Nitric Oxide Turn-on Fluorescent Probe Based on Deamination of Aromatic Primary Monoamines
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Supporting Information

ABSTRACT: The stable, water-soluble, and nonfluorescent FA-OMe can sense nitric oxide (NO) and form the intensely fluorescent product dA-FA-OMe via reductive deamination of the aromatic primary amine. The reaction is accompanied by a notable increase of the fluorescent quantum yield from 1.5 to 88.8%. The deamination mechanism of FA-OMe with NO was proposed in this study. The turn-on fluorescence signals were performed by suppression of photoinduced electron transfer (PeT), which was demonstrated by density functional theory (DFT) calculations of the components forming FA-OMe and dA-FA-OMe. Furthermore, FA-OMe showed water solubility and good stability at physiological pHs. Moreover, the selectivity study indicated that FA-OMe had high specificity for NO over other reactive oxygen/nitrogen species. In an endogenously generated NO detection study, increasing the incubation time of FA-OMe with lipopolysaccharide (LPS) pretreated Raw 264.7 murine macrophages could cause an enhanced fluorescence intensity image. In addition, a diffusion/localization cell imaging study showed that FA-OMe could be trapped in Raw 264.7 cells. These cell imaging results demonstrated that FA-OMe could be used as a turn-on fluorescent sensor for the detection of endogenously generated NO.

1. INTRODUCTION
Nitric oxide (NO), produced by nitric oxide synthases in the biological systems, has been known as a ubiquitous signaling molecule. Despite being a simple molecule, NO is involved in diverse physiological and pathological pathways. Although NO is a key vertebrate biological messenger, the mechanisms by which it performs its diverse biological roles remain elusive. In order to comprehend the diverse biological roles of NO, several techniques such as chemiluminescence, colorimetry, electron paramagnetic resonance, electrochemistry, and fluorimetry have been developed. In view of its sensitivity, selectivity, spatiotemporal resolution, and experimental feasibility, fluorimetry, which exploits fluorescent probes to monitor analytes of interest under a fluorescence microscope, has been regarded as a most promising method to detect endogenous NO. There have been a number of small-molecule organic or inorganic fluorescent NO sensors reported to date. A two-component system comprised of an NO-reactive moiety coupling to an organic fluorophore has been widely investigated to develop NO-responsive organic molecule based sensors. The NO-reactive moiety acts as a modulator in the photoinduced electron transfer (PeT) mechanism. After reaction with NO or the NO oxidized products, the PeT fluorescence quenching property is suppressed and the fluorescence of the probe is restored. The o-diaminophenyl group is the most commonly used NO-reactive moiety in organic fluorescent probes. The electron-rich o-diaminophenyl group can react with NO under aerobic conditions to produce fluorescent triazole derivatives. However, there are some undesired handicaps, such as complicated and low-yield synthetic procedures and blank fluorescence, existing in these kinds of NO probes.

In this study, the possibility of using an aromatic primary monoamine moiety as an NO-reactive site for the modulation of PeT was explored. The basic concept is outlined in Figure 1. The study was inspired by the genotoxic effects of NO. The NO-induced deamination of cytosine, adenine, guanine, and 5-methylcytosine indeed represents an important NO-induced genotoxic mechanism. In addition, previous research reported that some aromatic amines react with NO or N2O3 to give the
corresponding deaminated aromatics.\textsuperscript{11} In addition, it was reported that fluorescence of a fluorescent molecule could be adjusted by changing the electronic influence of the nitrogen electron lone pair.\textsuperscript{12} Therefore, we have investigated the ability of NO to deaminate the aromatic primary amino group, thereby altering the electronic influence of the aromatic moiety on the fluorophore. To the best of our knowledge, a NO turn-on fluorescent probe through reductive deamination of an aromatic primary monoamine has never been reported.

Among the various aromatic amines, we prefer aromatic amine esters to be the NO-reactive moieties. The ester functionality can enhance the cellular uptake of a probe and ensure its trappability within the plasma membrane.\textsuperscript{13} In the current study, 5-amino-2-(6-hydroxy-3-oxo-3H-xanthen-9-yl)-benzoic acid methyl ester (FA-OMe) was synthesized as an NO turn-on fluorescent probe. The reaction product (dA-FA-OMe) of FA-OMe with NO under aerobic conditions was identified, and a possible reaction mechanism was proposed. The DFT calculations of dA-FA-OMe relative to FA-OMe were undertaken to interpret the PeT phenomenon. For realizing the level of fluorescence enhancement, the fluorescent properties of FA-OMe and dA-FA-OMe were studied and compared. In order to confirm their stability, the fluorescence intensities of FA-OMe and dA-FA-OMe under various pH conditions were measured. To comprehend the NO-detecting ability, time- and concentration-dependent studies were also investigated. In addition, the selectivity of FA-OMe for NO against other reactive oxygen and nitrogen species was investigated. Furthermore, in vitro fluorescence images were obtained.

2. EXPERIMENTAL SECTION

2.1. Materials and Instruments. All chemicals were obtained from commercial suppliers and used as received without further purification. Hydrogen peroxide (H\textsubscript{2}O\textsubscript{2}), dehydroascorbic acid (DHA), ascorbic acid (AA), ethylenediaminetetraacetic acid (EDTA), lipopolysaccharide (LPS), and fluoresceinamine were purchased from Sigma-Aldrich. Sodium nitrite (NaNO\textsubscript{2}) and sodium pyruvate were purchased from Cellgro. Fetal bovine serum (FBS), 10% sodium bicarbonate, and lipopolysaccharide were purchased from J. T. Baker. Dulbecco’s Modified Eagle Medium (DMEM) and sodium pyruvate were purchased from Cellgro. Methanol was distilled over Mg/I\textsubscript{2} and subsequently dried over 0.3 nm molecular sieves. Preparative thin-layer chromatographic (TLC) plates were prepared from silica gel (Macherey-Nagel).\textsuperscript{1} H and \textsuperscript{13}C NMR spectra were obtained on a Varian VXR-300 spectrometer at 300 and 75 MHz and were referenced to the internal \textsuperscript{1}H and \textsuperscript{13}C solvent peaks. ESI-LRMS spectra were recorded on a Micromass Q-TOF mass spectrometer. EI-HRMS spectra were recorded on a Finnigan/Thermo Quest MAT mass spectrometer. UV/vis absorption spectra were recorded on a Hitachi U-3000 spectrophotometer. Bright-field and fluorescence images were recorded on an Olympus IX71 fluorescence microscope equipped with a 100 W mercury lamp, B-2A filters, and a color CCD camera system.

2.2. Preparation of S-Amino-2-(6-hydroxy-3-oxo-3H-xanthen-9-yl)benzoic Acid Methyl Ester (FA-OMe). A modification of the synthetic procedure of FA-OMe was employed.\textsuperscript{15} A mixture of fluoresceinamine (200 mg, 0.576 mmol), MeOH (10 mL), and concentrated H\textsubscript{2}SO\textsubscript{4} solution (0.2 mL) was heated for 48 h under reflux. The solution was poured on ice (300 g), and then a saturated NaHCO\textsubscript{3} solution was slowly added to adjust the pH to 6.5. An orange precipitate was separated by centrifugation and then subjected to TLC, with MeOH/CH\textsubscript{3}Cl\textsubscript{2} as eluant (1/12, v/v). A yellow band was isolated and identified as 5-amino-2-(6-hydroxy-3-oxo-3H-xanthen-9-yl)-benzoic acid methyl ester (FA-OMe; 184 mg, 88.5%). ESI-LRMS: m/z 361.1 ([M + H]+), M\textsuperscript{+} calculated 361.1. \textsuperscript{1}H NMR (DMSO-d\textsubscript{6}): δ 7.30 (s, 1H), 7.00 ([d, J = 9.0 Hz, 1H], 6.90 ([d, J = 9.0 Hz, 1H], 6.77 ([d, J = 9.0 Hz, 2H], 6.30 ([d, J = 9.0 Hz, 2H], 6.23 ([s, 2H]), 5.81 ([br s, 2H], 3.49 ([s, 3H], 1.4C NMR (DMSO-d\textsubscript{6}): δ 178.4, 166.2, 157.3, 152.9, 149.7, 131.5, 130.8, 129.8, 122.8, 120.7, 117.0, 114.7, 111.1, 103.1, 51.9.

2.3. Reaction of FA-OMe with NO/\textsubscript{O}_{2}. In 100 mL of an FA-OMe (30 mg, 0.083 mmol) solution in 100 mM HEPES buffer at pH 7.4, 10% NO gas was bubbled under aerobic conditions at room temperature. The reaction was terminated using TLC to ensure the consumption of all of FA-OMe. Then the solvent was removed on a rotary evaporator. The residue was subjected to TLC, with MeOH/CH\textsubscript{3}Cl\textsubscript{2} as eluant (1/10, v/v). The first bright yellow band was isolated and identified as 2-(6-hydroxy-3-oxo-3H-xanthen-9-yl)benzoic acid methyl ester (dA-FA-OMe; 224 mg, 78.0%). ESI-LRMS: m/z 347.1 ([M + H]+), M\textsuperscript{+} calculated 346.1. EI-HRMS: m/z 346.0836 (M\textsuperscript{+}), M\textsuperscript{+} calculated 346.0841. \textsuperscript{1}H NMR (DMSO-d\textsubscript{6}): δ 8.12 ([d, J = 7.8 Hz, 1H], 7.80 ([d, J = 7.5 and 7.8 Hz, 1H], 7.77 ([d, J = 7.5 and 7.8 Hz, 1H], 7.40 ([d, J = 7.5 Hz, 2H], 6.59 ([d, J = 9.0 Hz, 2H], 6.30 ([m, 4H], 3.54 ([s, 3H], 1.4C NMR (DMSO-d\textsubscript{6}): δ 187.4, 166.2, 157.3, 152.9, 149.7, 131.5, 130.8, 129.8, 122.8, 120.7, 117.0, 114.7, 111.1, 103.1, 51.9.

2.4. Reaction of Fluoresceinamine with NO/\textsubscript{O}_{2}. In 45 mL of a fluoresceinamine (10 mg, 0.029 mmol) solution in 100 mM HEPES buffer at pH 7.4, 10% NO gas was bubbled under aerobic conditions at room temperature. The reaction was terminated using TLC to ensure the consumption of all of fluoresceinamine. The solvent was removed on a rotary evaporator. The residue was subjected to TLC, with MeOH/CH\textsubscript{3}Cl\textsubscript{2} as eluant (1/10, v/v). The first bright yellow band was isolated and identified as fluorescein (7.54 mg, 78.3%).

2.5. Preparation of NO Stock Solutions. The preparation of NO and its stock solutions were carried out according to the reported methods.\textsuperscript{16} NO could be generated by slowly dropping 2.0 M H\textsubscript{2}SO\textsubscript{4}(aq) into a glass flask containing a saturated NaN\textsubscript{O}\textsubscript{2} water solution. Since O\textsubscript{2} will rapidly oxidize NO to form NO\textsubscript{2}, all apparatus was carefully degassed with argon for 30 min to remove O\textsubscript{2}. The forming gas was passed twice through a 30% NaOH solution and once through water to trap NO\textsubscript{2} generated from the reaction of NO with traces of O\textsubscript{2}. To produce a saturated NO solution (1.8 mM, at 20 °C) as a stock solution, 10% of deoxyglucose (dextrose) was bubbled with NO for 30 min and kept under an NO atmosphere. The saturation concentration was ascertained by using the Griess method.\textsuperscript{3a} Stock solutions were freshly prepared for each experiment and stored in a glass flask with a rubber septum.

2.6. Fluorometric Analysis. Fluorescence spectra were recorded by using a Hitachi F-7000 spectrophotometer. The slit widths were 5.0 and 2.5 nm for excitation and emission, respectively. The photon multiplier voltage was 500 V. Spectra were routinely acquired at 25.0 ± 0.1 °C in quartz cuvettes with a volume of 3.5 mL.

2.6.1. Fluorescence Quantum Yield Determination. Fluorescence quantum yields of FA-OMe and dA-FA-OMe were measured in 0.1 M NaOH(aq) by using fluorescein (Φ\textsubscript{f} = 0.95 in 0.1 M NaOH(aq)) as a standard.\textsuperscript{26} Fluorescence quantum yields were calculated by use of the following equation: Φ\textsubscript{unknown} = Φ\textsubscript{reference}(A\textsubscript{unknown}/A\textsubscript{reference})(I\textsubscript{unknown}/I\textsubscript{reference})\textsuperscript{2}, where
resceinamine were measured after 2 h of mixing (mM HEPES in the pH range 6–10 were prepared by mixing 0.2 mL of 500 μM solutions of FA-OMe in deionized water with 4.8 mL of 100 mM HEPES whose pH values were adjusted by hydrogen chloride and sodium hydroxide. Solutions of 20 μM FA-OMe and 20 μM fluoresceinamine in 100 mM HEPES in the pH range 6–10 were prepared as described above. The fluorescence intensities of FA-OMe, dA-FA-OMe, and fluoresceinamine were measured after 2 h of mixing (λ<sub>ex</sub> 460 nm, λ<sub>em</sub> 524 nm).]

2.6.3. Selectivity Studies. All the fluorescence tests were performed in 100 mM HEPES buffer at pH 7.4 and 25.0 ± 0.1 °C. The fluorescence intensities were monitored after reacting 3 mL of 20 μM FA-OMe (in 100 mM HEPES buffer at pH 7.4) with 100 equiv of reactive oxygen species (ROS), reactive nitrogen species (RNS), ascorbic acid (AA), and dehydroascorbic acid (DHA) for 2 h, respectively (λ<sub>ex</sub> 460 nm, λ<sub>em</sub> 524 nm). The concentration of the ONOO· stock solution was calculated by using its molar extinction coefficient of 1670 M<sup>−1</sup> cm<sup>−1</sup> at 302 nm. The degassed aqueous solutions of Angeli’s salt were freshly prepared under anaerobic conditions and used as a nitrite (NO<sub>2</sub><sup>−</sup>) source. Hydroxyl radical (·OH) was produced by reacting ferrous ammonium sulfate hexahydrate (Fe(NH<sub>4</sub>)<sub>2</sub>(SO<sub>4</sub>)<sub>2</sub>·6H<sub>2</sub>O) with H<sub>2</sub>O<sub>2</sub> under anaerobic conditions. Hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) was diluted promptly from the 30% H<sub>2</sub>O<sub>2</sub> solution and quantified by measuring its absorbance at 240 nm with a molar absorption coefficient of 43.6 M<sup>−1</sup> cm<sup>−1</sup>. The aqueous solutions of NaN<sub>3</sub> and NaNO were freshly prepared and used as nitrate (NO<sub>3</sub><sup>−</sup>) and nitrite (NO<sub>2</sub><sup>−</sup>) sources, respectively. The nitrogen-purged aqueous solutions of AA and DHA in 100 mM HEPES buffer at pH 7.4 containing 3 mM EDTA were freshly prepared before use.

2.7. Computational Methods. Density functional theory (DFT) calculations at the B3LYP/6-31++G** level were used to obtain optimized geometries and energy levels of frontier molecular orbitals for methyl 3-aminobenzoate, methyl benzoate, and xanthene, the components forming FA-OMe and dA-FA-OMe. The hydration effects were considered by performing single-point energy calculations with the polarizable continuum model on gas-phase optimized structures. All calculations were carried out with the Gaussian 09 program.

2.8. Cell Culture and In Vitro Cell Imaging. Raw 264.7 murine macrophage is organized from tumor ascites induced by intraperitoneal injection of Abselon Leukemia Virus (A-MuLV) in male mice. Raw 264.7 murine macrophages were obtained from the American Type Culture Collection (Manassas, VA). The cells were cultured in DMEM and supplemented with 10% FBS, 1% sodium pyruvate, and 1% MEM nonessential amino acids at 37 °C under a humidified 5% CO<sub>2</sub> atmosphere. For imaging studies, Raw 264.7 murine macrophages were passed and plated into poly-D-lysine coated plates containing 2 mL of DMEM and incubated at 37 °C with 5% CO<sub>2</sub>. For endogenously generated nitric oxide detection studies, iNOS (inducible nitric oxide synthase) was induced in Raw 264.7 murine macrophages with 0.5 μg mL<sup>−1</sup> of LPS for 4 h, and the cells were then coincubated with 10 μM of FA-OMe for 4 and 8 h. Prior to imaging, the cells were washed three times with 1 mL of PBS and then bathed in 2 mL of PBS. The fluorescence alterations were monitored by the fluorescence microscope. For FA-OMe diffusion/localization studies, Raw 264.7 murine macrophages were pretreated with 1.25 μg mL<sup>−1</sup> of LPS for 4 h, and then the cells were coincubated with 10 μM of FA-OMe for 8 h. The cells were washed three times with 1 mL of PBS prior to imaging and then bathed in 2 mL of PBS during the imaging procedure. To mimic media flow, the PBS was removed, and...
the cells were washed three times with 2 mL of fresh PBS and then bathed in 2 mL of fresh PBS while on the fluorescence microscope. This process was repeated every 10 min during imaging.

2.9. Cell Viability Studies. The Raw 264.7 murine macrophages were grown in 96-well plates at an initial density of 10^5 cells per well for 24 h. Subsequently, different concentrations of FA-OMe were incubated with Raw 264.7 murine macrophages at 37 °C for 24 h. After incubation, the supernatant was removed and the cells were washed three times with PBS buffer solution. Cell viability was evaluated using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) reduction assay. Briefly, MTT (20 μL, 5 mg mL⁻¹) was added to each well. After 4 h of incubation, each well was treated with dimethyl sulfoxide (100 μL) with pipetting. The absorbance at 570 nm was measured on a plate reader. Each treatment was done in six wells, and the experiments were repeated three times. Cell viability was calculated relative to the absorbance of the control for each treatment. The viability of untreated cells was assumed to be 100%.

3. RESULTS AND DISCUSSION

3.1. Synthesis, Characterization, and Proposed Mechanism. FA-OMe was easily prepared by refluxing the MeOH solution of fluoresceinamine with acid catalyst. To grossly observe the optical behavior of FA-OMe, NO gas was bubbled into an aqueous solution of FA-OMe. The solution immediately changed from yellow to light green. This indeed qualitatively establishes that FA-OMe is reactive toward NO. The reaction product dA-FA-OMe was purified, and the structure was elucidated using NMR spectroscopy and mass spectrometry. Thus, FA-OMe can detect NO and produce strongly fluorescent dA-FA-OMe through reductive deamination of FA-OMe.

The 1H NMR spectra of FA-OMe and dA-FA-OMe are shown in Figure 2. FA-OMe and dA-FA-OMe exhibit the expected idealized C_{2v} symmetry in solution due to the exchange process of the carbonyl and hydroxyl of the xanthene moiety. The lack of an apparent broad signal at around 11.0 ppm shows that the hydroxyls of FA-OMe and dA-FA-OMe may be deprotonated in the solution. Thus, eight types of protons in different chemical environments for FA-OMe (a–h) and dA-FA-OMe (a′–h′) are observed, respectively. As shown in Figure 2B, b′ and c′ signals of dA-FA-OMe present the splitting pattern of a doublet of doublets (dd, J = 7.5 and 7.8 Hz) at 7.80 and 7.71 ppm, respectively. In addition, the f′ signal merging with the g′ signal is a doublet signal, like the e signal of FA-OMe, as shown in Figure 2A. The primary amino group of FA-OMe, because of its electron-donating effect, increases the valence electron density around the protons of the phenyl ring. Therefore, the protons (a–c) are shielded higher than those of dA-FA-OMe (a′–d′) without an amino group. This result makes the a–c signals shift upfield in comparison to a′–d′ signals. The 13C NMR spectra of FA-OMe and dA-FA-OMe display 15 resonance signals, respectively, which indeed correspond with the C_{2v} symmetry (Figure S1 in the Supporting Information).

The reaction mechanism for deamination of FA-OMe is proposed as shown in Scheme 1. First, the primary amino group of FA-OMe is nitrosylated by N₂O₃, which forms by the autoxidation of NO,²³ to produce the N-nitrosoamine intermediate 1.⁹g Subsequently, nitrite ion reacts with 1 to produce the nitramine intermediate.²⁴ Afterward, the secondary amine of 2 is nitrosylated by N₂O₃ to form the N-nitroso nitramine intermediate 3, and then the N-nitroso nitramine moiety of 3 rearranges into the diazo nitrate moiety to form the diazo nitrate intermediate 5. It was reported that some diazo nitrates could decompose by homolytic bond rupture to form the corresponding radicals, nitrogen gas, and nitrate radical.²⁵ Thus, the diazo nitrate intermediate 5 homolytically decomposes to give the fluoresceinyl radical 6, which readily abstracts hydrogen from the solvent to give the deaminated product dA-FA-OMe. According to the ESI-mass spectrum of FA-OMe reacting with NO(aq) for 1.0 min under aerobic conditions.
(Figure S2 in the Supporting Information), the peaks corresponding to the proposed intermediates 1, 2, and 5 (or 3 and 4) were certainly observed. These results supported the proposed mechanism. However, further experimental studies are necessary to completely demonstrate our proposed mechanism. In other words, the proposed intermediates or their derivatives formed by a specific reagent (trap) should be isolated and fully characterized. The related experimental studies are currently under investigation.

3.2. Computations. A computational study was carried out using the Gaussian 09 program to understand the theoretical aspects of the change in fluorescence intensity. Density functional theory (DFT) calculations at the B3LYP/6-31+** level were applied to obtain the optimized geometries and energy levels of frontier molecular orbitals of the components forming FA-OMe and dA-FA-OMe (methyl 3-aminobenzoate, methyl benzoate, and xanthene). As shown in Figure 3, the highest occupied molecular orbital (HOMO) of the methyl 3-aminobenzoate moiety (electron donor) matches that of the xanthene moiety (electron acceptor); therefore, when the latter is photoexcited, the intramolecular electron transfer from the former to the latter is energetically favorable. In contrast, the HOMO of the methyl benzoate moiety is significantly lower than that of the xanthene moiety; consequently, the intramolecular electron transfer from the former to the latter is energetically unfavorable. The DFT calculations indicate that the behavior of the NO turn-on fluorescence of FA-OMe is associated with the PeT phenomenon.

3.3. Optical Properties of FA-OMe and dA-FA-OMe. The fluorescence properties of FA-OMe and dA-FA-OMe are summarized in Table 1. The ultraviolet-visible (UV/vis) spectrum profiles of FA-OMe and dA-FA-OMe are not significantly different, as shown in Figure 4. However, the quantum yield of dA-FA-OMe is approximately 59-fold higher than that of FA-OMe. Consequently, even a small amount of dA-FA-OMe formed in the reaction of FA-OMe with NO can produce remarkable fluorescence enhancement.

![Figure 3. Energy levels of the methyl 3-aminobenzoate, methyl benzoate, and xanthene moieties. The values in parentheses were derived from the calculations considering the aqueous environment.](image)

![Table 1. Fluorescence Quantum Yields (Φfl) and Absorption and Emission Maxima (λabs, λem) of FA-OMe and dA-FA-OMe in 0.1 M NaOH(aq).](image)

<table>
<thead>
<tr>
<th>compd</th>
<th>λabs (nm)</th>
<th>λem (nm)</th>
<th>Φfl (%)</th>
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<tr>
<td>FA-OMe</td>
<td>488</td>
<td>518</td>
<td>1.5</td>
</tr>
<tr>
<td>dA-FA-OMe</td>
<td>490</td>
<td>514</td>
<td>88.8</td>
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3.4. Stability of FA-OMe and dA-FA-OMe. Since the stability of probes under various pH conditions concerns the biological applicability, the effects of pH on the fluorescence intensities of FA-OMe and dA-FA-OMe were investigated at different pHs ranging from 6.5 to 10, as shown in Figure 5. A significant change in the fluorescence intensity of FA-OMe was not observed with an increase in pH from 6.6 to 10. In contrast, the gradually increasing fluorescence intensity of dA-FA-OMe reached a maximum value at pH 7.4. However, the fluorescence intensity ratio of dA-FA-OMe to FA-OMe shows high values between pH 6.5 and 10. This result indicates that FA-OMe can be used as an efficient fluorescent probe for the detection of NO under physiological conditions.

3.5. Concentration and Time Dependence of FA-OMe Reacting with NO. The concentration- and time-dependent reactions between FA-OMe and NO were carried out in 100 mM HEPES buffer at pH 7.4. To 50 μM FA-OMe solutions were added different amounts of the NO stock solution (1.8 mM). The fluorescence intensity at 524 nm is enhanced upon increasing the amount of NO(aq), as shown in Figure 6. Furthermore, the time-dependent fluorescence turn-on of FA-OMe shows a highly uniform pattern with different amounts of NO, as shown in Figure 7. The fluorescence intensities at 524 nm increase with time and reach a plateau within about 20 min. These results indicate that the NO concentration and reaction time influence proportionally the turn-on fluorescent intensity of FA-OMe. In addition, judging from Figure 6, the
enhancement in fluorescence intensity presents a linear correlation with concentration. The detection limit for NO, calculated as the concentration of NO at which the fluorescence signal is equal to the blank value plus a 3-fold standard deviation, was estimated as ~44 nM. Therefore, FA-OMe could be used for detecting endogenous NO in living cells.

3.6. Selectivity. The high specificity of the NO probe is imperative for its application in understanding the diverse biological roles played by NO. Therefore, the fluorescence intensities of FA-OMe in the presence of a series of possible competitive reactive oxygen species (ROS), reactive nitrogen species (RNS), AA, and DHA at up to 100-fold excess were examined under the same conditions. As shown in Figure 8, no detectable responses for these agents were observed under the same conditions. As shown in Figure 8, no detectable responses for these agents were observed. It was reported that the commercially available NO probe DAF-2 reacted with DHA or AA and turned on the probe under physiological conditions. Additionally, in the presence of H2O2/peroxidase, OONO−, or *OH, the o-diaminophenyl group is oxidized into an unstable arylaminyl radical, which can directly combine with NO to form triazole.

The better selectivity of FA-OMe for NO, in contrast to that of DAF-2, may be attributed to its less electron rich NO-reactive moiety. However, because of its highly electron rich o-diaminophenyl group, DAF-2 has a comparatively high reaction rate for NO according to the time-dependent experiments of DAF-2 and FA-OMe with 10 equiv of NO(aq) (Figure S3 in the Supporting Information).

3.7. Fluorescence Imaging of NO in Living Cells. The cytotoxicity of FA-OMe was investigated by incubating various concentrations of FA-OMe with Raw 264.7 murine macrophages. After 24 h, cell viability was examined by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay and the results demonstrated that FA-OMe showed low cytotoxicity to Raw 264.7 murine macrophages, as shown in Figure 9. To evaluate the utility of FA-OMe for the detection of endogenously generated NO, fluorescence microscopic imaging of biologically produced NO in Raw 264.7 murine macrophages was performed. The Raw 264.7 murine macrophages were stimulated by lipopolysaccharide (LPS; 0.5 μg mL−1) for 4 h and then incubated with 10 μM FA-OMe for an additional 4 and 8 h. In a control experiment, cells were incubated for 8 h with 10 μM FA-OMe in the absence of LPS. As shown in Figure 10B,C, a visible increase in fluorescence was observed over 4–8 h following FA-OMe incubation (8–12 h after induction of iNOS), which was not observed in the control cells (Figure 10A). These results indicate that FA-OMe can detect endogenously produced NO. To test the cell trappability of FA-OMe, Raw 264.7 murine macrophages prestimulated with 1.25 μg mL−1 LPS were incubated with 10 μM FA-OMe for 8 h and then washed and imaged by the fluorescence microscope four times over the course of 30 min, where a minimal change in fluorescence intensity was observed, as shown in Figure 11. Therefore, FA-OMe has cell-trappable characteristics. Since the ester groups can be cleaved by intracellular esterases to give carboxylates under physiological pH conditions to restrict recrossing of the cell membrane, the cell trappability may be caused by hydrolyzing FA-OMe into FA-OMe-carboxylate, which may react with NO/O2 to form dA-FA-OMe-carboxylate.

To prove that the turn-on fluorescence would be not achieved by FA-OMe-carboxylate in living cells, pH-dependent studies of FA-OMe and fluoresceinamine, which would transform into FA-OMe-carboxylate in aqueous solutions above pH 4,12a were undertaken in the pH range of 6.5–10, as shown in Figure 12. The fluorescence intensities are not apparently different at this pH range. These results show that FA-OMe-carboxylate cannot turn on the fluorescence. On the basis of the formation
of strongly fluorescent fluorescein, which would transform into dA-FA-OMe-carboxylate in aqueous solutions above pH 4.0, in the reaction of fluoresceinamine with NO/O₂ under neutral pH conditions, FA-OMe-carboxylate can react with NO/O₂ to generate dA-FA-OMe-carboxylate in living cells. Therefore, FA-OMe can be used as a fluorescent probe for the detection of NO formed in living cells.

4. SUMMARY
In this study, we have demonstrated the first example of an NO turn-on fluorescent probe based on reductive deamination of an aromatic primary monoamine. The DFT calculations of dA-FA-OMe relative to FA-OMe suggest that the fluorescence enhancement is associated with the PeT phenomenon. The high quantum yield of deaminated product shows that even a small deamination induced by NO can result in significant fluorescence enhancement. In addition, the use of a less
Figure 12. pH dependence of the fluorescence intensities of 20 μM fluoresceinamine (▲) and FA-OMe (◆) in 100 mM HEPES buffer at 25.0 ± 0.1 °C. The fluorescence intensities were detected at 524 nm with excitation at 460 nm. Inset: comparison to that of 20 μM dA-FA-OMe (●) in 100 mM HEPES buffer at 25.0 ± 0.1 °C.


