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A novel method for the synthesis of 1,2-benzisoxazoline-3-one and its application to hypochlorite recognition†

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The reaction of salicylhydroxamic acid with hypochlorite produces 1,2-benzisoxazoline-3-one, a heterocycle that contains a fluorophore. As a result, this reaction was used as the basis for a new, selective and sensitive fluorescence system for the recognition of hypochlorite. The effectiveness of the method was demonstrated by its use to detect hypochlorite in a disinfectant solution as well as to image hypochlorite in cells.

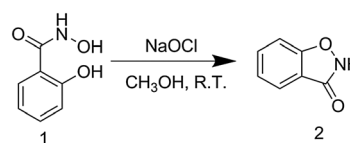
In modern-day organic chemistry, a growing trend exists to develop green and sustainable synthetic processes.^{1,2} The preparation of heterocyclic compounds has remained an important issue in synthetic chemistry because these substances play important roles in coordination chemistry, biochemistry, and pharmaceutical and materials science.³ Many biologically active natural products are heterocycles that are used in the agrochemical, pharmaceutical and flavorant industries.^{4,5} For example, 1,2-benzisoxazole derivatives have been widely utilized as anti-cancer, anti-microbial and anti-thrombotic agents.⁶ As a result, the observation that 1,2-benzisoxazoline-3-ones can be prepared from hydroxamic acids, using either thionyl chloride at low temperature, or 1,1'-carbonyldiimidazole (CDI) at high temperature, is relevant.⁷ The design and discovery of new methods to synthesize heterocyclic compounds, like 1,2-benzisoxazoline-3-ones, remain important.

Reactive oxygen species (ROS) are involved in a number of biological functions.⁸ Among ROS, the hypochlorite ion (ClO⁻) has gained importance because of the biological roles it plays.⁹ The hypochlorite ion is produced in living systems mainly by

myeloperoxidase (MPO) catalyzed peroxidation of chloride ions.¹⁰ However, because of its high and nonselective reactivity, abnormal production of hypochlorite can lead to various diseases including cardiovascular ailments,¹¹ arthritis,¹² neuronal degeneration¹³ and cancer.¹⁴ As a consequence, methods for sensitive and selective detection of the hypochlorite ion are highly significant. To date, a variety of fluorescent probes for ClO⁻ sensing, which are designed based on HOCl induced oxidation reactions, have been described.¹⁵ However, fluorescent probes that display a fluorescence enhancement (turn-on) response and that function in 100% aqueous media are scarce. In addition, tedious synthetic procedures for these probes can have serious disadvantages. In the current study, we report a simple, rapid and economical method to sense ClO⁻.

The method uncovered to prepare 1,2-benzisoxazoline-3-one (**2**), shown in Scheme 1, is both simple and efficient. Specifically, treatment of a methanol solution of salicylhydroxamic acid (**1**) at ambient temperature with excess sodium hypochlorite leads to rapid (30 s) formation of **2**. This heterocycle is isolated in 82% yield following implementation of a simple extraction procedure. Compound **2** was confirmed by ¹H-NMR, ¹³C-NMR and ESI-MS.

Because 1,2-benzisoxazoline-3-one is fluorescent and the oxidation reaction producing it is both rapid and efficient, we envisaged that the process would serve as the basis for a new method for HOCl detection and imaging the physiological generation of this ROS. In Fig. 1 emission spectra are displayed which show the fluorescence changes that occur when ClO⁻ (1.0 equiv.) and other ROS and RNS (10 equiv., H₂O₂, ¹O₂, •O²⁻, ROO•, NO• and •OH) are added to aqueous HEPES (10 mM, pH 7.4) solutions of **1**. The spectra demonstrate that only ClO⁻



Scheme 1 Synthesis of 1,2-benzisoxazoline-3-one (**2**) from salicylhydroxamic acid (**1**) using hypochlorite.

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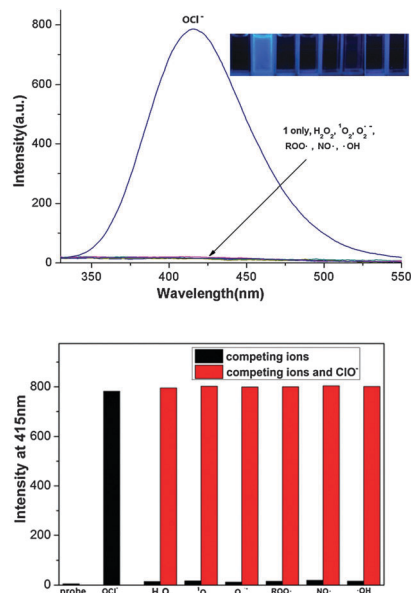


Fig. 1 Top: Fluorescence spectra of mixtures of **1** (10 μM) and other ROS and RNS (100 μM) including ClO^- , H_2O_2 , $^1\text{O}_2$, $\text{O}_2^{\bullet-}$, ROO^\bullet , NO^\bullet , $^\bullet\text{OH}$ in HEPES (10 mM, pH 7.4) ($\lambda_{\text{ex}} = 300$ nm). In the inset are photographic images of mixtures of **1** containing ClO^- (blue) and other anions (colorless) under illumination with a 365 nm UV lamp. Bottom: Fluorescence intensities (415 nm) of the probe (2.5 μM) upon the addition of 2.5 μM ClO^- in the presence of 25 μM of the other analytes in HEPES (10 mM, pH 7.4) solutions. Black bar: probe + various analytes. Red bar: probe + various analytes + ClO^- ($\lambda_{\text{ex}} = 300$ nm, $\lambda_{\text{em}} = 415$ nm).

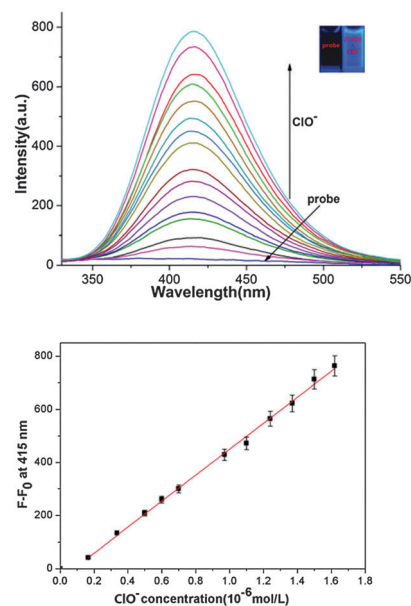


Fig. 2 Top: Fluorescence spectra of **1** (2.5 μM) in the presence of various concentrations of ClO^- (0–1.8 μM) in HEPES (10 mM, pH 7.4) ($\lambda_{\text{ex}} = 300$ nm). Bottom: Plot of intensities of fluorescence at 415 nm as a function of the concentrations of ClO^- .

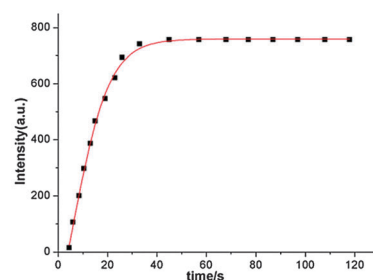


Fig. 3 Reaction time profile of probe **1** and ClO^- .

causes production of **2** in association with a large increase in the intensity of the fluorescence band with a maximum at 415 nm (300 nm excitation). UV-Vis spectroscopic monitoring of the reaction between **1** and ClO^- shows that an increase in absorption occurs at 265 nm as the reaction proceeds (Fig. S3, ESI[†]) and fluorescence titration data show that a ClO^- concentration dependent fluorescence enhancement occurs at 415 nm (Fig. 2a).

Analysis of the fluorescence titration profile of the reaction of **1** (2.5 μM) with ClO^- demonstrates that the limit of detection of the system for ClO^- is 49 nM (Fig. 2b). As shown in Fig. S5 (ESI[†]), probe **1** shows fluorescence enhancement at pH in the range of 6–12 upon the addition of HOCl.

Time-dependent changes in the fluorescence spectra, brought about by the addition of 10 equiv. of ClO^- to a solution of **1**, were monitored. The results (Fig. 3) show that reaction to form **2** is completed within 30 s, indicating that the probe reacts rapidly with this ROS under the conditions employed. This fast response suggests that the process can be possibly employed directly for the quantitative detection of ClO^- .

The ability of **1** to be used to visualize ClO^- within living cells was also evaluated. For this purpose, laser confocal fluorescence imaging was carried using a Leica TCS SP5 laser scanning microscope, with an optical window in the blue channel (400–450 nm) as a signal output. As the images given in Fig. 4a show, HepG2 cells incubated with the 2.5 μM probe for 30 min at 37 $^\circ\text{C}$ do not fluoresce when selectively excited at

300 nm. In contrast, HepG2 cells do display blue fluorescence after they are incubated with 2 μM of **1** for 30 min at 37 $^\circ\text{C}$ and then treated with 5 μM ClO^- (Fig. 4b). The results of these

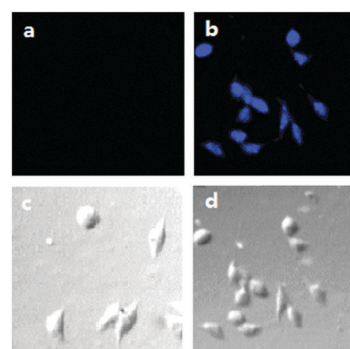


Fig. 4 Confocal fluorescence microscope (a) and bright field (c) images of HepG2 cells in the absence of **1**. Confocal fluorescence microscope (b) and bright field (d) images of HepG2 cells following incubation with 2.5 μM **1** for 10 min followed by addition of 5 μM ClO^- .



Table 1 Determination of HOCl concentration in a disinfectant

| Solution added ^a (μL) | Relative fluorescence ^b (ΔF) | [OCl ⁻] ^c (mmol L ⁻¹) |
|----------------------------------|---|--|
| 1 7 | 142 | 106 ± 3 |
| 2 12 | 301 | 114 ± 2 |
| 3 29 | 640 | 95 ± 6 |

^a Diluted 1000 fold. ^b 2.5 μM probe in 10 mM HEPES, pH 7.4. ^c Hypochlorite concentration of sodium hypochlorite disinfectant ($n = 5$).

whole cell experiments demonstrate that the new probe is cell permeable and that it can be used to visualize ClO⁻ within living cells.

To investigate the applicability of the new probe for monitoring ClO⁻ in real samples, the levels of this ROS in a common disinfectant were examined (Table 1). Aliquots of a commercial bleach solution, without pretreatment, were added to a solution of **1** in buffer. By comparing the observed fluorescence intensities to those of a standard curve (Fig. S4, ESI[†]), the concentration of ClO⁻ in the bleach sample was determined to be 105 ± 3 mM (Table S1, ESI[†]).

In summary, in this study we uncovered a simple, rapid and economical method for synthesis of the interesting heterocycle, 1,2-benzisoxazoline-3-one, using ClO⁻ promoted oxidation of salicylhydroxamic acid at room temperature for 30 s. Compared with traditional protocols used to prepare this substance,⁷ the new method is more environmentally benign. In addition, we demonstrated that this process can be used as a fluorescent probe for ClO⁻. The application of this method to the synthesis of other heterocyclic compounds and to the development of new fluorescent probes with longer wavelength emission characteristics is under current investigation.

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