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Unique assembly of carbonylpyridinium and chromene reveals mitochondrial thiol starvation under ferroptosis and novel ferroptosis inducer†

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To reveal the delicate function of mitochondria, spatiotemporally precise detection tools remain highly desirable. However, current probes with positively charged warheads for targeting mitochondria diffuse out of the mitochondria as the potential of the mitochondrial membrane changes, which directly influences the accuracy of the detection. Herein, we assembled carbonylpyridinium and chromene to afford the probe **CM-Mit**. Following the ultrafast response to thiol and the dissociation of carbonylpyridinium, the formation of *o*-quinone methide from **CM-Mit** was proposed to label proteins, thus avoiding diffusion out of the mitochondria. Therefore, the accurate spatiotemporal detection of thiol in mitochondria was realized. With this excellent probe, ferroptosis inducers were proved to stimulate thiol starvation in mitochondria for the first time in cancer cells. Moreover, **CM-Mit** was used to screen a compound library developed in-house and the stemona alkaloid analog **SA-11** was shown to induce ferroptosis in various cancer cell lines, including a drug-resistant one.

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Introduction

Mitochondrial thiol systems are central to protecting mitochondria against oxidative damage and are assumed to be the key means through which potential redox signals are transmitted, modulated and sensed.^{1,2} Activity-based sensing probes have been considered as desirable and indispensable tools for studying dynamic processes in the organelle.³ Thus, various mitochondrial-targeted fluorescent probes have been developed to understand the roles and functions of the analytes in mitochondria.⁴ Due to strong negative membrane potential, cations with strong lipophilicity tend to penetrate the membrane and accumulate inside the mitochondrial matrix.⁵ However, current positively charged warheads for targeting mitochondria, such as TPP (triphenylphosphine) and alkylpyridinium cations, can affect the membrane potential, triggering membrane rupture and disturbance of the microenvironment.⁶ Moreover, the probes with positive charge for targeting mitochondria can diffuse out of mitochondrial as the mitochondrial membrane potential changes, which could influence the accuracy of the

detection. Notably, Zhang and co-workers reported their elegant mitochondria-targetable probe for accurate detection of Cys without interference from other sites.⁷ However, the fluorescence was re-distributed in the cell with the extension of the incubation time following the response to the analytes. Despite the solid-state organic fluorophores (SSOFs) being successfully constructed to avoid the diffusion out of mitochondria,⁸ the aqueous dispersibility and low biocompatibility of the SSOFs are currently unsatisfactory.⁹

We recently took note of the thiol–chromene “click” nucleophilic reaction and developed a sensing probe for thiol with prominent selectivity, which enables us to monitor thiol fluctuations in various physiological and pathological processes (Fig. 1a and b)^{10–14}. Ferroptosis is a unique type of non-apoptotic-regulated cell death culminating in overwhelming lipid peroxidation downstream of metabolic dysfunctions^{15,16} and holds promise as a novel therapeutic approach.^{17,18} Recent research indicated that cysteine deprivation could induce the ferroptosis of pancreatic ductal adenocarcinoma. However, the mitochondrial membrane potential (MMP) fluctuates in ferroptosis, which influences the accuracy of current mitochondrial-targeting probes.¹⁹ Therefore, this novel strategy for accurate visualization of thiol-mediated oxidative stress in mitochondria remains highly desirable but challenging. Moreover, the developed sensing probe could be further used for rapid screening of efficient ferroptosis-inducing compounds.^{20,21} It is noteworthy that an acrylate-based fluorescent probe was developed for monitoring intracellular cysteine in ferroptosis.²² However, the relatively low reaction

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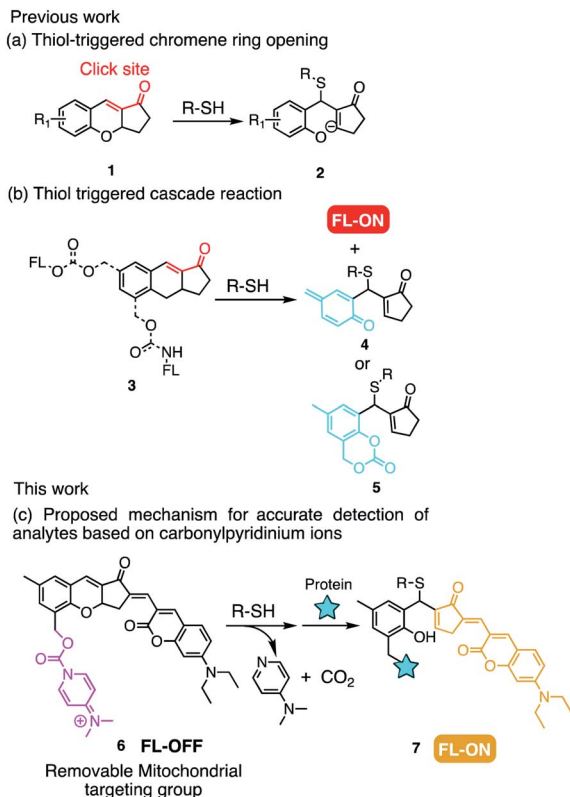


Fig. 1 Previous work: (a) thiol-mediated nucleophilic attack on unsaturated α,β -ketone and (b) self-immolation to release the fluorophore. (c) Design strategy of this work.

rate and no organelle-targeting ability limited its use for further research on the role of thiol in ferroptosis.

With these considerations in mind, we envisioned the possibility of utilizing the thiol–chromene “click” nucleophilic reaction to provide the probe with a high signal to background ratio (Fig. 1c). Besides, the formation of *o*-quinone methide after recognizing the analytes could label the proteins in the mitochondria, which may lead to mitochondrial targeting with high accuracy.²³ Although alkylpyridinium ions has extensively been employed for mitochondrial targeting,^{8,24,25} there have been no reports concerning carbonylpyridinium as a mitochondrial-targeting group in the probe. Carbonylpyridinium cation is also assumed to accelerate the recognition reaction due to its excellent leaving ability.²⁶

In this work, a novel fluorophore was constructed based on the chromene skeleton, which was integrated with the carbonylpyridinium ions to afford the ultrafast selective probe **CM-Mit** for accurate detection of thiol in mitochondria. The formation of the *o*-quinone methide enables it to label the proteins in the mitochondria, thus avoiding diffusion out of the mitochondria. With the probe **CM-Mit**, the ferroptosis inducer erastin and RSL3 were demonstrated for the first time to trigger thiol starvation in mitochondria. The screening approach was developed for discovering the potential thiol starvation modulator in mitochondria as a ferroptosis inducer. Notably, a stemona alkaloid analog (**SA-11**) with an α -methylene- γ -lactone moiety

was first proved to induce thiol starvation and ferroptosis in various cancer cells, including drug-resistant ones. Overall, this work provides a robust and reliable tool for ferroptosis research.

Results and discussion

Probe design and synthesis

In our previous study, the chromene structure (**8**) was synthesized from (2-hydroxy-5-methyl-1,3-phenylene)dimethanol in two steps.¹⁰ To realize the condensation of compounds **8** and **9**, mild conditions need to be identified. When catalytic amounts of acetic acid and piperidine were added with toluene as the solvent, the reaction was quite messy, with only a small amount of product **10** obtained (Scheme S1†). Using just piperidine as the catalyst with molecular sieves and ethanol as the solvent gave a better result. The crude product of the Knoevenagel condensation was submitted for the subsequent esterification and compound **11** was obtained with 32% yield for two steps, followed by reacting with DMAP to afford the probe **CM-Mit** with moderate yield.

Spectral properties

The capability of **CM-Mit** to detect thiols was evaluated using UV-vis absorption and fluorescence spectra (Fig. 2a and b). The selectivity of **CM-Mit** for thiols was investigated in DMSO/PBS (5 μ M, pH 7.4, v/v, 1 : 9). The probe responded to various other amino acids, such as Ala, Arg, Asn, Asp, Cys, Gln, Glu, Gly, His, Ile, Leu, Lys, Met, Phe, Pro, Ser, Thr, Tyr, Val, Cys, and Hcy, and GSH was measured. Upon the addition of these analytes, only thiols caused significant changes in fluorescence emission at 578 nm with excitation at 475 nm. However, no obvious changes

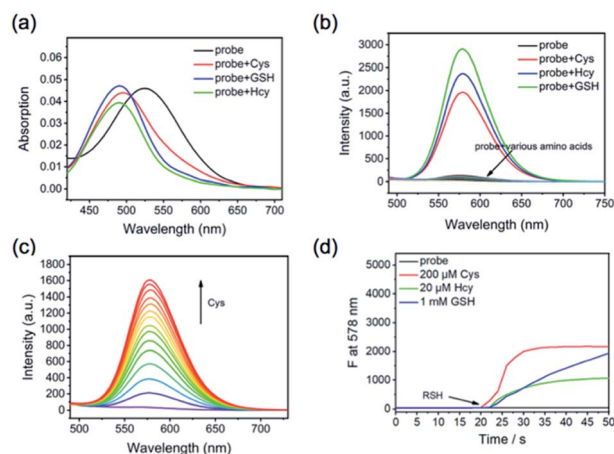


Fig. 2 (a) Absorption changes of **CM-Mit** (5 μ M, DMSO : PBS, 1 : 9, v/v) after adding 200 μ M Cys, Hcy, or GSH. (b) Fluorescence intensity changes of probe **CM-Mit** (5 μ M, DMSO : PBS, 1 : 9, v/v) with the addition of 200 μ M of various amino acid after 5 min; (λ_{ex} = 475 nm, slit: 5 nm/5 nm). (c) Fluorescence spectral changes of **CM-Mit** (5 μ M, DMSO : PBS, 1 : 9, v/v) as the concentration of Cys (0–200 μ M) increases. (d) The kinetic curve of **CM-Mit** (5 μ M) in the mixed solvent of DMSO and PBS (1 : 9, v/v) without or with 200 μ M Cys, 20 μ M Hcy, and 1 mM GSH at 578 nm (λ_{ex} = 475 nm, slit: 5 nm/5 nm).

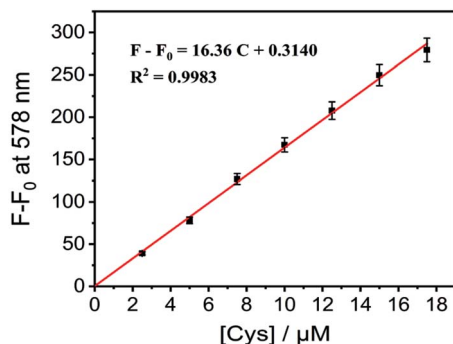


Fig. 3 The working curve of **CM-Mit** (5 μM) in the presence of different concentrations of Cys. Each point represents the average and standard deviation of three independent repeated experiments.

were observed for other amino acids (0.2 mM), anions and cations (Fig. S1†). Those results displayed that probe **CM-Mit** could detect thiols specifically. The addition of Cys at different pH values was also tested (Fig. S2†). The results indicated that probe **CM-Mit** showed almost no fluorescence emission in acidic and alkaline solutions.

Based on the above result, we selected pH 7.4 as the measurement system in our following experiments in terms of the body pH value. The fluorescence spectrum of **CM-Mit** was monitored in the presence of **CM-Mit** (5 μM , DMSO/PBS, 1 : 1, v/v) with increasing concentration of Cys (1–40 μM) (Fig. 2c). The kinetic study showed that the reaction was completed within 10 s when Cys (200 μM) was added, while the reactions for GSH (1 mM) and Hcy (20 μM) were completed in 200 and 30 s (Fig. 2d and S3†). As we expected, the introduction of the carbonylpyridinium ion significantly improved the reaction rate compared with the probe (Cys: 7 min, Hcy: 3 min, GSH: 7 min) in our previous work¹⁰ (Table S2†).

A significant increase in the fluorescence emission intensity at 575 nm was measured in DMSO/PBS (v/v, 1 : 9, pH 7.4) after the addition of Cys (0–3.6 equiv.) into the mixture. Meanwhile, a linear relation was observed between the fluorescence intensity and the concentration of Cys (Fig. 3). Furthermore, the detection limit of **CM-Mit** for Cys based on the IUPAC definition ($\text{CDL} = 3\text{Sb}/m$) is 0.49 μM , which is comparable to that for NIR-HMPC reported by our group.¹³ The above results demonstrated that **CM-Mit** is an ultrafast fluorescent probe for thiol detection with excellent selectivity and sensitivity.

Proposed mechanism

Originally, a photoinduced electron transfer (PET) mechanism was assumed to be responsible for the fluorescence quenching of **CM-Mit**, due to the existence of the carbonylpyridinium moiety. The S1 photoexcitation of **CM-Mit** mainly involves the transition from the highest occupied molecular orbital (HOMO) to the lowest unoccupied molecular orbital (LUMO). Between these two orbitals, no other molecular orbitals from a quencher are available. HOMO–1 and LUMO+1 are 0.971 eV and 0.878 eV away from the HOMO and LUMO, respectively. These energy

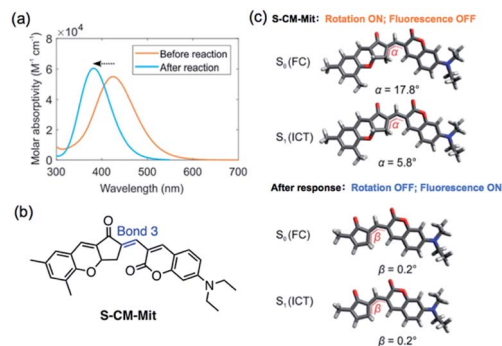


Fig. 4 (a) Calculated UV-vis absorption spectra of **CM-Mit** and the product after the reaction of **CM-Mit** and thiol. (b) The simplified structure of **CM-Mit** (**S-CM-Mit**). (c) **S-CM-Mit** and the product after the response in the ground state (the Franck–Condon state) and the ICT state. The insets show the dihedral angles (α and β) along with bond 3.

levels do not meet the requirement of PET²⁷ (Fig. S5†). Further computational analysis showed that two factors contribute to the low quantum yield of the **CM-Mit** and subsequent fluorescence turn-on after recognizing the analytes (Fig. 3). A hypsochromic shift in the UV-vis absorption spectrum occurred upon the reaction of **CM-Mit** with biological thiols and the distinct difference in the absorbance could effectively contribute to the fluorescence intensification of the product.²⁸ The second factor for modulating the fluorescence intensity is related to intramolecular rotations upon photoexcitation in probe **CM-Mit**. To reduce the computational load, we analyzed the intramolecular rotations in a simplified **CM-Mit** structure (**S-CM-Mit**) (Fig. 4b). In the most stable structure of **S-CM-Mit**, the dihedral angle along bond 3 is 17.8° in the ground state (the Franck–Condon state) but reduced to 5.8° in the ICT state in water, which could activate low-frequency vibrations and reduce the fluorescence intensity of **S-CM-Mit**, while the most stable molecular structure after **S-CM-Mit** recognizing thiol is planar in both the ground state and the ICT state (Fig. 4c).²⁹ Our viscosity-dependence study supports the intramolecular rotations of **CM-Mit** (Fig. S6†). As the volume ratio of glycerol increased, the emission intensity of **CM-Mit** was enhanced by four times.

The thiol–chromene “click” nucleophilic mechanism was proposed *via* the ¹H NMR titration experiment (Fig. S4†). The signal of the original proton (H-3) at 5.32 ppm shifted low-field to 6.85. Meanwhile, the signal of the proton (H-4) on the double bond shifted from 7.25 to 5.53 ppm. These NMR data delivered direct proof for the thiol–chromene-based ring-opening reaction and the formation of quinone methide as the intermediate.

Imaging in cells and tumors

Since probe **CM-Mit** showed an increased fluorescence emission response to thiol *in vitro* and the detection limits of probe **CM-Mit** for thiol was micromolar, bioimaging detection of thiol-containing species *in vivo* may become possible. Before



bioimaging assays, MTT assays were performed to check the cytotoxicity of **CM-Mit** against HeLa cells. As shown in Fig. S7,† the HeLa cells treated with various concentrations of **CM-Mit** (1–50 μM) for 5 and 10 h showed high cell viability, indicating that the agent had fairly low toxicity to living cells. Then, the ability of probe **CM-Mit** to detect thiol in living cells was evaluated (Fig. S8†). The results indicated that probe **CM-Mit** penetrated the cell membrane well and selectively monitored the level of thiols in the living cells. The feasibility for real-time visualization of endogenous thiols in HeLa tumor-bearing mice was also investigated. As shown in Fig. S9 and S10,† intratumor injection with **CM-Mit** resulted in a significant fluorescence signal in the tumor only 2 min after injection, which indicated that **CM-Mit** could be activated rapidly by endogenous thiol in the tumor.

Subsequently, the colocalization experiment was conducted and the results showed that probe **CM-Mit** selectively localized in mitochondria (Fig. 5) and the Pearson's coefficient is around 0.98. This result indicated that probe **CM-Mit** could selectively detect thiols in mitochondria without interference from the cytoplasmic thiols. Moreover, the fluorescence reached a high level after incubation with the probe for only 2 min (Fig. S11†). Previous research into accurate detection of thiols in mitochondria indicated that the product of the probe and the analytes may escape from mitochondria with extended incubation time.⁷ To test whether the probe could accurately monitor the thiol level in mitochondria over a longer time, the Pearson's coefficient was utilized to evaluate. In the colocalization experiment, it is interesting to find that the Pearson's coefficient is around 0.97 even with another 25 min incubation time, which was probably due to the formation of the *o*-quinone methide after response to the thiol, followed by labeling the protein in the mitochondria.²³ To examine this hypothesis, the cells were

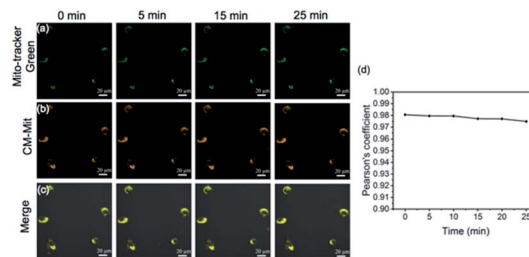


Fig. 6 Fluorescence images of **CM-Mit** responding to thiols in living HeLa cells along with reaction time by confocal fluorescence imaging. (a) Images of Mito-Tracker Green; (b) images of **CM-Mit**. (c) Merged images of (a) and (b). (d) Time-dependent Pearson's coefficient. The HeLa cells were first incubated with **CM-Mit** for 20 min, followed by incubation with Mito-Tracker Green for 20 min. Fluorescence images were captured from the green channel of 500–540 nm with excitation at 488 nm (first row) and orange channel of 563–593 nm (second row) with excitation at 475 nm.

incubated with **CM-Mit**, followed by analyzing the lysate by SDS-PAGE. The result indicated that certain proteins were indeed stained compared to the lysate with no probe (Fig. S12e†). Moreover, the fluorescence improved significantly when the cells were treated with Cys and **CM-Mit** (Fig. S12f†), which was probably due to the formation of *o*-quinone methide promoted by the thiol. Overall, the above data confirmed that probe **CM-Mit** could firmly monitor the mitochondrial thiols (Fig. 6).

CM-Mit-based fluorescence assays for discovery of potential ferroptosis inducers

We further attempted to assess the potential use of probe **CM-Mit** to visualize the thiol profile of ferroptosis. We applied two ferroptosis inducers, including erastin and RSL3, which are respectively selective inhibitors of system X_c^- and glutathione peroxidase 4 (GPX4). Cells incubated with probe **CM-Mit** displayed bright fluorescence in the orange channel, whereas pre-treatment with erastin and RSL3 significantly decreased the fluorescence intensity (Fig. 7), indicating that probe **CM-Mit** could specifically monitor mitochondrial thiols of ferroptosis and the ferroptosis inducer could trigger thiol starvation in mitochondria.

Stemona alkaloids possess diverse structures and interesting biological activities.^{30,31} In our previous work,^{32–35} we synthesized several stemona alkaloids and their derivatives. To identify stemona alkaloids that induce ferroptosis, we performed **CM-Mit**-based fluorescence assays to detect the effects of these stemona alkaloids on mitochondrial thiol levels. The live HeLa cells were pretreated with 15 μM of different compounds as well as the positive control erastin, followed by imaging and analysis of the endogenous thiol levels in mitochondria with the probe **CM-Mit**. The structure of the stemona alkaloids and their derivatives is shown in Table S1.† The results indicated that **SA-11** induced a significant decrease in the fluorescence signal, which suggested that **SA-11** significantly reduced the mitochondrial thiol level (Fig. 8). Previously, **SA-11** was proved to overcome the resistance to 5-fluorouracil in colorectal cancer cells.³⁶ However, the exact underlying mechanism is still

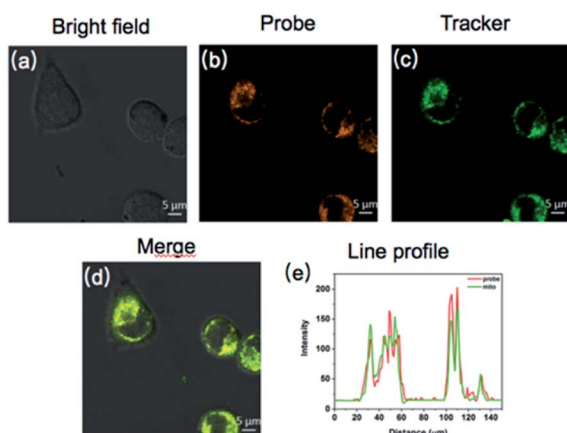


Fig. 5 Confocal fluorescence images of HeLa cells co-stained with **CM-Mit** and a tracker. (a–c) Cells pretreated with **CM-Mit** (10 μM) for 20 min and subsequently Mito-Tracker Green (1 μM) at 37 $^{\circ}\text{C}$ for 20 min. (a and d) Images from **CM-Mit**, λ_{ex} = 475 nm and λ_{em} = 563–593 nm. (b) Fluorescence images from Mito-Tracker Green, λ_{ex} = 488 nm and λ_{em} = 500–540 nm. (d) Overlay of the orange and green channels. (e) Line profile: intensity profile of the white line in image overlap. Scale bars: 5 μm .



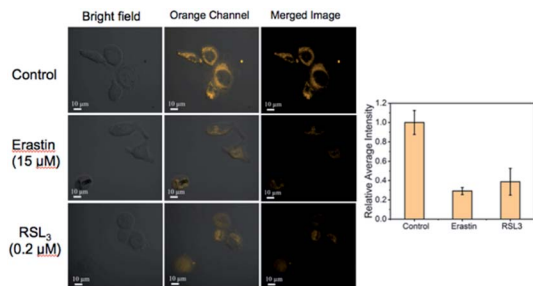


Fig. 7 Imaging of endogenous thiols in live HeLa cells treated with ferroptosis inducer. (Above) Cells were pretreated without other reagents. (Middle and below) Cells were pretreated with ferroptosis inducer, erastin (15 μ M) and RSL3 (0.2 μ M). All of the above were cultured at 37 $^{\circ}$ C for 24 h, followed by exchange into PBS (pH = 7.4) with 10 μ M CM-Mit and incubation for another 15 min at 37 $^{\circ}$ C. Data are represented as mean $\pm$ SD (n = 3).

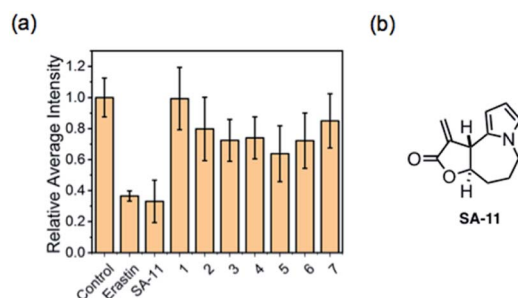


Fig. 8 Screening of stemona alkaloids and their derivatives for controlling the thiol level. (a) HeLa cells were pretreated with ferroptosis inducer erastin (5 μ M), SA-11 and its analogs (15 μ M) in a 5% carbon dioxide incubator at 37 $^{\circ}$ C for 24 h, and the cells were incubated with 10 μ M CM-Mit for 15 min. The mean intensity from Fig. S9,† where the control group is defined as 1.0. Data are represented as mean $\pm$ SD (n = 3); (b) the structure of SA-11.

unknown. Given that SA-11 triggered thiol starvation in mitochondria, we propose that SA-11 could induce ferroptosis of cancer cells to overcome drug resistance.

To examine this hypothesis, ES-2 cells were first treated with SA-11 (30 μ M) and the apoptosis inhibitor z-VAD or the necroptosis inhibitor necrostatin-1 (Nec-1) (Fig. 9a). The cytotoxicity of SA-11 was not abrogated by z-VAD and Nec-1, indicating that SA-11 had no effect on apoptosis and necroptosis. When ES-2 cells were treated with various concentrations of SA-11, the ES-2 cells were inhibited in a dose-dependent manner (Fig. 9b). In contrast, when ES-2 cells were incubated with SA-11 and the ferroptosis inhibitor ferrostatin-1 (fer-1), the cell death was significantly rescued. When ES-2 cells were treated with SA-11 combined with erastin, the cell viability was further reduced compared to the effect of erastin alone, which suggested that SA-11 and erastin could collaboratively induce cell death. Meanwhile, the lethality induced by SA-11 was also suppressed by *N*-acetyl cysteine (NAC), which also served as an antioxidant (Fig. 9c). A further experiment was conducted to monitor the effect of SA-11 in HCT-116 and the drug-resistant HCT-116p53^{-/-}

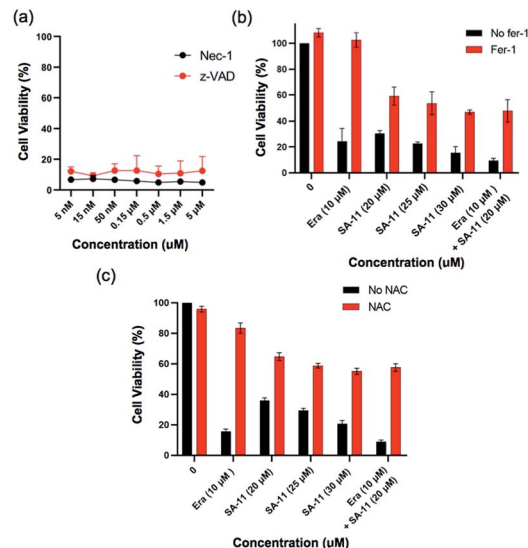


Fig. 9 SA-11 induces cancer cell death through ferroptosis. (a) Ability of apoptosis inhibitor z-VAD or necroptosis inhibitor Nec-1 to protect cells against SA-11 (30 μ M) treatment after 24 hours in ES-2 cells. For (b) and (c) effect of ferroptosis Fer-1 (2 μ M) and NAC (1 mmol) on the viability of ES-2 cells after treated with SA-11 (20 μ M, 25 μ M, 30 μ M), erastin (10 μ M) and erastin (10 μ M) combined with SA-11 (20 μ M) for 24 hours. Data are represented as mean $\pm$ SD (n = 3).

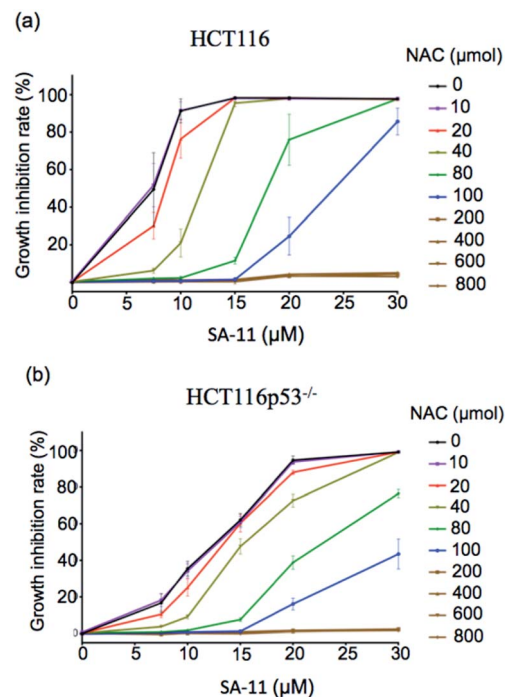
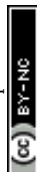


Fig. 10 Effect of NAC on viability of (a) HCT-116 and (b) HCT116p53^{-/-} after incubation with different concentration of SA-11. Data are represented as mean $\pm$ SD (n = 3).

The results indicated that both cell lines treated with different concentrations of NAC could be rescued from the death induced by SA-11 (Fig. 10). These results clearly indicate that SA-11 triggers ferroptotic cell death in cancer cells.



Conclusion

In this work, the novel probe **CM-Mit** was constructed for accurate detection of thiol in mitochondria based on the chromene skeleton and carbonylpyridinium structure. Extensive computational studies indicated that the hypsochromic shift after the response to analytes and the intramolecular rotations between the ground state and the ICT state contribute to the fluorescence turn-on. **CM-Mit** displays excellent mitochondrial-targeting ability and could accurately detect the thiols in mitochondria without interference from other sites. Moreover, the formation of the *o*-quinone methide after the response to thiols could label the proteins, which enables it to localize in mitochondria with the extension of incubation time. With the probe **CM-Mit**, the ferroptosis inducers erastin and RSL3 were demonstrated for the first time to trigger thiol starvation in mitochondria. Additionally, the screening approach was successfully developed for discovering potential thiol-level modulators as ferroptosis inducers. The stemona alkaloid derivative **SA-11** with an α -methylene- γ -lactone moiety was found to trigger thiol starvation in mitochondria and demonstrated to induce the ferroptosis of cancer cells, including drug-resistant ones. This work not only provides carbonylpyridinium as a robust mitochondrial-targeting group for mitochondrial function research but also illustrates a versatile strategy for the discovery of small molecules as ferroptosis inducers.

Author contributions

K. M. and C. Y. conceived the project. K. M., H. Y. and L. Z. synthesized the dyes and conducted photophysical characterization and analysis. H. Y. and L. Z. performed the fluorescence imaging experiments. T. R. and X. L. conducted the theoretical calculations. F. H. characterized the structures. All authors participated in writing the manuscript. We thank Xingkang Wu for the discussion about the biological assays, and Zhihong Wei for the help in computational analysis.

Conflicts of interest

There are no conflicts to declare.

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