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Imaging of hypochlorous acid in mitochondria using an asymmetric near-infrared fluorescent probe with large Stokes shift†

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Small-molecule near-infrared (NIR) imaging facilitates deep tissue penetration, low autofluorescence, non-invasive visualization, and a relatively simple operation. As such it has emerged as a popular technique for tracking biological species and events. However, the small Stokes shift of most NIR dyes often results in a low signal-to-noise ratio and self-quenching due to crosstalk between the excitation and emission spectra. With this research, we developed a NIR-based fluorescent probe **WD-HOCl** for hypochlorous acid (HOCl) detection using the NIR dye **TJ730** as the fluorophore, which exhibits a large Stokes shift of 156 nm, with no crosstalk between the excitation and emission spectra. It contains acyl hydrazide as the responsive group and a pyridinium cation as the mitochondria-targeting group. The fluorescence intensity of **WD-HOCl** was enhanced by 30.1-fold after reacting with HOCl. Imaging studies performed using BV-2 cells indicated that **WD-HOCl** could be used for endogenous HOCl detection and imaging in living cells exposed to glucose and oxygen deprivation/reperfusion. Finally, we demonstrated that inhibiting the expression of NOX2 reduced the HOCl levels and the severity of oxidative stress during stroke in a mouse model.

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Introduction

Near-infrared small-molecule fluorescent probes have been extensively developed due to their enhanced properties including deep tissue penetration, low autofluorescence, non-invasive visualization, and relatively simple operation,^{1–3} and as such have emerged as a popular technique for the real-time sensing and tracking of biological species and events.^{4,5} However, to improve their internal conversion efficiency, the HOMO–LUMO band gap of NIR dyes must be reduced. This task is difficult and often decreases the Stokes shift.⁶ Efforts to extend the Stokes shift include nitrogen insertion *via* cyanine or pyronine groups, twisted intramolecular charge transfer, and solvent cage relaxation. Unfortunately, none of these techniques can eliminate crosstalk between the excitation and

emission spectra.^{7–9} Inducing a pseudo-Stokes shift based on energy transfer is complicated, is limited by the energy transfer efficiency, and dual emission peaks in the NIR region are difficult to achieve.^{10,11} These issues result in many disadvantages such as a low signal-to-noise ratio and self-quenching, making it difficult to increase the sensitivity and accuracy of imaging techniques.^{12–14} In contrast, the group of Yuan¹⁵ introduced asymmetric electronic structures to fluorescein, squaraine, cyanine, rhodamine, and oxazine dyes to improve their photophysical properties. Which was achieved by enhancing the internal conversion, reducing emission energy, and increasing the Stokes shift of fluorophores using vibronic contributions to the lowest unoccupied molecular orbital (LUMO). Inspired by this principle, we designed and synthesized a NIR dye containing a strong electron-donating heteroatom which increased the asymmetry of the electronic structure through vibronic contributions. Therefore, the NIR dye, **TJ730**, was used to prepare a novel NIR probe which eliminated crosstalk between excitation and emission spectra and improved the sensitivity and accuracy of imaging analysis.

Specifically, a novel heteroatom-substituted rhodamine-based NIR fluorescent probe for HOCl, named **WD-HOCl**, was developed. The probe consists of a fluorophore **TJ730** decorated with acyl hydrazide as HOCl recognition group.¹⁶ **TJ730** exhibited a large Stokes shift (156 nm in acetonitrile) and achieved almost complete separation of its excitation and emission peaks. The fluorescence intensity of the NIR fluorescent probe

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Next, we evaluated the chemical stability of **TJ730** to ensure its suitability for the high-fidelity imaging of hypochlorous acid (HOCl). Several previous studies have demonstrated that S, as the most important component of mercaptan derivatives, can maintain the redox balance of cells by reducing oxidizing species.¹⁸ The methyl thioether group has frequently been used as a modulator of the redox cycle between HOCl and GSH.¹⁹ Therefore, we suspected that **TJ730** could be easily destroyed by the endogenous ROS produced, leading to poor reliability and stability. As illustrated in Fig. S10A and B,† we evaluated the stability of **TJ730** against various oxidants and nucleophiles including H₂O₂, O₂^{•-}, H₂S, HSO₃⁻, NO, ONOO⁻, HOCl, and •OH. The results indicated that **TJ730** had good stability and is suitable for HOCl detection. We hypothesize that the extended π -conjugated system decreased the activity of S and increased the chemical stability of the dye.

Design and synthesis of WD-HOCl

Fluorometric detection of HOCl using WD-HOCl

The photophysical properties of **TJ730** in 10 mM PBS and organic solvents with different polarities were investigated to optimize the detection system. Fig. S10C and D[†] shows the absorption and fluorescence spectra of **TJ730**, and the corresponding photophysical data are presented in Table S1.[†] **TJ730** exhibits an absorption band between 606 nm and 615 nm, which was attributed to the π - π^* transition of the ring-opened rhodamine conjugate. Upon excitation at 606 nm, **TJ730** presented a characteristic NIR fluorescence emission $\lambda_{\text{em}} \geq 745$ nm. The absorption and fluorescence spectra revealed a weak bathochromic shift upon increasing solvent polarity due to the high polarity of the molecule resulting from its D- π -A structure. The results also illustrated that **TJ730** exhibited the best photophysical properties in acetonitrile, with an absorption maximum at 606 nm ($\epsilon = 0.35 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$) and an emission maximum at 762 nm ($\Phi = 4.7\%$). The Stokes shift

Dye	λ_{abs}^a (nm)	λ_{em}^b (nm)	ϵ^c (M ⁻¹ cm ⁻¹)	Φ^d	Stokes shift (nm)	$\epsilon \times \Phi^e$ (M ⁻¹ cm ⁻¹)
RhB	553	567	109 100	0.7	14	76 370
DQF-RB	597	682	127 000	0.358	85	45 466
TJ730	606	762	628 000	0.047	156	29 516

^a Maximum absorption wavelength. ^b Maximum fluorescence emission wavelength. ^c Molar absorptivity. ^d Quantum yield. ^e Brightness.



Fig. 1 (A) Chemical structures and normalized absorption (black line)/emission (red line) spectra of RhB (B), DQF-RB (C), and TJ730 (D) in acetonitrile. (E) Density functional theory-optimized molecular orbital plots (lowest occupied molecular orbital and highest occupied molecular orbital) of RhB, DQF-RB, and TJ730. (F) Calculated absorption and emission wavelengths and oscillator strengths of the dyes (PCM solvation model with acetonitrile as the solvent).

between the excitation and emission peaks exceeded 156 nm. We then decorated **TJ730** with acyl hydrazide (which can be oxidized and hydrolyzed specifically by HOCl) and a pyridinium cation (which can anchor in the cell mitochondria due to the positive charge). Therefore, we obtained a novel heteroatom-substituted rhodamine-based NIR fluorescent probe for HOCl which we called **WD-HOCl**. **TJ730** and **WD-HOCl** were readily synthesized, and their synthetic routes and characterization details are given in Scheme S1 and Fig. S1–S9.†

The proposed recognition and imaging mechanism of HOCl in mitochondria is given in Fig. 2A. **WD-HOCl** was sufficiently water-soluble for solution-phase detection and cell staining (Fig. S11†). The absorption and fluorescence spectra of **WD-HOCl** with and without HOCl in 10 mM PBS containing 33% acetonitrile were recorded. **WD-HOCl** displayed a negligible absorption peak at 596 nm owing to the closed-ring form of the probe. However, upon adding HOCl (2 eq.), a new absorption maximum appeared at 606 nm ($\epsilon = 1387 \text{ M}^{-1} \text{ cm}^{-1}$) (Fig. S12A†), indicating the ring-opening of **TJ730**. **WD-HOCl** is

non-fluorescent however a strong emission peak at approximately 762 nm appeared immediately upon the addition of HOCl (2 eq.) (Fig. S12B†), indicating a large Stokes shift of about 156 nm. **WD-HOCl** was then used to quantify HOCl concentrations *in vitro*. Upon the addition of 2 eq. HOCl, a 30.1-fold enhancement of the fluorescence intensity was observed (Fig. 2B and C). Furthermore, a linear correlation between the fluorescence intensity of **WD-HOCl** at 762 nm and HOCl concentrations was observed (Fig. 2D). The detection limit of **WD-HOCl** for HOCl was calculated to be 1.3 nM, demonstrating its high sensitivity for the determination of HOCl. As presented in Fig. 2E, the reaction of **WD-HOCl** with HOCl (3, 10, and 16 μM) was nearly complete at 100 s, confirming that **WD-HOCl** exhibited a rapid response to HOCl. The mechanism for the interaction of **WD-HOCl** with HOCl was confirmed using high-resolution mass spectrometry (HRMS). The peak at $m/z = 521.1863$ was ascribed to the characteristic peak of the reaction product **TJ730**, and the peak at $m/z = 654.2538$ (representing the reactant) disappeared (Fig. S13†).

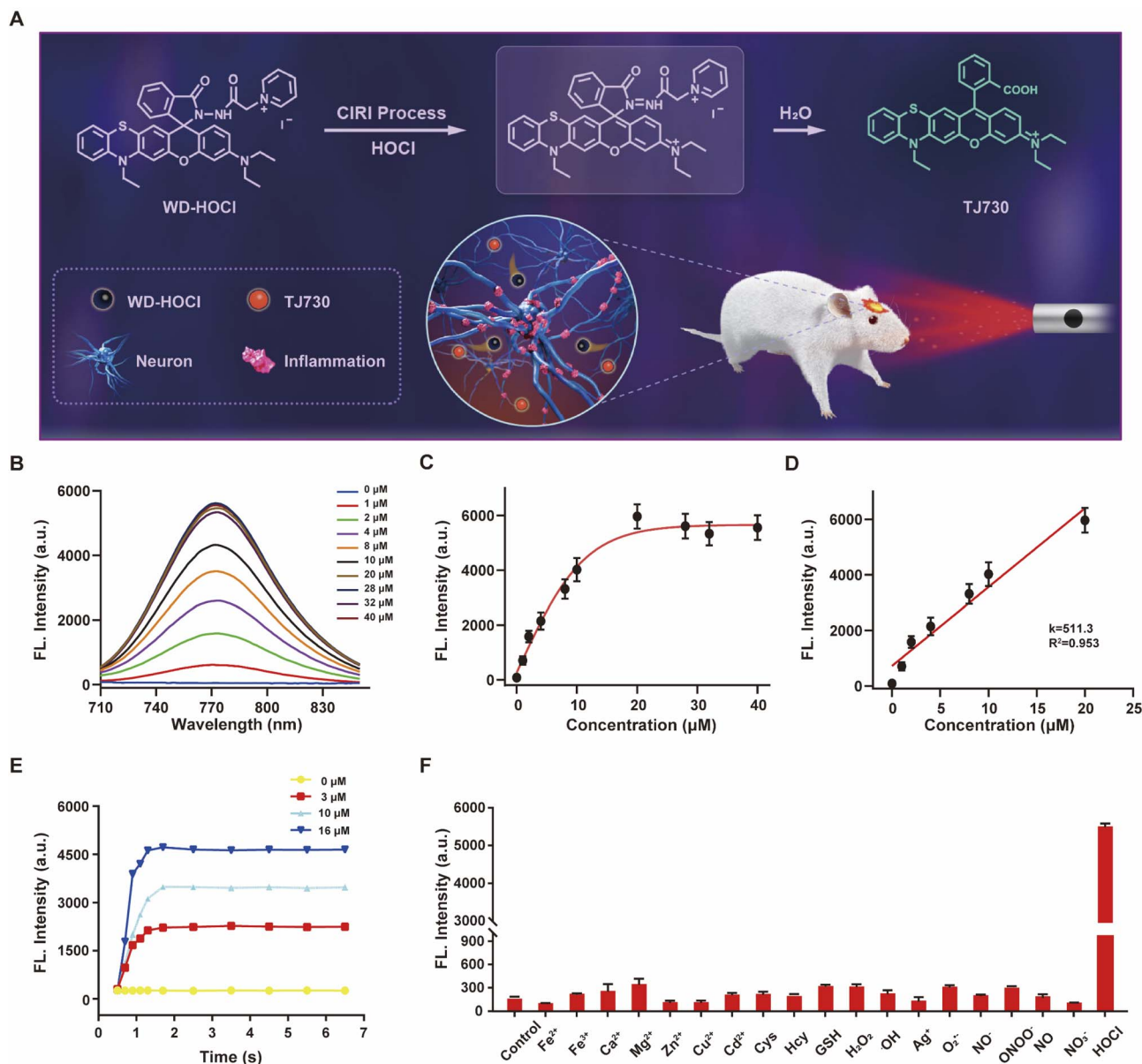


Fig. 2 (A) Recognition mechanism of **WD-HOCl**. (B) Fluorescence spectra for **WD-HOCl** upon exposure to different amounts of **HOCl** (0–40 μM). (C) Fluorescence intensity of **WD-HOCl** at 762 nm as a function of the **HOCl** concentration. (D) Linear fluorescence response of **WD-HOCl** toward **HOCl** (0–20 μM). (E) Time-dependent reaction profiles of 10 μM **WD-HOCl** and (0, 3, 10, and 16 μM) **HOCl**. (F) Fluorescence responses of **WD-HOCl** to a blank control (a), 20 μM metal ions ((b–i) Fe^{2+} , Fe^{3+} , Ca^{2+} , Mg^{2+} , Zn^{2+} , Ag^{+} , Cu^{2+} , and Cd^{2+} , respectively), 1.0 mM biomolecules ((j–l) Cys, Hcy, and GSH, respectively), 100 μM ROS ((m–o), H_2O_2 , $\cdot\text{OH}$, and $\text{O}_2^{\cdot-}$, respectively), 100 μM reactive nitrogen species (RNS, (p–s) NO_2^- , ONOO^- , NO , and NO_3^- , respectively), and 20 μM **HOCl**.

Selectivity is a critical factor for improving the performance of fluorescent probes for bioimaging applications. Thus, we investigated the fluorescence responses of **WD-HOCl** towards **HOCl** and 20 μM of potential interfering metal ions (Fe^{2+} , Fe^{3+} , Ca^{2+} , Mg^{2+} , Zn^{2+} , Ag^{+} , Cu^{2+} , and Cd^{2+}), 1.0 mM biomolecules (Cys, Hcy, and GSH), 100 μM ROS (H_2O_2 , $\cdot\text{OH}$, and $\text{O}_2^{\cdot-}$, respectively), 100 μM reactive nitrogen species (RNS, NO_2^- , ONOO^- , NO , and NO_3^-). As illustrated in Fig. 2F, only negligible enhancements in the fluorescence intensity was caused by non-target species. However, the addition of **HOCl** remarkably enhanced the fluorescence, indicating that **WD-HOCl** was selective towards **HOCl** over other interferents. In addition, the

fluorescence response of **WD-HOCl** towards **HOCl** was evaluated at different pH values (Fig. S14[†]). In the absence of **HOCl**, **WD-HOCl** exhibited no obvious fluorescence intensity changes over the pH range of 4.2–8.0, however, the fluorescence intensity fluctuates slightly after reaction with **HOCl**.

Bioimaging applications of **WD-HOCl** in living cells

Potential applications of **WD-HOCl** for monitoring fluctuations in **HOCl** levels in living cells were then evaluated. Using a 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay, the cytotoxicity of **WD-HOCl** towards BV-2 cells was



examined after being incubated with different concentrations (0, 2, 5, 10, and 15 μM) of **WD-HOCl** for 24 h. The resultant cell viabilities all exceeded 90%, confirming that **WD-HOCl** exhibited negligible cytotoxicity, even at a concentration of 10 μM (Fig. S15[†]). Furthermore, the photostability of **WD-HOCl** during the imaging of BV-2 cells was also investigated. The results indicate that the fluorescence intensity of **WD-HOCl** remained nearly constant for 20 min, confirming its photostability (Fig. S16[†]).

In order to more accurately reflect the fluctuation of HOCl in pathological process, it is necessary for the probe to anchor at the cell mitochondria. As an organelle with a double-membrane structure, the negative charge of the inner mitochondrial membrane enables positively charged molecules to be specifically enriched and anchored to the mitochondria. Since the

pyridinium cation group of **WD-HOCl** has a positive charge, it should exhibit mitochondrial localization ability. To this end, **WD-HOCl** was mixed with a commercial mitochondrial localization dye, 100 nM Mito-Tracker Green, a commercial lysosome localization dye, 100 nM Lyso-Tracker Green, and a commercial endoplasmic reticulum localization dye, 100 nM ER-Tracker Green. These were co-incubated with BV-2 cells for 30 min and then imaged using a confocal microscope. The results in Fig. S17[†] indicate that **WD-HOCl** was mainly distributed in the mitochondria (the Pearson colocalization coefficient reached 0.94). The Pearson colocalization coefficients in the lysosome and endoplasmic reticulum were 0.31 and 0.33, respectively, indicating that **WD-HOCl** has a good mitochondrial localization ability, providing a good basis for studying ROS produced during respiration. To explore the biological applications of

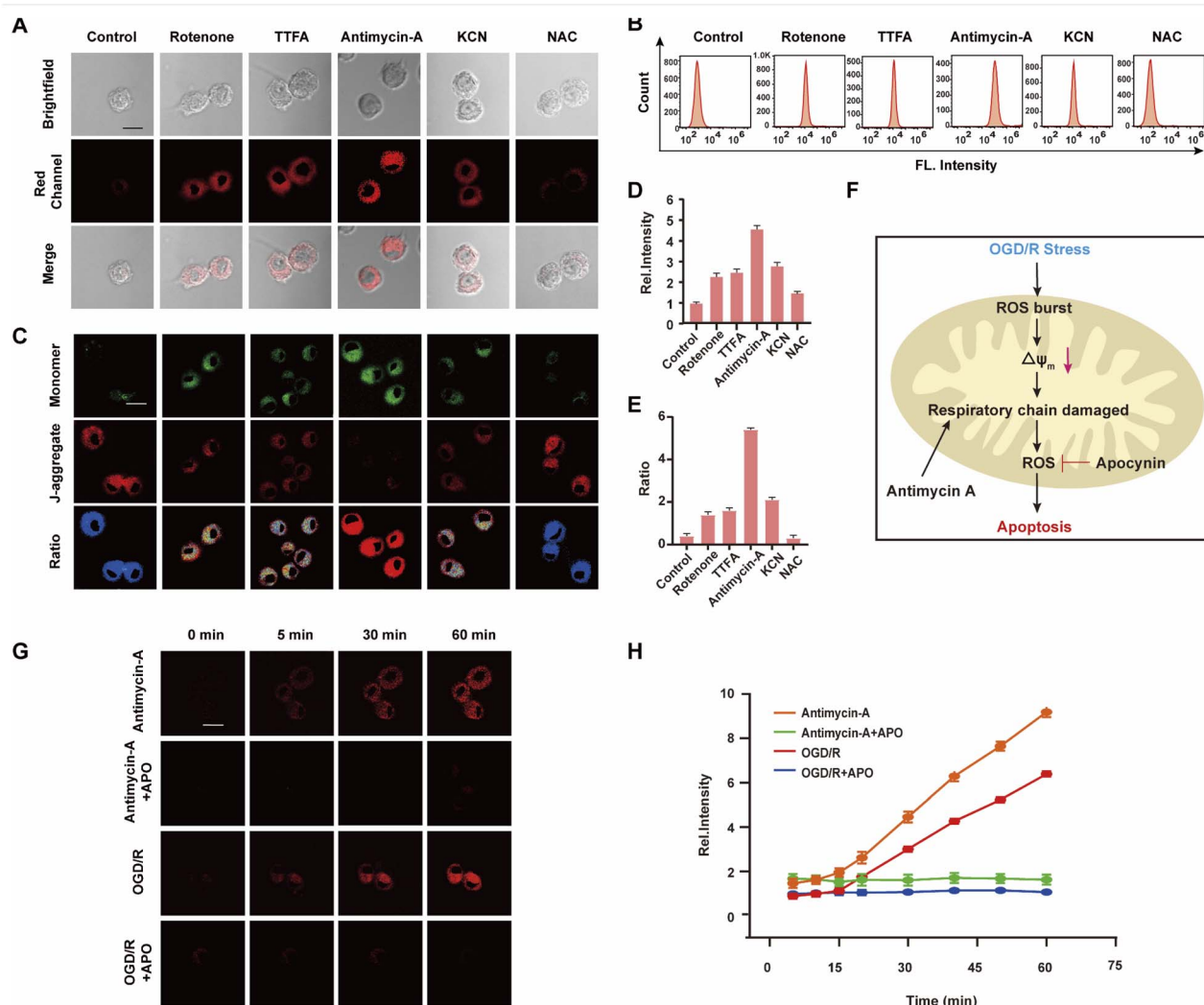


Fig. 3 (A) Images of **WD-HOCl** (10 μM) loaded BV-2 cells with the mitochondrial respiratory inhibitor including control, 5 μM of rotenone, 5 μM of TTFA, 5 μM of antimycin A and 500 μM of malonic for 30 min. (B) Flow cytometry analysis of (A). (C) Images of mitochondrial membrane potential analyzed by JC-1. (D) Relative fluorescence intensities of BV-2 cells as used in (A). (E) Ratio intensities of BV-2 cells as used in (C). (F) Scheme of OGD/R-induced signal transduction pathways. Imaging (G) and time course of fluorescence ratio changes (H) of BV2 cells. Cells incubated with 10 μM **WD-HOCl** with the first line: antimycin A (5 μM); the second line: antimycin A + APO (10 mM); the third line: OGD/R and the fourth line: OGD/R + APO. For the fluorescent images, the experiment was repeated using three cultures; similar results were obtained each time. Scale bar = 20 μm . Emissions were collected at the red channel (680–738 nm) with 606 nm excitation. The error bars mean SD ($n = 33$ cells).

WD-HOCl, exogenous HOCl in BV-2 cells was determined. BV-2 cells were incubated with **WD-HOCl** (5 μM) for 30 min, washed with PBS, treated with different concentrations of NaOCl (0, 5, 10, and 20 μM) for another 30 min, and immediately imaged using a confocal fluorescence microscope (Fig. S18†). The fluorescence intensity in the red channel (I_{red}) increased in a concentration-dependent manner, indicating that **WD-HOCl** was membrane-permeable and could be used to detect exogenous HOCl in living cells.

To detect endogenous HOCl using **WD-HOCl**, BV-2 cells were preconditioned with the ROS stimulant phorbol 12-myristate-13-acetate (PMA, 2 $\mu\text{g mL}^{-1}$) for 1 h and then treated with **WD-HOCl** for another 30 min. As illustrated in Fig. S19,† I_{red} of the PMA-stimulated cells was substantially higher than that of cells treated with only the probe. Furthermore, I_{red} for the cells pretreated with HOCl scavengers including lipoic acid (5 $\mu\text{g mL}^{-1}$), *N*-acetyl-L-cysteine (NAC, 2 $\mu\text{g mL}^{-1}$) and L-methionine (1 $\mu\text{g mL}^{-1}$) exhibit weak fluorescence. The experimental results confirm that **WD-HOCl** can be used to monitor endogenous HOCl in living cells.

The mitochondrial respiratory chain is the chief source of HOCl generation.²⁰ Complex I (NADH:ubiquinone oxidoreductase), complex II (succinate dehydrogenase), complex III (ubiquinol-cytochrome c reductase) and complex IV (cytochrome c oxidase) are all redox centers for HOCl generation. Mitochondrial respiratory inhibitors, such as rotenone (a complex I inhibitor), thenoyltrifluoroacetone (TTFA, a complex II inhibitor), antimycin A (a complex III inhibitor) and KCN (a complex IV inhibitor), were employed to block the respiratory chain selectively and induce mitochondrial HOCl overproduction. Remarkable I_{red} enhancement were observed for the four inhibitors *versus* that in the control group (Fig. 3A and D). In addition, the cells were incubated for 30 min with NAC to eliminate HOCl. The weak fluorescence in Fig. 3A and D illustrates a very low endogenous HOCl concentration in BV-2 cells. Flow cytometry analysis also indicated fluorescence intensity variations. Compared with the control group, significant fluorescence enhancement was observed with incubation of BV-2 cells with the four mitochondrial respiratory inhibitors, and reduced by NAC, which confirms that these four respiratory inhibitors of the respiratory chain result in outbursts of HOCl (Fig. 3B). Then, the mitochondrial membrane potential ($\Delta\psi_{\text{m}}$) as an indicator of mitochondrial function, was used to evaluate the origin of HOCl. In the subsequent research, we measured changes in $\Delta\psi_{\text{m}}$ using the red/green fluorescence ratio (JC-1 assay, Fig. 3C and E). The same trend as Fig. 3A was observed, indicating that the destruction of the cell respiratory chain may damage the cell mitochondria, resulting in the formation of HOCl. In addition, confocal imaging indicated that maximal HOCl production was induced by antimycin A (Fig. 3A and D), which prompted us to determine the dynamic real-time monitoring ability of antimycin A (5 μM) to produce HOCl. As shown in Fig. 3G, H and S20,† the fluorescence signals of both groups appeared in the first 5 min and increased gradually for the subsequent 10 min (5–15 min). This phenomenon indicates that the probe fully entered the cell after 5 min and that HOCl production began after 5 min. In addition, compared with the

antimycin A + APO (apocynin, 10 mM, NADPH oxidase 2 inhibitor²⁰) group, a significant increase in I_{red} was observed for antimycin A, indicating that the fluorescence signal changes during antimycin A-induced HOCl production were mainly attributed to changes in oxidative stress.

Strokes are usually caused by the natural reperfusion or iatrogenic reperfusion of ischemic brain tissue, which further damages the structure and functional metabolism of the nervous system, *i.e.*, cerebral ischemia-reperfusion injury (CIRI), which results from prolonged ischemic insult followed by the restoration of blood perfusion, which can induce tissue damage and initiate a cascade of deleterious cellular responses, leading to cell death and ultimately organ failure. Under hypoxia, cells consume several intracellular reducing substances, especially reductases, to maintain normal metabolic activity. During reperfusion, the ROS byproducts produced by mitochondria when the cells return to normal respiration cannot be cleared sufficiently quickly, resulting in oxidative stress damage to cells and tissues).²¹ Based on the above experimental results, we speculate that HOCl can be used to characterize CIRI, however this connection remains largely unexplored because of the strong oxidizing properties, short lifetime, and low concentration of HOCl during this process.²² To confirm this assumption, **WD-HOCl** was used to monitor HOCl changes with real-time tracking (0–60 min) during oxygen-glucose deprivation/reperfusion (OGD/R) to simulate the occurrence of CIRI in cells. Briefly, cells were kept in sugar-free DMEM and a three-gas incubator for 12 h without oxygen, and then the normal state of cultured cells was resumed. As presented in Fig. 3G, H and S20,† the relative fluorescence intensity increased dramatically upon increasing the OGD/R time, indicating that the HOCl content gradually increased upon prolonging the OGD/R. To further confirm the source of HOCl during the OGD/R process, we used APO, which can reduce ROS generation, in an attempt to ameliorate cell death and cerebral injuries induced by CIRI. Cells were incubated with APO (10 mM) and subjected to OGD/R over different time periods. Compared with the OGD/R group, the fluorescence signals were reduced after the cells were co-incubated with APO. This suggests that APO can reduce ROS production during CIRI and therefore inhibit apoptosis caused by oxidative stress (Fig. 3F).

NOX2 is a key enzyme that produces cellular reactive oxygen species (ROS) that can induce oxidative stress. To further elucidate the relationship between oxidative stress and cerebral strokes, the NOX2 gene of BV-2 cells was removed (NOX2KD, operational approach according to the reported methods²³) and cultured in a sugar-free and oxygen-free environment for 12 h (Fig. 4A). Then, the cells were incubated in normal medium for 1 h and then incubated with **WD-HOCl**. Fig. 4B and F indicated that the I_{red} value obtained from the OGD/R group was much higher than that from the control group, but the I_{red} value of the BV-2 cells changed slightly after removal of the NOX2 gene (NOX2KD). The I_{red} value of the negative control group (performed according to a previous report²³) was significantly enhanced. Flow cytometric analysis also indicated changes in the fluorescence intensity, confirming that the OGD/R process results in oxidative stress of the cells (Fig. 4C). Apoptosis in



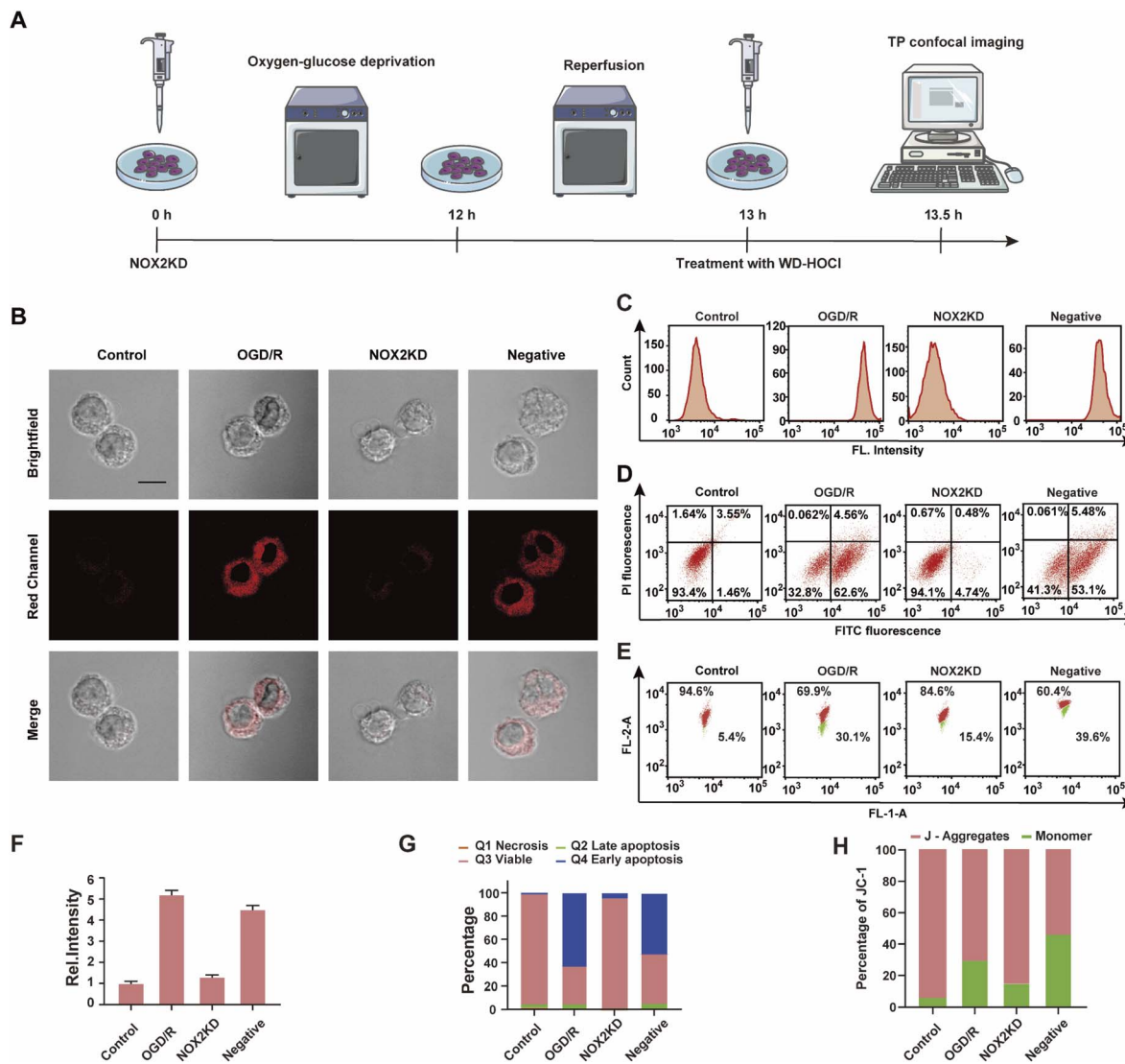


Fig. 4 (A) Schematic illustration of the OGD/R modelling and probe treatment; (B) imaging of WD-HOCl-loaded BV-2 cells. Control, untreated cells; OGD/R, OGD/R treated cells; NOX2KD, NADPH oxidase 2 gene knockout; negative, negative control of NOX2KD. $n = 48, 61, 55$ and 65 cells, respectively. Emissions were collected in the red channel (680–738 nm) with 606 nm excitation. Scale bar: 20 μ m. (C) Flow cytometry analysis of (B); (D) apoptosis assay of cells in (B); viable cells (FITC–/PI–), early apoptosis (FITC+/PI–), late apoptosis, and necrosis (FITC+/PI+); (E) mitochondrial membrane potential analyzed by JC-1; (F) histograms of the average fluorescent intensities in (B); (G) quantification data of the apoptosis assay results in (D); (H) quantification data of JC-1 assay results in (E). In (F)–(H), error bars represent SD.

these processes was assayed using an Annexin V-FITC/PI apoptosis assay kit, as shown in Fig. 4D and G. Cells in the control and NOX2KD groups maintained similar and high survival rates, but the apoptosis rates of cells in the OGD/R and negative groups were 62.6% and 53.1% respectively and the cell survival rates were only 32.8 and 41.3%. These results suggest that the oxidative stress levels in the OGD/R cells can regulate the degree of apoptosis. $\Delta\psi_m$ was also measured using the JC-1 assay, and the results indicated that the change was consistent with that of the Annexin V-FITC/PI apoptosis assay, confirming that apoptosis induced by OGD/R was caused by mitochondrial damage (Fig. 4E and H). The released ROS could induce the overexpression of the inflammatory vesicle, NLRP3 (NOD-like receptor thermal protein domain associated protein 3) and

COX-2 (cyclooxygenase 2), which results in cellular inflammation. The above process also induced significant oxidative stress. Therefore, the expression levels of the cellular oxidative stress marker (NOX2) and inflammatory factors (NLRP3 and COX-2) in the above process were evaluated. As shown in Fig. S21,† the concentration trends of the mentioned proteins in cells under different treatments were detected using western blotting, and the relationship between oxidative stress and inflammatory response during OGD/R was confirmed.

In vivo imaging of WD-HOCl during stroke

Based on the good results observed for the cell OGD/R models, we decided to use WD-HOCl to monitor HOCl fluctuations during *in vivo* stroke processes. The *in vivo* cytotoxicity of WD-

HOCl was evaluated using the hematoxylin and eosin staining (H&E) of organs after the intravenous injection of the probe. As shown in Fig. S22,† no necrosis, edema, inflammatory infiltration, or hyperplasia were observed in any of the tissues, indicating that WD-HOCl (200 μ L, 200 μ M) exhibited good biocompatibility and could be used for *in vivo* imaging. Combined with 2,3,5-triphenyltetrazolium chloride (TTC, measures tissue viability and is used to evaluate the infarct size)

staining (Fig. 5C and F), the mouse cerebral stroke model was established using the middle cerebral artery occlusion (MCAO) method. *In situ* detection results indicated that the I_{red} values 1 and 3 days after a cerebral stroke were significantly higher than the 0 day group. This indicates that the probe can monitor oxidative stress during a cerebral stroke *in vivo* (Fig. 5B and C). However, the fluorescence intensity of mice after intraperitoneal injection of APO (2.0 mg kg^{-1}) was almost unchanged,



Fig. 5 (A) Schematic illustration of the middle cerebral artery occlusion (MCAO) modeling method and probe treatment. (B) *In vivo* imaging of HOCl in the brain during middle cerebral artery occlusion (MCAO) at different times when subjected to different treatments: sham group (mice not undergoing MCAO), MCAO group (mice undergoing MCAO), vehicle group (injection of saline to mice tail veins), and APO (2.0 mg kg^{-1}) group (intraperitoneally injection with apocynin in mice). Emissions were collected at the red channel (680–738 nm) with 606 nm excitation. Scale bar = 2 cm. (C) 2,3,5-Triphenyltetrazolium chloride (TTC) staining images of infarct regions in slices of the ipsilateral hemisphere undergoing different treatments. (D) ROS revealed by dihydroethidium (DHE) staining from samples in panel (B). (E) Histogram of the average fluorescence intensities in panel (B). (F) Percentages of infarct size for the contralateral hemispheres shown in (B). (G) Rotarod test. (H) Neurological scores. (I) Quantification of ROS-positive cells in the ischemic border. Scale bar, 50 μ m. (J) Western blotting illustrating the expression of NOX-2, NLRP-3, IL-1 β , and GAPDH from the samples in (B); quantification data of the western blot results of NOX-2 (K), NLRP-3 (L), and IL-1 β (M) from the samples in (J). In (E)–(I), and (K)–(M), the error bars indicate the SD.

which again suggests that APO can inhibit brain oxidative stress and reduce the concentration of HOCl in mice. Additionally, the rotarod test was used to evaluate the effect of APO on the motor coordination of mice and results similar to the neurological scores, with both behavioral tests indicated that APO improved behavioral performance after 3 days (Fig. 5G and H). In addition, we evaluated the NOX2 expression and ROS levels using a western blot and DHE staining, respectively (Fig. 5D and I–K). The results indicated that the production of both NOX2 and ROS were significantly higher three days after a stroke. Furthermore, APO injection significantly suppressed the production of ROS while also inhibiting NOX2 expression. The western blot results indicated that the inflammatory-related proteins, NLRP3 and IL-1 β (interleukin-1 beta), exhibited similar expression levels 3 days after a stroke and were significantly inhibited by APO. These results confirm that the onset of stroke was accompanied by neuroinflammation (Fig. 5J, L and M).

Conclusions

With this research we have developed a NIR fluorescent probe **WD-HOCl** that exhibits a large Stokes shift of 156 nm, which enables almost complete separation of its excitation and emission peaks. The probe was able to target mitochondria and was suitable for bioimaging and the evaluation of HOCl levels during stroke. **WD-HOCl** rapidly reacted with HClO and could be used for the selective and sensitive detection of HOCl without interference from other ROS or RNS. **WD-HOCl** specifically accumulated in the mitochondria and highlighted changes in endogenous HOCl levels. Using cell OGD/R models, **WD-HOCl** was used to monitor HOCl levels. Based on HOCl imaging in a mouse stroke model using **WD-HOCl**, we were able to determine that a stroke causes extreme damage to brain tissue in the early stage, and APO can reduce the production of HOCl by inhibiting oxidative stress and neuroinflammation *in vivo*. In summary, **WD-HOCl** is a promising and effective tool for bioimaging and medical diagnosis.

Ethical statement

Wild-type C57BL/6J mice ($n = 300$; 25–30 g) were purchased from Hubei Experimental Animal Research Center (Hubei, China; no. 43004700018817, 43004700020932). All animal experimental protocols were approved by the Animal Experimentation Ethics Committee of South-Central University for Nationalities (No. 2020-scuec-043) and were conducted according to the Animal Care and Use Committee guidelines of South-Central University for Nationalities. Animals were housed in a room with controlled humidity ($65 \pm 5\%$) and temperature ($25 \pm 1^\circ\text{C}$), under a 12/12 hour light/dark cycle with free access to food and water for at least 1 week before the experiments.

Data availability

The datasets supporting this article have been uploaded as part of the ESI.†

Author contributions

W. Hu, T. Qiang and T. D. James conceived the project. W. Hu, F. Cheng and X. Song synthesized the dyes and conducted photophysical characterization and analyses. W. Hu, C. Li and B. Wang performed fluorescence imaging experiments. L. Ren, and M. Li performed the OGD/R and MCAO experiments. T. Qiang characterized all the compounds. All authors participated in writing the manuscript.

Conflicts of interest

TDJ acts as an academic consultant for TQ as part of a guest professorship at SUST.

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