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1. Introduction

During the past decade, social instability, terrorist activities and other criminals have made the enormous harm to human health and social stability. A major concern in the field of public safety today is to continue to develop high sensitive sensors for a fast and accurate identification of trace explosives. The technology of fluorescence sensing based on fluorescent sensors has been intensively investigated recently because it has the advantages of high sensitivity, good specificity, rapid response, portability and simple operation. Many studies have been carried out on this technology which is widely used in trace detection of ions, biological molecules, explosives and environmental pollutants.

Peroxide-based explosives (PEs) such as triacetone triperoxide (TATP) are increasingly used to make improvised explosive devices in some crime and terrorism activities because of their simple preparation from readily available materials (TATP, just three ingredients: hydrogen peroxide (H$_2$O$_2$), an acid and acetone), poor chemical stability and easy decomposition leading to large explosive power. Compared to the nitroaromatic explosives, e.g. 2,4,6-trinitrotoluene (TNT), peroxide explosives vapor is difficult to be detected with fluorescence probe because PEs are not typical quenching agents without nitro groups or aromatic units, which can easily interact with electron-rich chromophores by intermolecular π-π stacking interaction. In addition, sensitivity to mechanical stress, low stability, limited solubility and so on also make the detection of PEs become more challenging.

For example, Germain et al. introduced a fluorescence detection method targeted H$_2$O$_2$ by using a chelator formed via reaction with H$_2$O$_2$, which was only limited in liquid phase. For an efficient detection, both molecular structure and morphology of the film are critical. In molecular structure aspect, both pyrene and fluorene derivatives are important building blocks in fluorescent materials for their relatively simple molecular structure, easy modification and high fluorescence quantum efficiency. And borate ester units, as the chain end groups, were introduced to selectively detect...
peroxides. In addition, amplified fluorescence signal could improve the sensing properties via the “molecular wires” or the hyperbranched conjugated structure of polymers. In particular, properties associated with the hyperbranched polymers such as modifiable surface functionality, available internal cavities, uniform film and good solubility make them attractive for biological, chemical and medical field 28. In morphology aspect, the microstructure of films, which determined the quenching efficiency, response time etc has an important effect on the sensing performance. We find that using ZnO nanorods array as the substrate of the sensing film can effectively increase the signal strength, reaction rate and sensitivity due to high area to volumn ratio for efficient analyte permeability and catalytic ability of light oxidation 30. Herein, we systematically discussed that the effects of the configuration of polymer, chain end location and number of boron ester unit, as well as the polymers/ZnO nano rod array composite on the sensing performance of the peroxide vapor detection.

2. Experimental Section

2.1 Instruments

The 1H-NMR spectra were recorded on a Brucker DRX500 instrument, and tetramethyldisilane (TMS) was used as an Internal standard. UV-Vis absorption and fluorescence analysis were obtained from a Jasco V-670 spectrophotometer and a Jasco FP 6500 spectrometer, respectively. High vacuum infrared spectra were performed on Brucker VERTEX 70v via surface reflection-absorption model. Cyclic voltammetry (CV) experiments were performed with a CH instruments electrochemical analyzer. The electrochemical behaviors of S1, S2 and S3 were investigated in a standard three electrode electrochemical cell (a glassy carbon working electrode, a platinum counter electrode and a silver chloride electrode as a reference electrode) with 0.1 M tetra-n-butylammonium hexafluorophosphate (Bu4NPF6) in acetonitrile solution, and the scanning rate was 100 mV/s under nitrogen atmosphere at room temperature.

2.2 Synthesis

All the chemicals and solvents were obtained from commercial sources and used as received. The synthetic procedures were illustrated in Figure 1. All the polymers were synthesized via Suzuki-Miyaura cross-coupling reaction with yields of 34%, 41% and 48%, respectively. The raw materials, 1, 3, 6, 8-Tetrabromopyrene, 9, 9-diocetyl-2, 7-bis (boronic acid pinacol ester) fluorene, 1, 6-Dibromopyrene, 2, 7-dibromo-9, 9-didodecylfluorene and 2, 7- bis (4, 4, 5, 5-tetramethyl-1, 3, 2-dioxaborolan-2-yl) - pyrene were obtained from commercial resources.

2.2.1 Polymerization of S1

A mixture of 1, 3, 6, 8-Tetrabromopyrene (0.156 g, 0.301 mmol), 9, 9-diocetyl-2, 7-bis (boronic acid pinacol ester) fluorene (0.951 g, 1.480 mmol), Pd(PPh3)4 (98.6 mg, 0.085 mmol), K2CO3 (aq, 2 M/10 mL), dioxane (20 mL), ethanol (10 mL) and aliquat 336 (two drops) was charged under nitrogen and stirred for 24 h at 100 °C. After cooling to room temperature, the solution was washed with water/CH2Cl2 solution and dried with MgSO4. The organic phase solution was evaporated under reduced pressure to get crude solid product. The crude polymer was then dissolved with least amount of toluene and precipitated in methanol for three times. The polymer was then further purified by a short silica column with THF as eluent and precipitated in MeOH again. The final product was dried to afford the polymer as a light green solid (234 mg, 34%).1H-NMR (500 MHz, CDCl3, 25 °C, TMS): δ= 8.20 (s, 4H), 8.07 (s, 2H), 7.85 (d, 4H), 7.83 (d, 4H), 7.78 (t, 8H), 7.67(d, 8H), 2.04 (s, 20H), 1.26 (m, 30H), 1.12 (m, 90H), 0.83 (m, 50H). GPC (THF vs. PS): Mn= 2875, Mw= 3514, PDI=1.22.

2.2.2 Polymerization of S2

Following the same polymerization procedure as S1, with 1,6-Dibromopyrene (277.2 mg, 0.77 mmol), 9, 9-diocetyl-2, 7-bis (boronic acid pinacol ester) fluorene (0.73 g, 1.136 mmol), Pd(PPh3)4 (89 mg, 0.077 mmol), K2CO3 (aq, 2 M/10 mL), tetrahydrofuran (20 mL) and aliquat 336 (four drops). The final product was dried to afford S2 as a celadon solid (250 mg, 41%).1H-NMR (500 MHz, CDCl3, 25 °C, TMS): δ= 8.34 (m, 3H), 8.15 (m, 3H), 8.09 (m, 1H), 7.83 (m, 2H), 7.73 (m, 2H), 7.65(m, 2H), 2.16 (s, 2H), 2.08 (s, 2H), 1.43 (s, 2H),1.25 (m, 18H) 0.97 (m, 14H), 0.84 (m, 6H). GPC (THF vs. PS): Mn= 4819, Mw= 6981, PDI=1.45.

2.2.3 Polymerization of S3

Following the same polymerization procedure as S1, with 2, 7-dibromo-9, 9-didodecylfluorene (330 mg, 0.5 mmol), 2, 7-bis(4, 4, 5, 5-tetramethyl-1, 3, 2-dioxaborolan-2-yl)-1, 6-dihydropyrene (340 mg, 0.75 mmol), Pd(PPh3)4 (60 mg, 0.05 mmol), K2CO3 (aq, 2 M/15 mL), toluene (15 mL) and aliquat 336 (four drops). The final product was dried to afford the polymer as a light brown solid (170 mg, 48%).1H-NMR (500 MHz, CDCl3, 25 °C, TMS): δ= 8.53 (m, 3H), 8.25 (m, 4H), 7.95 (m, 8H), 2.18 (m, 4H), 1.25 (s, 8H), 1.22 (s, 48H), 0.88 (m, 18H). GPC (THF vs. PS): Mn= 8278, Mw= 27905, PDI=3.371.
time-course responses were recorded as soon as the film was exposed to analyte vapor and ended at 300 s.

3. Result and discussion

3.1 Optical and electrochemical properties

Figure 2 shows the absorption and emission spectra of S1, S2 and S3 in films and in THF solution. Their photophysical results are summarized in Table 1. On the one hand, it can be found that the maximum absorption peak of linear polymers S2 and S3 in THF are 383 and 346 nm, respectively. Compared with that of S2, the maximum absorption peak of S3 is blue-shifted by 37 nm and the similar result also appears in the film state. This proves that a different connection position of borate ester unit has a different effect on the spectral characteristics. On the other hand, the emission peak of hyperbranched polymer S1 in THF is 465 nm, red-shifted by 28 nm compared to that of S2. The same result was also observed in film state (red-shifted by 20 nm). The borate ester number of S1 in one molecule unit is more than that of S2, and there are different configurations in S1 (hyperbranched polymer) and S2 (linear polymer), which leads to more efficient conjugation in S1. It is easy to find that the number of borate esters on the chain end and configuration of polymer have different impact on the spectral property.

The emission peak of S2 and S3 films are red-shifted by 9 and 10 nm from solution to film state respectively, while that of S1 film is almost the same (a red shift of only 1 nm). The result illustrates the hyperbranched structure of S1 has larger steric hinderance and it is beneficial for preventing the π-π stacking caused self-aggregation effect in solid state and hence improving the sensing performance. In addition, both S1 and S2 have very high fluorescent quantum yield (Φ) in their THF solution, which are nearly 1. It is also an important factor as sensing materials. Moreover, Stokes shifts of S1 and S3 are larger than that of S2 in both solution state and solid state, which can avoid the self-absorption by the material and reduce the interference of the background to improve the sensitivity. Last but not the least, there are plenty of internal cavities 31 and external boron ester groups in hyperbranched configuration of S1 so that it makes important contributions to increasing permeability of analytes and response rate of the sensing process.

The energy level is related to the probe’s reactivity to the oxidation reagents. The electrochemical results (Table 1) reveal that the HOMOs of S1, S2 and S3 are -5.49, -5.99 and -6.04 eV, respectively, suggesting a stepwise decreased HOMO from S1 to S3. Their band gaps are 2.80, 3.21 and 3.25 eV and S1 is the least one among them. These features mean that S1 will be likely the most reactive polymer toward oxidation reaction. The hyperbranched molecular structure and more borate ester groups on the periphery structure of S1 make it more efficient conjugation and more reactive than the linear polymers S2 and S3. At the same time, narrow band gap means that S1 needs an exciting light with longer wavelength. Thus, it can not only reduce the cost of the detector but also improve the light stability of the material. All these optical and electrochemical conditions above suggest the hyperbranched polymer S1 is likely a promising candidate for fluorescent sensing application.

3.2 Detection of peroxide in the vapor phase

The sensing performances of the three polymers to peroxide vapor were monitored by fluorescent spectroscopy. The films were fabricated by dip-coating their toluene solution with a concentration of 10−4 g/mL onto quartz plate. The normalized peak emission intensity change in air or explosive vapour with time is shown in Figure 3. It exhibits that the quenching efficiency of three probes towards the saturated vapor of 30% H2O2 aqueous are 40% (S1), 24% (S2) and 23% (S3) respectively. As can be seen, the hyperbranched polymer S1 shows a much better sensing performance than that of the linear polymer S2 and S3.

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The fluorescence quantum yields of S1, S2 and S3 compounds in THF dilute solution were measured using a dilute solution of 9,10-diphenylanthracene (Φ ~ 1) in THF solution as the standard.

Figure 3. (a-c) Stability and sensing properties of S1-S3 films on different substrates exposed to air and H2O2 saturated vapor respectively within 300s. (a: S1; b: S2; c: S3); (d) Quenching efficiency of S1-S3 films on different substrates in H2O2 saturated vapor within 300s.

Since the sensing performance is also related to their film morphology. SEM was used for the morphology characterization. Figure 4 shows that more uniform film is formed for hyperbranched polymer S1, while S2 and S3 suffered a heavy aggregation. SEM results suggest that the linear polymer structure with polar borate ester end groups will lead to serious aggregation, which not only prevents the penetration of H2O2 vapor but also reduces the specific surface area of films resulting in a lower sensitivity.

Figure 4. SEM images of S1-S3 films prepared with toluene solvents (all concentrations, 10^3 g/ml) and ZnO nanorod array as film substrate (a: S1; b: S2; c: S3; d: ZnO nanorod array).

The morphology of the substrate may also greatly influence the sensitivity, since a high specific surface area and surface activity of the films will contribute to the sensing performance. Our group has successfully developed ZnO nanorod arrays with different surface morphology and proved to improve the sensing efficiency by a catalytic functionality to accelerate the reaction rate and evanescent effect to increase the emission intensity. Therefore, we coated the fluorescent materials on the surface of ZnO nanorod arrays (Figure 4(d)) vertically arranged on quartz plate to make it a composite. Figure 3 (a) presents that the S1/ZnO nanorod arrays composite produces enhanced fluorescence quenching with a response rate of about 30% within 50 s and finally 60% within 300 s upon exposure to a vapor of 30% H2O2 aqueous, corresponding to a 20% increase relative to that on quartz plate. Under the same condition, the quenching efficiency of hyperbranched polymer S1 to H2O2 vapor is also much better than that of linear polymer S2 and S3 (43% and 30% quenching efficiency within 300 s, respectively). It demonstrates hyperbranched polymer S1 with more external boron ester end groups and internal cavities is conducive to peroxide vapor penetration and contributes to more effective detection.
In order to interpret the sensing mechanism, both fluorescence and high vacuum reflection-absorption infrared spectra (IRAS) were used for monitoring the changes of S1/ZnO nanorod array composite film before and after the H$_2$O$_2$ vapor exposure. Figure 5 (a) shows that emission changes of S1 by dissolving the composite film into THF after being exposed to H$_2$O$_2$ in a period of 20 minutes. The emission peak of S1 declines gradually with a slight red shift from 461 to 465 nm. Figure 5 (b) exhibits the IRAS changes that, after the exposure, the band at 1343 cm$^{-1}$ (B-O characteristic peak) disappears along with the appearance of new band at 3253 cm$^{-1}$ (O-H characteristic peak). It proves that the deboronation reaction occurred between external boron ester end groups of S1 and peroxide vapor leading to the fluorescence turn-off as shown as Figure 5 (c). In addition, the fluorescent conjugated structure of hyperbranched polymer S1 can amplify fluorescence signal due to its effective conjugation and multiple borate esters structure. On the other hand, ZnO nanorod arrays not only change the surface morphology of the film but also increase the reactivity to H$_2$O$_2$ vapor, which will accelerate the deboronation reaction of S1 under UV light.

Figure 5. (a) Changes in the emission spectral pattern of S1 film on ZnO nanorod array during reaction with H$_2$O$_2$ saturated vapor followed by dissolving the film into THF solvent within the reaction of 20 minutes. (b) High vacuum infrared spectra of S1/ZnO nanorod array composite dissolved into THF solution before (red) and after exposure to hydrogen peroxide (black) vapor for 1h. (c) Deboronation reaction of boron ester group by H$_2$O$_2$.

The light stability will determined the lifetime of the sensing devices. The photo bleaching of S1 on quartz plate is measured to be ~7% in 300 s under air condition. However, the light bleaching of S1/ZnO nanorod arrays composite can be well controlled within 2% suggesting that ZnO nanorod arrays are also favorable for enhancing the stability of material. Nevertheless, compared with the good stability of S1 and S2, the photo bleaching rate of S3 film either on quartz plate or on ZnO nanorod arrays is not ideal (~9%). The results illustrate that the connection position of borate ester groups has obvious influence on light stability and the sensing performance of sensing films. A direct connecting borate ester to fluorene causes better light stability and sensing performance than that to pyrene. This is likely relevant to the increased intermolecular interaction among pyrene molecules due to the polar borate ester units. S3 film is very difficult to be used as efficient sensor due to worse photo stability which leads to a short lifetime and false alarm signals of the detection device. The selectivity is another important issue for the sensing performance. Figure 6 shows fluorescence responses of S1/ZnO composite to different analytes saturated vapor. S1 film presents obvious fluorescent quenching response to H$_2$O$_2$ (~60%) and TATP (~30%). It is mentionable that compared with H$_2$O$_2$, TATP is hard to be detected with high sensitivity owing to the lower saturation vapor pressure (78 ppm). Hence S1 will be a promising fluorescent probe for TATP vapor detection. The other common used solvents vapor such as methylene chloride, acetone and ethyl alcohol show very low quenching efficiencies, which will not interfere with the sensing process. It is important to note that some solvents such as toluene can lead to enhanced fluorescence intensity due to swelling effect. But it will not produce interference to the peroxide detection based on fluorescence quenching process. In addition, the quenching efficiencies of S1 to some saturated vapor mixtures of different analytes after an exposure of 300 s are all less than 15% so that it will not interfere with the sensing process (As shown as Figure S4).

Figure 6. Fluorescence responses of S1 film on ZnO nanorod array to different compounds saturated vapor after an exposure of 300s.
In order to determine the detection limit of the S1/ZnO nanorod array composite film, diverse concentrations of H$_2$O$_2$ solution were prepared by diluting the 30 wt% H$_2$O$_2$ solution with deionized water to produce a corresponding equilibrium vapor pressure (1, 19, 38 and 225 ppm, corresponding to 90, 10, 5, and 1 times dilution of the commercial 30 wt% H$_2$O$_2$ solution) at room temperature (25 °C) 32. The quenching efficiencies towards different concentration of H$_2$O$_2$ vapor are shown in Figure 7 (a). Then the different quenching rates and corresponding equilibrium vapor pressures data were made a linear fitting, which is well-fitted to the Langmuir equation. As shown as Figure 7 (b), the detection limit of H$_2$O$_2$ could be as low as 1.6 ppb if the triple multiple signal to noise ratio of the fluorescent detection device was considered as 0.01.

4. Conclusions

In summary, we reported a relatively simple, sensitive and selective borate ester endcapped pyrene-fluorene hyperbranched copolymer for fluorescent detection of peroxide explosive vapor. The sensing performance is based on a deboronation reaction under peroxide and signal amplification effect of polymer resulting in fluorescence quenching, which is related to the molecular structure, substitution position on the aromatic ring and the numbers of the borate units. A composite of the sensory polymer/ZnO nanorod array can improve the stability and sensibility of material because of the catalytic ability of light oxidation and surface morphology with high area to volum ratio for efficient analyte permeability. The results show that compared with two linear analogues S2 and S3, the hyperbranched polymer S1 demonstrates more sensitive response toward peroxide (~60% quenching and 30% for TATP within 300 s) due to larger steric hindrance, higher HOMO level, more internal cavities and external boron ester groups of hyperbranched structure. The detection limit to H$_2$O$_2$ vapor is estimated to be 1.6 ppb. This method provides a new way for trace and on-site chemical detection of peroxide-based explosives (PEs) for human health and public safety.

Acknowledgements

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Notes and references

† Electronic Supplementary Information (ESI) available: [Fourier transform infrared spectra of S3, S2, S1 and the contrast, The GPC and CV curves of S1-S3 polymer, The quenching efficiency (1-I/I$_0$) of S1 film on ZnO nanorod array to different vapor pressure of H$_2$O$_2$ after an exposure of 300s]. See DOI: 10.1039/b000000x/