## Fullerene sensors based on calix[5]arene

## Takeharu Haino, $^a$ Hiromi Araki, $^a$ Yoshihisa Fujiwara, $^b$ Yoshifumi Tanimoto $^b$ and Yoshimasa Fukazawa\* $^a$

- <sup>a</sup> Department of Chemistry, Graduate School of Science, Hiroshima University, 1-3-1 Kagamiyama, Higashi-Hiroshima 739-8526, Japan. E-mail: fukazawa@sci.hiroshima-u.ac.jp
- <sup>b</sup> Department of Mathematical and Life Sciences, Graduate School of Science, Hiroshima University, 1-3-1 Kagamiyama, Higashi-Hiroshima 739-8526, Japan

Received (in Cambridge, UK) 27th May 2002, Accepted 2nd August 2002 First published as an Advance Article on the web 22nd August 2002

## A new class of fullerene sensors based on calix[5]arenes has produced the highly sensitive detection of $C_{60}$ and $C_{70}$ .

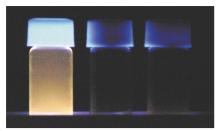
The development of chemical sensors has received broad attention in recent years. One of the most appealing approaches involves the combination of supramolecular concepts and luminescence techniques. A number of sensors<sup>1</sup> have been developed with a focus upon selective binding for metal ions, phosphates and neutral molecules coupled with luminescence signal modulation.

During our investigation on fullerene receptors, we have developed calix[5]arene-based fullerene receptors, which show high selective binding for  $C_{60}$  and/or  $C_{70}$ .<sup>2</sup> Among numerous studies on developing fullerene receptors reported so far,<sup>3</sup> the fullerene sensor is limited. We envisioned developing fullerene sensors. Our strategy for the fullerene sensor is based on the introduction of a covalently attached lumophore onto the fullerene binding site. Since it is well known that fullerene is a good acceptor in an energy transfer reaction, it might act as a good luminescence quencher. The luminescence of the sensor can be quenched when fullerene is situated in close proximity to the lumophore. This optical response should produce highly sensitive detection for fullerenes. Along this concept, sensors 1 and 2 have been designed.

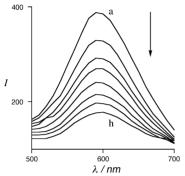
The synthesis of sensors 1 and 2 was started according to Scheme 1. Coupling reaction of iodocalix[5]arene 3<sup>4</sup> with methyl ester 4<sup>5</sup> through palladium catalysis, followed by hydrolysis of ester groups produced double-calix[5]arene carboxylic acid 5. Bipyridine 6<sup>6</sup> was introduced to 5 to give double-calix[5]arene 7. Reaction of 7 with Re(CO)<sub>5</sub>Cl in toluene at 100 °C produced sensor 1 in 60% yield. Bromobipyridine 8<sup>4</sup> reacted with 9<sup>7</sup> through Suzuki conditions to give 10. Coupling reaction of 10 with 3, followed by basic hydrolysis of acetyl groups furnished calix[5]arene derivative 11, which reacted with Re(CO)<sub>5</sub>Cl in toluene at 100 °C, produced sensor 2 in 46% yield.

Sensors 1 and 2 are soluble in toluene. Strong orange luminescence showed up when the solution of 1 was exposed to 365 nm UV light (Fig. 1). Addition of  $C_{60}$  or  $C_{70}$  to the solution of 1 caused a dramatic change. The luminescence was immediately extinguished upon the addition of  $C_{60}$  or  $C_{70}$ . The steady-state phosphorescence spectral change of 1 in toluene  $(\lambda_{\rm exc} = 400 \text{ nm})$  is shown in Fig. 2. The characteristic luminescence band appearing around 600 nm was assigned to the metal-to-ligand charge transfer (MLCT) band.8 Upon the addition of  $C_{60}$  to the solution of 1 in toluene, the luminescence intensity gradually decreased (Fig. 2). In order to discuss the quenching quantitatively, Stern-Volmer analysis was carried out for 1, 2 and Re(bpy)(CO)<sub>3</sub>Cl 12 with C<sub>60</sub> or C<sub>70</sub> in toluene. Steady-state luminescence quenching studies with C<sub>60</sub> yielded two linear Stern-Volmer plots for 2 and 12, as shown in Fig. 3. The results were interpreted in terms of diffusion-controlled intermolecular quenching between  $C_{60}$  and the lumophore. For 1 with  $C_{60}$  or  $C_{70}$ , the quenching is much more efficient than the others. The quenching of 1 with  $C_{60}$  or  $C_{70}$  produced curved Stern-Volmer plots (Fig. 3). There should be two luminescence quenching processes associated with the energy transfer from the lumophore to the fullerene: one is an intermolecular quenching process arising from the simple collision between the unbound fullerene and the lumophore, the other is an intra-

Scheme 1



**Fig. 1** Luminescence changes induced on sensor **1**  $(1.0 \times 10^{-5} \text{ mol L}^{-1})$ : (left) in the absence of fullerenes, (middle) with  $C_{60}$ , (right) with  $C_{70}$ .



**Fig. 2** Steady-state luminescence spectra ( $\lambda_{\rm exc} = 400$  nm) of **1** (5.6  $\times$  10<sup>-6</sup> mol L<sup>-1</sup>) in toluene upon the addition of C<sub>60</sub>: a; 0, b; 0.67, c; 1.3, d; 2.0, e; 3.3, f; 4.7, g; 6.0, h; 8.7 ( $\times$  10<sup>-5</sup> mol L<sup>-1</sup>).

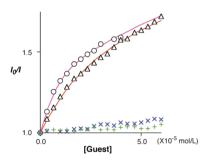


Fig. 3 Stern–Volmer plots of sensors 1, 2 and 12  $(5.6 \times 10^{-6} \text{ mol L}^{-1})$  in the presence of fullerenes in toluene. The symbols represent the experimental data and the solid lines are given by curve-fitting analysis ( $\circ$ :  $C_{70}$   $\nu s$ . 1;  $\triangle$ :  $\nu s$ . 1;  $\times$ :  $C_{60}$   $\nu s$ . 2; +:  $C_{60}$   $\nu s$ . 12).

molecular quenching process between the bound fullerene and the lumophore of the complex. It is known that the receptor bearing two calix[5]arenes shows remarkably strong binding ability to the fullerenes in toluene.<sup>2</sup> Although these plots qualitatively account for the effect of the fullerene complexation, more detailed understanding is required to discuss the quenching properties. For this, luminescence lifetime measurements and the determination of the quenching rate constants were carried out.

The luminescence lifetimes ( $\tau_0$ ) of 1, 2 and 12 in toluene were determined to be 41.4 ns (1), 39.3 ns (2), and 34.5 ns (12), respectively, by nanosecond laser spectroscopy. The quenching rate constants ( $k_q$ )<sup>9</sup> on 2 and 12 for  $C_{60}$  or  $C_{70}$  were determined on the basis of the Stern–Volmer plots and the luminescence lifetimes (2:  $k_q$ ( $C_{60}$ ) = 37.4 × 10<sup>9</sup> M<sup>-1</sup> s<sup>-1</sup>, 12:  $k_q$ ( $C_{60}$ ) = 12.7 × 10<sup>9</sup> M<sup>-1</sup> s<sup>-1</sup> and  $k_q$ ( $C_{70}$ ) = 8.8 × 10<sup>9</sup> M<sup>-1</sup> s<sup>-1</sup>). On the other hand, the non-linear curve fitting analysis of the Stern-Volmer plots for sensor 1 with  $C_{60}$  or  $C_{70}$  yielded both the intermolecular ( $k_q$ ) and the intramolecular quenching rate constants ( $k_q$ ') in the bound state, and the binding constants ( $K_a$ ) ( $k_q$ ( $C_{60}$ ) = 5.7 × 10<sup>9</sup> M<sup>-1</sup> s<sup>-1</sup>,  $k_q$ '( $C_{60}$ ) = 0.0020 × 10<sup>9</sup> s<sup>-1</sup>,  $K_a$  = 71000 M<sup>-1</sup>;  $k_q$ ( $C_{70}$ ) = 12 × 10<sup>9</sup> M<sup>-1</sup> s<sup>-1</sup>,  $k_q$ '( $C_{70}$ ) = 0.0012 × 10<sup>9</sup> s<sup>-1</sup>,  $K_a$  = 360000 M<sup>-1</sup>).

Slight enhancement of the quenching rate was observed between  $\bf 2$  and  $\bf 12$  for  $C_{60}$ . The relative quenching rate constant

of **2** for  $C_{60}$  is only 3 times as large as that of **12**. This can be rationalized by the complexation of **2** to  $C_{60}$ . It appears that the quenching is mainly due to the intermolecular collision; however, a small portion of the intramolecular quenching process should contribute to the quenching. A more obvious contribution of the intramolecular quenching process in the bound state is seen in the luminescence quenching of **1** with the fullerenes. The apparent quenching rate constants  $(k_q^{app})$  are expressed as follows:

$$k_{\mathbf{q}}^{\mathrm{app}} = k_{\mathbf{q}}[\mathbf{G}]_0$$
 for  $\mathbf{2}$  and  $\mathbf{12}$   
 $k_{\mathbf{q}}^{\mathrm{app}} = k_{\mathbf{q}}[\mathbf{G}] + k_{\mathbf{q}}'$  for  $\mathbf{1}$ 

where  $[G]_0$  and [G] denote the concentration of total and unbound guests, respectively. The apparent quenching rate constants  $(k_q^{\rm app})$  were calculated using the  $K_a$  value and the total fullerene concentration of  $5.6 \times 10^{-6}$  mol L<sup>-1</sup> ( $12: k_q^{\rm app}(C_{60}) = 7.1 \times 10^4 \, {\rm s}^{-1}, k_q^{\rm app}(C_{70}) = 4.9 \times 10^4 \, {\rm s}^{-1}; 2: k_q^{\rm app}(C_{60}) = 21 \times 10^4 \, {\rm s}^{-1}; 1: k_q^{\rm app}(C_{60}) = 200 \times 10^4 \, {\rm s}^{-1}, k_q^{\rm app}(C_{70}) = 120 \times 10^4 \, {\rm s}^{-1}$ ). The apparent luminescence quenching rate constants of 1 for  $C_{60}$  and  $C_{70}$  are over 20 times higher than that of 12 at that concentration. This suggests that the apparent quenching rate constant is strongly associated with the concentration of the bound fullerene; thus, the high binding ability of 1 toward  $C_{60}$  and  $C_{70}$  brings about the extremely high sensitivity and selectivity 11 to them even at concentrations of less than  $10^{-5}$  mol  $10^{-5}$  mol  $10^{-5}$  mol  $10^{-5}$  mol  $10^{-5}$ 

We have demonstrated the first example of the highly sensitive detection of fullerenes using calix[5]arene-based sensors produced by the combination of the supramolecular concept and the luminescence technique.

## Notes and references

- P. D. Beer and P. A. Gale, *Angew. Chem., Int. Ed.*., 2001, 40, 486; E. V.
  Anslyn and J. J. Lavigne, *Angew. Chem., Int. Ed.*, 2001, 40, 3118.
- T. Haino, M. Yanase and Y. Fukazawa, *Angew. Chem., Int. Ed.*, 1998,
  997; T. Haino, M. Yanase and Y. Fukazawa, *Tetrahedron Lett.*,
  1997, 38, 3739; T. Haino, M. Yanase and Y. Fukazawa, *Angew. Chem., Int. Ed. Engl.*, 1997, 36, 259.
- 3 D. Sun, F. S. Tham, C. A. Reed, L. Chker, M. Burgess and P. D. W. Boyd, J. Am. Chem. Soc., 2000, 122, 100704; K. Tashiro, T. Aida, J.-Y. Zheng, K. Kinbara, K. Saigo, S. Sakamoto and K. Yamaguchi, J. Am. Chem. Soc., 1999, 121, 9477; J. L. Atwood, L. J. Barbour, P. J. Nichols, C. L. Raston and C. A. Sandoval, Chem. Eur. J., 1999, 5, 990; M. J. Hardie and C. L. Raston, Chem. Commun., 1999, 1153; H. Matsubara, T. Shimura, A. Hasegawa, M. Semba, K. Asano and K. Yamamoto, Chem. Lett., 1998, 1099; K. Tsubaki, K. Takayoshi, T. Kinoshita and K. Fuji, Chem. Commun., 1998, 895; N. S. Isaacs, P. J. Nicholls, C. L. Raston, C. A. Sandova and D. J. Young, Chem. Commun., 1997, 1839; A. Drljaca, C. Kepert, L. Spiccia, C. L. Raston, C. A. Sandoval and T. D. Smith, Chem. Commun., 1997, 195; T. Andersson, G. Westman, G. Stenhagen, M. Sundahl and O. Wennerström, Tetrahedron Lett., 1995, 36, 597; J. L. Atwood, G. A. Koutsantonis and C. L. Raston, Nature, 1994, 368, 229; T. Suzuki, K. Nakashima and S. Shinkai, Chem. Lett., 1994, 699; Z. Yoshida, H. Takekuma, S. Takekuma and Y. Matsubara, Angew. Chem., Int. Ed. Engl., 1994, 33, 1597.
- 4 T. Haino, H. Araki, Y. Yamanaka and Y. Fukazawa, *Tetrahedron Lett.*, 2001, 42, 3203.
- 5 B. Förster, J. Bertran, F. Teixidor and C. Viñas, J. Organomet. Chem., 1999, 587, 67.
- 6 I. Hamachi, S. Tanaka, S. Tsukiji, S. Shinkai and S. Oishi, *Inorg. Chem.*, 1998, 37, 4380.
- 7 M. H. Todd, S. Balasubramanian and C. Abell, *Tetrahedron Lett.*, 1997, 38, 6781.
- 8 K. A. Walters, L. L. Premvardhan, Y. Liu, L. A. Peteanu and K. S. Schanze, Chem. Phys. Lett., 2001, 339, 255.
- 9 The quenching rate conctants of 12 for  $C_{60}$  and  $C_{70}$  were determined by Stern–Volmer plots of the fullerene concentration  $vs. \tau_0/\tau$ .
- 10 **2**:  $K_a(C_{60}) = 1100 \text{ M}^{-1}$ ; **11**:  $K_a(C_{60}) = 1060 \text{ M}^{-1}$ ,  $K_a(C_{70}) = 170 \text{ M}^{-1}$ .
- 11 A control experiment with anthracene as a small guest was carried out to test the selectivity of 1. A Stern–Volmer plot of 1 produced  $k_{\rm q}$  (21.2  $\times$  109 M $^{-1}{\rm s}^{-1}$ ) and  $k_{\rm q}^{\rm app}$  (11.9  $\times$  104 s $^{-1}$  at a total guest concentration of 5.6  $\times$  10 $^{-6}$  mol L $^{-1}$ ) is 17 times as small as that of 1 with C $_{60}$ . This indicates that sensor 1 shows selective detection of fullerenes even if a small aromatic guest is present.