

Cite this: *Chem. Sci.*, 2017, 8, 2150

# Photoacoustic probes for real-time tracking of endogenous H<sub>2</sub>S in living mice†

Ben Shi,<sup>a</sup> Xianfeng Gu,<sup>b</sup> Qiang Fei<sup>a</sup> and Chunchang Zhao<sup>\*a</sup>

H<sub>2</sub>S is a key chemical mediator that exerts a vital role in diverse physiological and pathological processes. However, *in vivo* tracking of endogenous H<sub>2</sub>S generation still remains difficult due to the lack of reliable analytical methods. Herein, we present the first example of activatable photoacoustic probes for real-time imaging of H<sub>2</sub>S in living mice through the full utilization of the superiority of photoacoustic imaging modality at fine spatial resolution during deep tissue penetration. The designed probe can generate high NIR absorption at 780 nm in the presence of H<sub>2</sub>S, thus producing a strong photoacoustic signal output in the NIR region. Furthermore, this probe exhibits extremely fast and highly selective responsiveness, good water-solubility and excellent biocompatibility. In light of these outstanding features, this probe realizes the direct photoacoustic trapping of endogenous H<sub>2</sub>S generation in a HCT116 tumor-bearing mouse model. These preliminary imaging studies show that HCT116 colon tumors exhibit CBS upregulation activity, resulting in an increased rate of H<sub>2</sub>S generation.

Received 21st October 2016  
Accepted 26th November 2016

DOI: 10.1039/c6sc04703c

www.rsc.org/chemicalscience

## Introduction

Mounting studies have shown that H<sub>2</sub>S plays a vital role in diverse physiological and pathological processes. It has been identified as a physiological gasotransmitter with robust cytoprotective actions in multiple organ systems, including regulatory roles in the cardiovascular system and modulation of the central nervous, respiratory, and gastrointestinal systems.<sup>1,2</sup> Nevertheless, abnormal production of H<sub>2</sub>S can contribute to a variety of diseases ranging from Down's syndrome and Alzheimer's disease to diabetes and hypertension.<sup>3</sup> As such, it is of great scientific interest to develop selective methods for the real-time tracking of this small molecule in living systems. To date, a number of reaction-based fluorescent probes have been designed as attractive molecular imaging techniques for monitoring H<sub>2</sub>S in living cells.<sup>4–7</sup> However, few probes have been employed for *in vivo* H<sub>2</sub>S imaging due to the fact that fluorescence imaging suffers from limitations such as shallow tissue penetration and poor spatial resolution inside deep biological tissue. Therefore, development of reaction-based probes with new imaging modality is highly demanded in order to address these challenges.

Photoacoustic (PA) imaging is a state-of-the-art imaging modality which relies on the translation of excitation light into ultrasonic waves based on the PA effect, and provides deeper tissue imaging penetration and higher *in vivo* spatial resolution when compared with traditional optical imaging techniques.<sup>8</sup> As a result of these significant advantages, PA imaging has found broad applications in biology and medicine as a non-invasive imaging tool.<sup>9,10</sup> However, it is still a big challenge to devise activatable probes for the photoacoustic imaging of molecular targets of interest. Given that H<sub>2</sub>S is a key chemical mediator, the full utilization of the superiority of PA imaging modality for the construction of H<sub>2</sub>S-activatable photoacoustic probes should be imperative to fully explore the biological roles of H<sub>2</sub>S *in vivo*. To our knowledge, there are no probes available for the PA imaging of this important small molecule. Herein, we present the first example of reaction-based PA probes that show a fast response to H<sub>2</sub>S accompanied with “turn-on” PA signals.

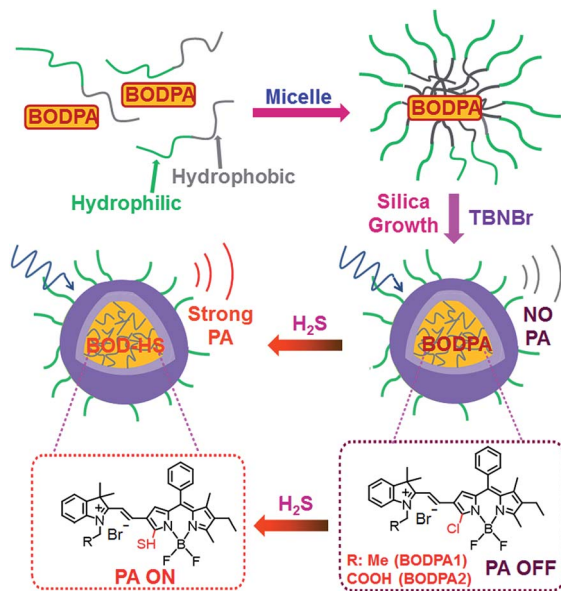
In this contribution, a H<sub>2</sub>S-activatable probe was fabricated by encapsulating semi-cyanine-BODIPY hybrid dyes (BODPA) into the hydrophobic interior of core-shell silica nanocomposites for *in vivo* photoacoustic imaging (Scheme 1). Our design strategy is based on modulating the electronic nature of the substituents to generate a near-infrared absorbing dye to produce PA signals, employing the feasible thiol-halogen nucleophilic substitution of a monochlorinated hybrid dye with H<sub>2</sub>S. Indeed, the presence of H<sub>2</sub>S transformed BODPA into BOD-HS inside nanoparticles through aromatic nucleophilic substitution, which resulted in high NIR absorption around 780 nm. Thus, the Si@BODPA probe generated a strong photoacoustic signal output in the NIR region. As shells of these nanoparticles were composed of poly(ethylene glycol) (PEG) and

<sup>a</sup>Key Laboratory for Advanced Materials, Institute of Fine Chemicals, School of Chemistry and Molecular Engineering, East China University of Science & Technology, Shanghai 200237, P. R. China. E-mail: zhaocchang@ecust.edu.cn

<sup>b</sup>Department of Medicinal Chemistry, School of Pharmacy, Fudan University, Shanghai, 201203 China

† Electronic supplementary information (ESI) available: Procedures for the synthesis of BODPA, living mice imaging, characterization data, and supplementary figures. See DOI: 10.1039/c6sc04703c





Scheme 1 Schematic illustration of the construction of activatable photoacoustic probes for H<sub>2</sub>S.

positively charged silica units, Si@BODPA possessed good water-solubility and excellent biocompatibility as well as extremely fast responsiveness, enabling real-time imaging of H<sub>2</sub>S-related biological processes. In light of these outstanding features, Si@BODPA allowed the direct photoacoustic tracking of endogenous H<sub>2</sub>S generation in a HCT116 (human colon cancer cells) tumor-bearing mouse model.

## Results and discussion

Two monochlorinated BODIPY-based hybrid dyes (BODPA1 and BODPA2) were chosen to prepare core-shell silica nanocomposites. Both dyes responded to H<sub>2</sub>S readily and were efficiently converted to BOD-HS within 50 minutes in CH<sub>3</sub>CN/PBS buffer mixtures (1 : 1, v/v, 20 mM, pH 7.4, room temperature), which was demonstrated by HPLC and HRMS (Fig. S1†). Specifically, a strong NIR absorption band around 735 nm was activated, accompanied by a decrease of absorption at 540 nm (Fig. S2†), indicative of BODPAs as building blocks for the construction of activatable photoacoustic probes. Notably, BOD-HS is non-fluorescent which facilitates thermal deactivation to generate much stronger PA signals.<sup>10b</sup> Considering the limited solubility of BODPAs as well as their slow reaction with H<sub>2</sub>S in aqueous solution, we here intended to apply hydrophilic nanoparticles for the encapsulation of BODPAs to overcome this issue. Firstly, BODPAs and mPEG-DSPE were mixed in a 0.85 N hydrochloride solution under rigorous stirring, leading to the formation of self-assembled micelles with hydrophobic inner cores and hydrophilic PEG chains as outer arms.<sup>11</sup> The hydrophobicity of BODIPY dyes trap them in the interior core. Following the *in situ* growth of silica shells under acidic conditions with *N*-trimethoxysilylpropyl-*N,N,N*-tri-*n*-butylammonium bromide (TBNBr) as silica agents for cross-linking, water-dispersible nanoparticles (Si@BODPA) were afforded,

which were dialyzed for 48 h to remove unreacted chemicals. Additional characterization using TEM and DLS were carried out, which were identified in Fig. S3.†

The response of Si@BODPA toward H<sub>2</sub>S was then evaluated in PBS buffer solutions (pH 7.4) using UV-vis spectroscopy. The addition of NaHS induced the generation of the NIR absorption at 780 nm and 777 nm for BODPA1 and BODPA2, respectively. The reduction of the original peak at around 540 nm was also noted (Fig. 1A). This observation was in good agreement with that of BODPAs when interacted with NaHS in CH<sub>3</sub>CN/PBS, which demonstrated the occurrence of a reaction between BODPAs and H<sub>2</sub>S within the water-dispersible silica core-shell nanoparticles. Notably, BOD-HS in the hydrophobic interior of the nanocomposites showed an obvious red-shift of 45 nm in its absorption compared to that in CH<sub>3</sub>CN/PBS buffer mixtures (1 : 1, v/v). This variation could be rationalized as the effect of H-bonding interactions between the sulfur atom and H atom of H<sub>2</sub>O. Such interactions existing in aqueous solutions rather than in nanocomposites reduced the electron-donating ability

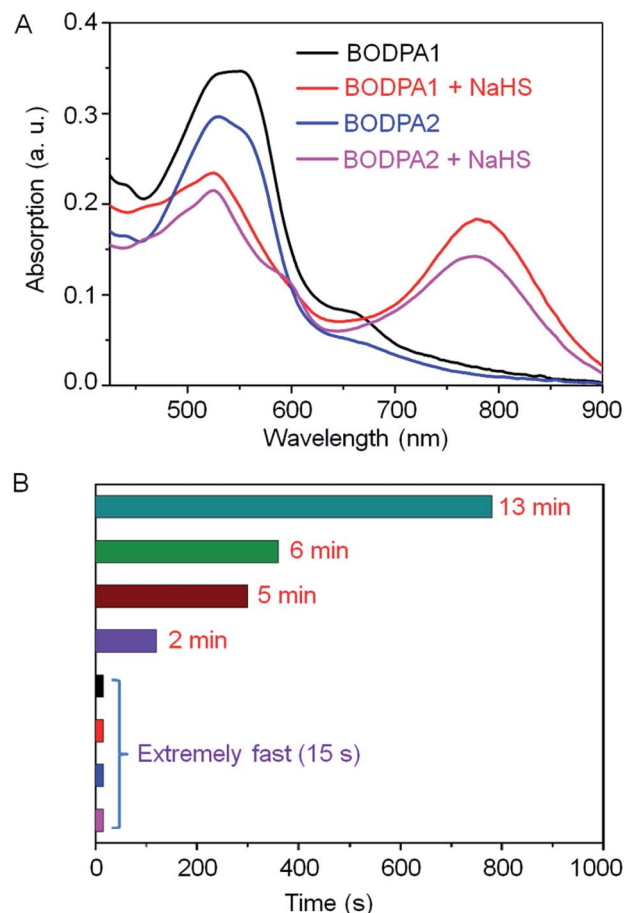


Fig. 1 (A) The absorption changes of Si@BODPAs (10  $\mu$ M BODPA1 or BODPA2) in the absence and presence of NaHS (100  $\mu$ M). (B) The dynamics of the reaction between BODPAs (10  $\mu$ M) and H<sub>2</sub>S (100  $\mu$ M) in Si@BODPAs (13 min and 6 min for BODPA2 in Si@BODPA30 and Si@BODPA90, respectively; 5 min and 2 min for BODPA1 in Si@BODPA30 and Si@BODPA90, respectively). Si@BODPA180 and Si@BODPA270 afforded rapid detection of H<sub>2</sub>S within 15 s.



of sulfur which ultimately led to the shift of the absorption to shorter wavelengths. Impressively, the dynamics of the reaction between BODPAs and  $H_2S$  in Si@BODPA was Si dose-dependent (Fig. 1B and S4†). In general, increasing the amount of Si in the shell, as determined by energy dispersive X-ray (EDX) spectroscopy (Fig. S5†), caused a much faster reaction. The process of absorption change in Si@BODPA30 could reach completion in about 5 and 13 minutes for BODPA1 and BODPA2, respectively. In the case of Si@BODPA90, such change was promoted to be about 2 min and 6 min for BODPA1 and BODPA2, respectively. Gratifyingly, the responsive reaction was dramatically accelerated when further increasing the Si content, resulting in ideal probes, Si@BODPA180 and Si@BODPA270, which could rapidly detect  $H_2S$  within 15 s. Given the transient nature of  $H_2S$  which is catabolized rapidly in biological systems, one of the most important prerequisites for a probe is the extremely fast reactivity toward  $H_2S$ . Thus, the nanoprobe established here possess the superiority and capability of trapping transient  $H_2S$  and monitoring of  $H_2S$ -related biological processes in real-time. As an example, Si@BODPA180 was chosen as a model probe for studying its potential applications. Undoubtedly, the entrapment of BODPAs into the interior of silica nanocomposites enables the advantage of rapid reaction, which can be attributed to the transfer of  $H_2S$  via the hydrophilic assembly from water into its polar interior, which facilitates nucleophilic substitution.<sup>12</sup> Such a polar microenvironment within the core is due to the positively charged ammonium functionality in TBNBr. More importantly, these nanoparticles are smaller than 100 nm and therefore are feasibly endocytosed by cells.

The photoacoustic responsiveness to  $H_2S$  was then established. In the absence of  $H_2S$ , Si@BODPA180 showed no detectable photoacoustic signals in PBS buffer solutions (pH 7.4). When  $H_2S$  was introduced to the solution, Si@BODPA180 efficiently produced bright photoacoustic signals following excitation at 780 nm and ultimately led to a 44-fold turn-on response (Fig. 2A), which was in agreement with  $H_2S$ -triggered formation of BOD-HS within nanoparticles. These results infer that this nanoprobe could serve as an activatable photoacoustic probe.

$H_2S$  triggered photoacoustic amplitudes of the nanocomposites upon excitation at 780 nm were examined at various concentrations. A linear correlation could be obtained between the PA signal and concentration (Fig. 2B). To explore the effect of local concentrations of BODPA within the interior of the nanoparticles on the activated PA properties, we also prepared Si@BODPA with different ratios of BODPA/mPEG-DSPE in weight. It was found that, at the same concentration of BODPA, nanoparticles Si@BODPA180(7) with the ratio of 7/100 exhibited higher PA signals than those of Si@BODPA180(1) and Si@BODPA180(4) obtained from the ratio of 1/100 and 4/100, inferring that Si@BODPA180(7) is a superior system for the establishment of photoacoustic imaging probes. These results reflect that elevated local concentrations of BODPA benefit the thermal deactivation of the excited chromophores, thereby affording stronger PA signals.

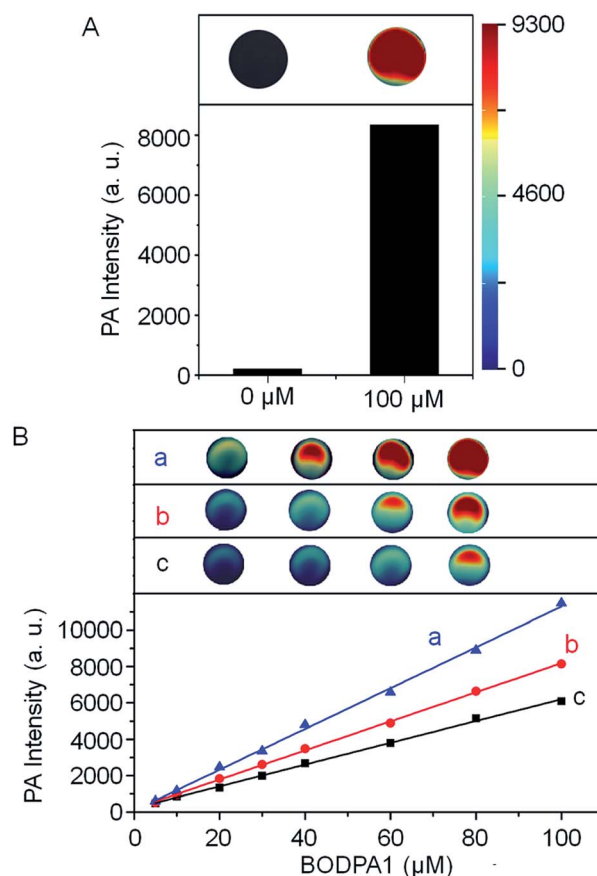


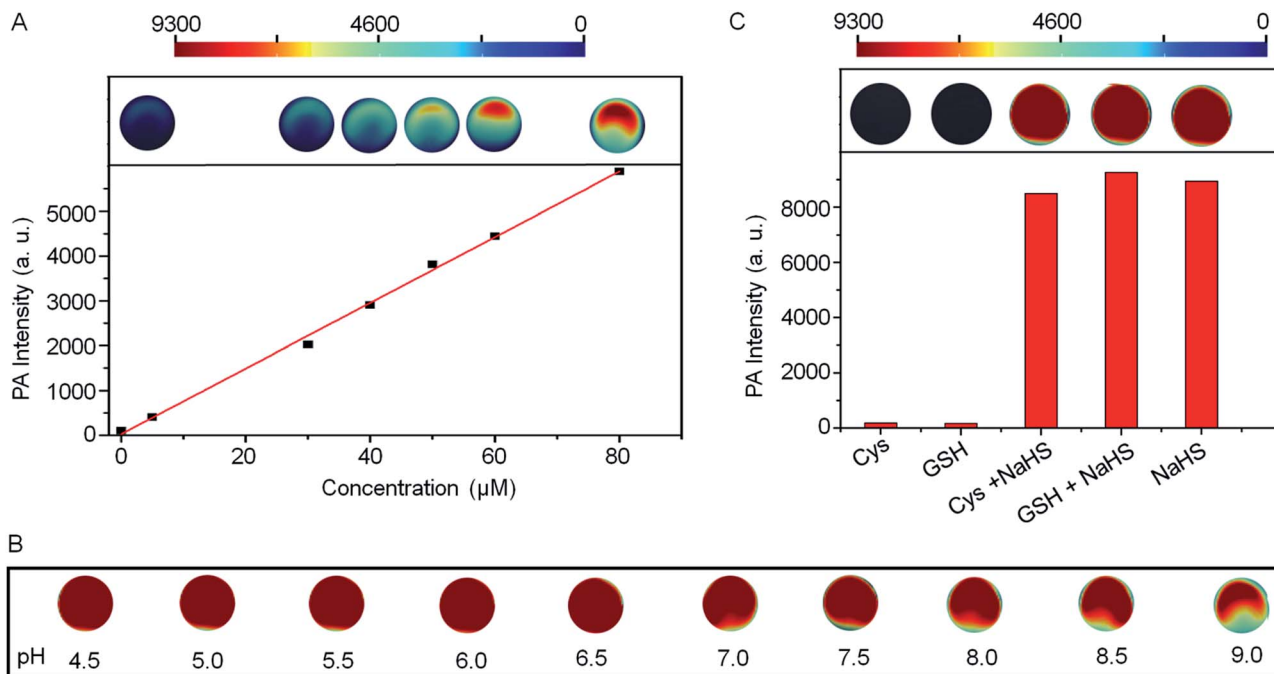
Fig. 2 (A) Representative photoacoustic images and amplitudes of Si@BODPA180 (80 μM BODPA1) in the absence and presence of NaHS (100 μM). (B) Representative photoacoustic images and amplitudes of Si@BODPA180 as a function of BODPA1 concentration upon addition of two molar equivalents of  $H_2S$ . (a): Si@BODPA180(7); (b): Si@BODPA180(4); (c): Si@BODPA180(1); these nanoparticles were obtained by employing the weight ratio (BODPA1/mPEG-DSPE) of 7/100, 4/100 and 1/100, respectively.

The PA intensities of Si@BODPA180(7) showed a linear and dose-dependent correlation with the concentrations of NaHS in the range of 0–80 μM (Fig. 3A). The detection limit was determined to be as low as 53 nM, suggesting sufficient sensitivity to detect endogenous  $H_2S$  in living systems. Furthermore, Si@BODPA180(7) was found to give excellent PA responsiveness to  $H_2S$  within a physiological range from pH 9 to approximately 4.5 (Fig. 3B).

Si@BODPA180(7) also showed a highly selective PA response to  $H_2S$  over various potentially interfering species (Fig. 3C and S6†). It was noticeable that only  $H_2S$  induced a bright PA signal upon excitation at 780 nm, while other analytes, including reactive sulfur (RSS), oxygen (ROS), and nitrogen (RNS) species triggered no obvious PA response. Of note, the strong PA signal elicited by  $H_2S$  was not affected by the presence of 1 mM GSH and 1 mM Cys in competition assays, highlighting the potential applications in a complex biological system.

With these promising results in hand, we next employed Si@BODPA180(7) for the *in vivo* photoacoustic imaging of endogenous  $H_2S$  generation in a HCT116 tumor-bearing mouse





**Fig. 3** (A) The linear plot of PA intensities of Si@BODPA180(7) (80 μM BODPA1) as a function of H<sub>2</sub>S concentrations. (B) The pH effects on the PA responsive process of Si@BODPA180(7) toward H<sub>2</sub>S. (C) Photoacoustic amplitude changes of Si@BODPA180(7) (80 μM BODPA1) in the presence of NaHS and other biologically relevant competing analytes in PBS (pH 7.4) at room temperature. Data shown are for 1 mM glutathione, 1 mM cysteine, and 100 μM NaHS.

model. It has been suggested that the H<sub>2</sub>S-producing enzyme cystathionine-β-synthase (CBS) is overexpressed in colon cancer, resulting in increased H<sub>2</sub>S production.<sup>13</sup> In this study, we employed Si@BODPA180(7) as an imaging tool to provide direct evidence for supporting such a functional relationship. In these experiments, saline or Si@BODPA180(7) were injected subcutaneously into tumor regions and normal regions, and images were recorded at various times after probes injection (Fig. 4). To our delight, robust activation of PA signals could be observed specifically in the tumor region in the probe-treated mice within 10 min post-injection (Fig. 4C), indicative of a rapid activation of the probe *in vivo*. Notably, minimal PA signals were observed in the normal site injected with the probe within 2 h (Fig. 4B). Furthermore, no detectable PA signals in the tumor and normal sites could be observed in the saline-treated mice (Fig. 4A). These results indicated that Si@BODPA180(7) can serve as a promising probe for imaging markedly enhanced H<sub>2</sub>S production in colon tumors.

In our study, we also found that the PA signals were greatly attenuated by the CBS inhibitor aminooxyacetic acid (AOAA) (Fig. 4D). In sharp contrast, addition of an allosteric CBS activator *S*-adenosyl-L-methionine (SAM) afforded dramatic elevation of PA signals (Fig. 4E). These preliminary imaging studies suggested that HCT116 colon tumors exhibited CBS upregulation activity which resulted in an increased rate of H<sub>2</sub>S generation. Undoubtedly, this activatable probe can be explored for trapping transient H<sub>2</sub>S and for the real-time monitoring of H<sub>2</sub>S-related biological processes *in vivo*.

## Experimental

### Live subject statement

All animal experiments were performed in compliance with the relevant laws and institutional guidelines for the Care and Use of Research Animals established by Fudan University Animal Studies Committee, and the experiments were approved by the committee.

### General procedure for preparation of Si@BODPAs

BODPAs (0.3 μmol) were rapidly poured into a solution of mPEG-DSPE (19.1 mg) in 3 mL 0.85 N hydrochloride solution under rigorous stirring, resulting in the formation of nanosized micelles. The *in situ* shell cross-linking under acidic conditions with desired amount of silylation agent was carried out at room temperature while kept stirring for 24 h. Then the aqueous solution was dialyzed for 2 days, and the stock solution was thus obtained, which can be diluted to the desired concentration for further studies. The concentration of BODPA1 or BODPA2 in Si@BODPAs was determined according to their standard UV-vis absorption.

The *in situ* shell cross-linking under acidic conditions with TBNBr of 5.73 mg (TBNBr/mPEG-DSPE = 30/100, w/w), 17.19 mg (TBNBr/mPEG-DSPE = 90/100, w/w), 34.38 mg (TBNBr/mPEG-DSPE = 180/100, w/w) and 51.57 mg (TBNBr/mPEG-DSPE = 270/100, w/w), afforded water dispersible nanocomposites Si@BODPA30, Si@BODPA90, Si@BODPA180 and Si@BODPA270, respectively.



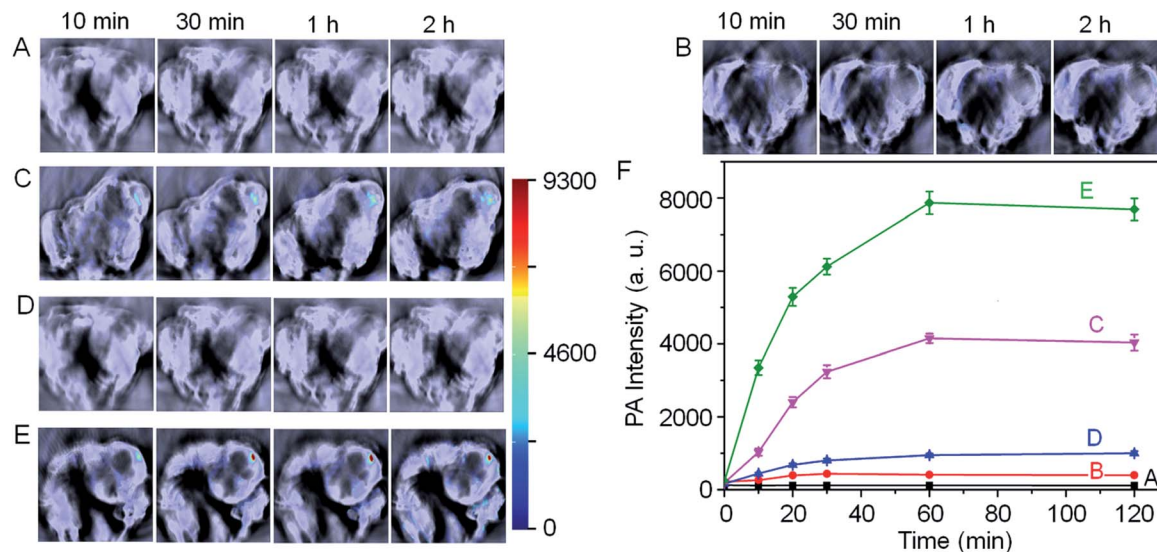


Fig. 4 *In vivo* photoacoustic imaging of tumor-bearing mice using Si@BODPA180(7) (10 nmol BODPA1). Saline or the probe in PBS was injected subcutaneously into the tumor regions and normal regions in the dorsal side of living mice, and images were taken at various time points after injection. (A) Saline-treated mice in the tumor regions; (B) probe-treated mice in the normal sites; (C) probe-treated mice in the tumor regions; mice pretreated with (D) 100 nmol AOAA, (E) 300 nmol SAM for 12 h, were subcutaneously injected Si@BODPA180(7) in the tumor regions. (F) PA intensities as a function of time post-injection of Si@BODPA180(7).

BODPA1 (0.3  $\mu\text{mol}$ ), mPEG-DSPE (19.1 mg) and TBNBr (34.38 mg) afforded Si@BODPA180(1); BODPA1 (1.2  $\mu\text{mol}$ ), mPEG-DSPE (19.1 mg) and TBNBr (34.38 mg) afforded Si@BODPA180(4); BODPA1 (2.1  $\mu\text{mol}$ ), mPEG-DSPE (19.1 mg) and TBNBr (34.38 mg) afforded Si@BODPA180(7).

## Conclusions

In summary, we have successfully presented an activatable photoacoustic probe for imaging endogenous  $\text{H}_2\text{S}$  in living mice. The probe was fabricated by encapsulating semi-cyanine-BODIPY hybrid dyes into the hydrophobic interior of core-shell silica nanocomposites, thus endowing the probe with good water-solubility and excellent biocompatibility. In the presence of  $\text{H}_2\text{S}$ , a high NIR absorption around 780 nm was triggered, ultimately leading to a strong photoacoustic signal output in the NIR region. Furthermore, the designed probe showed a highly selective PA response to  $\text{H}_2\text{S}$  with a favourable detection limit of 53 nM. In particular, this probe displayed an extremely fast response, permitting the trapping of transient  $\text{H}_2\text{S}$  and real-time monitoring of  $\text{H}_2\text{S}$ -related biological processes. More importantly, this probe could serve as a promising tool for the direct photoacoustic imaging of endogenous  $\text{H}_2\text{S}$  generation in a HCT116 tumor-bearing mouse model, verifying the activating effect of CBS upregulation on the elevated level of  $\text{H}_2\text{S}$ . To the best of our knowledge, this work represents the first example of NIR-activatable photoacoustic probes capable of reporting the endogenous production of  $\text{H}_2\text{S}$  in real time, which may facilitate further exploration of the complex roles of  $\text{H}_2\text{S}$  in living systems.

## Acknowledgements

We gratefully acknowledge the financial support from the National Science Foundation of China (grant numbers 21172071, 21190033, 21372083, 21572039), NSFC for Creative Research Groups (21421004), National 973 Program (2013CB733700) and the Programme of Introducing Talents of Discipline to Universities (B16017). We thank Prof. Ping Shi and Yajing Liu for assistance with *in vivo* imaging.

## Notes and references

- (a) H. Kimura, *Antioxid. Redox Signaling*, 2010, **12**, 1111; (b) N. Skovgaard, A. Gouliarov, M. Aalling and U. Simonsen, *Curr. Pharm. Biotechnol.*, 2011, **12**, 1385; (c) R. Wang, *Physiol. Rev.*, 2012, **92**, 791.
- (a) G. Tang, G. Yang, B. Jiang, Y. Ju, L. Wu and R. Wang, *Antioxid. Redox Signaling*, 2013, **19**, 1634; (b) K. Kondo, S. Bhushan, A. L. King, S. D. Prabhu, T. Hamid, S. Koenig, T. Murohara, B. L. Predmore, G. Gojon Sr, G. Gojon Jr, R. Wang, N. Karusula, C. K. Nicholson, J. W. Calvert and D. J. Lefler, *Circulation*, 2013, **127**, 1116.
- (a) P. Kamoun, M.-C. Belardinelli, A. Chabli, K. Lallouchi and B. Chadeaux-Vekemans, *Am. J. Med. Genet.*, 2003, **116A**, 310; (b) M. Lee, C. Schwab, S. Yu, E. McGeer and P. L. McGeer, *Neurobiol. Aging*, 2009, **30**, 1523; (c) W. Yang, G. Yang, X. Jia, L. Wu and R. Wang, *J. Physiol.*, 2005, **569**, 519; (d) B. D. Paul, J. I. Sbodio, R. Xu, M. S. Vandiver, J. Y. Cha, A. M. Snowman and S. H. Snyder, *Nature*, 2014, **509**, 96.
- (a) V. S. Lin, W. Chen, M. Xian and C. J. Chang, *Chem. Soc. Rev.*, 2015, **44**, 4596; (b) X. Zhou, S. Lee, Z. Xu and J. Yoon,



- Chem. Rev.*, 2015, **115**, 7944; (c) J. Chan, S. C. Dodani and C. J. Chang, *Nat. Chem.*, 2012, **4**, 973.
- 5 (a) A. R. Lippert, E. J. New and C. J. Chang, *J. Am. Chem. Soc.*, 2011, **133**, 10078; (b) H. Peng, Y. Cheng, C. Dai, A. L. King, B. L. Predmore, D. J. Lefer and B. Wang, *Angew. Chem., Int. Ed.*, 2011, **50**, 9672; (c) W. Xuan, C. Sheng, Y. Cao, W. He and W. Wang, *Angew. Chem., Int. Ed.*, 2012, **51**, 2282; (d) S. K. Bae, C. H. Heo, D. J. Choi, D. Sen, E.-H. Joe, B. R. Cho and H. M. Kim, *J. Am. Chem. Soc.*, 2013, **135**, 9915; (e) V. S. Lin, A. R. Lippert and C. J. Chang, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 7131; (f) M. D. Hammers, M. J. Taormina, M. M. Cerda, L. A. Montoya, D. T. Seidenkranz, R. Parthasarathy and M. D. Pluth, *J. Am. Chem. Soc.*, 2015, **137**, 10216; (g) D.-W. Li, L.-L. Qu, K. Hu, Y.-T. Long and H. Tian, *Angew. Chem., Int. Ed.*, 2015, **54**, 12758; (h) H. A. Henthorn and M. D. Pluth, *J. Am. Chem. Soc.*, 2015, **137**, 15330; (i) W. Chen, A. Pacheco, Y. Takano, J. J. Day, K. Hanaoka and M. Xian, *Angew. Chem., Int. Ed.*, 2016, **55**, 9993.
- 6 (a) C. Liu, J. Pan, S. Li, Y. Zhao, L. Y. Wu, C. E. Berkman, A. R. Whorton and M. Xian, *Angew. Chem., Int. Ed.*, 2011, **50**, 10327; (b) Y. Qian, J. Karpus, O. Kabil, S. Y. Zhang, H. L. Zhu, R. Banerjee, J. Zhao and C. He, *Nat. Commun.*, 2011, **2**, 495; (c) Y. Qian, L. Zhang, S. T. Ding, X. Deng, C. He, X. E. Zheng, H.-L. Zhu and J. Zhao, *Chem. Sci.*, 2012, **3**, 2920; (d) F. Yu, P. Li, P. Song, B. Wang, J. Zhao and K. Han, *Chem. Commun.*, 2012, **48**, 2852; (e) X. Wang, J. Sun, W. Zhang, X. Ma, J. Lv and B. Tang, *Chem. Sci.*, 2013, **4**, 2551; (f) Y. Chen, C. Zhu, Z. Yang, J. Chen, Y. He, Y. Jiao, W. He, L. Qiu, J. Cen and Z. Guo, *Angew. Chem., Int. Ed.*, 2013, **52**, 1688; (g) L. Zhang, H. Zhu, M. Li and X. Gu, *Chem. Commun.*, 2015, **51**, 13135; (h) C. Liu, W. Chen, W. Shi, B. Peng, Y. Zhao, H. Ma and M. Xian, *J. Am. Chem. Soc.*, 2014, **136**, 7257; (i) W. Chen, E. W. Rosser, T. Matsunaga, A. Pacheco, T. Akaike and M. Xian, *Angew. Chem., Int. Ed.*, 2015, **54**, 13961; (j) S. Wang, S. Xu, G. Hu, X. Bai, T. D. James and L. Wang, *Anal. Chem.*, 2016, **88**, 1434; (k) C. Zhao, X. Zhang, K. Li, S. Zhu, Z. Guo, L. Zhang, F. Wang, Q. Fei, S. Luo, P. Shi, H. Tian and W.-H. Zhu, *J. Am. Chem. Soc.*, 2015, **137**, 8490.
- 7 (a) K. Sasakura, K. Hanaoka, N. Shibuya, Y. Mikami, Y. Kimura, T. Komatsu, T. Ueno, T. Terai, H. Kimura and T. Nagano, *J. Am. Chem. Soc.*, 2011, **133**, 18003; (b) X. Gu, C. Liu, Y.-C. Zhu and Y.-Z. Zhu, *Tetrahedron Lett.*, 2011, **52**, 5000; (c) S. Sarkar, Y. S. Ha, N. Soni, G. Il An, W. Lee, M. H. Kim, P. T. Huynh, H. Ahn, N. Bhatt, Y. J. Lee, J. Y. Kim, K. M. Park, I. Ishii, S.-G. Kang and J. Yoo, *Angew. Chem., Int. Ed.*, 2016, **55**, 9365.
- 8 L. V. Wang and S. Hu, *Science*, 2012, **335**, 1458.
- 9 (a) K. Pu, A. J. Shuhendler, J. V. Jokerst, J. Mei, S. S. Gambhir, Z. Bao and J. Rao, *Nat. Nanotechnol.*, 2014, **9**, 233; (b) J. Levi, S. R. Kothapalli, T.-J. Ma, K. Hartman, B. T. Khuri-Yakub and S. S. Gambhir, *J. Am. Chem. Soc.*, 2010, **132**, 11264; (c) A. Dragulescu-Andrasi, S.-R. Kothapalli, G. A. Tikhomirov, J. Rao and S. S. Gambhir, *J. Am. Chem. Soc.*, 2013, **135**, 11015; (d) H. Li, P. Zhang, L. P. Smaga, R. A. Hoffman and J. Chan, *J. Am. Chem. Soc.*, 2015, **137**, 15628; (e) Q. Chen, X. Liu, J. Chen, J. Zeng, Z. Cheng and Z. Liu, *Adv. Mater.*, 2015, **27**, 6820; (f) Q. Miao, Y. Lyu, D. Ding and K. Pu, *Adv. Mater.*, 2016, **28**, 3662.
- 10 (a) J. F. Lovell, C. S. Jin, E. Huynh, H. Jin, C. Kim, J. L. Rubinstein, W. C. W. Chan, W. Cao, L. V. Wang and G. Zheng, *Nat. Mater.*, 2011, **10**, 324; (b) K. Pu, J. Mei, J. V. Jokerst, G. Hong, A. L. Antaris, N. Chattopadhyay, A. J. Shuhendler, T. Kurosawa, Y. Zhou, S. S. Gambhir, Z. Bao and J. Rao, *Adv. Mater.*, 2015, **27**, 5184; (c) C. Zhu, Y. Yang, M. Luo, C. Yang, J. Wu, L. Chen, G. Liu, T. Wen, J. Zhu and H. Xia, *Angew. Chem., Int. Ed.*, 2015, **54**, 6181; (d) L. Wang, P.-P. Yang, X.-X. Zhao and H. Wang, *Nanoscale*, 2016, **8**, 2488.
- 11 (a) X. Wu, S. Chang, X. Sun, Z. Guo, Y. Li, J. Tang, Y. Shen, J. Shi, H. Tian and W. Zhu, *Chem. Sci.*, 2013, **4**, 1221; (b) X.-D. Wang, J. A. Stolwijk, T. Lang, M. Sperber, R. J. Meier, J. Wegener and O. S. Wolfbeis, *J. Am. Chem. Soc.*, 2012, **134**, 17011; (c) Y. Li, A. Shao, Y. Wang, J. Mei, D. Niu, J. Gu, P. Shi, W. Zhu, H. Tian and J. Shi, *Adv. Mater.*, 2016, **28**, 3187; (d) B. M. Weckhuysen and J. Yu, *Chem. Soc. Rev.*, 2015, **44**, 7022.
- 12 S. Bai, H. Yang, P. Wang, J. Gao, B. Li, Q. Yang and C. Li, *Chem. Commun.*, 2010, **46**, 8145.
- 13 (a) C. Szabo, C. Coletta, C. Chao, K. Módis, B. Szczesny, A. Papapetropoulos and M. R. Hellmich, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 12474; (b) K. Módis, C. Coletta, A. Asimakopoulou, B. Szczesny, C. Chao, A. Papapetropoulos, M. R. Hellmich and C. Szabo, *Nitric Oxide*, 2014, **41**, 146.

