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Wei Li, Jing Yu, Jing Wang, Xuejing Fan, Ximing Xu, et al.. How Does Ferrocene Correlate with Ferroptosis? Multiple Approaches to Explore Ferrocene-Appended GPX4 Inhibitors as Anticancer Agents. *Chemical Science*, 2024, 15, pp.10477-10490. 10.1039/D4SC02002B . hal-04601570

HAL Id: hal-04601570

<https://hal.science/hal-04601570v1>

Submitted on 5 Jun 2024

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Chemical Science

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How Does Ferrocene Correlate with Ferroptosis? Multiple Approaches to Explore Ferrocene-Appended GPX4 Inhibitors as Anticancer Agents

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Keywords bioorganometallic chemistry; ferrocene; ferroptosis; antitumour agents; bioinorganic chemistry.

Abstract

Ferroptosis has emerged as a form of programmed cell death and exhibits remarkable promise for anticancer therapy. However, it is challenging to discover ferroptosis inducers with new chemotypes and high ferroptosis-inducing potency. Herein, we report a new series of ferrocenyl-appended GPX4 inhibitors rationally designed in a "one stone kills two birds" strategy. Ferroptosis selectivity assays, GPX4 inhibitory activity and CETSA experiments validated the inhibition of novel compounds on GPX4. In particular, the ROS-related bioactivity assays highlighted the ROS-inducing ability of **17** at the molecular level and their ferroptosis enhancement at the cellular level. These data confirmed the dual role of ferrocene



as both the bioisostere motif maintaining the inhibition capacity of certain molecules with GPX4 and also as the ROS producer to enhance the vulnerability to ferroptosis of cancer cells, thereby attenuating tumor growth *in vivo*. This proof-of-concept study of ferrocenyl-appended ferroptosis inducers via rational design may not only advance the development of ferroptosis-based anticancer treatment, but also illuminate the multiple roles of the ferrocenyl component, thus opening the way to novel bioorganometallics for potential disease therapies.

Introduction

Ferroptosis was defined in 2012 as an iron-dependent form of programmed cell death caused by increased cellular reactive oxygen species (ROS) and lipid peroxidation (LPO).¹ Lipid metabolism, ROS biology, and iron regulation have been summarized as intersections from the centrality of the framework of ferroptosis.² In the past decades, ferroptosis has been proven to engage in human diseases including cancer, neurodegeneration and ischemic disease.³⁻⁹ Notably, accumulating evidence shows that ferroptosis plays important roles in the cancer occurrence, development and metastasis, which highlights its great potential for the treatment of refractory tumors.¹⁰ An deeper understanding of the epithelial–mesenchymal transition (EMT) in metastasis underscores the critical role of glutathione peroxidase 4 (GPX4) in cancer cells undergoing EMT; it is the only peroxidase in mammals capable of reducing lipid hydroperoxides.¹¹⁻¹² Indeed, when GPX4 function is impaired, lipid peroxidation causes ferroptosis which is a tumor-suppressive process. Moreover, further research suggested that blocking GPX4 function or decreasing the protein level of GPX4 can suppress tumor growth.¹³ Thus, aggressive neoplastic diseases might be treated through the use of GPX4 inhibitors.

Iron homeostasis composes the essential part of ferroptosis which is mainly mediated via the Fenton chain reaction.¹⁴⁻¹⁵ Therefore, elevated levels of iron can increase the vulnerability to ferroptosis. The traditional Fenton system composed of H₂O₂ and Fe²⁺ has been intensively investigated and used in cancer treatment. The ferrocenyl group is perhaps the most widely-studied organometallic moiety and enables the generation of ROS, which has contributed to its wide-spread exploration for use in medicinal chemistry.¹⁶ The electron donor–acceptor ability of the central iron atom endows ferrocene with good redox characteristics and high catalytic capacity as an excellent Fenton catalyst, which effectively enhances the biological behavior of the original drugs.¹⁷⁻²⁰ Considering the high level of interest in ferroptosis, whose impetus towards the formation of lipid hydroperoxides arises from the iron-catalyzed Fenton reaction, exploration of the effect of ferrocene in the framework of ferroptosis (Figure 1A)



could be a valuable contribution. Thus far, however, comprehensive investigation of ferrocene in terms of ferroptosis inducers has been rather limited, and reported only for iron/ferrocene-based inorganic complexes or nanomaterials.²¹⁻²⁴

In recent years, transition metal bioorganometallic chemistry has gradually come to the fore regarding its potential for the development of metallodrugs to circumvent the drawbacks²⁵⁻³¹ associated with the platinum complexes.³²⁻³⁷ Amongst the organometallic complexes that can be used as antitumoral agents, ferrocene occupies a privileged position as a compact, stable, non-toxic metallocene not only acting as a bioisostere for aryl/heteroaryl rings but also interfering the cellular redox homeostasis.^{17-18, 38-41} Initially, a library of ferrocene-based compounds (>100 entries) was established and then used to screen for ferroptosis inducers. Unfortunately, none of these random ferrocenyl-bearing entities could be characterized as ferroptosis inducers, although some have moderate cytotoxicities. These results indicated that regular ferrocenyl entities may not be able to induce ferroptosis significantly, which contradicted the conservative perception of ferrocene for ferroptosis and drove us towards the rational design of ferrocene-based ferroptosis inducers.

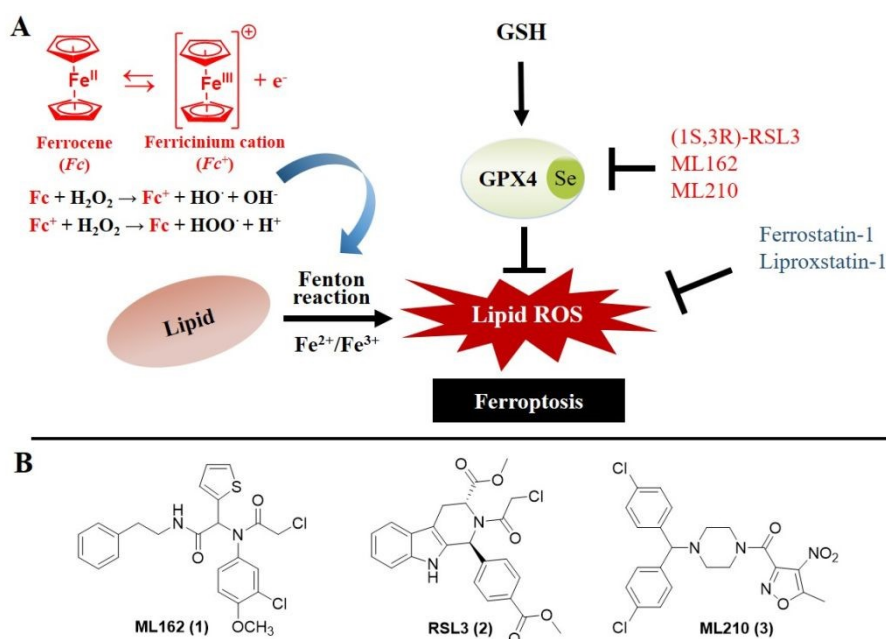
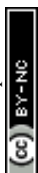


Figure 1. A) Illustration of the basis of developing ferrocene-based ferroptosis inducers. B) Typical small molecule ferroptosis inducers.

Thus, a “one stone kills two birds” strategy has been developed in this paper integrating the ferrocene fragment into the typical ferroptosis inducers, such as the widely-used GPX4 inhibitors (ML162, RSL3 and ML210, Figure 1B). We here describe multiple approaches towards the development of ferrocene-appended GPX4 inhibitors as anticancer agents,



including structure-based drug design (SBDD), ferroptosis-targeted phenotypic evaluation and other molecular and cellular investigations. The ferroptosis selectivity assays, GPX4 inhibitory activity and CETSA experiments validated the inhibition of these novel compounds on GPX4. The ROS-related bioactivity assays emphasized the ROS-inducing ability of selected compounds at the molecular level and their ferroptosis enhancement at the cellular level. These data confirmed the dual role of ferrocene as both the bioisostere fragment maintaining the binding capacity of certain molecules with GPX4 and also as the ROS producer to enhance the vulnerability to ferroptosis of cancer cells, thereby attenuating tumor growth *in vivo*. This work describes a proof-of-concept study of ferrocene-appended dual-function ferroptosis inducers for cancer therapies via a rational design procedure, and consummated the potential application of ferrocene in ferroptosis-targeted drug development.

Results and Discussion

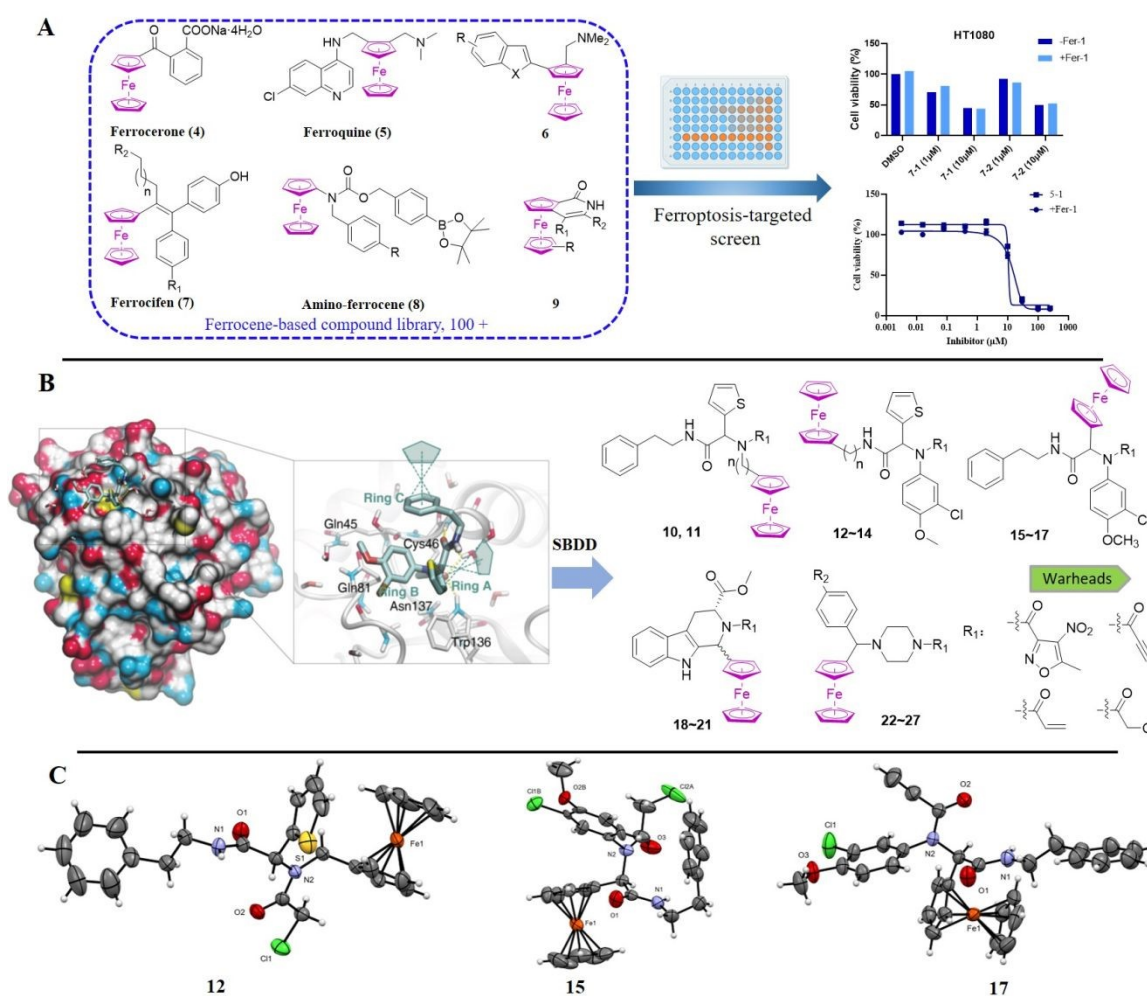


Figure 2. A) Illustration of the ferrocene-based organometallic compound library construction and screening. B) Structure-based drug design (SBDD) of ferrocene-appended GPX4 inhibitors. C) Molecular structures of the ferrocene complexes **12**, **15** and **17** with thermal ellipsoids shown at 50% probability.

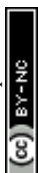


Ferrocene-based organometallic compound library construction and screening

The bioorganometallic community has explored the use of ferrocenyl compounds as inhibitors or as auxiliary groups to augment the efficacy of existing organic moieties, and some of them, such as ferroquine,⁴² have entered clinical trials. In an initial approach to explore the potential of ferrocenyl compounds as ferroptosis inducers, an in-house ferrocene-based compound library was constructed, consisting of simple mono- or di-substituted ferrocenes, known ferrocene-based drugs candidates, such as the ferrocifen derivatives developed in our own laboratories,⁴³⁻⁴⁶ aminoferrocene analogues from the Mokhir group^{40, 47} and ferroquine, as well as several chiral ferrocene derivatives reported by the You group⁴⁸⁻⁵⁰ (Figure 2A). Approximately 10% of the library was purchased from commercial sources and used without further modification, while the remaining majority of complexes were either generous gifts from the original laboratory, or were prepared according to literature procedures. Initially, a primary screening was carried out for ferrocene complexes at two concentrations (1 and 10 μM) on ferroptosis-sensitive HT1080 cells, with or without ferrostatin-1 (fer-1), a known specific ferroptosis inhibitor.¹ For the complexes having moderate antiproliferative activities, their IC_{50} values were further tested to double check their ferroptosis selectivity. Unfortunately, although some of these organometallics can inhibit proliferation of HT1080 cells at the micromolar level, these cytotoxicities cannot be reversed significantly by the ferroptosis inhibitor fer-1, as shown in Figure S1. These results indicated that the antiproliferative effect of the tested ferrocene complexes may not be relevant to ferroptosis, even though some molecules have been claimed to produce ROS significantly at both the molecular and cellular levels including ferrocifen and aminoferrocene derivatives.^{40, 51} These observations suggested there is no immediate connection between ferrocene and ferroptosis, which highlights its unique mechanism.

Rational design and synthesis of ferrocene-appended GPX4 inhibitors

Apparently, the currently available ferrocene-based chemical entities could not induce ferroptosis, which prompted us to explore the effect of ferrocene in terms of ferroptosis via a rational design strategy. Inhibition of GPX4 with small molecules has been established as a pharmacological stimulator of ferroptosis, that can achieve durable responses in a range of cancer types supported by evidence both *in vitro* and *in vivo*.⁵²⁻⁵³ The most well-known and potent GPX4 inhibitors involve two covalent binders RSL-3 and ML162 having chemically reactive chloroacetamides as warheads, as well as an additional prodrug, ML210, that is converted to a nitrile oxide reactant in cells.⁵⁴ In a preliminary analysis of the cocrystalline

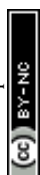


structure of GPX4 with its covalent inhibitor ML162 in a monomer model,⁵⁵ we noted that the solvent-exposed area of the ligand ML162 exceeds 60%, especially in terms of rings A and C (Figure 2B). Assuming that the under-occupied lipophilic pocket around rings A and C may be able to accommodate the ferrocenyl sandwich motif, this could provide an approach towards targeting GPX4 by ferrocene-appended GPX4 inhibitors.

To this end, 18 covalent GPX4 inhibitors with a ferrocenyl substituent incorporated at varied positions were synthesized so as to replace the aromatic ring of ML162, RSL3 and ML210, as shown in Figure 2B. Efficient syntheses of the designed ferrocenyl dipeptide analogues **10** - **17** proceeded in moderate to good yields, each via a Ugi four-component reaction from a diversity of reactants, aryl aldehydes, amines, carboxylic acids and isocyanides (Scheme S2). The synthesis of ferrocenyl RSL3 derivatives **18** - **21** was accomplished by Pictet–Spengler condensation of tryptophan analogues with formylferrocene, followed by an amide coupling (Scheme S3). It took several steps to prepare the third series, ferrocenyl compounds **22** - **27** were each obtained in a multi-step process (Scheme S4), Benzoylferrocene derivatives were first prepared by a Friedel-Crafts acylation and then reduced, using sodium borohydride, to the corresponding alcohols **FC-2a/2b**. Treatment of **FC-2a/2b** with acetic anhydride provided the ester analogues, and was followed by treatment with excess of piperazine in refluxing acetonitrile to afford **FC-3a/3b**. Coupling of secondary amines **FC-3a/3b** with carboxylic acids or acyl chlorides in dichloromethane delivered the final compounds **22** - **27**. The molecular structures of **12**, **15** and **17** were unequivocally established by X-ray crystallography, as illustrated in Figure 2C and Table S5-S8.

In Vitro Anticancer Evaluations and Cellular Uptake Measurements

As mentioned above, a preliminary screening on ferroptosis-sensitive HT1080 cells, with or without fer-1, has been routinely used to evaluate the anticancer activity of our molecules (as shown in Table 1). Encouragingly, in the case of ML162 analogues, the designed compound **15** with a ferrocenyl group instead of 2-thienyl showed comparable antitumor activity and ferroptosis selectivity to that of ML162. However, introducing the ferrocenyl group at either the R₂ or R₄ position on the dipeptide scaffold of ML162 seems to have a negative influence on both the antitumor activity and the ferroptosis selectivity. Compounds **10-14** displayed only moderate cytotoxicity on HT1080 cells without significant ferroptosis selectivity. More interestingly, the ferroptosis-inducing ability of dipeptide analogues bearing a ferrocenyl group can be modulated by changing the covalent warhead.



Compound **17** possessing a propiolamide as its warhead showed a large improvement in antitumoral activity on HT1080 with IC₅₀ values at nanomolar level as well as excellent ferroptosis selectivity. However, replacing chloroacetamide with acrylamide, as in **16**, resulted in a very dramatic loss of activity relative to ML162. The best combination therefore appears to be a ferrocenyl group at the R₃ position and a propiolamide as the warhead.

Table 1. Antitumor activities and ferroptosis selectivity of target compounds towards HT1080 cells.

Comp.	R ₁	R ₂	R ₃	R ₄	HT1080, IC ₅₀ (μM) ^a		Selectivity ^b
					-fer-1	+fer-1	
10					1.044±0.201	1.699±0.014	1.6
11					4.980±0.954	6.732±1.228	1.4
12					2.520±0.290	1.749±0.238	0.7
13					4.866±1.087	2.982±0.667	0.6
14					1.449±0.069	4.068±0.520	2.8
15					0.043±0.007	2.772±0.231	64.5
16					8.510±1.371	9.391±0.459	1.1
17					0.007±0.001	1.486±0.193	212.3
ML162					0.036±0.013	2.200±1.320	61.1

(a) Values are expressed as the mean of three independent experiments ± SD; (b) Selectivity: HT1080 IC₅₀ with fer-1/ HT1080 IC₅₀ without fer-1.

In contrast to the diverse role of ferrocene on the antitumor behavior of ML162, incorporation of the ferrocenyl unit in RSL3 and ML210 by replacing the phenyl with ferrocenyl generally triggers a global loss of activity (as shown in Table S9). In particular, in the analogues of ML210, all synthesized compounds bearing ferrocene show diminished ferroptosis-inducing activity, even with varied covalent warheads. One should note that, in the RSL3 series, only ferrocenyl complex **21** possessing a propiolamide warhead exhibited approximately two-fold decreased antitumor activity, and five-fold reduced ferroptosis



selectivity, compared to that of RSL3. These data emphasized the complexity of the effect of ferrocene on these three classical ferroptosis inducers in terms of their antitumoral activities and ferroptosis selectivity.

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Table 2. Antitumor activities and ferroptosis selectivity of selected compounds on other cell lines: human renal cell carcinoma (OS-RC2), triple negative mouse breast cancer (4T1), normal liver cells (L-02) and normal breast cells (MCF-10A).

Comp.	OS-RC2, IC ₅₀ (μM) ^a		Selectivity ^b	GPX4 Inhibition(%) at 1.0 μM	4T1 IC ₅₀ (μM)	L02 IC ₅₀ (μM)	MCF-10A IC ₅₀ (μM)
	-fer-1	+fer-1					
12	0.055±0.017	10.16±0.28	185	25.40±4.24	0.634±0.074	4.98±0.87	NT
13	0.027±0.003	2.42±0.70	89.6	42.37±1.93	0.192±0.008	14.87±5.78	7.81±0.49
15	0.023±0.005	7.687±0.030	334.2	34.02±4.94	0.169±0.008	15.38±4.35	7.79±0.44
17	0.009±0.001	6.917±1.211	768.6	49.16±4.60	0.095±0.006	3.54±0.56	7.32±0.30
18	4.805±0.364	4.281±0.275	0.9	17.88±1.03	NT	4.04±0.79	NT
19	6.584±0.442	4.023±0.307	0.6	12.80±2.67	NT	9.98±2.69	NT
20	26.320±0.930	25.560±0.515	0.9	17.60±11.13	12.930	15.76±2.78	NT
21	0.028±0.003	2.909±0.589	103.9	15.74±0.34	2.63±0.6	4.27±1.55	6.77±0.11
ML162	0.021±0.009	2.574±0.49	122.3	45.02±1.40	0.516±0.059	1.20±0.26	2.93±0.12
RSL3	0.024±0.004	0.69±0.32	28.8	46.14±5.44	0.34±0.062	0.89±0.13	16.93±1.40

(a) Values are expressed as the mean of three independent experiments ± SD); (b) Selectivity: OS-RC2 IC₅₀ with fer-1/ OS-RC2 IC₅₀ without fer-1.

Inspired by the above results, we next attempted to evaluate the cytotoxicity effects of certain molecules on more cancers including ferroptosis-susceptible human renal cell carcinoma (OS-RC2), triple negative mouse breast cancer (TNBC 4T1), as well as two non-tumorigenic cell lines (L02 and MCF-10A), as shown in Table 2. OS-RC2 cells display high vulnerability to ferroptosis and are widely used to evaluate the specificity of compounds to ferroptosis, and the inhibitory activities of compounds against the GPX4 enzyme were evaluated on this cell line at 1 μM. Gratifyingly, ferrocene-appended compounds **15** and **17** exhibited outstanding ferroptosis selectivity and potent inhibition on GPX4 function, that was validated upon the exceptional reversal of cytotoxicity by ferroptosis inhibitor. These data were in line with their antitumoral behaviors on the aforementioned ferroptosis-sensitive HT1080 cells. Its interesting to see that compounds **12** and **13**, each with a ferrocene group at



the R₄ position on the ML162 scaffold, bring about more vulnerability on OS-RC2 cells than HT1080 cells, and these two compounds also possess certain inhibitory activity on the GPX4 enzyme. It indicated that the growth of OS-RC2 may be more dependent on the function of GPX4. This phenomenon has also been observed for the ferrocene-appended RSL analogue **21** which has remarkable ferroptosis-inducing ability on OS-RC2 with a selectivity of 103.9 superior to that of HT1080. An obvious decrease in the cytotoxic activity was observed in the case of active derivatives on TNBC 4T1 cells, which indicated the distinctive damage wrought by ferroptosis on multiple cancer cells. On the other hand, compounds **15** and **17** display weaker cytotoxicity on human normal cells of L02 and MCF-10A compared to that of ML162, and this criterion indicates that they may be good candidates for drug development.

To comprehensively understand the synergistic effects between ferrocene and GPX4 inhibitors in these dual-function inducers, we then explored whether a simple combination of ferrocene compound and GPX4 inhibitors could enhance ferroptosis induction and antitumor effects. As shown in Figure S10, the combination of ferrocene with ML162 only resulted identical inhibition compared with ML162 alone on HT1080 cells; those antitumor effects are obviously weaker than the ferrocene-appended GPX4 inhibitor, **17**. These results acknowledged the synergies of ferrocene and GPX4 inhibitors to exert stronger ferroptosis inducing activities. In addition, ferrocenyl groups have been recognised as important hydrophobic organometallic units to improve the penetration of cell membrane in terms of biologically active compounds,⁵⁶ the improvement of this effect is of interest to investigate for the compounds herein. The cellular uptake efficiency of ML162 and **17** were examined by LC-MS/MS method (Figure S11), which demonstrates that both complexes accumulate rapidly within cells on a time scale of hours. The internalization of **17** is significantly fast compared with ML162 and leads to higher concentration inside of cells. As a consequence, the higher cytotoxicity of **17** may at least partly, emerge from its high cellular accumulation due to the lipophilic ferrocenyl group.

Cell death and EMT process studies

To investigate the ferroptosis selectivity of ferrocene-appended GPX4 inhibitors, their modes of cell death were explored by employing the apoptosis inhibitor Z-VAD-FMK, the necrosis inhibitor necrostatin-1 (Nec-1), the autophagy inhibitor Wortmannin and the typical ferroptosis inhibitors fer-1 and deferoxamine (DFO). As shown in Figure 3A, the percentage of cell viability of OS-RC2 upon combining the ferrocene-appended molecule **17** and the apoptosis inhibitor Z-VAD-FMK remained unchanged, indicating that **17** could not induce



cell apoptosis at current concentration. Moreover, the co-treatment of **17** with Nec-1 and Wortmannin, respectively, did not significantly thwart the cellular death of OS-RC2 induced by these two compounds. It demonstrated that the cellular death caused by **17** may not be attributable to necrosis and autophagy. On the contrary, when combined with typical ferroptosis inhibitors fer-1 and DFO, the cytotoxicity of **17** was obviously reversed. Of particular note are the higher EC₅₀ values of ferroptosis inhibitors to rescue the cellular death posed by **17** compared to that of ML162 (Table S10); these observations confirmed that the potent antiproliferative activities of **17** were achieved mainly by inducing ferroptosis of cancer cells, and also underscored its remarkable ferroptosis selectivity.

EMT is connected to carcinogenesis, invasiveness, metastasis, and therapeutic resistance in cancer. Growing evidence shows that cancer cells undergoing EMT are vulnerable to ferroptotic cell death.¹¹⁻¹² Thus, the effect of the ferroptosis-inducing agent **17** on the migration and invasion of cancer cells was examined. Through a wound-healing assay, it was found that those compounds significantly prevented the migration of 4T1 cells time-dependently, especially in the case of **17** (Figure S11). Similarly, transwell assays revealed that this molecule disrupts the invasion of tumor cells dose-dependently (Figure 3B). These results indicated the superior capacity of ferrocene-appended ferroptosis inducers to block the migratory and invasive properties of 4T1 cells, denoting their enormous promise for the treatment of refractory cancer metastasis, taking into account that TNBC 4T1 cells have high invasiveness, and approximately half of TNBC cases will have distant metastasis.⁵⁷

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DOI: 10.1039/C4CY02002B



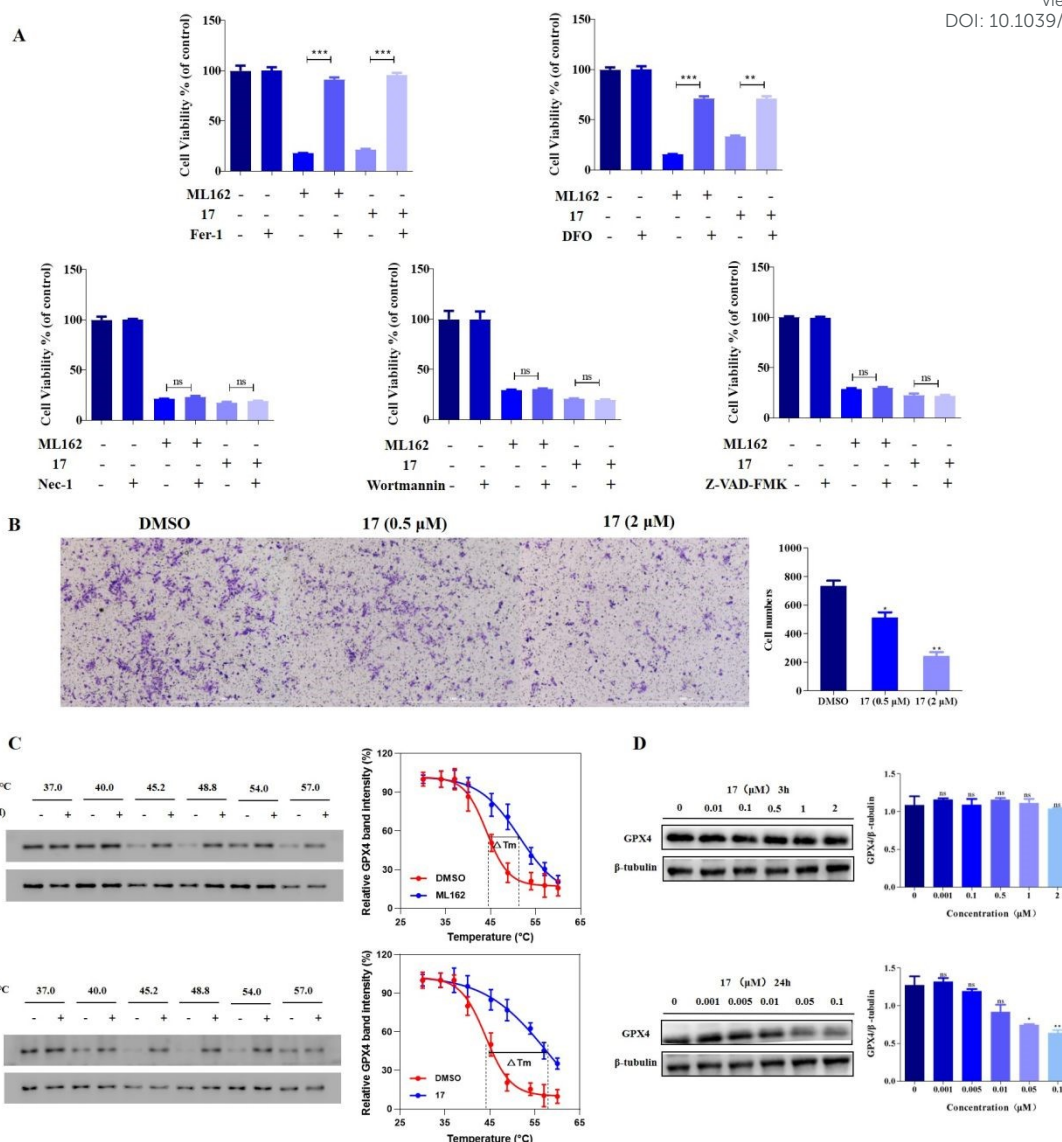


Figure 3. A) The percentage of cell viability after treatment of ML162 or **17** (0.4 μM) with Z-VAD-FMK (10 μM), Nec-1 (10 μM), Wortmannin (30 μM), Fer-1 (2 μM), and DFO (30 μM), analysis results represented the mean ± SD. B) Transwell assays were used to detect the invasion of cancer cells after treatment of **17** (0.5 and 2 μM). C) CETSA experiments were performed on OS-RC-2 cells treated with DMSO, ML162 and **17** (5 μM) for 4 h. The protein levels were analyzed by western blotting after heating at different temperatures (30–60 °C), the apparent T_{agg} values for GPX4 in OS-RC-2 cells in the absence and existence of ML162 and **17** were 44.09 °C to 51.42 °C, and 44.05 to 58.00 °C, respectively. D) Western blot analysis of GPX4 levels in OS-RC-2 cells following treatment of **17** in 3 and 24 h. * $P < 0.05$, ** $P < 0.01$, and *** $P < 0.001$ versus the control groups.

GPX4 CETSA and western blotting evaluations

Since certain ferrocene-appended molecules exhibited high selectivity for ferroptosis and remarkable GPX4 inhibition activities, the effect of **17** on intact GPX protein was further determined, and the results are shown in Figures 3C and 3D. The ability of **17** for binding GPX4 was first validated by the cellular thermal shift assay (CETSA) considering the



stability of the protein is changed when it is bound with the molecule. It was found that the aggregation temperature (T_{agg}) of GPX4 was improved by ~ 14 °C in OS-RC2 cells after treatment of **17**, whereas the T_{agg} in the case of ML162 increased by ~ 7 °C, indicating the stronger interaction of **17** with GPX4 in the cells. The increased thermal stability on GPX4 is in line with that of ML210 and other alkyne analogues on the stabilization of GPX4 reported previously,⁵⁴ suggesting that they may bind to GPX4 in a similar way. Meanwhile, the expression level of GPX4 was further investigated by western blot, and it is noteworthy that there was no obvious difference in the protein level of GPX4 compared to that of the control group within 3 h of **17** treatment (Figure 3D). However, the down-regulation of GPX4 was evident after 24 h treatment. One could speculate that the potent ferroptosis-inducing ability of ferrocene-based ML162 analogues may result from both the inhibition and downgrading of GPX4 in the long-term cellular incubation. These results accentuate the complex effects of covalent inhibitors to GPX4.

LPO related bioactivity evaluations

It is well-known that certain ferrocene complexes can induce the formation of ROS via a Fenton type reaction. Therefore, the ability of ferrocene-appended GPX4 inhibitor **17** to yield ROS was first evaluated in physiological conditions by the widely used fluorescent ROS indicator, 2',7'-dichlorofluorescein diacetate (DCFH-DA). As shown in Figure 4A, a known H₂O₂-induced activation method was carried out to monitor the ROS generation by selected molecules in buffer,⁴⁷ containing FeSO₄ as a positive control. It is evident that **17** can facilitate the conversion of H₂O₂ to OH• and increase the production of ROS in a time-dependent manner, reflected by the OH•-mediated oxidation of nonfluorescent 2',7'-dichlorofluorescein (DCFH) to fluorescent 2',7'-dichlorofluorescein (DCF). In contrast, the pure organic compound ML162 did not exhibit obvious catalytic efficiency for yielding ROS. Unrestricted lipid peroxidation is a hallmark of ferroptosis, which can be evaluated in cells by multiple approaches. As a metabolite of LPOs, the level of MDA is commonly used as a characteristic index for the determination of LPOs. As illustrated in Figure 4B, **17** improved the contents of MDA in cells dose-dependently and more prominently than that of ML162 at the same concentration, which corresponds to the phenomenon observed at the molecular level.



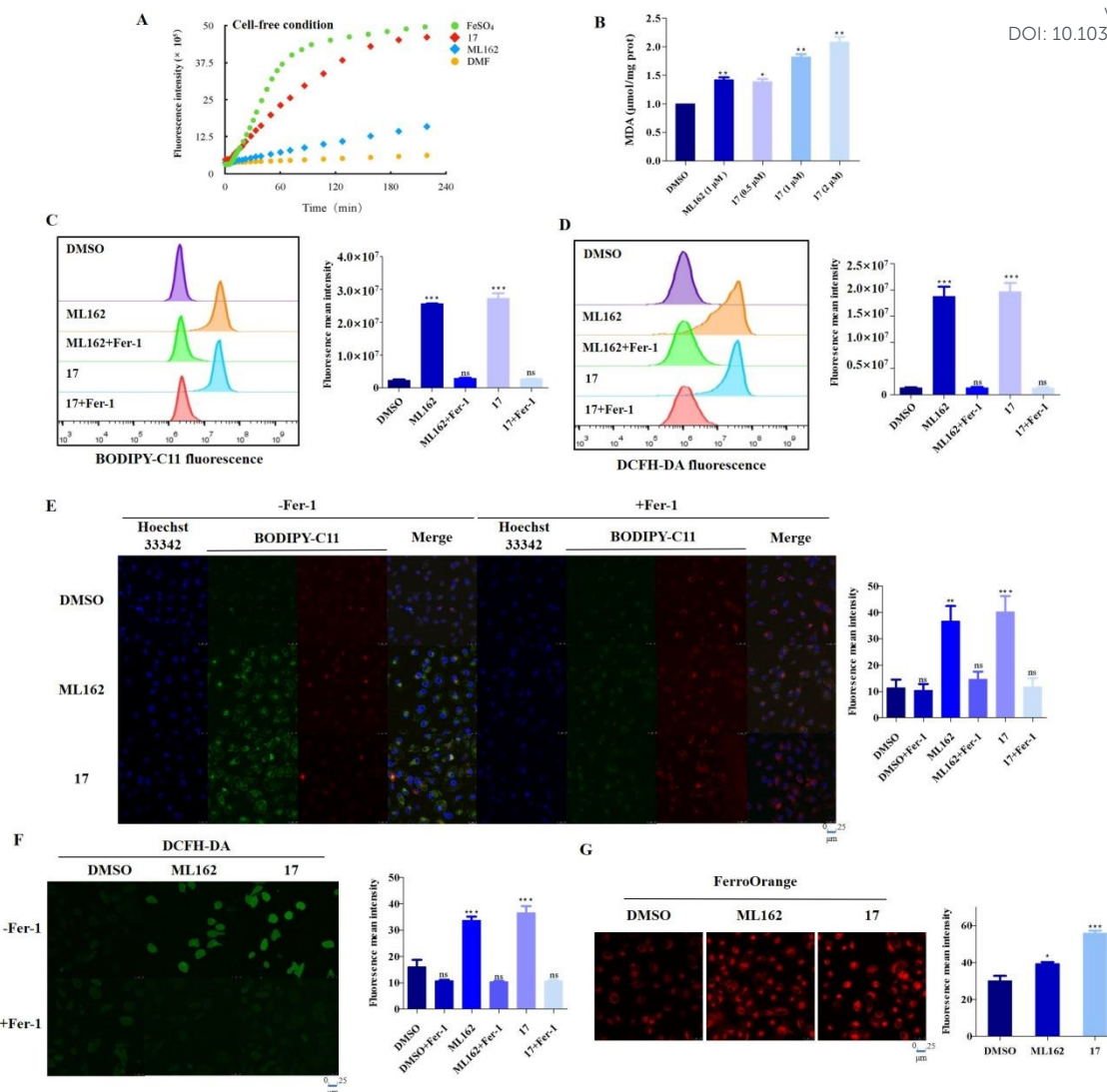


Figure 4. A) Monitoring of the generation of ROS of corresponding compounds (100 μM) using fluorescence spectroscopy in combination with ROS-sensitive DCFH-DA ($\lambda_{\text{ex}} = 501 \text{ nm}$, $\lambda_{\text{em}} = 531 \text{ nm}$). B) OS-RC2 cells were treated with ML162 (1 μM), and **17** (0.5, 1, and 2 μM) for 4 h, and then the content of MDA was determined. C) Flow cytometry analysis for intracellular LPO by C11-BODIPY of OS-RC-2 cells treated with DMSO, ML162 and **17** (0.5 μM) with and without fer-1 (1.5 μM) respectively. D) Flow cytometry analysis for intracellular ROS by DCFH-DA of OS-RC-2 cells treated with DMSO, ML162 and **17** (0.5 μM) with and without fer-1 (1.5 μM) respectively. E) Representative confocal images and analysis of C11-BODIPY staining of OS-RC-2 cells after incubation with ML162 and **17** (0.5 μM) with and without fer-1 (1.5 μM) respectively. F) Representative confocal images and analysis of DCFH-DA staining of OS-RC-2 cells after incubation with ML162 and **17** (0.5 μM) with and without fer-1 (1.5 μM) respectively. G) Intracellular Fe^{2+} was visualized with the FerroOrange fluorescence probe. OS-RC-2 cells were treated with ML162 and **17** (0.5 μM) for 4 h, and the content of intracellular Fe^{2+} was determined. * $P < 0.05$, ** $P < 0.01$, and *** $P < 0.001$ versus the control groups.

Accordingly, the fluorescent probes C11-BODIPY and DCFH-DA were employed to investigate the intracellular ROS and LPOs via flow cytometry and confocal laser scanning microscopy (CLSM). As illustrated in Figure 4C-4D, **17** could significantly induce the



accumulation of LPOs slightly superior to that of ML162 and this trend could be reversed by fer-1 totally according to the flow cytometry analysis, which emphasized its remarkable ferroptosis selectivity. In agreement with this, the oxidative fluorescence intensity (green) of C11-BODIPY and DCFH-DA in OS-RC2 cells significantly increased upon treatment with **17** (Figure 4E-4F). Considering the iron dependence which can make cancer cells more vulnerable to ferroptosis, the overall level of cellular ferrous iron was detected by Fe²⁺-sensitive FerroOrange staining. As shown in Figure 4G, the images of FerroOrange staining intuitively illustrated that a notable fluorescent intensity of the cells after treatment with **17** is found to be significantly higher than that of cells incubated with ML162, suggesting that **17** can increase the level of cellular ferrous iron. Overall, these results verified by the increasing of LPOs and Fe²⁺, lead to a consistent conclusion that ferrocene-appended GPX4 inhibitors could significantly induce the accumulation of LPOs and subsequently engender severe ferroptosis of cancer cells.

Molecular Docking

The molecular simulation of small molecular ligands with GPX4 has been challenging due to the flat surface surrounding the active site of GPX4,⁵⁸ thus a template-based docking by Watvina was carried out. According to the atomic types and their contribution to ligand-receptor interaction, we first generate a template-containing mapping of heavy atoms, aromatic atoms and hydrogen bond receptors (Figure S12). Using the genetic algorithm for global searching and BFGS for local refinement in Watvina, the conformation with the highest template matching score of docked ferrocene derivatives was used for further analysis. As shown in Figure 5, the selected ligands **12**, **13**, **15** and **17** docked well to the ML162 space with paired atoms RMSD less than 0.5 Å. Those four organometallics enabled the formation of two key hydrogen bonds with residues of GPX4 like that of ML162. One of the cyclopentadienyl rings of **15** or **17** occupied the same hydrophobic space as that of ML162, and the resulting conformation of **17** revealed that it filled the active pocket better than compound **15**. In the case of **12** and **13**, the ferrocene groups are placed in lieu of the benzene (Ring C) part in ML162 (Figure 2B, 5C and 5D). Since ring C occupied by ferrocene is flexible and distal from the binding pocket, **12** and **13** may exhibit weak binding activity with GPX4. As for compounds **10** and **11** with the ferrocenyl group sited at the very cramped ring B part (Figure 2B), they were expected to lose the capacity of conjugating GPX4. Indeed, these docking data, which held the binding mode of covalent inhibitors with GPX4, concur well with their GPX4 inhibitory activities and thus provided a reasonable interpretation.



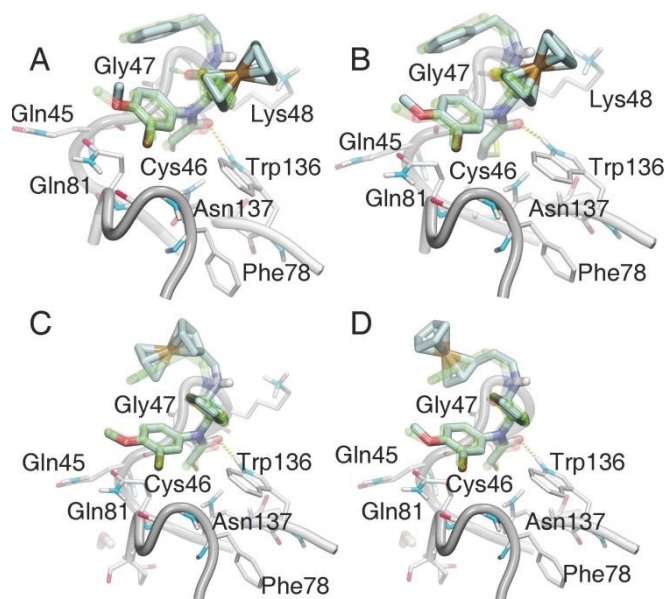


Figure 5. The binding models of ferrocenyl ligands in the GPX4 crystal structure (PDB code 6HKQ). Compounds **15** (A), **17** (B), **12** (C), **13** (D) were docked to GPX4 by Watvina with template-based method, then all the structures were minimized with xtb program. The original ligand ML162 was shown in green transparent stick mode, while ferrocenyl ligands in cyan stick mode. The hydrogen bonds were represented with yellow dashed line by VMD software.

In vivo profiling of ferrocene-based ferroptosis inducers

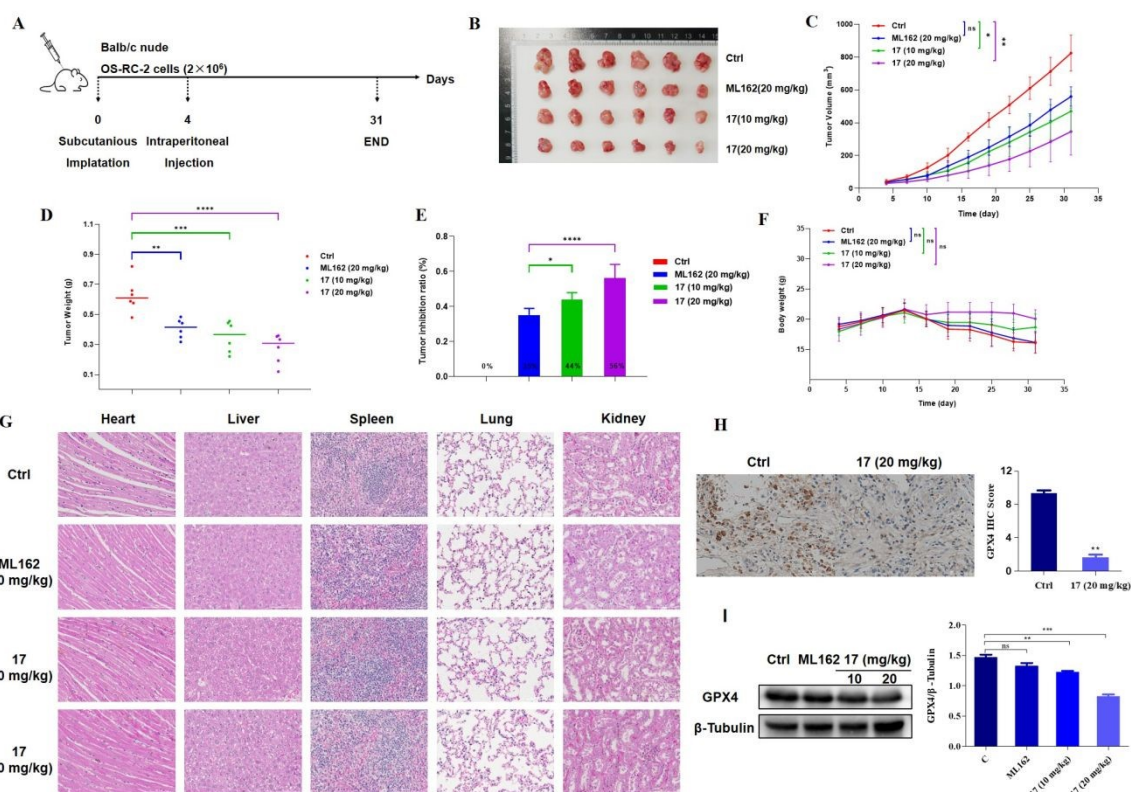


Figure 6. A) *In vivo* antitumor activity of **ML162** and **17** in a human OS-RC2 renal carcinoma model (BALB/c nude mice, given a medicine every three days, n = 6). B) Picture



of excised tumors from the different groups after the treatment; C) Tumor volume curves during the treatment period; D) Tumor weights after treatment; E) Tumor inhibition ratio after the treatment period. F) Body weight curves of different groups after the treatment; G) H&E staining of the major organs of mice after treatment with compounds. H) anti-GPX4 immunohistochemistry images and analysis from the OS-RC-2 xenograft model. I) Western blot analysis of GPX4 levels in tumor tissue following treatment of corresponding compounds. *P < 0.05, **P < 0.01, ***P < 0.001 and ****P < 0.0001 versus the control groups.

Considering the excellent antitumor activities of **17** on two ferroptosis sensitive cell lines, the OS-RC2 xenograft model was established in *BALB/C* nude mice to assess its *in vivo* efficacy. The tested compounds were administered by intraperitoneal injection every three days for approximately 4 weeks (Figure 6). The tumor growth inhibition (TGI) values of **17**-treated groups were 44%@10 mg/kg and 56%@20 mg/kg, which showed that **17** could suppress the progression of the renal tumor with a significant dose-dependent effect. It is noteworthy that **17** displayed prominent antitumor activities *in vivo* compared to ML162 in the same case of delivering a high dose of drugs. Additionally, there was no obvious difference in body weight of the **17** treatment groups compared with the control group. Furthermore, hematoxylin and eosin (H&E) staining analysis of the major organs were examined to investigate the safety profile of ML162 and **17**, which indicated that there was no apparent toxic damage observed in major organs such as the heart, liver, spleen, lung, and kidney. Subsequent immunohistochemical analysis and western blot studies showed that **17** evidently reduced the level of GPX4 protein in tumor tissues, revealing that ferrocenyl complex **17** effectively induced ferroptosis *in vivo*. These results emphasized the superior potential of ferrocene-based ferroptosis inducers as anticancer drug candidates for further research.

RNA-seq analysis

To further illuminate the impact of **17** on the gene expression pattern, we employed the NovaSeq 6000 PE150 platform (Illumina Inc., CA, USA) to analyze gene expression differences (DEGs) between the extracts of tumor tissues from OS-RC-2 xenograft model rats treated with **17** (20 mg/kg) and normal control rats. A volcano plot illustrated significant differences in mRNA expression patterns between the two groups (Figure 7A and Table S13). Utilizing the Limma package, we identified differential genes under the condition: p-value < 0.05 & |logFC| > 1, resulting in 790 differentially expressed genes (324 up-regulated and 466 down-regulated).



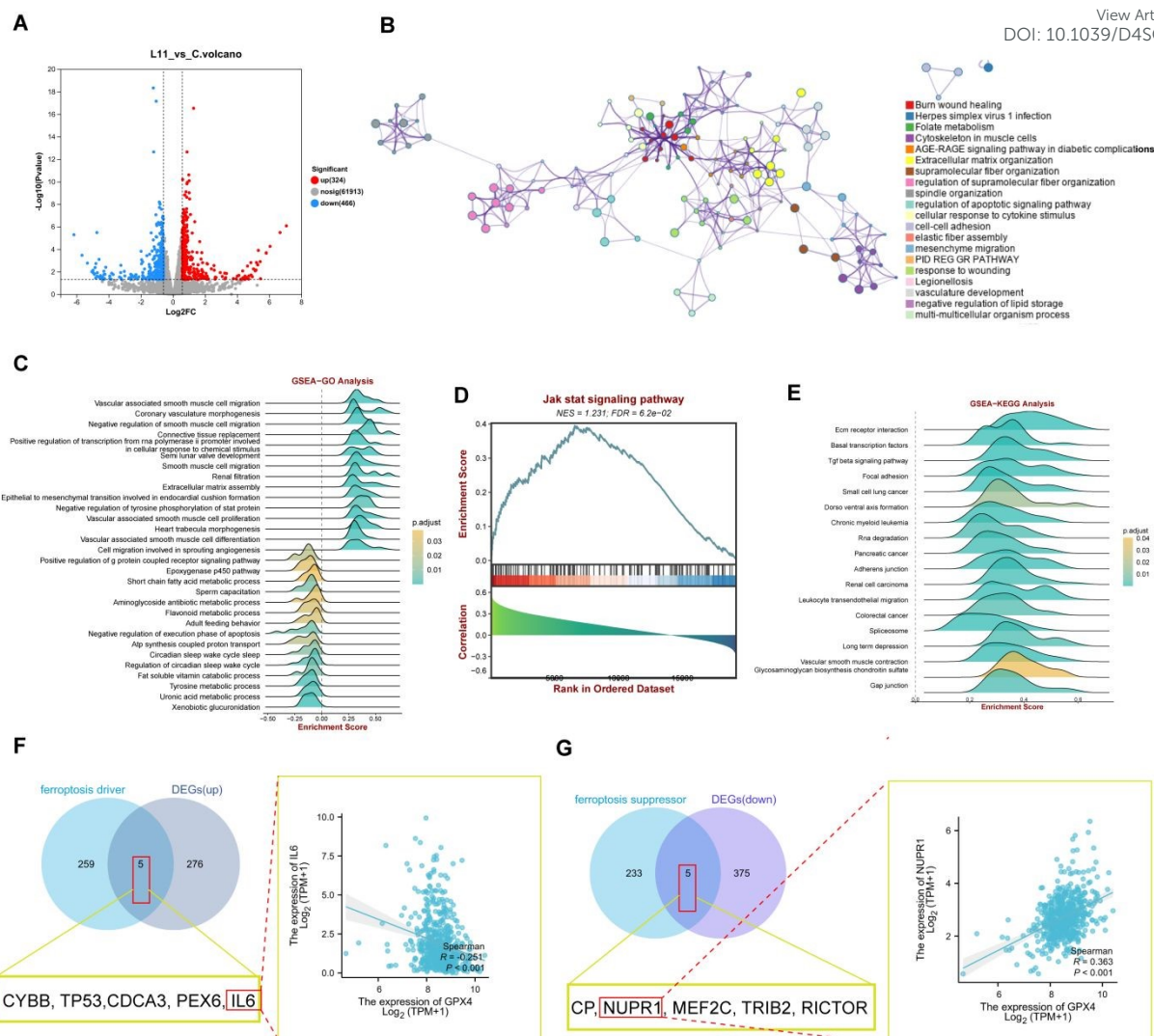


Figure 7. RNA-seq analysis of samples from OS-RC-2 xenograft model rats treated with **17** (20 mg/kg) and control. A) Volcano plot of all DEGs, red indicates upregulation, whereas blue indicates downregulation. B) Protein-protein interaction (PPI) networks constructed by Metascape and the protein function were enriched. C) GO term enrichment analysis of DEGs from **17** (20 mg/kg) treatment group. D) GSEA analysis of DEGs in JAK-STAT signaling pathway. E) KEGG enrichment analysis of DEGs from **17** (20 mg/kg) treatment group. F, G) The genes as the driver and suppressor of ferroptosis overlap with DEGs that are up-regulated and down-regulated by Venn diagram.

Subsequently, we constructed a PPI network of the 790 DEGs using Metascape. This analysis revealed enrichment in ferroptosis-related signaling pathways and cell phenotypes, including folate metabolism, AGE-RAGE signaling pathway in diabetic composites, negative regulation of lipid storage, and mesenchymal migration (Figure 7B). Further insight into the biological functions and signaling pathways associated with ferroptosis was gained through GO and KEGG analyses (Figure 7C-7E). These analyses demonstrated the involvement of DEGs in multiple biological processes related to iron homeostasis, such as cyclooxygenase and G protein-coupled receptor signaling pathways. Notably, the activation of multiple inflammation-related signaling pathways leading to ferroptosis was observed, with DEGs



enriching in the inflammatory signaling pathway Jak-stat, as indicated by GSEA analysis (Figure 7D). KEGG analysis also highlighted involvement in pathways related to iron and lipid metabolism, including short-chain fatty acid metabolism and tyrosine metabolism (Figure 7E). To explore the correlation of **17** with ferroptosis-related pathways, we examined the overlapping genes between ferroptosis driver and DEGs, or ferroptosis suppressor genes and DEGs after **17** treatment. Results revealed that **17** upregulated ferroptosis driver genes (CYBB, TP53, CDCA3, PEX6, IL6) and downregulated suppressor genes (CP, NUPR1, MEF2C, TRIB2, RICTOR, including GPX4). The correlation analysis showed a significant negative correlation between the driving gene IL6 and GPX4 (Figure 7F) and a significant positive correlation with UPR1 and GPX4 (Figure 7G), indicating the targeted inhibition of GPX4 function by **17** to induce ferroptosis in renal cell carcinoma *in vivo*.

Conclusions

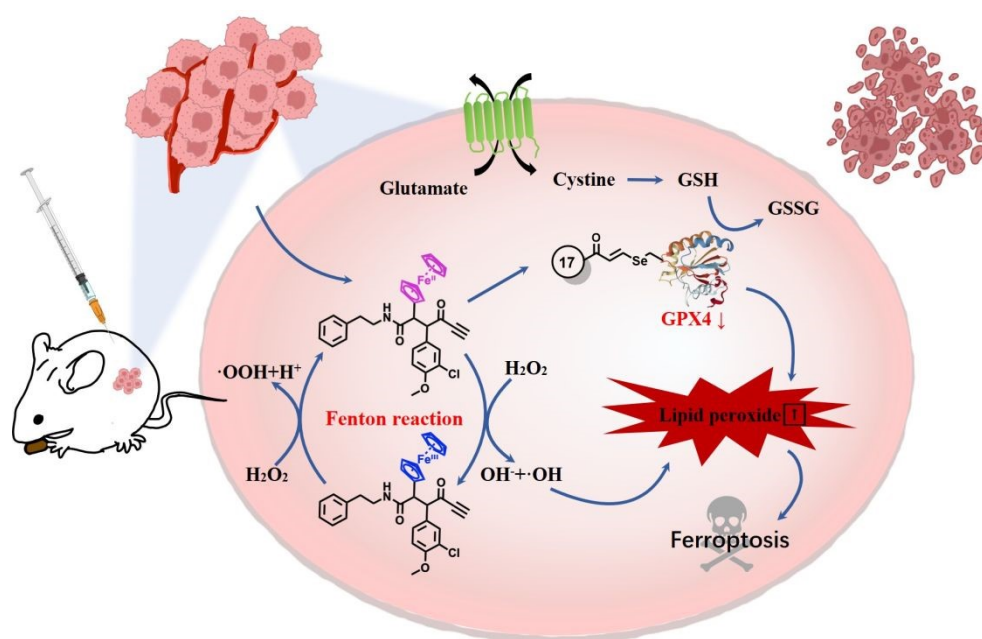
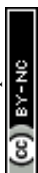


Figure 8. Proposed mechanisms of ferrocene-appended ferroptosis inducers.

According to the ferroptosis-based phenotypic screening of our in-house organometallic compound library, the usual ferrocene complexes did not possess visible ferroptosis-inducing activities, which contradicted the conventional understanding of ferrocene for ferroptosis. As a result, a series of ferrocene-appended GPX4 inhibitors has been rationally designed by a one-stone-two-birds strategy for the first time. Distinct from the known bioferrocene species, only using the ferrocenyl group as an auxiliary fragment to existing organic moieties, or merely taking advantage of the redox behaviors of ferrous iron to generate ROS, complex **17** can both inhibit GPX4 effectively and catalyze the Fenton-like reaction to yield ROS, and



consequently result in stronger antiproliferative activity and higher ferroptosis selectivity than the corresponding organic compound ML162 (Figure 8). Therefore, as a proof-of-concept study, we successfully achieved the goal to use a single molecule to modulate the two main pathways of ferroptosis, iron homeostasis and the antioxidant defense system. In particular, cellular CETSA experiments confirmed the strong interaction of **17** with intact GPX4 protein, and this result was in accord with its capability to disrupt the function of GPX4 in OS-RC2 cells. Moreover, the significant ROS-inducing ability of **17** at both the molecular level and at the cellular level acknowledged the ferroptosis enhancement of the ferrocene-appended GPX4 inhibitor. Subsequent *in vivo* experiments demonstrated that **17** is a potent attenuator of the growth of renal tumors and, and ferroptosis is pronounced to be involved in this inhibition process, supported by the GPX4 content analysis and RNA-seq study of tumor tissues. In conclusion, we present here a comprehensive study of rationally designed ferrocene-appended ferroptosis inducers, not only to promote the development of ferroptosis-based anticancer treatments, but also to provide new insight into the design of novel bioorganometallics based on ferrocene for potential disease therapies.

Supplementary Information. Experimental procedures for syntheses and biological evaluation, supplementary Figures, Schemes and Tables, computational methods and details, cif files and X-ray data of **12**, **15** and **17**, these material is available free of charge via the Internet at [xxx](#).

Author contributions

W. L., J. Y., J. W. and X. F. contributed equally to this manuscript. Y. W. and J. L. designed the study. W. L., J. W., H. W. and Q. Z. performed the synthesis and characterization of chemical compounds. J. Y., X. F., Y. X., X. Z. and X. Qi performed the *in vitro* and *in vivo* anticancer evaluation of the compounds. X. X. performed the docking calculation work. X. L. and X. Qiu contributed to the target protein binding affinity evaluation. P. P., Q. G., G. J. and S. Y. contributed to the construction of ferrocene-based compounds library. J. B-C and M. J. M. prepared the crystal figure. Y. W., J. L. and X. Qiu supervised the study. Y. W., J. L. and M. J. M. wrote the manuscript. All authors participated in valuable discussion and manuscript correction.

Declaration of competing interest

Y. W., J. L., J. W., W. L., H. W. and X. Z. are inventors on a patent application related to this work.



Acknowledgements

View Article Online
DOI: 10.1039/D4SC02002B

Y. W. received funding from Qingdao Marine Science and Technology (2022QNLM030003-2), Taishan Scholar Youth Expert Program in Shandong Province (tsqn202103035) and Distinguished Young Scholars of Shandong Provincial Natural Science Foundation (Overseas, 2022HWYQ-069), X. Q. thanks the support of the Fundamental Research Funds for the Central Universities of Ocean University of China (No. 202262015).

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