.8, 129.2 (2C), 129.3, 133.8, 134.1, 140.6, 140.8, 141.0, 141.7, 142.3, 142.7.

**Cryptand 9.** Eluent: CH\(_2\)Cl\(_2\)/MeOH 3:1. Yield 18 mg (13 %), yellowish solid. (MALDI-TOF) found: 661.4068 C\(_{26}\)H\(_{32}\)N\(_8\)O\(_4\) requires 661.4019 [M+H]+. \(^{1}H\) NMR (CDCl\(_3\), 298 K) \(\delta\) ppm: 1.79–2.80 m (2H), 2.81–2.91 m (4H), 2.90–2.92 m (6H), 2.97–2.98 m (6H), 3.09 (2H, \(J = 13.0\) Hz), 3.66 d (2H, \(J = 8.1\) Hz), 5.67 s (2H, \(J = 8.1\) Hz), 5.89 d (2H, \(J = 8.1\) Hz), 7.02 d (2H, \(J = 8.1\) Hz), 7.09 d (2H, \(J = 8.1\) Hz), 7.22–7.35 m (8H), 7.40–7.48 m (4H), 7.50–7.58 m (4H), 7.67–7.76 m (2H), 7.85–7.94 m (4H), two NH protons were not assigned.

Dansyl-substituted cryptands:

**Cryptand 12.** A flask equipped with a magnetic stirrer was charged with the cryptand 6 (0.03 mmol, 19 mg), dansyl chloride 11 (0.066 mmol, 19 mg), K\(_2\)CO\(_3\) (0.12 mmol, 18 mg) and 1 ml MeCN. The reaction mixture was stirred at ambient temperature for 24 h, the residue was filtered, washed with 5 ml dichloromethane, combined organic fractions were evaporated in vacuo, the residue was dissolved in dichloromethane (3 ml), washed with water (3×5 ml), dried over molecular sieves 4 Å, evaporated in vacuo, and the target compound 12 was obtained as a yellow glassy compound. Yield 30 mg (90 %). (MALDI-TOF) found: 1099.4802 C\(_{66}\)H\(_{67}\)N\(_8\)O\(_3\) requires 1099.4727 [M+H]+. \(^{1}H\) NMR (CDCl\(_3\), 298 K) \(\delta\) ppm: 1.60–1.68 m (2H), 2.00–2.05 m (2H), 2.14 dt (2H, \(J = 13.4\) Hz, \(J = 4.4\) Hz), 2.29–2.36 m (2H), 2.40 s (12H), 2.52 d (2H, \(J = 12.9\) Hz), 2.56 d (2H, \(J = 12.9\) Hz), 2.83–2.90 m (4H), 3.32–3.37 m (4H), 3.40–3.49 m (2H), 3.64–3.73 m (2H), 5.67 s (2H), 5.89 d (2H, \(J = 7.6\) Hz), 6.25–6.29 m (4H), 6.43 d (2H, \(J = 7.6\) Hz), 7.01 t (2H, \(J = 7.6\) Hz), 7.11–7.17 m (4H), 7.27–7.33 m (6H), 7.77 d (2H, \(J = 9.0\) Hz), 7.82 d (2H, \(J = 7.6\) Hz), 7.91–7.96 m (4H), 8.02 d (2H, \(J = 8.9\) Hz), 8.06 d (2H, \(J = 8.3\) Hz).

The investigations of the spectral properties of the cryptands 6-9 and 13 in the presence of different chiral molecules were carried out in a following manner: 3 ml of the solution of 6 (C = 31.0 μM), or 7 (C = 16.0 μM), or 8 (C = 15.0 μM), or 9 (C = 12.1 μM), or 13 (C = 12.5 μM), in MeCN was placed in a spectrofluorimeter cuvette, solutions of appropriate chiral compounds (S)- and (R)-leucinol, (S)- and (R)-tert-leucinol, (S)- and (R)-phenylglycinol, (S)- and (R)-2-amino-propan-1-ol, (S)- and (R)-2-amino-1,2-diphenylethanol, in MeCN (C = 0.2 M) were added sequentially (100, 200, 500, 1000 equiv.) and after each addition UV-Vis and fluorescence spectra were recorded. Also the solutions of Li4, Na4, K4, Ag4, Mg4, Ca4, Ba4, Al4, Fe4, Mn4, Co4, Ni4, Cu4, Zn4, Cd4, Pb4 perchlorates and Ga4, In4, Y4 nitrates in MeCN (C = 0.01 M) were added sequentially to the solutions of the cryptands 6-9 and 12 in MeCN (1, 2, and 5 equiv.) and after each addition UV-Vis and fluorescence spectra were recorded.

Results and Discussion

The synthesis of the target cryptands was accomplished using Buchwald-Hartwig amination applied to the macrocyclization reactions. They were successfully employed by us earlier to obtain various macrobicyclic and macrotricyclic derivatives of cyclo(1,4,7,10-tetraazaacyclodecane) and cyclam(1,4,8,11-tetraazaacyclotetradecane).[24-31] The starting compounds, N,N′-dibromomethyl) substituted cyclens 1, 2 and cyclams 3, 4 were synthesized in high yields from free tetraazaacmacrocycles by three-step procedures described previously.[24-25] The reactions of these compounds with (S)-BINAM (5) were carried out in the presence of the catalytic system Pd(dba)3/BINAP (8:9 mol%) in boiling dioxane at the concentration of the starting compounds 0.02 M using sodium tert-butoxide as a base (Scheme 1). After the reactions were over, target cryptands were isolated from the reaction mixtures using column chromatography. This approach proved to be quite efficient in the present case as it provided the desired cryptands 6-9 in yields up to 34 %. It is to be noted, however, that the steric constraint in this cyclization reaction is quite strict due to the rigid structure of BINAM with a short distance between the two amino groups. It is not surprising that for different bromobenzyl derivatives of tetraazaacmacrocycles the outcome of the reaction was different. As usual, macrocyclization reaction

Cycloadition reaction was charged with the cryptand 8 (0.026 mmol, 17 mg), dansyl chloride 11 (0.130 mmol, 33 mg), K2CO3 (0.2 mmol, 28 mg) and 1 ml MeCN. The reaction mixture was stirred at ambient temperature for 24 h, the residue was filtered, washed with 5 ml dichloromethane, combined organic fractions were evaporated in vacuo, the residue was dissolved in dichloromethane (3 ml), washed with water (3×5 ml), dried over molecular sieves 4 Å, evaporated in vacuo, and the target compound was synthesized in high yields from free tetrazaacmacrocycles by three-step procedures described previously.[24-25] The reactions of these compounds with (S)-BINAM (5) were carried out in the presence of the catalytic system Pd(dba)3/BINAP (8:9 mol%) in boiling dioxane at the concentration of the starting compounds 0.02 M using sodium tert-butoxide as a base (Scheme 1). After the reactions were over, target cryptands were isolated from the reaction mixtures using column chromatography. This approach proved to be quite efficient in the present case as it provided the desired cryptands 6-9 in yields up to 34 %. It is to be noted, however, that the steric constraint in this cyclization reaction is quite strict due to the rigid structure of BINAM with a short distance between the two amino groups. It is not surprising that for different bromobenzyl derivatives of tetraazaacmacrocycles the outcome of the reaction was different. As usual, macrocyclization reaction
proceeds in poorer yields in the cyclam series. In one case, namely with $N,N'$-di(3-bromobenzyl)cyclen, we managed to isolate the cyclic dimer 10, actually a macrotricycle, as a by-product in 13% yield. In the same reaction a mixture of cyclic trimers and tetramer was also obtained as a separate fraction (yield 20%). This compound is of interest as it possesses a larger macrocyclic cavity compared to the cryptands 6-9. In other cases, only fractioned mixture of cyclic and linear oligomers was obtained by chromatography.

Next, we modified the synthesized macrobicycles with dansyl fluorophore groups. For this purpose the cryptands 6-9 were reacted with 2.2 equiv. of 5-(dimethylamino)naphthalene-1-sulfonyl chloride (dansyl chloride 11) in MeCN in the presence of $K_2CO_3$ at ambient temperature (Scheme 2). Macrobicycles 6-8 produced corresponding didansyl derivatives 12-14 in yields from 30 to 90%, while compound 9 was unable to afford the target product. Such difference in the reactivity of the cryptands is supposedly due to the difference in the steric hindrances at the secondary amino groups of the tetraazamacrocycles.

Indeed, quite close position of the binaphthalene moiety to the tetraazamacrocycle may hinder the alkylation of these amino groups with naphthalene-1-sulfonyl chloride. Didansyl derivatives 13 and 14 were isolated using column chromatography on silica gel.

NMR spectra of the obtained cryptands possess the following peculiarities. In all macrobicycles protons in $NCH_2Ph$ methylene group are diastereotopic. Cyclen fragment in compounds 6, 7, 12 and 13 is characterized by four different carbon atoms and not two as was usual for many other achiral macrobicycles synthesized by us earlier. However, macrotricyclic dimer 10 displays only two different carbon atoms assigned for cyclen moiety which can be explained by a different reciprocal position of cyclen and BINAM fragments. In the case of cyclam derivative 8 the asymmetrical effect is even more pronounced as all protons and carbons in this molecule are inequivalent thus leading to a complicated spectral pattern. On the other hand, in other cyclam derivatives (9 and 14) symmetry is not disturbed, benzyl spacers and naphthalene moieties...
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Obtained cryptands were tested for their ability to perform as fluorescent enantioselective detectors using optically pure amino alcohols (7 pairs of enantiomers). The analytes structures are given on Figure 1. Experiments were carried out with the cryptands 6-9, and 13 in MeCN, corresponding analytes were added stepwise (100, 200, 500, 1000 equiv.) and fluorescent spectra were measured in each case. Compounds 6-9 possess an absorption band near $\lambda_{\text{max}} = 350$ nm, and being excited at this wavelength they emit at 414 nm (6), 432 nm (7), 422 nm (8), 435 nm (9). The cryptand 6 did not change notably its fluorescence in the presence of both isomers of such amino alcohols like lecinol, tert-leucinol, 2-aminopropan-1-ol and 2-amino-1,2-diphenyl ethanol. In the case of 2-aminobutan-1-ol, the addition of its (S)-isomer (here and after 1000 equiv. of each enantiomer are considered) quenched emission by 20 % while (R)-isomer quenched it 1.5 times. Similarly, both isomers of phenylglycinol led to fluorescence quenching, (S)-isomer was more efficient (1.5 times) than (R)-isomer (by 40 %), moreover, in both cases bathochromic shift of the emission maxima by 20-25 nm was observed. Thus, compound 6 cannot be used for distinguishing between selected amino alcohols.

As for 7, the addition of enantiomers of the majority of amino alcohols did not change notably the fluorescence intensity. In one case, with both (S)- and (R)-isomers of 2-amino-1,2-diphenylethanol tiny quenching of emission was noted, and only the addition of the enantiomers of 2-phenylglycinol led to different results: with (S)-isomer the emission enhanced by ca 20 % while with (R)-isomer its intensity did not change (Figure 2). It makes possible to claim that the cryptand 7 is an efficient chemosensor for this amino alcohol.

The investigation of the compound 8 showed that the addition of both isomers of valinol resulted in the emission quenching: with L-isomer by 15 %, with D-isomer by 40 %. On the other hand, in the presence of both enantiomers of phenylglycinol or 2-amino-1,2-diphenyl ethanol emission enhancement was observed, more pronounced in the second case. The cryptand 8 turned to be unable to discriminate any of amino alcohols enantiomers.

The fluorescence of the cryptand 9 was susceptible to the presence of phenylglycinol enantiomers: the addition of its (R)-isomer led to an increase in the fluorescence intensity by 15 % while (S)-isomer increased it by 40 %; simultaneously weak bathochromic shifts of the maxima (ca 5–10 nm) were noted. The addition of L-valinol caused a tiny emission quenching, and with D-valinol it was somewhat more pronounced. In fact, the compound 9 was found to be able to distinguish between 2-aminobutan-1-ol enantiomers: the presence of its (S)-isomer does not change the spectrum at all and its (R)-isomer causes 15 % emission quenching (Figure 3). Thus, both cryptands 7 and 9 which were found to be more selective towards certain amino alcohols, comprise para-phenylene spacer.

Unfortunately, the modification of the macrobicycles with two dansyl fluorophores was not helpful and all amino alcohols were detected by the cryptands 7 and 9.
alcohols under investigation did not cause significant changes in the fluorescence intensity of compound 13. Probably, this is due to a strong shielding of the coordination sites with dimethylaminonaphthalene groups preventing the formation of the molecular complexes with even small organic compounds.

The fluorimetric investigations were continued with metal salts (also in MeCN). Totally 21 metal cations were used for this purpose: Li⁺, Na⁺, K⁺, Mg²⁺, Ca²⁺, Ba²⁺, Al³⁺, Cr³⁺, Mn²⁺, Fe²⁺, Co³⁺, Ni²⁺, Cu²⁺, Zn²⁺, Cd²⁺, Pb²⁺, Ag⁺, Hg²⁺ (perchlorates), Ga³⁺, In³⁺, Y³⁺ (nitrates). The addition of the metal salts led to much more pronounced changes in fluorescence with all cryptands tested. Some metals led to emission quenching, the most important effect was noted for Hg²⁺ (by 70 %) and especially by Cu²⁺ which produced full quenching (Figure S1, see Supporting Information at https://macroheterocycles.isuct.ru/en/mhc190337a). It was shown that the addition even of the first equivalent of this cation led to a dramatic decrease in the intensity of the emission (Figure S2). Some metals caused tiny emission enhancement, only in the presence of In³⁺ it increased substantially accompanied by a hypsochromic shift of the maximum by 10 nm. It was shown that this shift was observed with the addition of the first equivalent of indium, and the enhancement was notable after the second equivalent was added (Figure S3). The maximal effect was achieved with 15 equiv. and further input of the salt did not change the spectrum. As for Ga³⁺, the maximal enhancement was noted for 5 equiv. of this metal while with 10 equiv. the intensity of fluorescence decreased to some extent (Figure S4). This effect may be explained by the formation of a new complex with different structure and spectroscopic features. Thus, the cryptand 6 can be considered as a molecular sensor for Cu²⁺ and In³⁺.

The fluorescence of the cryptand 7 also changes in the presence of most of the cations investigated (Figure S5): the addition of Co³⁺ and Ni³⁺ led to ca twofold decrease in the emission intensity while In³⁺, Pb²⁺, Cr³⁺, Al³⁺, Ga³⁺ and Zn²⁺ induced about twofold enhancement of the fluorescence intensity. These changes in emission were accompanied with small hypsochromic shifts (by less than 10 nm) in the majority of cases. However, the addition of only 2 equiv. of Cu²⁺ quenched totally the fluorescence (Figure S6), this can be helpful in the selective detection of this metal.

The addition of the metal salts to the macrobicycle 8 generally results in more or less pronounced emission quenching (Figure S7). The most efficient quenching was noted for Hg²⁺ (by 45 %), Co³⁺ (by 60 %) and Cu²⁺ (by 80 %). It is clearly seen from Figure S8 that unlike in the case of the two previous ligands 6 and 7, here the addition of even 10 equiv. of copper was not enough to achieve full emission quenching. However, with this metal cation the hypsochromic shift of the emission maximum from 422 to 406 nm was observed. This effect can distinguish copper among other metal cations. No enhancement of the fluorescence intensity was noted in any case, unlike it was common for cyclen-based cryptands 6 and 7.

As for the cryptand 9, most of metal cations more or less diminished the intensity of fluorescence. Three of them, Hg²⁺, Cd²⁺ and Cu²⁺ led to hypsochromic shifts of the emission maxima (by 5, 10 and 25 nm, respectively) (Figure S9). In the presence of these metals fluorescence quenched by 70, 65 and 85 %, respectively. Almost full emission quenching with Cu²⁺ was achieved with 10 equiv. (Figure S10). The macrobicycle 9 can be judged as a molecular probe for the above-mentioned metals. To note, cyclen-containing cryptands 6 and 7 undergo almost full emission quenching in the presence of 2–5 equiv. of Cu²⁺ cations, while the fluorescence of cyclam-containing macrobicycles 8 and 9 is quenched only partially by this metal added in comparable amounts. This fact is in a good correspondence with the known better stability of the cyclen-copper complex compared to cyclam analogue.

Quite different behavior of fluorescence in the presence of the studied metals was observed in the case of the dansyl-decorated cryptand 13 (Figure S11). Excitation of the ligand at 340 nm led to intensive emission at 512 nm. No enhancement of the emission was noted in the presence of any ligand, and many metals caused partial quenching of fluorescence. Its intensity diminished by 20–40 % in the presence of Al³⁺, In³⁺, Y³⁺, Fe²⁺, Ni²⁺, Zn²⁺, Pb²⁺, and Cu²⁺ led to 5-times decrease in the emission intensity. Moreover, generally bathochromic shifts of the emission maxima were observed (525–535 nm) after the addition of metal salts which caused quenching. To note, full emission quenching by the copper cations was achieved after adding 10 equiv. of the corresponding salt (Figure S12).

Conclusions

To conclude, in the course of the present work we used our previously elaborated method for the synthesis of BINAM-containing macrobicycles based on disubstituted cyclen and cyclam. The compounds were decorated with dansyl fluorophore groups. Spectrofluorimetric investigations of synthesized cryptands demonstrated the possibility of the cyclen-based compound 7 to distinguish between two enantiomers of 2-phenylglycinol and of cyclam-based macrobicycle 9 to detect enantiomers of 2-aminobutan-1-ol by different changes in the emission caused by the presence of enantiomers of said compounds. Spectrofluorimetric titrations of the ligands revealed the ability of the cryptand 7...
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to act as a selective fluorescent detector of Cu\textsuperscript{II} cations, while macrobicycles 6, 8, 9, and 13 were shown to be molecular sensors for Cu\textsuperscript{II}, Hg\textsuperscript{II}, Cd\textsuperscript{II} and In\textsuperscript{III}.

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