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Rational Designed Sonocatalyst-Enhanced Supramolecular Ferroptosis Inducers for Effective Cancer Therapy

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Ferroptosis is a promising strategy against apoptosis-resistant tumors, yet traditional iron-induced approaches face safety issues and unsatisfied efficacy within complex tumor microenvironments, highlighting the need for biocompatibility and highly effective ferroptosis inducers. Herein, we rationally constructed a series of supramolecular ferroptosis inducers (**Ru1-Ru3**) with sonosensitivity and sonocatalytic properties via molecular engineering, designed for cancer treatment through near-infrared fluorescence-guided sonodynamic therapy. Among them, **Ru3** exhibited highly ultrasound-triggered ¹O₂ generation efficiency ($\Phi_A=0.89$) owing to its extended π -conjugation system and enhanced intramolecular charge transfer effect. Moreover, **Ru3** possessed catalase mimic and peroxidase mimic activities, significantly improving ROS generation and diversifying ROS species. Further studies revealed that **Ru3** localized predominantly in lysosomes, where it induced lysosomal membrane permeabilization and activated ferritinophagy under US irradiation, leading to the release of iron ions into the cytosol and triggering a Fenton reaction. Furthermore, **Ru3** catalyzed the depletion of GSH and oxidation of NADPH, disrupting redox homeostasis. These effects collectively suppressed GPX4 activity, promoted lipid LPO accumulation, and ultimately enhanced ferroptosis. In vivo experiments confirmed that US-activated **Ru3** effectively inhibited 4T1 tumor growth with favorable biosafety. This work provides a research framework for the design of the next generation ferroptosis inducers.

Introduction

Despite significant advances in cancer treatment, malignancies with high incidence and mortality rates continue to pose a major threat to human health.¹⁻³ Sonodynamic therapy (SDT) has emerged as a promising anticancer approach, leveraging sonosensitizers to generate cytotoxic singlet oxygen (¹O₂) upon ultrasound (US) irradiation, effectively eliminating cancer cells. SDT offers several advantages, including precise spatiotemporal control, non-invasive application, and deep tissue penetration (> 10 cm).⁴ The primary mechanism underlying its anticancer effects involves caspase-dependent apoptosis.⁵⁻⁶ However, the overexpression of anti-apoptotic proteins in malignant cells fosters resistance to apoptosis-inducing agents, thus

diminishing the therapeutic efficacy of SDT.⁷⁻⁹ This underscores the urgent need to explore novel, non-apoptotic mechanisms of cell death to further enhance the effectiveness of SDT.

Ferrous accumulation and lipid peroxidation (LPO)-mediated ferroptosis, a non-apoptotic cell death pathway, has garnered increasing attention.¹⁰⁻¹² However, the direct delivery of iron may induce severe side effects, such as hypersensitivity reactions in normal tissues. Therefore, employing non-iron-dependent agents to induce LPO could be safer for triggering ferroptosis in tumor cells. As the primary 'executor' in SDT, reactive oxygen species (ROS) can oxidize polyunsaturated fatty acids to lipid peroxides, ultimately leading to ferroptosis.¹³⁻¹⁷ The key factor promoting ferroptosis is the elevation of intracellular ROS levels. Recent studies have shown that organic sonosensitizers, with well-defined structures and flexible designs, can effectively generate ROS under US irradiation.¹⁸⁻²² However, most traditional sensitizers (e.g., cyanine and porphyrin) exhibit low ROS generation efficiency and are prone to ROS quenching.²³⁻²⁶ On the other hand, the hypoxic tumor microenvironment (TME) and the overexpression of reductive substances, such as nicotinamide adenine dinucleotide phosphate (NADPH) and glutathione (GSH), can directly or indirectly reduce ROS levels.²⁷ NADPH plays a key role in the recycling of GSH by donating electrons to glutathione reductase, which reduces oxidized glutathione (GSSG) back to GSH. GSH plays a critical role in supporting the antioxidant function of glutathione peroxidase 4 (GPX4), which converts harmful lipid peroxides into benign lipid alcohols, thereby protecting cells.

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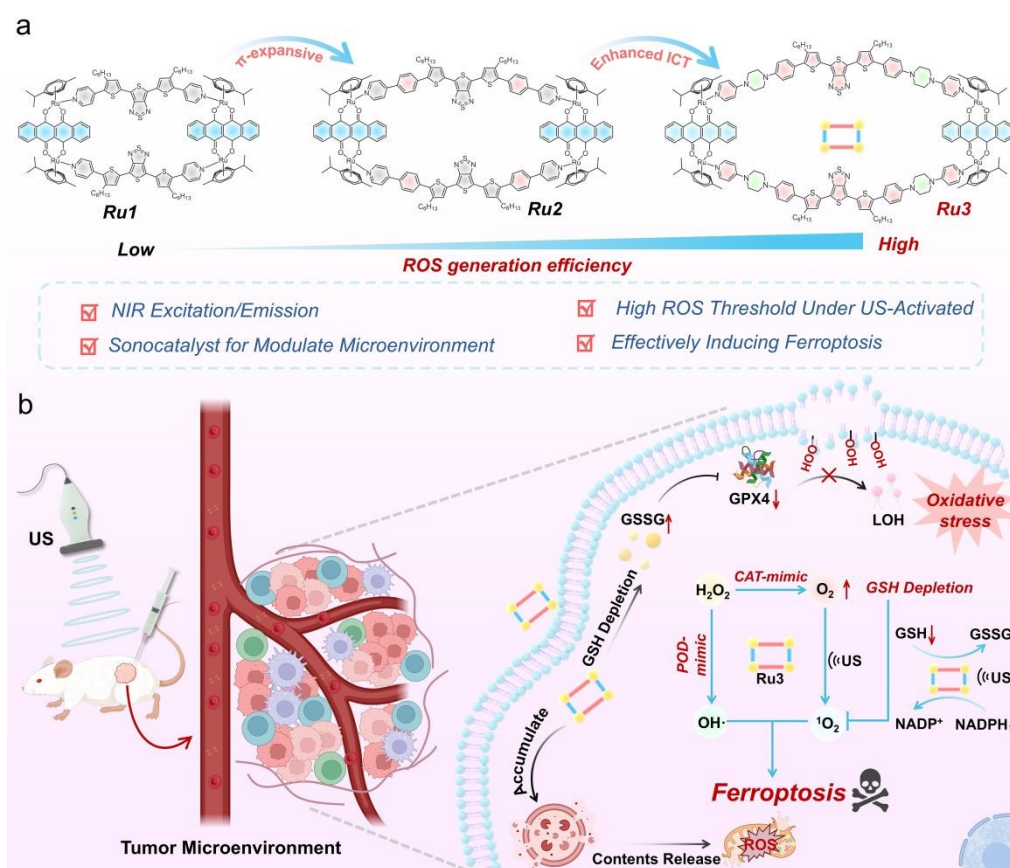
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Specifically, by directly affecting NADPH and depleting GSH, the activity of GPX4 can be inhibited, disrupting the redox balance in tumor cells and promoting ROS-mediated ferroptosis.²⁸ Therefore, in the development of efficient sonosensitizers, it is crucial to consider their inherent multifunctionality, including alleviating hypoxia and disrupting antioxidant defense systems, to better induce ferroptosis.

Recent studies have shown that supramolecular coordination complexes (SCCs), including Pt(II)- or Ru(II)-based SCCs *et al.*, have demonstrated distinct advantages over small molecular precursors in applications such as bioimaging, photodynamic therapy, and SDT.^{29–33} The formation of SCCs effectively confines sonosensitizers within a rigid structure, thereby minimizing the aggregation-induced ROS quenching effects. Furthermore, the incorporation of heavy metals such as the Ru atom into SCCs enhances intersystem crossing (ISC) and intramolecular charge transfer (ICT) processes, promoting the generation of sonosensitizer-induced ROS.^{34–36} More importantly, the presence of Ru(II) metal centers in SCCs, with their rich redox, optoelectronic, and sonocatalytic properties, may endow these complexes with significant potential for mimicking enzymatic and sonocatalytic activities, particularly in adapting to the complex tumor microenvironment.^{37–40} However, the use of SCCs as potential US-activated ferroptosis inducers remains an underexplored area.

In this study, we successfully designed and synthesized a series of supramolecular ferroptosis inducers (**Ru1–Ru3**) via molecular engineering, which possess sonosensitizing and sonocatalytic properties for near-infrared (NIR) fluorescence-guided SDT. Among them, **Ru3** exhibited superior ¹O₂ generation capability ($\Phi_{\Delta}=0.89$), attributed to its larger π -conjugation system and enhanced ICT effect, compared to homologues **Ru1** ($\Phi_{\Delta}=0.64$) and **Ru2** ($\Phi_{\Delta}=0.82$). In vitro studies demonstrated that the **Ru3** triggered a cascade of molecular events under US activation: First, **Ru3** displayed catalase (CAT)-mimic and peroxidase (POD)-mimic catalytic activities, significantly enhancing ROS generation and expanding their diversity. And the **Ru3** mainly accumulated in lysosomes, where the ROS generated in it increased lysosomes' membrane permeability (LMP), activated ferritinophagy, and released Fe²⁺ into the cytoplasm, where it catalyzed the Fenton reaction, further amplifying oxidative stress. More importantly, **Ru3** significantly induced the disruption of cellular redox homeostasis by catalyzing GSH depletion and promoting NADPH oxidation. These effects collectively suppressed GPX4 activity, promoted lipid LPO accumulation, and ultimately enhanced ferroptosis. Finally, in vivo experiments demonstrated that US-activated **Ru3** can effectively and safely ablate tumors in 4T1 tumor-bearing mice under the NIR fluorescence-guided assistance. Therefore, this study provides a promising strategy for the development of long-wavelength-emitting supramolecular ferroptosis inducers with potential clinical applications.



Scheme 1 Schematic illustration of the supramolecular ferroptosis inducers **Ru3** to modulate the tumor microenvironment under US irradiation for tumor therapy. (a) Molecular engineering strategy used for the synthesis of **Ru3**. (b) The potential anti-tumor mechanisms of **Ru3**.

Results and discussion

Design, synthesis, and characterization of Ru(II) metallacycles

To develop supramolecular ferroptosis inducers with high ROS generation efficiency and sonocatalytic capability, a series of novel



sonosensitizers were synthesized through rational molecular design and structural modifications. In this design, thienothiadiazole (TTD) was selected as an electron-deficient core to construct a donor–acceptor–donor (D-A-D) conjugated structure, considering that the strong push-pull electron effect would favor enhanced ROS generation.^{41–42} First, **L-1** was developed by using sterically hindered 3-hexylthienyl-substituted pyridine as the electron donor. Based on the thienyl donor engineering, **L-2** was obtained by introducing a phenyl ring structure to further expand the π -conjugation. To further enhance the ICT effect, **L-3** was synthesized by incorporating a piperazine structure with a stronger electron-donating ability. The chemical structures of **L-1** to **L-3** were comprehensively characterized using ¹H NMR, ¹³C NMR, and MALDI-TOF MS (Fig. S1–S24). Furthermore, density functional theory (DFT) calculations were performed using the Gaussian 09 to determine the highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) of **L-1**, **L-2**, and **L-3** (Fig. S25). The results showed that the energy gaps of **L-1** (1.74 eV), **L-2** (1.66 eV), and **L-3** (1.59 eV) gradually decreased, validating their potential as sonosensitizers.^{43–44} To further enhance the ROS generation efficiency and impart catalytic properties to these sonosensitizers, we chose a half-sandwich Ru(II) acceptor, with low dark toxicity and high ROS yield,^{45–46} to self-assemble with **L-1**, **L-2**, and **L-3** at a 1:1 molar ratio to form SCCs, named **Ru1**, **Ru2**, and **Ru3** (Scheme 1). Compared to the individual small-molecule precursors, the rigid structure and incorporation of heavy metal Ru in SCCs may enhance ROS yield under US activation. More critically, the presence of Ru(II) metal centers in SCCs, with their rich redox, optoelectronic, and catalytic capabilities, it could be that these SCCs are endowed with significant potential for mimicking enzymatic and sonocatalytic activities, particularly in adapting to the complex tumor microenvironment.

The SCCs, **Ru1**, **Ru2**, and **Ru3**, were synthesized through the reaction of **L-1**, **L-2**, and **L-3** with Ru(II) acceptor in a 1:1 methanol/chloroform mixture, stirred at room temperature for 12 hours, affording total yields of 6.48% (**Ru1**), 4.10% (**Ru2**), and 2.43% (**Ru3**), respectively, as detailed in the supplementary information. To characterize the synthesized SCCs, we initially employed ¹H NMR and 2D COSY (Fig. S26–S37). As shown in Fig. 1a and 1b, a downshift of the pyridine proton peak (H_a) was observed compared to the corresponding ligands, with chemical shifts of 8.69–8.83 ppm for **Ru1**, 8.74–8.87 ppm for **Ru2**, and 8.35–8.71 ppm for **Ru3**. For **Ru1**, **Ru2**, and **Ru3**, the proton H₁ of the SCCs was upshifted by approximately 0.20, 0.40, and 0.24 ppm, respectively, relative to the proton H₁ of the free Ru(II) receptor, while proton H₂ was upshifted by approximately 0.16, 0.17, and 0.21 ppm, respectively. These observed chemical shift changes during the self-assembly process are primarily put down to the decrease of electron density when the TTD-based precursor coordinate to the electron-deficient Ru(II) centers. The 2D COSY spectra further confirmed the correct assignment of each proton in **Ru1**, **Ru2**, and **Ru3**. The clear spectra and distinctive NMR signals from both ¹H NMR and 2D COSY spectra support the formation of these discrete SCCs. To further confirm the assembly of the SCCs, electrospray ionization time-of-flight mass spectrometry (ESI-TOF-MS) was carried out (Fig. 1c). Characteristic peaks were observed due to the elimination of trifluoromethanesulfonate (OTf[−]) counterions, indicating successful formation of the [2+2] rectangular SCCs, with *m/z* values of 975.21 for [**Ru1**-3OTf]³⁺, 1076.35 for [**Ru2**-3OTf]³⁺, and 1189.69 for [**Ru3**-3OTf]³⁺. And all assigned isotope peaks closely

matched the theoretical distribution, confirming that **Ru1**, **Ru2**, and **Ru3** have the anticipated [2+2] assembly structure (Figs S29, S33, and S37). Finally, DFT calculations were performed to optimize the most stable conformations of **Ru1**, **Ru2**, and **Ru3**. These calculations revealed that all three SCCs adopt a planar rectangular geometry (Fig. 1d and S38), with approximate cavity dimensions of 7.84/21.64 Å, 7.78/29.36 Å, and 7.75/37.31 Å (width/length) for **Ru1**, **Ru2**, and **Ru3**, respectively. These combined results confirm the successful self-assembly of the designed SCCs, **Ru1**, **Ru2**, and **Ru3**.

Photophysical properties, sonodynamic performance, and sonocatalytic activity studies

Based on these molecules, we subsequently investigated the photophysical and sonodynamic properties of **Ru1**, **Ru2**, and **Ru3**. Ultraviolet-visible (UV-Vis) absorption and photoluminescence (PL) spectroscopy were employed to study the optical properties of these molecules in dimethyl sulfoxide (DMSO). **Ru1**, **Ru2**, and **Ru3** exhibited maximum absorption wavelengths at 632 nm, 664 and 712 nm, respectively, which were similar to the absorption bands of precursors **L-1** to **L-3** (Fig. 2a and S39). Additionally, the maximum emission wavelengths of these molecules were observed at 944 nm, 959 nm and 1032 nm (with 808 nm as the excitation light), indicating their potential application in NIR fluorescence imaging. Notably, compared to **Ru1** and **Ru2**, **Ru3** exhibited a significant redshift in both absorption and emission wavelengths, ascribed to the expanded π -conjugation system and enhanced ICT in the molecular backbone.

Next, the chemical and sonostability of **Ru1**, **Ru2**, and **Ru3** were evaluated by monitoring changes in their absorption spectra. As shown in Fig. 2b and S40–S41, after incubating in phosphate-buffered saline (PBS) or 10% fetal bovine serum (FBS) for 6 days, no significant degradation in their absorption spectra was observed, indicating strong stability under physiological conditions. To assess the sonodynamic performance of these sonosensitizers, the ROS generation efficiency of **Ru1**, **Ru2**, and **Ru3** was evaluated using 2',7'-dichlorofluorescein (DCFH) as an indicator. Previous literature reports indicated that SDT commonly employed low-intensity focused US with a frequency range of 0.5–2 MHz. Among these, 1 MHz offered a more suitable balance between spatial resolution, controllability and ROS production efficiency, providing sufficient tissue penetration depth while minimizing tissue damage.^{4, 7–9} Upon US irradiation (1 W/cm², 1 MHz), DCFH alone did not produce fluorescence. Interestingly, after 5 minutes of irradiation, the DCF fluorescence intensity increased by 7.21-fold, 10.14-fold, and 14.31-fold for **Ru1**, **Ru2**, and **Ru3**, respectively (Fig. 2c). These results demonstrated that **Ru3** has significantly higher ROS generation efficiency than **Ru1** and **Ru2**. Furthermore, electron paramagnetic resonance (EPR) spectroscopy was conducted to assess the types of ROS generated by **Ru1**, **Ru2**, and **Ru3** under US irradiation. Among them, ¹O₂ was detected using 2, 2, 6, 6-tetramethylpiperidine (TEMP) as the indicator, while hydroxyl radicals (•OH) and superoxide anions (O₂^{•−}) were detected using 5,5-dimethyl-1-pyrroline N-oxide (DMPO). As shown in Fig. S42, under US irradiation, **Ru1**, **Ru2**, and **Ru3** primarily generated ¹O₂, with **Ru3** exhibiting the highest efficiency. The ¹O₂ quantum yields (Φ_Δ) of **Ru1**, **Ru2**, and **Ru3** under US irradiation were determined to be 0.64, 0.82, and 0.89, respectively, using methylene blue (MB, Φ_Δ = 0.52) as a reference sonosensitizer (Fig. S43). These three sonosensitizers produced negligible amounts of •OH and O₂^{•−} (Fig. S44–S45).



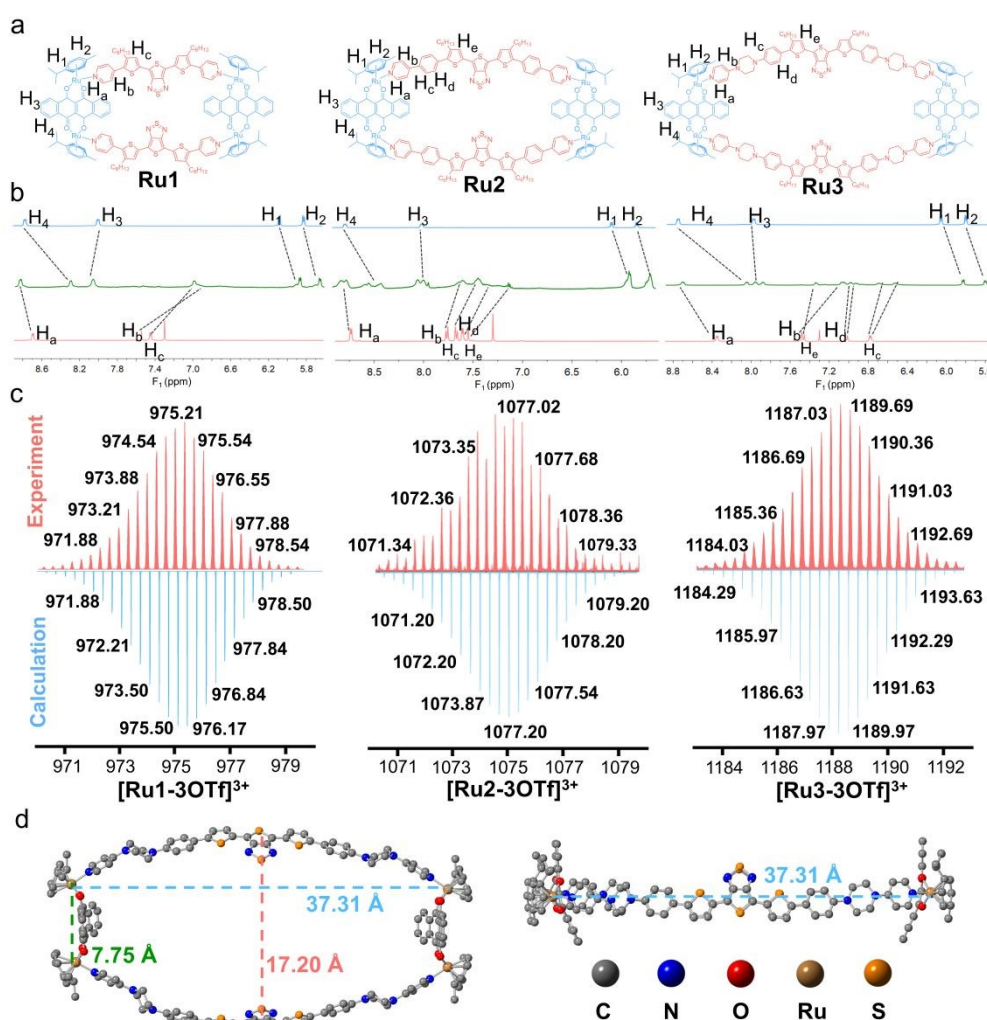


Fig. 1 The characterization of SCCs **Ru1**–**Ru3**. (a) The chemical structure of **Ru1**, **Ru2** and **Ru3**. (b) The partial ¹H NMR (400 MHz, 298 K) spectra of metallacycles **Ru1**–**Ru3**. (top: acceptor, middle: SCCs, bottom: ligand) (c) Calculated and experimental ESI-TOF-MS spectra of **Ru1** ([**Ru1**-3OTf]³⁺), **Ru2** ([**Ru2**-3OTf]³⁺), and **Ru3** ([**Ru3**-3OTf]³⁺). (d) Optimized molecular model of **Ru3**, top view (left), side view (right). For clarity, counterions and hydrogen atoms are omitted.

Previous studies have demonstrated that the incorporation of heavy atoms such as Ru in SCCs enhances ISC, thereby promoting ROS generation under external energy stimuli.⁴⁷ To explore the underlying mechanisms, we performed geometry optimization and molecular orbital analysis of **Ru1** to **Ru3** using the Gaussian 09. The results indicate that, compared to **Ru1** and **Ru2**, **Ru3** has the smallest singlet-triplet energy gap (Δ_{EST}) (**Ru1** = 0.46 eV, **Ru2** = 0.40 eV, and **Ru3** = 0.31 eV, Fig. S46). Additionally, the formation of SCCs effectively confines the sonosensitizer within a rigid structure, minimizing the aggregation-induced ROS quenching effect. To evaluate the ROS quenching resistance, we assessed the ROS generation of **Ru1**, **Ru2**, and **Ru3** in various DMF/water mixtures (Fig. 2d). The results showed that, with increasing water content in the

DMF/water mixture, the ROS generation of **Ru3** decreased dramatically due to strong intermolecular π - π stacking interactions. In contrast, **Ru3** exhibited superior ROS quenching resistance, attributed to its rigid structure and the steric hindrance of the Ru acceptor.^{29–35} Furthermore, to investigate whether SCCs can effectively generate ROS in deeper tissues, we studied the ROS penetration depth of **Ru3** in 1% lipid (solidified by 1% agarose) using a tissue model. The results revealed that **Ru3** achieved a penetration depth exceeding 10 cm (Fig. S47). In addition to the excellent sonodynamic performance exhibited by **Ru3** under US, its sonothermal properties have also been further evaluated (Fig. S48). Overall, **Ru3** demonstrated efficient ROS generation under US conditions, deep tissue ROS penetration, and excellent resistance to



ROS quenching, underscoring its potential as a ferroptosis inducer candidate.

Considering the redox enzyme-like activity demonstrated by **Ru3**-based agents, we next examined the catalytic activity of **Ru3** under conditions mimicking the intracellular environment. To evaluate the CAT-mimic activity of **Ru3**, we assessed its ability to decompose H_2O_2 into O_2 . Initially, $\text{Ru}(\text{dpp})_3\text{Cl}_2$ (RDPP) was used as an O_2 indicator to probe its CAT-mimic performance. RDPP is a well-known luminescent O_2 probe, widely used for O_2 detection and quantification.⁴⁸ The fluorescence of the RDPP (with an emission maximum of 613 nm) was strongly quenched by molecular O_2 due to dynamic quenching (λ_{max} at 455 nm). As shown in Fig. 2e, under US conditions, **Ru3** induced a decrease in the fluorescence intensity of RDPP in the

buffer solution. Similarly, the increase in dissolved O_2 and the formation of gas bubbles supported substantial O_2 generation (Fig. S2f and S49). We also confirmed this by observing an increase in ROS level upon the addition of H_2O_2 (Fig. S50). To assess the POD-mimic activity of **Ru3**, we measured its ability to induce $\bullet\text{OH}$ generation. As illustrated in Fig. 2g and S51, the characteristic absorption peak of MB at 665 nm underwent a pronounced reduction, coupled with visible color alterations following the addition of **Ru3**, thereby confirming the generation of $\bullet\text{OH}$. Similarly, by using tetramethylbenzidine (TMB) as a probe, which reacted with $\bullet\text{OH}$ to generated blue oxidized TMB with a characteristic absorption peak at 652 nm,⁴⁹ we observed a clear stronger absorption when the addition of **Ru3** (Fig. S52).

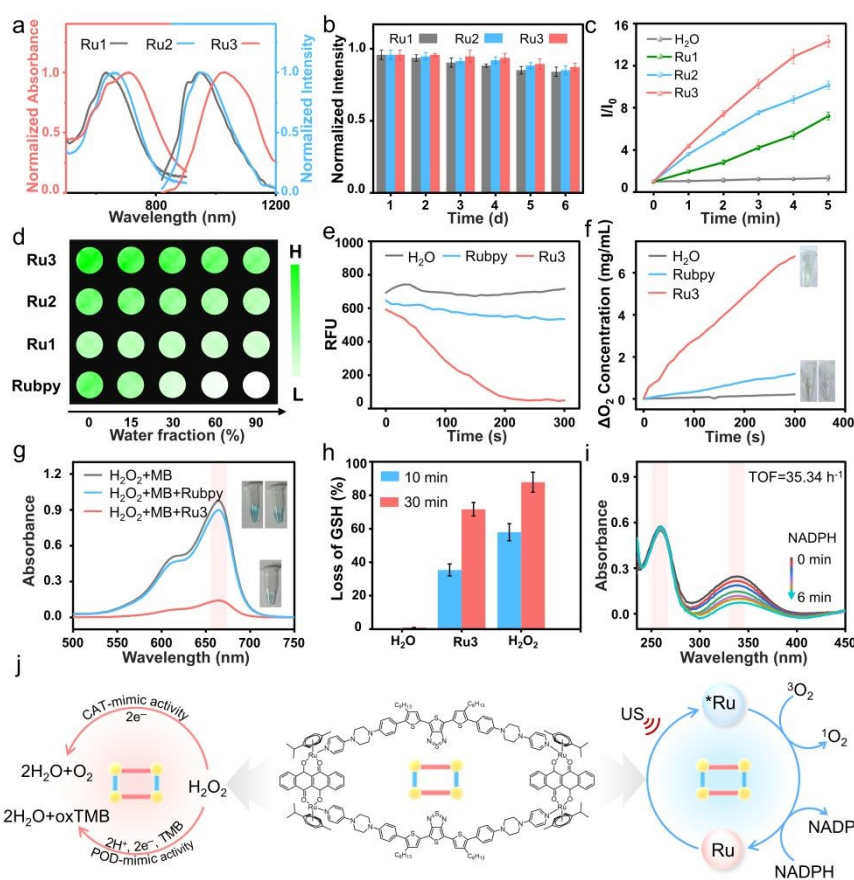


Fig. 2 The in vitro photophysical properties, sonodynamic performance, and sonocatalytic activity of SCCs **Ru3**. (a) Normalized absorption spectra and the emission spectra ($\lambda_{\text{ex}}=808$ nm) of **Ru1–Ru3** in DMSO. (b) Normalized absorption spectra of **Ru1–Ru3** after incubation in PBS for 1–6 d. (c) Normalized absorption spectra of **Ru1–Ru3** in DMF under US irradiation (1 W/cm²). (d) Fluorescent images of DCFH for detecting ROS generation of **Ru1–Ru3** in different ratios of DMF and H₂O during US irradiation. (e) Average fluorescence intensity of RDPP induced by **Ru3** and Rubpy. (f) The DO (ΔO_2) production after H₂O₂ co-incubation with **Ru3**. (g) UV/Vis absorption spectra of MB after co-incubation with or H₂O₂ and **Ru3**. (h) Time-dependent GSH consumption perform after co-incubation with **Ru3** or H₂O₂ using DTNB as indicator (n=3, mean±SD). (i) Average absorption spectra of NADPH after co-incubation with **Ru3** at different time points in PBS solution. (j) Schematic illustration of the sonocatalytic capability of **Ru3**.

GSH is a major endogenous antioxidant that plays a key role in maintaining redox homeostasis, capable of scavenging potential ROS bursts that may subsequently impair therapeutic outcomes.^{50–51} Given the redox properties of Ru(II), we further investigated the depletion of GSH in the presence of **Ru3**. To evaluate **Ru3**'s ability to consume GSH, we used 5, 5'-dithiobis(2-nitrobenzoic acid) (DTNB) as a GSH indicator. DTNB can react with GSH to form 5-thio-2-nitrobenzoic acid (TNB).⁵² As shown in Fig. 2h, **Ru3** could induce a considerable amount of GSH depletion over time, fully demonstrating the effective depletion effect of **Ru3** on GSH. Then, ¹H

NMR spectroscopy further verified the successful conversion of GSH to its oxidized form GSSG (Fig. S53). Also, cellular GSH can be rejuvenated from oxidized GSSG through a key NADPH-dependent process.⁵³ Therefore, we proceeded to investigate the sonocatalytic oxidation of NADPH by **Ru3**. The results showed that, upon addition of **Ru3**, the NADPH absorption at ~336 nm was significantly reduced with extended US exposure, and the NADPH oxidation turnover frequency (TOF) was 35.34 h⁻¹ (Fig. 2i and S54). Similarly, ¹H NMR spectroscopy further validated the transformation of NADPH into its oxidized form, NADP⁺ (Fig. S55). These observations suggested that



Ru3 exhibited CAT/POD-mimic multi-enzyme activity and sonocatalyst-enhanced GSH and NADPH depletion, highlighting its significant potential in sonocatalyst-enhanced SDT for tumor cells ferroptosis (Fig. 2j).

In vitro cell uptake, localization, cytotoxicity and sonocatalytic activity studies

Given the excellent SDT performance demonstrated by the physicochemical properties of supramolecular sonosensitizers, we subsequently conducted in vitro experiments in 4T1 cells. Firstly, inspired by the high-resolution imaging capabilities of **Ru3**, we evaluated its cellular uptake and localization. As shown in Fig. 3a and S56, after incubation with **Ru3**, 4T1 cells exhibited NIR-II fluorescence, which increased with incubation time and peaked at 24 hours. And the NIR-II fluorescence intensity of **Ru3** remained high during the 48-hour monitoring period. After determining the cellular uptake and retention efficiency of **Ru3**, we examined its subcellular localization to reveal its distribution within organelles. After incubated of 4T1

cells with **Ru3** and the LysoSensor Red probe (a commercial probe for lysosome imaging) for 30 min revealed that the NIR fluorescence signals from **Ru3** overlapped with the red fluorescence from LysoTracker Red, showing a Pearson correlation coefficient (PCC) of 0.81 (Fig. 3b). Likewise, **Ru3** could well overlap with a commercial MitoTracker Red imaging probe (PCC=0.72). These results suggested that **Ru3** effectively entered into 4T1 cells and primarily concentrated in lysosomes and mitochondria, which was critical for its anticancer efficacy.⁵⁴ Then, we further used the MTT assay to assess the cell viability of 4T1 cells after treatment with **Ru1-Ru3**. As shown in Fig. 3c and S57, the results indicated that **Ru3** exhibited negligible dark toxicity at low concentrations in the absence of US irradiation. When incubated with 10 μ M of Rubpy, **Ru1**, **Ru2**, and **Ru3**, and conducted US irradiation, the cell viability of Rubpy, **Ru1**, **Ru2**, and **Ru3** decreased to 90 %, 78 %, 57 %, and 35%, respectively, confirming that **Ru3** demonstrated significantly better performance in SDT compared to the other treatment groups.

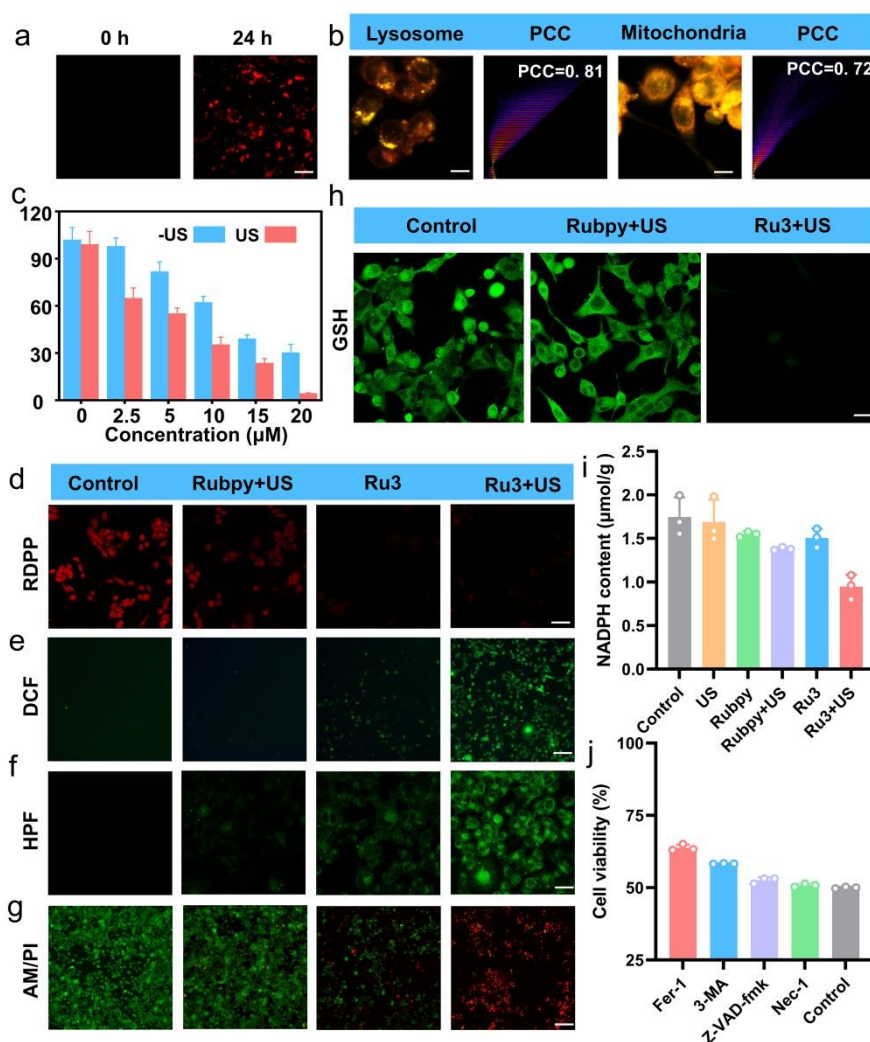


Fig. 3 In vitro cell uptake, localization, cytotoxicity and sonocatalytic activity of SCCs **Ru3** in 4T1 cells. (a) Fluorescence images of 4T1 cells treated with **Ru3** (10 μ M) at different time points (0 h and 24 h). Scale bar: 10 μ m. (b) Colocalization assay of **Ru3** (10 μ M) in 4T1 cells by using LysoTracker Red and MitoTracker Red. Scale bar: 10 μ m. (c) Cell viabilities of 4T1 cells after incubation with different concentrations of **Ru3** under US power (1 W/cm²). Error bars are mean \pm SD (n=3). (d) Fluorescence images of 4T1 cells co-incubation with RDPP and different treatments (Rubpy: 10 μ M, **Ru3**: 10 μ M, US power: 1 W/cm²). Scale bar: 10 μ m. (e) Fluorescence images of 4T1 cells co-incubation with DCF under different treatments. Scale bar: 25 μ m.



μm . (f) Fluorescence images of 4T1 cells stained by HPF incubated with **Ru3** (10 μM) and irradiated by US (1 W/cm^2). Scale bar, 10 μm . (g) Calcein-AM and PI-stained images of 4T1 cells incubated with different treatments (Rubpy: 10 μM , **Ru3**: 10 μM , US power: 1 W/cm^2). Scale bar, 25 μm . (h) The fluorescence images of GSH levels within the 4T1 cells after co-incubation with different treatment groups. (Rubpy: 10 μM , **Ru3**: 10 μM , US power: 1 W/cm^2). Scale bar, 10 μm . (i) Relative NADPH activity in 4T1 cell under various treatments. Error bars are mean \pm SD (n=3). (j) Cell viabilities of 4T1 cells co-incubated (24 h) with **Ru3** (10 μM) and various inhibitors, Fer-1 (10 μM), 3-MA (10 μM), Z-VAD-FMK (10 μM) and Nec-1 (10 μM) under US irradiation (1 W/cm^2). Error bars are mean \pm SD (n=3).

To assess the potential of **Ru3** as a sonosensitizer adaptable to the tumor microenvironment, we then evaluated the O_2 levels within 4T1 cells using the RDPP staining method. As shown in Fig. 3d and S58, in the **Ru3**-treated group, a significant reduction in the red fluorescence of RDPP was observed, confirming an elevation in O_2 levels within the 4T1 cells. Conversely, no substantial change in RDPP fluorescence was detected after Rubpy treatment, suggesting its limited O_2 production ability. These results indicated that **Ru3** may generate O_2 through the CAT-mimic process, alleviating hypoxia in 4T1 cells. Then, we incubated the SCCs **Ru3** together with the DCFH probe to validate the effective ROS generation of **Ru3** in 4T1 cells. As shown in Fig. 3e, after incubation with **Ru3** and 4T1 cells without irradiation, a visible green fluorescence was observed, which may suggest that **Ru3**-mediated POD-mimic catalysis induced the generation of $\bullet\text{OH}$ within the cells. Upon US treatment, the **Ru3** displayed a marked enhancement in green fluorescence compared to the Rubpy group, indicating US-triggered enhanced ROS generation. We further evaluated the POD-mimic activity by employing HPF as a $\bullet\text{OH}$ probe, and the results revealed a notable increase in HPF fluorescence intensity after **Ru3** incubation, indicating the production of $\bullet\text{OH}$ via the POD-mimic mechanism in 4T1 cells (Fig. 3f). Finally, we used Calcein AM and propidium iodide (PI) staining to distinguish live cells (green) from dead cells (red) (Fig. 3g). These results demonstrate that the **Ru3** plus US group effectively generated ROS in 4T1 cells, showing enhanced sonodynamic performance.

Next, we investigated the consumption of GSH and the sonocatalytic oxidation of NADPH by **Ru3** in 4T1 cells. ThiolTrace Violet 500 was used as an indicator for intracellular GSH levels. As shown in Fig. 3h and S59, the green fluorescence of ThiolTrace Violet 500 in the **Ru3**-treated group was significantly diminished, indicating a reduction in GSH levels in 4T1 cells, which confirmed **Ru3**'s consumption of intracellular GSH. In contrast, no significant change in fluorescence was observed in the Rubpy-treated group, suggesting a weaker GSH depletion. After US irradiation, the green fluorescence in the **Ru3** group almost completely disappeared, indicating that US-triggered ROS generation further depleted intracellular GSH. Subsequently, the sonocatalytic oxidation of NADPH by **Ru3** in 4T1 cells was studied (Fig. 3i). After **Ru3** treatment and US irradiation, significant oxidation of NADPH was observed, consistent with our sonocatalytic studies.

Cell Death Mechanisms studies

To explore the potential mechanisms of cell death under US irradiation, we assessed the cell viability of 4T1 cells in the presence of various cell death pathway inhibitors (Fig. 3j), including ferroptosis, autophagy, apoptosis and necrosis. After treatment with z-VAD-fmk (an apoptosis inhibitor) or necrostatin-1 (Nec-1, a necrosis inhibitor), cell viability almost remained unchanged, indicating non-apoptotic

and non-necrotic cell death. On the contrary, the addition of the ferroptosis inhibitor Ferrostatin-1 (Fer-1) effectively increased the survival of 4T1 cells, suggesting that US-activated **Ru3** may induce ferroptosis. Furthermore, the autophagy inhibitor 3-methyladenine (3-MA) was found to partially improve cell viability, indicating that autophagic processes in lysosomes possibly contribute to this cell death mode.

Increasing evidence suggests that lysosome-targeting sonodynamic materials exhibit ferroptosis effects in tumor cells, which are more prominent than ROS-induced cell damage. Lysosomal membrane permeabilization (LMP) induces the release of various substances into the cytoplasm, such as protons and iron, which activate or amplify cell death signaling under iron depletion conditions.⁵⁵⁻⁵⁷ Similarly, mitochondrial dysfunction is a significant marker of ferroptosis. Cells undergoing ferroptosis often show dissipation of mitochondrial membrane potential (MMP), increased mitochondrial membrane permeability, and structural damage to mitochondria. MMP is essential for maintaining mitochondrial function and energy metabolism.⁵⁸⁻⁶¹ Given that **Ru3** was primarily concentrated in both lysosomes and mitochondria, we then investigated the sonodynamic damage to lysosomes and mitochondria in situ. The JC-1 probe was used as a probe to detect the changes in MMP in cells. As shown in Fig. 4a, the **Ru3** plus US treatment group exhibited the strongest green fluorescence, indicating synergistic depolarization and mitochondrial membrane damage. Subsequently, using acridine orange (AO) as an indicator, the red fluorescence of AO in 4T1 cells treated with **Ru3** plus US disappeared dramatically compared to other treatments, indicating a loss of lysosomal integrity and dysfunction (Fig. 4b).

Previous studies have shown GSH depletion can inactivate GPX4, thereby upregulating ROS-dependent LPO, which is a crucial factor in ferroptosis-induced cell death.^{62, 10-13} Considering the oxidative stress imbalance and GPX4 inactivation, we assessed the changes in LPO levels during the US-induced tumor cell death process. As shown in Fig. 4c and S60, the **Ru3** plus US treatment group exhibited significant LPO accumulation, as evidenced by the red fluorescence in the control group and the decrease in red fluorescence and increase in green fluorescence in the **Ru3** plus US treatment group. Moreover, since malondialdehyde (MDA) is a key end product of LPO, we measured the MDA concentration in 4T1 cells treated with different conditions. The results showed that the **Ru3** plus US group exhibited the highest MDA levels compared to other control groups (Fig. 4d). Additionally, transmission electron microscopy (TEM) analysis revealed morphological changes in 4T1 cells mitochondria, including mitochondrial shrinkage, increased membrane density, and decreased cristae, which are typical of ferroptosis-induced mitochondrial dysfunction (Fig. 4e).



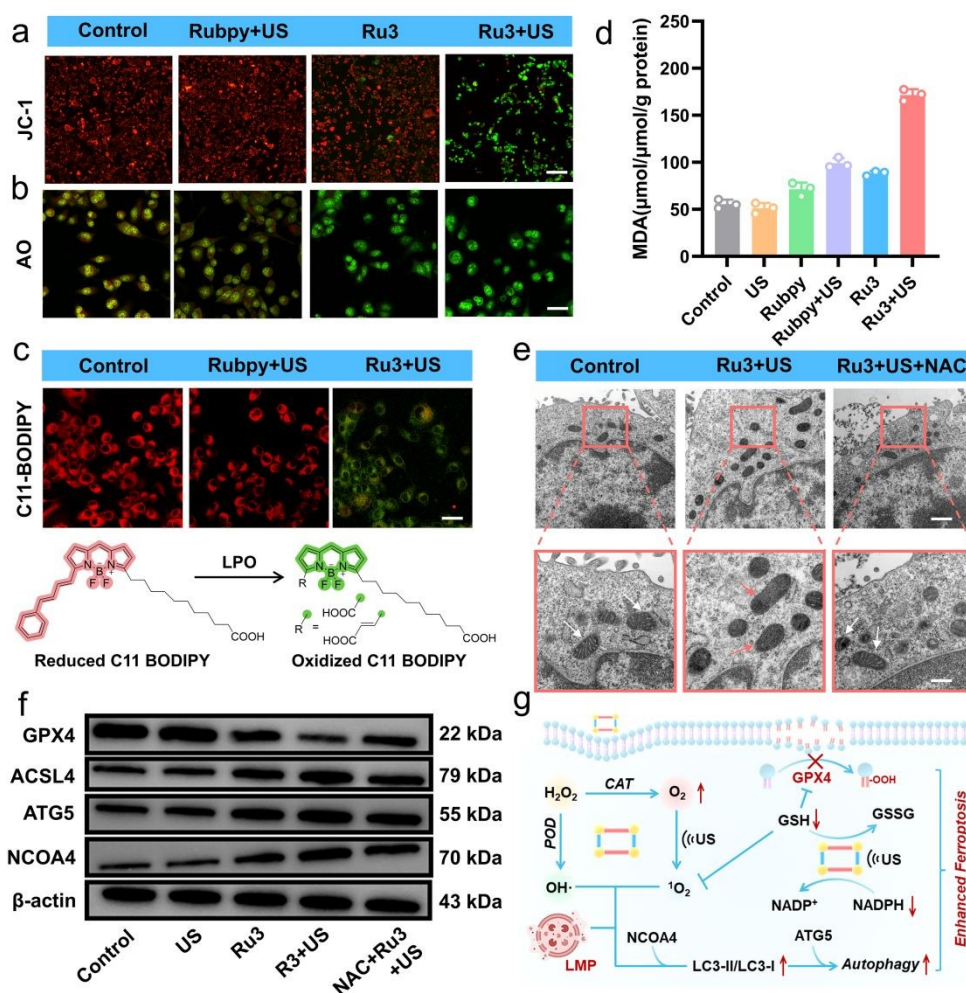


Fig. 4 Ferroptosis mechanism of 4T1 cells under **Ru3** plus US treatment. Fluorescence images of 4T1 cells co-incubation with Rubpy/**Ru3** (10 μM) and probe (a) JC-1 (b) AO, with or without US irradiation. Scale bar JC-1: 25 μm; scale bar AO: 10 μm. (c) Fluorescence images of C11-BODIPY-stained 4T1 Cells with the red and green channels indicating reduced C11-BODIPY and oxidized C11-BODIPY, respectively (top) and the mechanism of C11-BODIPY ox/red. Scale bars: 10 μm. (d) Relative MDA activity in 4T1 cells under various treatments. Error bars are mean ± SD (n=3). (e) Bio-TEM images showing the existence of ferroptosis after being treated with various treatments. Scale bar (top image): 5 μm; Scale bar (below image): 1 μm. (f) Western blot assay of GPX4, ACSL4, ATG5 and NCOA4 levels in 4T1 cell treated with various treatments. (g) Cartoon illustration of the ferroptosis mechanism induced by **Ru3** under US irradiation in 4T1 cells.

To further investigate the ferroptosis induced by **Ru3** under US irradiation in 4T1 cells, we performed western blot (WB) analysis to detect the expression of ferroptosis-related proteins. As shown in Fig. 4f and S61, the protein expression of GPX4 was visibly reduced, which can be attributed to **Ru3**'s ability to deplete GSH, while US irradiation further exacerbated this depletion. These results suggest that the generation of multiple ROS, including $\bullet\text{OH}$ and $^1\text{O}_2$ and the depletion of GSH may together lead to irreversible GPX4 inactivation, consistent with the classical hallmark of ferroptosis. Additionally, we examined ferroptosis-related pathway markers, including Acyl-CoA synthetase long-chain family member 4 (ACSL4), which facilitates the

esterification of polyunsaturated fatty acids (PUFAs) into PUFA-phospholipids (PUFA-PLs), which are prone to lipid peroxidation.⁶³⁻⁶⁶ WB analysis revealed that **Ru3** plus US treatment significantly upregulated the expression of ACSL4, further confirming that **Ru3** can induce ferroptosis in tumor cells. Previous reports have shown that oxidative stress can induce ferritinophagy and increase intracellular iron levels. This prompted us to explore whether ferritinophagy occurs in the cells. Nuclear receptor coactivator 4 (NCOA4) is known to be a key regulator of ferritinophagy, as it binds to ferritin and facilitates its transport to lysosomes for degradation.⁶⁷⁻⁷² Notably, WB results showed a significant increase



in NCOA4 expression, as well as in the LC3II/LC3I ratio and autophagy-related protein 5 (ATG5) under the treatment of **Ru3** plus US, which could be reversed by NAC treatment. Additionally, changes in the levels of the relevant proteins GPX4, ASCL4, and ACOA4 were further monitored through Raman signal variation, and the results

were consistent with the ferroptosis outcomes (Fig. S62). Overall, the enzyme-like activity and sonocatalytic performance of **Ru3** enhanced ROS generation, induced ferritinophagy, depleted GSH and NADPH, and ultimately inhibited GPX4 activity, significantly promoting ferroptosis in tumor cells (Fig. 4g).

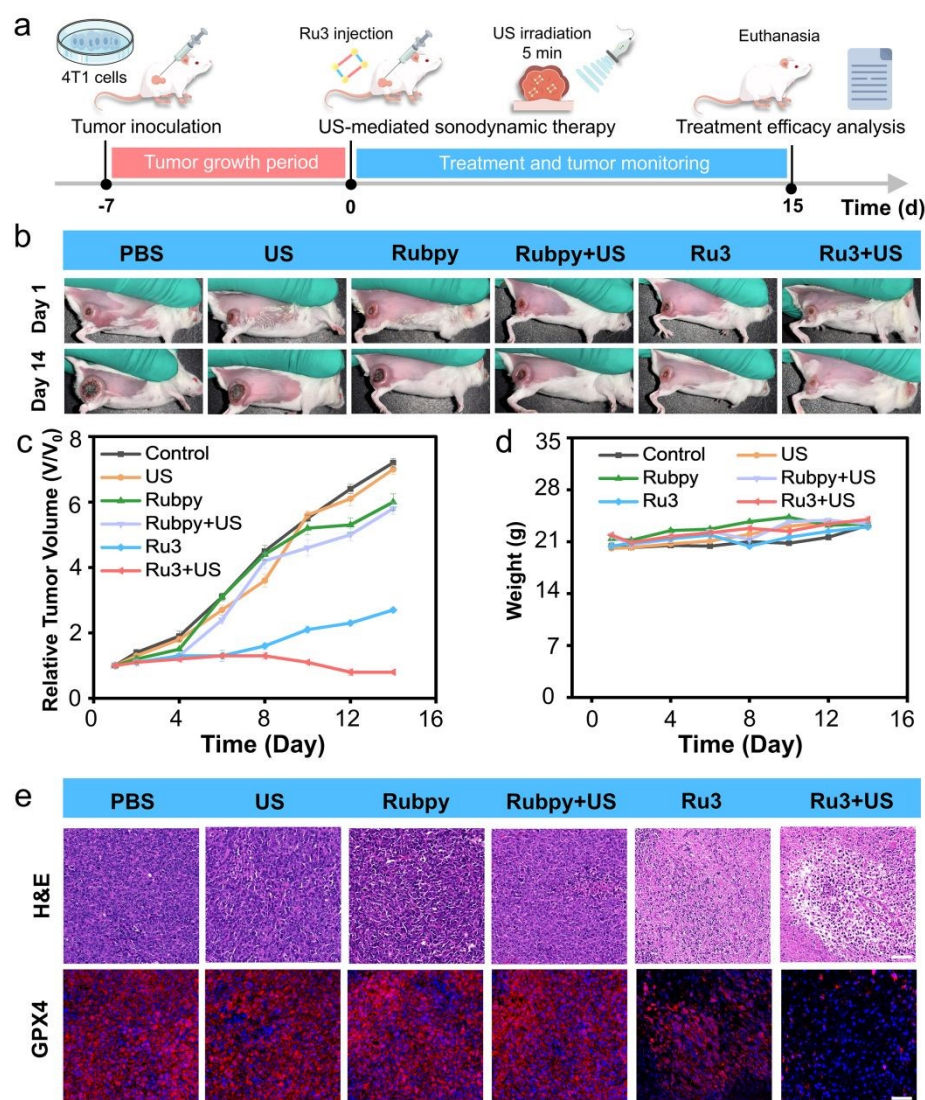


Fig. 5. In vivo antitumor experiments. (a) The protocol of the experiment. (b) Representative image of 4T1 tumor-bearing mice under different treatments at day 1 and day 14. (c) The linear curve depicts the 4T1 tumor suppression in different treatment groups. Error bars are mean \pm SD (n=5). (d) Representative image depicts the body weight change of mice after various treatments. Error bars are mean \pm SD (n=5). (e) H&E and GPX4 staining of tumor slices collected from tumor mouse models. Scale bar: H&E, 100 μ m; GPX4, 50 μ m.

In vivo anticancer application

Given the excellent in vitro synergistic therapeutic effects of **Ru3**, we evaluated its in vivo antitumor efficacy using a 4T1 tumor-bearing mouse model. Before conducting in vivo experiments, we first performed hemolysis tests to ensure the biocompatibility of **Ru3** in biological systems (Fig. S63). Next, a total of 30 mice were randomly divided into 6 groups (5 mice per group): (1) control group, (2) US group, (3) Rubpy group, (4) Rubpy + US, (5) **Ru3** group, and (6) **Ru3** + US group. Mice in the Rubpy and **Ru3** groups were intratumorally

injected with Rubpy (1 mg Ru/kg) and **Ru3** (1 mg Ru/kg), respectively, while the other groups received PBS. Following intratumoral injection of PBS, Rubpy or **Ru3**, the mice were subjected to US irradiation (1.0 MHz, 1.0 W/cm², 50% duty cycle, 5 min) at 12 hours post-injection (US, Rubpy + US, **Ru3** + US groups), or no US treatment (PBS, Rubpy, **Ru3** groups) (Fig. 5a). Mouse body weight and tumor volume were recorded every other day. We then visualized the tumor after injection with **Ru3** by using NIR fluorescence imaging, which guided subsequent sonodynamic therapy (Fig. S64). After US irradiation, the results showed complete tumor eradication in the



Ru3 plus US group after 14 days of treatment. In contrast, the Rubpy plus US group exhibited only mild tumor suppression (Fig. 5b and 5c). Furthermore, to assess the acute systemic toxicity of **Ru3**, we monitored body weight changes in the mice (Fig. 5d). The results indicated that the weight loss observed during treatment was negligible. Finally, on day 20, normal organs from each group of mice were collected for hematoxylin and eosin (H&E) staining. After 20 days of treatment, no significant organ damage was observed in any of the treatment groups (Fig. S65). It is noteworthy that the tumor tissue from the **Ru3**+US group exhibited a significantly lower cancer cell density compared to the other control groups, indicating the significant effectiveness of **Ru3**-mediated sonodynamic therapy. GPX4 immunofluorescence staining showed a marked downregulation in the **Ru3**+US group, suggesting that **Ru3**-induced ferroptosis plays a key role in its antitumor activity (Fig. 5d). These preliminary but comprehensive evaluations suggest that the sonosensitizer **Ru3** exhibits excellent biocompatibility and is suitable for potential in vivo therapeutic applications.

Data availability

All the data supporting this article have been included in the main text and the ESI.†

Author contributions

Y. Sun conceived the project and designed the experiments. Y. Pang, C. Li and J. Zhang designed, synthesized and characterized the materials. L. Mei carried out theoretical calculation. Y. Luo and Q. Li performed the *in vitro* and *in vivo* study. Y. Sun, Y. Pang, C. Li and J. Li wrote the manuscript. All authors analyzed and discussed the results and have given approval to the final version of the manuscript.

Data availability

There are no conflicts to declare.

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Conclusions

In conclusion, we successfully developed a series of supramolecular ferroptosis inducers (**Ru1–Ru3**) through a molecular engineering approach for NIR fluorescence-guided ferroptosis-enhanced SDT. Under US irradiation, **Ru3**, with its extended π -conjugation system and enhanced ICT effect, demonstrated superior ROS generation. In vitro studies revealed that **Ru3** exhibited both CAT-mimic and POD-mimic catalytic activities, further boosting ROS production and broadening their diversity. Furthermore, **Ru3** catalyzed the depletion of GSH and promoted NADPH oxidation, disrupting cellular redox homeostasis, inducing autophagy, and triggering ferroptosis in 4T1 tumor cells. In vivo, **Ru3** effectively inhibited 4T1 tumor growth upon US activation with negligible side effects. This work offers a promising strategy for developing long-wavelength-emitting supramolecular sonosensitizers, particularly for catalytically enhanced ferroptosis therapy with potential clinical applications.

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Data availability

All the data supporting this article have been included in the main text and the ESI.†

