Make Caffeine Visible: a Fluorescent Caffeine ‘‘Traffic Light’’ Detector

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Caffeine has attracted abundant attention due to its extensive existence in beverages and medicines. However, to detect it sensitively and conveniently remains a challenge, especially in resource-limited regions. Here we report a novel aqueous phase fluorescent caffeine sensor named Caffeine Orange which exhibits 250-fold fluorescence enhancement upon caffeine activation and high selectivity. Nuclear magnetic resonance spectroscopy and Fourier transform infrared spectroscopy indicate that \( \pi \)-stacking and hydrogen-bonding contribute to their interactions while dynamic light scattering and transmission electron microscopy experiments demonstrate the change of Caffeine Orange ambient environment induces its fluorescence emission. To utilize this probe in real life, we developed a non-toxic caffeine detection kit and tested it for caffeine quantification in various beverages. Naked-eye sensing of various caffeine concentrations was possible based on color changes upon irradiation with a laser pointer. Lastly, we performed the whole system on a microfluidic device to make caffeine detection quick, sensitive and automated.

Caffeine, although only first isolated in 1819, has been extensively consumed as caffeinated nutrients throughout eastern and western cultures for over a thousand years. Its widespread consumption is attributed to its stimulating effects in the nerve cells. Caffeine from beverages such as Cola could enhance the performance of athletes, as well as reduce the possibility of type 2 diabetes. However, caffeine is yielding either positive or negative effects in various types of cancer. It may also cause headache, abnormal heart rhythms, especially in children and pregnant women with caffeine allergy. In addition, due to its abundant existence in domestic wastage, caffeine was found to be an important indicator of natural water system pollution by domestic drain. Therefore, it is urgent to develop a convenient sensor tool to detect caffeine.

Traditional caffeine detection methods include thin-layer chromatography (TLC), high-performance liquid chromatography-mass spectrometry (HPLC-MS), quartz balance, and immunoassay. However, these methods require expensive instruments and sophisticated handling that are not convenient for public usage. Chemosensors provide great prospects for caffeine identification. Recently, Waldvogel group reported the first artificial caffeine receptor, which is based on hydrogen bonding. Subsequent research modified this receptor to achieve better binding affinity and selectivity. Nevertheless, these interactions only occur in organic solvent which is not eligible in real life usage. The Reinholdt group then reported an aqueous phase caffeine sensor based on the chelation between its metal centre and the 5-nitrogen of caffeine. But its absorbance shift is not visible through naked eyes, thus requiring an absorbance detector. On the other hand, the Severin group reported an aqueous phase fluorescence turn-off caffeine sensor, HPTS, which binds caffeine based on \( \pi \)-stacking. The authors tested this sensor in different samples by extraction method, yet the incorporation of toxic organic solvent (chloroform) and the fluorescence turn-off feature renders difficulties in real-life application. Although they further developed a similar-structure ratiometric caffeine fluorescence sensor and incorporated it into a caffeine detection strip, it would still require more than 20 minutes for caffeine detection in beverages. Moreover, the relatively low fluorescence response adds difficulties to direct caffeine observation.

To achieve fast, sensitive and convenient caffeine detection, we developed an aqueous phase fluorescence turn-on caffeine sensor based on the screening of diversity-oriented fluorescence libraries (DOFL) and rational modification of the selected hit compound. This sensor exhibits tremendous fluorescence enhancement towards caffeine, which can be clearly seen even with naked eyes. To make caffeine visible for common people, we further
develop a caffeine detection kit which is able to purify and concentrate caffeine from different samples. The sensor could respond to caffeine and show bright orange-red colour like a warning sign. Combining both fluorescent sensor and the detection kit, customers can easily tell the caffeine content in their beverages, in less than one minute.

Results

Design of a BODIPY-based caffeine sensor. Figure 1 exhibits the structure of our caffeine sensor. It was achieved from screening of diversity-oriented fluorescence libraries (DOFL) and rational modifications31. One compound (BD-185) which exhibits fluorescence increase to caffeine has been discovered by an unbiased DOFL screening system. BD-185 contains a bromo-substituted indole motif conjugating to the BODIPY core and exhibits remarkably bright emission upon treatment with caffeine. To figure out the role of bromine-containing indole, we synthesized 4 derivatives of BD-185, which only modify the indole moiety. Among them, the BODIPY compound bearing a fluorine atom yielded highest fluorescent intensity towards caffeine (Fig. S1 to S5). Therefore, it was selected as the final caffeine sensor and named as Caffeine Orange (CO). CO exhibits dissociation constant (K_D) of 16.81 mM and detection limit of 50 μM (Fig. S6). The concentration-dependent colour changes could be observed under the irradiation of a green laser pointer at 532 nm by naked eyes (Fig. 1b). We then tested this interaction under different conditions. In organic solvents there was no caffeine dose-dependent fluorescence responses since (Fig. S7). However, when we dried the organic solutions, the fluorescence responses recovered and the fluorescence intensity was even stronger than the aqueous solution (Fig. S8). To our knowledge, this is the first report on BODIPY-based fluorescence turn-on caffeine sensor that works both in aqueous solution and in dry state.

Determination of the binding event. To elucidate how CO and caffeine interact, we first injected a mixture of CO and caffeine into the HPLC-MS. Two separate peaks were observed (Fig. S9), which indicates that there is no covalent interaction between CO and caffeine. Next, we performed Fourier transformed infrared spectroscopy (FT-IR) (Fig. S10a) in an attempt to identify the likely non-covalent interaction between CO and caffeine. Figure S10b shows the peak at 3,379 cm⁻¹ shifts to 3,289 cm⁻¹ and broadens. It refers to hydrogen bond between N–H group of CO and caffeine32. The shifted peaks of 790 cm⁻¹, 1,165 cm⁻¹ and 1,504 cm⁻¹ are due to π stacking, which causes bending vibration and ring torsions (Fig. S10c).33–35.

To further confirm this hydrogen bond and π stacking interactions, we performed NMR titration experiments. Figure 2a shows that the indole N–H proton shifts from 8.60 ppm to 9.58 ppm, which indicates the formation of hydrogen bond between CO and caffeine22. Majority of the aromatic protons of CO exhibit upfield shift of around 0.1 ppm as caffeine amount increases (Fig. 2b). In fluorine NMR spectra, the upfield chemical shift of indole F from 212.5 ppm to 212.6 ppm demonstrates the involvement of F atom in the interaction (Fig. S11a). Figure 2b & Figure S11b exhibit that caffeine aromatic proton and three methyl protons all show upfield shift of around 0.05 ppm. The NMR titration results confirm the mutual shielding effect of caffeine rings on CO molecule. Hence, we can infer that these two molecules interact by π stacking and hydrogen bond interactions.

BODIPY dyes are known to self-assemble and self-quench in aqueous solutions36,37. To understand how CO fluoresces upon addition of

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**Figure 1** | **Fluorescence responses of CO against caffeine.** (a) Fluorescence spectra of CO (10 μM) with different concentration of caffeine in water under excitation of 532 nm green light. Inset is the structure of CO. (b) photographs of CO (10 μM) aqueous solutions containing caffeine concentrations from 0 mM to 50 mM under the irradiation of a 532 nm green laser beam.
Caffeine, we first conducted dynamic light scattering (DLS) analysis. Figure S12 indicates that CO aggregates in aqueous phase with an approximate radius of 20 nm, which corresponds with previously published results. Next, to confirm the existence of CO aggregates and to analyse the condition of CO in the presence of caffeine, we performed imaging by transmission electron microscopy (TEM). As shown in Fig.S13, the size of CO-caffeine complex increases with the addition of caffeine. This indicates that the ambient environment of CO is changed with the addition of caffeine. CO-caffeine interaction disassembles the aggregates of CO molecules and leads to the formation of larger complexes. Thus, the self-quench of CO in aqueous solution is reverted and its intrinsic fluorescence shows up.

**Selectivity test and development of caffeine detection kit.** To test the possibility of developing CO into a feasible caffeine detection kit, we tested its selectivity against other bio-related molecules and caffeine analogs, which may interfere with its fluorescence response. Through in-house screening, we found that CO exhibits superior selectivity towards caffeine, which rules out the likelihood of interference during detection (Fig. S14 & Fig. S15, Table S1). Moreover, CO demonstrates good selectivity for various caffeine analogs, such as theophylline and theobromine, which have been difficult to differentiate by previously reported caffeine receptors (Fig. 3a, Fig. S16). Caffeine yielded a 2-fold and 5-fold fluorescence enhancement as compared to theophylline and theobromine, respectively, while none of the other caffeine analogs exhibit substantial fluorescence response towards CO. Thus, we can safely differentiate caffeine from theophylline and theobromine, based on their remarkably different fluorescence responses. Furthermore, the caffeine amount in beverages significantly outnumbers theophylline and theobromine, which means that CO can be reliably developed into a selective and sensitive caffeine detection kit.

To make caffeine visible, we developed a caffeine detection kit to extract caffeine from beverages with complicated impurities and tested its eligibility to differentiate normal coffee and decaffeinated coffee. A syringe inserted with reverse phase materials was used for the elimination of auto-fluorescent impurities. A short C4 column rinsed with 75% ethanol (EtOH)/H2O was first loaded with coffee samples, and then washed with K2CO3 (1 mM) and deionized water, followed by eluting with 15% EtOH/H2O. The eluent was collected into a selective and sensitive caffeine detection kit.

**Development of automated caffeine detection system.** To fully utilize our traffic-light caffeine sensor, we setup an automated system by incorporating microfluidics technique. The whole extraction process can be fully automated using a centrifugal system by incorporating microfluidics technique. The whole extraction and detection process can be completed within a minute, making it highly practical in real-life caffeine detection.

Samples can also be accurately measured with the help of fluorescence standard curve (Fig. 3b, Fig. S19 & S20). Furthermore, the whole extraction and detection process can be completed within a minute, making it highly practical in real-life caffeine detection. This further demonstrates the feasibility of developing CO into a portable caffeine detection kit and quantifying the caffeine amount in various beverages (Details in Methods and movie supporting information).

To test the applicability of our caffeine detection kit, we estimated the caffeine amounts of various beverages and compared them with the actual caffeine amounts acquired from HPLC measurements (Fig. 4). As shown in the Fig. 4, results from the HPLC and the fluorescence data correlate very well, thus we confirm that this caffeine detection kit could achieve rapid, as well as sensitive measurement of caffeine. For the convenience of general users, we constructed a caffeine concentration-dependent colour bar with high amount of caffeine exhibiting red colour and low amount of caffeine showing green colour (Fig. 4). Since different amounts of caffeine show different colours in our detection kit, by comparing the observed colour with the reference bar, we can easily estimate the amount of caffeine present in the sample. This is referred to as a “traffic-light caffeine amount designator”, with the reddish orange colour indicating a stop sign for people who cannot uptake caffeine, Yellow colour as a warning and green colour indicating a safe zone. Thus, users could rapidly make a decision as to enjoy the beverage or not.
the C4 column chamber and the input channel is blocked by closing valve #2. Then, the C4 column is washed by K2CO3 and DI water by actuation of valves #3 and #4, respectively. Then, the channel to the waste chamber is closed by laser irradiation on valve #5 and the caffeine is eluted and transferred to the detection chamber by actuation of the valves #5 and #6. The eluted caffeine is mixed with pre-stored CO and the final concentration is measured under excitation at 532 nm with an optical fiber-coupled spectrophotometer. The total spin process takes about 5 minutes and can be further reduced by modifying the spinning velocity (Table S3). Figure 3b shows CCD images at each step (movie supplementary information 2). The calibration curve obtained using samples with known caffeine concentration showed good dynamic range and reproducibility (Fig. 5c). Figure 5d shows data measured with real beverage samples using the lab-on-a-disc. Hence, using this lab-on-a-disc device, caffeine extraction and detection can be fully automated and users can easily measure caffeine amount.

**Discussion**

DOFL has shed light on sensor development in the past decade\(^42\–44\). Through the high-content screening of more than 8,000 fluorescent compounds that cover the whole UV-Vis spectrum, we found the prototype caffeine sensor (BD-185). The structural diversity of our fluorescent compounds renders us confidence that the interaction between BD-185 and caffeine is specific and selective, yet we would like to investigate more deeply into the binding event. First we ruled out the possibility of single BODIPY interaction, since this fluorescent core has been widely explored at most of the positions in our DOFL and only the indole-bound dye shows response towards caffeine. Thus we next selected out all of our indole-bound BODIPY dyes and test their responses. Figure S4 clearly exhibits that any modification to the effective structure, such as when the Br atom of indole is replaced with methoxy, methyl or simply removed, will result in disappearance of fluorescence responses. Though limited by the indole diversity, we have already learned that Br atom plays an important role in the interaction. As is already known, heavy atoms lead to low fluorescence quantum yield\(^45\). Furthermore, if Br participates into the interaction through intermolecular binding, it is reasonable to infer that lighter atoms could achieve stronger interactions. Thus we tried to replace Br with Cl and F and successfully observed stronger fluorescence responses and binding affinity (\(K_D(\text{Br}) > K_D(\text{Cl}) > K_D(\text{F})\), Fig. S6).

Two aqueous sensors were reported by Severin group, both of which utilize \(\pi\)-stacking as the driving force for binding caffeine\(^29\–30\). CO, on the other hand, interacts with caffeine not only through \(\pi\)-stacking, but also through hydrogen bonding and electrostatic inter-actions, as exhibited by FT-IR and NMR. However, \(\pi\)-stacking and other non-covalent forces are well-known to cause fluorescence quenching due to the formation of low-lying dark excited states\(^46\). Thus it is extremely difficult to induce substantial fluorescence turn-on phenomenon based on \(\pi\)-stacking. BODIPY dyes are well-known to exhibit self-assembling/self-quenching phenomenon in the aqueous phase. Combining these two properties, we designed this turn-on sensor. The interaction of CO and caffeine is strong enough to
disassemble CO aggregate and force large excess of caffeine to circle around the CO molecules as shown by TEM. Inside caffeine protective layer which is highly hydrophobic, CO molecules could fully stretch and maximally reduce inter-/intramolecular energy loss and thus its fluorescence emission could be observed.

The centrifugal microfluidic discs has emerged as a promising tool for point-of-care diagnostics devices to miniaturize and automate biochemical assays. For example, “Sample-in and answer-out” type of fully automated immunoassays and blood chemistry analysis have been demonstrated on a centrifugal microfluidic disc starting from whole blood. It has been also applied to environmental applications to test water and soil quality. Lafleur et al. measured organic pollutants directly on the sorbent materials after pre-concentration in a solid phase extraction (SPE) microcolumn on a disc. However, to the best of our knowledge, a fully automated integration of SPE protocols including sample injection, binding, washing, elution, and detection has never been demonstrated before. In the lab-on-a-disc used in this study, the sorbent materials are packed not in the radially located microcolumn, but between supporting frits located between the top and the bottom discs, which provide reduced flow velocity for the efficient SPE extraction. In addition, the outlet from the SPE chamber has the serpentine channel to control the flow resistance. With the lab-on-a-disc developed in this study, users can have on-site information about the caffeine amount simply by spinning the disc with just one manual step of injecting the beverage sample on a disc.

Prior to this caffeine “traffic-light” designator, no practically applicable and customer-friendly caffeine detection methods have been reported. The detection kit developed in our group has several advantages: (I) it is easy to construct and simple to handle. The whole kit requires just one syringe equipped with reverse-phase materials and several washing solutions. Its incorporation into automated system has enhanced the handling even greater. (II) The detection process is safe and customer-friendly. No organic solvent is involved in the extraction process. (III) The caffeine extraction procedure is very fast, taking less than one minute. The customers can apply the kit as soon as they wish to enjoy the beverages or measure any sample. (IV) The detection kit shows great adaptability. It has been proven that the detection could extract caffeine from different beverages that are both chemically and physically complicated. Partial purification could not only remove most of the auto-fluorescent impurities, but also is very timesaving. The remarkable selectivity of CO could greatly reduce the efforts to purify the samples. (V) The dose-dependent colour bar renders direct visualization of caffeine amount in the sample. Thus the detection could function like a pH paper and clearly guide the customers to make decision. (VI) The automated microfluidics system serves as an upgraded version of portable caffeine detector and produces an even safer and convenient approach for caffeine monitoring.

In summary, we report the first BODIPY-based fluorescence turn-on caffeine sensor that works in aqueous solution. The fluorescence colour changes with caffeine concentrations and shows a traffic-light distinction. The interaction properties of CO with caffeine has been systematically explored with LC-MS, FT-IR spectroscopy, NMR, DLS and TEM approaches. The results demonstrate that the change of CO ambient environment driven by hydrogen-bond and π-stacking between CO and caffeine molecules renders CO a remarkable fluorescence turn-on feature and good selectivity. A caffeine detection kit was developed in order to purify beverage samples and achieve the quantitative measurement of caffeine. To facilitate the usage of our caffeine detection kit in real life, we tested its eligibility for numerous beverages and constructed a “traffic-light caffeine amount designator”, which could help estimate caffeine amount even faster. Furthermore, the whole process can be automated using microfluidics devices. The caffeine sensor and detection kit not only can help

Figure 5: Microfluidics device that can be utilized for automated caffeine amount determination in beverages. (a) The photograph of a disc for fully automated caffeine detection. The disc design shows detailed microfluidic layout. The number indicates the order of the valve operation and arrows indicate the flow of reagents. The red circles with numbers are normally open-laser irradiated ferrowax microvalves and the blue circles with numbers are normally open-laser irradiated ferrowax microvalve. (b) The CCD images of the spinning disc at each reaction steps. (c) The calibration curve obtained using the lab-on-a-disc and caffeine solution with known concentration. Each data points are an average of 4 samples tested with 4 different discs. (d) Caffeine concentration measured by fully automated lab-on-a-disc with real beverage samples; Decaf (Caffe Vergnano 1882), Coca-Cola, Red Bull (energy), Coffee (Angelinus Americano), and Espresso (Nespresso Roma). Values are represented as means and error bars as standard deviations (n = 3).
enhance product safety during the extensive consumption of caffeine, but also serve as a practical path of uniting science with real life.

Methods

Synthesis of (E)-5,5-difluoro-3-(2-(5-fluor-o-1H-indol-3-yl)vinyl)-1-methyl-5H-
5,4,6-dipyrrolo[1,2-c:2′,1′-d][1,2,3]diazaborinine (Caffeine Orange). Caffeine was first dissolved in dichloromethane and then dried, hence well-dispersed mixture of caffeine was formed. The mixture was visualized with a green laser pointer (5 mW, 532 nm, 17. Pavlik, J. W. Tlc detection of caffeine in commercial products. J Chromatogr A 881, 307–316 (2002).

Reversal phase-syringe based caffeine extraction procedure. Reverse phase solid phase extraction (SPE) was prepared by breaking two OROCHEN 3 mL C4 SPE cartridges (200 mg, 3.5 mL size) and inserting the reverse phase gel into a BRAUN Injet 5 mL/Luer Solo syringe. The mixture was filtered through a 0.22 μm filter and eluted with 15% EtOH in water to obtain 1 mL of EtOH in water. The eluent was collected in a glass tube containing 15 μL 1 mM dye solution and 100 μL EtOH. The mixture was visualized with a green laser pointer (5 mW, 532 nm, Aurora).

Caffeine extraction procedure on a disc. Reverse phase SPE was prepared by packing 40 mg C4 gel on a disc. The reverse phase SPE was rinsed with 0.4 mL of 75% EtOH in H2O and caffeine was adsorbed on SPE by transferring 1 mL beverage through the cartridge. The SPE was washed in sequence with 1 mL K2CO3 (0.2 mL) and H2O (0.2 mL), followed by elution with 15% EtOH in H2O (0.2 mL) and mixed with 22 μL 0.1 mM dye solution.

1. Joossef, M. R., Beral, V., Rolfs, R. T., Aral, S. O. & Cramer, D. W. Are caffeinated beverages risk-factors for delayed conception. Lancet 335, 136–137 (1990).
2. Hieshima, Y., Sutherland, D. M. & Smith, A. P. Effect of caffeine following a day of normal consumption of caffeinated beverages. J Psychopharmacol 16, 458–468 (2002).
3. Lindskog, M. et al. Involvement of DARPP-32 phosphorylation in the stimulant action of caffeine. Nature 418, 774–778 (2002).
4. Bruning, R., Luethke, M. J. & Davis, J. E. Caffeinated and decaffeinated beverages equally restore fluid balance after exercise in the heat. Med Sci Sport Exerc 42, 112–113 (2010).
5. van Dam, R. M. & Feskens, E. J. M. Coffee consumption and risk of type 2 diabetes mellitus. Lancet 360, 1477–1478 (2002).
6. Valcic, S. et al. Inhibitory effect of six green tea catechins and caffeine on the growth of four selected human tumor cell lines. Anti-Cancer Drug Dev 7, 461–468 (1994).
7. Smith, S. J. et al. Alcohol, smoking, passive smoking and caffeine in relation to breast-cancer risk in young-women. Br J Cancer 70, 112–119 (1994).
8. Page, R. M. Perceived consequences of drinking caffeinated beverages. Percept Motor Skill 65, 765–766 (1987).
9. Chan, J. Y., Vip, T. T. & Jeng, J. H. The role of caffeinated beverages in dental caries. Med Hypotheses 33, 22–29 (1990).
10. Caan, B. J. & Goldhaber, M. K. Caffeinated beverages and low-birth-weight - a case-control study. Am J Public Health 79, 1299–1300 (1989).
11. Pastore, L. M. & Savitz, D. A. Case-control study of caffeinated beverages and preterm delivery. Am J Epidemiol 141, 61–69 (1995).
12. Savoca, M. R., Evans, C. D., Wilson, M. E., Harshfield, G. A. & Ludwig, D. A. The association of caffeinated beverages with blood pressure in adolescents. Arch Pediatr Adolesc Med 158, 473–477 (2004).
13. Al-Wadai, H. A. N., Takahashi, T. & Schuller, H. M. Caffeine stimulates the proliferation of human lung adenocarcinoma cells and small airway epithelial cells via activation of PKA, CREB and ERK1/2. Oncol Rep 15, 431–435 (2006).
14. Takahashi, J. L., Bulky, A. M., Briatico, L. & Dewey, A. M. Sex differences in reinforcing value of caffeinated beverages in adolescents. Behav Pharmacol 21, 731–741 (2009).
15. Daneshvar, A. et al. Evaluating pharmaceuticals and caffeine as indicators of fecal contamination in drinking water sources of the Greater Montreal region. Chemosphere 88, 131–139 (2012).
16. Saufe, S. et al. Fecal coliforms, caffeine and carbamazepine in stormwater collection systems in a large urban area. Chemosphere 86, 118–123 (2012).
17. Pavlik, J. W. Tlc detection of caffeine in commercial products. Journal of Chemical Education 50, 134–134 (1973).
18. Zuo, Y. G., Chen, H. & Deng, Y. W. Simultaneous determination of catechins, caffeine and gallic acids in green, Oolong, black and pu-erh teas using HPLC with a photodiode array detector. Talanta 57, 307–316 (2002).
19. Kobayashi, T., Murawaki, Y., Reddy, P. S., Abe, M. & Fujii, M. Molecular imprinting of caffeine and its recognition assay by quartz-crystal microbalance. Anal Chim Acta 435, 141–149 (2001).
20. Carvalho, J. J., Weller, M. G., Panne, U. & Schneider, R. J. A highly sensitive caffeine immunoassay based on a monoclonal antibody. Anal Bioanal Chem 396, 2617–2628 (2010).
21. Waldvogel, S. R. Caffeine - A drug with a surprise. Chemreacpt Com 360, 65–67 (2006).
22. Waldvogel, S. R., Frohlich, R. & Schalley, C. A. First artificial receptor for caffeine - A new concept for the complexation of alkylated oxopurines. Angew Chem Int Ed 39, 2472–2475 (2000).
23. Ballester, P. et al. Binding of caffeine by a synthetic co-receptor. Tetrahedron Lett 41, 3849–3853 (2000).
24. Goswami, S., Mahapatra, A. K. & Mukherjee, R. Molecular recognition of xanthine alkaloids: First synthetic receptors for theobromine and a series of new receptors for caffeine. J Chem Soc Perkin Trans 1, 2717–2726 (2001).
25. Wei, Y. L., Ding, L. H., Dong, C., Niu, W. P. & Shuang, S. M. Study on inclusion complex of cyclodextrin with methyl xanthine derivatives by fluorometry. Spectrochim Acta A 59, 2697–2703 (2003).
26. Siering, C., Kerschbaumer, H., Niegier, M. & Waldvogel, S. R. A supramolecular fluorescence probe for caffeine. Org Lett 8, 1471–1474 (2006).
27. Mahapatra, A. K., Sahoo, P., Goswami, S., Fun, H. K. & Yeap, C. S. First artificial acridine fluorescent receptors for caffeine and other xanthine alkaloids. J Incl Phenom Macroc 67, 99–108 (2010).
28. Fiammengo, R., Crego-Calama, M., Timmerman, P. & Reinhoudt, D. N. Recognition of caffeine in aqueous solutions. Chem–Eur J 19, 784–792 (2003).
29. Rochat, S., Steimann, S. N., Corminboeuf, C. & Severin, K. Fluorescence sensing of caffeine in water with polysulfonated pyrenes. Chem Commun 47, 10584–10585 (2011).
30. Luisier, N. et al. A ratimetric fluorescence sensor for caffeine. Org Biomol Chem 10, 7487–7490 (2012).
31. Lee, J. S. et al. Synthesis of a bodipy library and its application to the development of alkylated oxopurine receptors. J Am Chem Soc 131, 10077–10086 (2009).
32. Gorman, T. The evidence from infrared spectroscopy for hydrogen bonding: A case history of the correlation and interpretation of data. Journal of Chemical Education 34, 3 (1957).
33. Tamam, L. K. & Tatulian, S. A. Infrared spectroscopy of proteins and peptides in lipid bilayers. Q Rev Biophys 30, 365–429 (1997).
34. Eronoto, S., Kawai, Y. & Sugita, M. Infrared spectrum of poly(vinylidine fluoride). J Polym Sci A2 13, 461 (1968).
35. Xu, Y. J. & McKellar, A. R. W. The infrared spectrum of the Ar-CO complex - Comprehensive analysis including vanderWaals stretching and bending states. Mol Phys 88, 859–874 (1996).
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Author contributions

Y.T.C. conceived and supervised the overall project. W.X. performed all the compound syntheses, spectra measurement, FT-IR & NMR measurement, TEM measurement, caffeine detection kit development. T.H.K. & Y.K.C. conceived and performed the microfluidics caffeine extraction. D.T.Z. helped spectra measurement. J.C.E. helped NMR measurements and data analysis. L.Y.Z. helped FT-IR & NMR measurements. A.A.K. helped TEM measurements. B.K.A. helped caffeine detection kit development. W.X., T.H.K., D.T.Z., J.C.E., L.Y.Z., A.A.K., Y.K.C. and Y.T.C. co-wrote the paper.

Additional information

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