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Design, synthesis and biological evaluation of naphthalene-1,4-dione analogues as anticancer agents†

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The increased metabolism of glucose *via* aerobic glycolysis, known as the Warburg effect, is a hallmark of most cancers. Identifying molecules that disrupt the Warburg effect may allow for selective cytotoxicity towards cancer cells and reduce side effects compared to current chemotherapy agents. Our initial hit compound, **BH10**, which potentially targets Kelch-like ECH-associated protein 1 (Keap1), increased oxygen consumption rate and displayed increased cytotoxicity towards cancer cells over normal cells *in vitro*. In this project, a library of analogues based on the **BH10** scaffold was prepared with the aim of improving potency and cancer-cell specificity. Among these analogues, several compounds showed notable potency, with activity (IC₅₀) observed around 1 μM. However, when considering selectivity, the imidazole derivative, compound **44**, exhibited the most optimal balance, achieving an IC₅₀ of 6.4 μM and selectivity ratio of 3.6 which indicates greater toxicity to cancer cells vs. normal cells.

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1. Introduction

A major drawback of many cancer therapies is their non-selective toxicity to healthy cells and tissues.^{1,2} The development of new drugs with better cancer-specific cytotoxicity would provide better treatment options and quality of life to cancer patients. A potential way to improve the selectivity of therapeutics is to target a pathway that is altered in cancer cells. One such potential target is glucose metabolism, which Hanahan and Weinberg described as a new ‘hallmark of cancer’.³ In healthy cells, glucose metabolism begins with glucose transporters (GLUTs) moving glucose from outside the cell and into the cytoplasm. Once glucose molecules are inside the cell, each glucose molecule is metabolised into two pyruvate molecules *via* glycolysis. Pyruvate can then be metabolised by one of two pathways depending on oxygen availability. If oxygen is present, pyruvate enters the mitochondria where it is metabolised through the tricarboxylic acid (TCA) cycle to produce reducing equivalents that feed electrons into the electron transport chain. This is required for the process of oxidative

phosphorylation (aerobic respiration), which generates approximately 36 ATP molecules per glucose molecule.⁴ In contrast, when oxygen is absent, pyruvate remains in the cytoplasm where it is converted to lactate *via* anaerobic respiration, yielding only 2 ATP molecules per glucose molecule. Hence, aerobic respiration is far more efficient for cellular energy production.⁴ In cancer cells, however, glucose metabolism is altered significantly.³ Cancer cells primarily metabolise glucose *via* glycolysis and convert pyruvate to lactate even in the presence of oxygen, and this metabolic preference is known as the “Warburg effect”. Although it appears to be an inefficient means of producing cellular energy (only 4 ATP molecules produced per molecule of glucose), it can provide a selective advantage for cancer cells by multiple mechanisms (Fig. 1a). For example, the replenishment of NAD when pyruvate is converted to lactate is essential for rapidly dividing cells.^{4–6}

We recently developed a screening platform to identify drug-like compounds that were more toxic to cancer cells than normal cells. From this screen, we identified a quinone molecule, 2-chloro-3-[(2-morpholin-4-ylethyl)amino] naphthalene-1,4-dione (referred to as **BH10** (Fig. 1b)) that showed greater cytotoxicity towards cancer cells.⁷ Furthermore, **BH10** alters cellular glucose metabolism and increases cellular oxygen consumption rate (OCR) by targeting mitochondrial redox defence in cancer cells. This alteration increases glucose oxidation and inhibits glycolysis, which in turns induces cancer cells necrosis.⁷ We also identified Kelch-like ECH-

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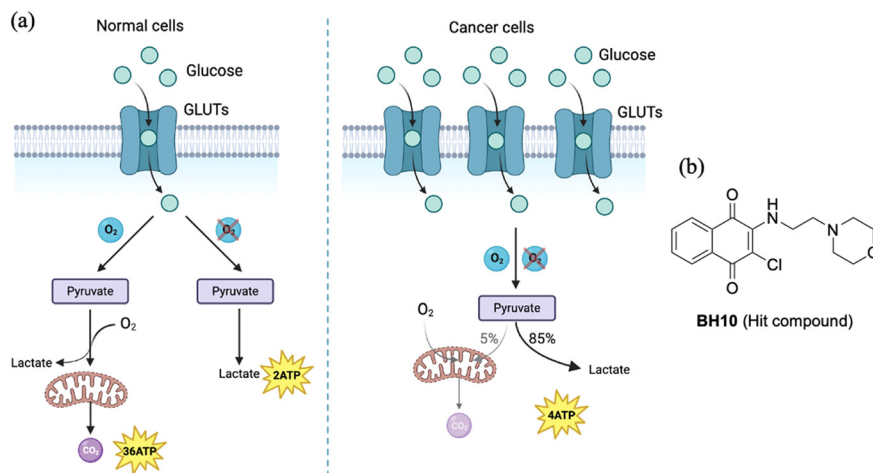


Fig. 1 (a) Glucose metabolism in differentiated tissues: glucose is metabolized through oxidative phosphorylation or anaerobic glycolysis. In cancerous tissues, glucose is predominantly metabolized through aerobic glycolysis (Warburg effect), resulting primarily in lactate fermentation.⁴ (b) The chemical structure of **BH10**.

associated protein 1 (Keap1) as a potential protein target of **BH10** via avidin pull-down assays with biotinylated-**BH10**.⁸ Keap1 is a critical regulator of cellular defence against oxidative stress.⁹

BH10 is a 1,4-naphthoquinone, and this structure is important in medicinal chemistry. Furthermore, the quinone scaffold is present in many frontline chemotherapy drugs.^{10,11} One of the best-known examples of quinones that has been used to treat cancer is doxorubicin.¹¹ Importantly, our study demonstrated that **BH10** had better cancer cell-specific cytotoxicity than 7 other well-known anticancer quinones including doxorubicin, CDC25 phosphatase inhibitor II, vitamin K3, mitomycin C, β -lapachone, tanespimycin (17-AAG) and

streptonigrin, even though these compounds show higher potency (Fig. 2).⁷ Thus, **BH10** represents a lead molecule for future drug development.

In this study, various functional groups were introduced to the 1,4-naphthoquinone scaffold to replace the morpholine chain and aimed to increase the scaffold's rigidity by modifying the 1,4-naphthoquinone into a naphthoimidazole. Therefore, the objectives of this study were to design, synthesize and screen **BH10**-like molecules to identify those that are more potent than **BH10** and have improved selectivity compared to **BH10**. Specifically, the comprehensive structure-activity relationship, and the potential anti-cancer mechanism as well as the pharmacokinetic profile of the best drug candidates were explored in this study.

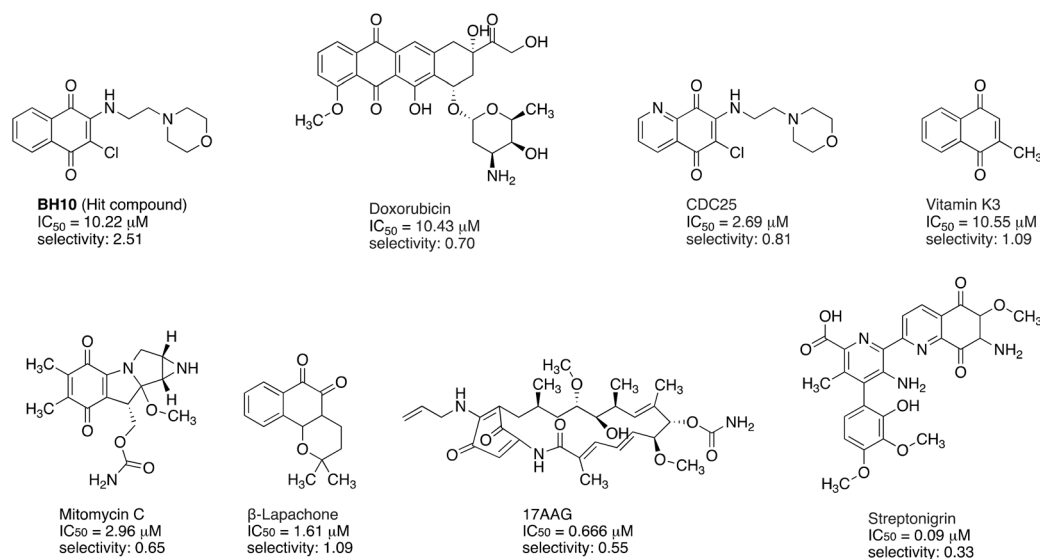


Fig. 2 Chemical structure and biactivity of **BH10** and seven other well-known anticancer compounds. IC₅₀ values (μM) indicate the cytotoxicity towards HEC1A endometrial cancer cells, while selectivity is calculated as the IC₅₀ value for non-cancerous MAD11 cells divided by the IC₅₀ value for HEC1A cells.⁷



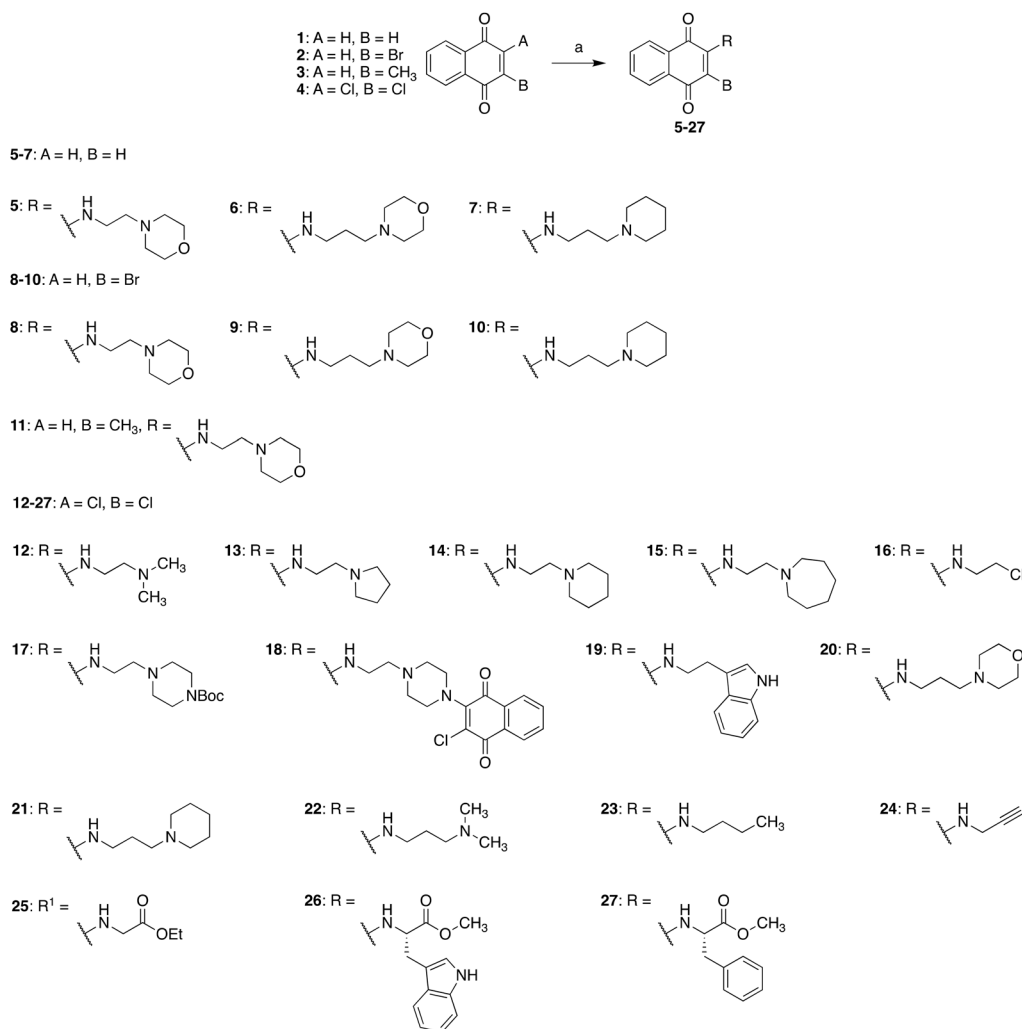
2. Result and discussion

2.1. Rational design and chemical synthesis

Firstly, 3-unsubstituted naphthoquinones 5–7 were synthesized to investigate whether the chlorine atom is necessary for its bioactivity. To achieve this, naphthalene-1,4-dione **1** was reacted with different amines with the presence of Et₃N or K₂CO₃ as the base to accelerate the reaction (Scheme 1). This provided naphthoquinones 5–7 in 25–80% yield. Moreover, the effect of replacing the 3-chloro group with a bromo or methyl group on the bioactivity of the analogues was investigated. The corresponding analogues **8–11** were synthesized *via* the forementioned reaction condition using 2-bromonaphthalene-1,4-dione **2** or 2-methylnaphthalene-1,4-dione **3** as the starting material. Subsequently, the effect of varying the steric nature and the electronic properties of the pendent amine group on biological activity was explored. As the morpholine ring in **BH10** contains a tertiary amine and an ether functional group, altering the oxygen atom in the ring could change the

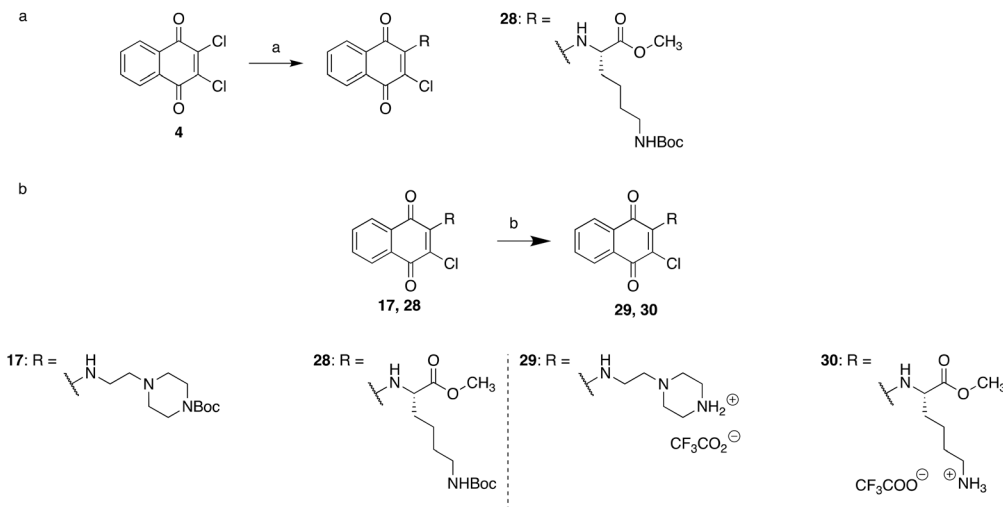
hydrophilic characteristic of the ring, which might provide an insight on the importance of a hydrogen acceptor of hydrogen bonding at that position. Furthermore, modifications to the ring size or replacing the entire ring with other groups could provide information about the steric requirements of the biological target. Hence, derivatives **12–25** were synthesized using the same reaction conditions mentioned above. Finally, to explore the effect of the introduction of an amino acid ester instead of the amine side chain of the hit compound on activity and specificity, tryptophan, and phenylalanine were used in the substitution reaction to obtain analogues **26–27**.

There have been reports in the literature that salt compounds such as ammonium salts possess good anticancer potency, therefore salt products **29** and **30** were synthesized.^{12–14} Using similar conditions in Scheme 1, dichloronaphthoquinone **4** was reacted with *N*-Boc-L-lysine methyl ester hydrochloride to obtain **28** (Scheme 2a). Deprotection of analogues **17** and **28** was achieved using TFA in DCM at room temperature for 3 h to afford final compounds **29** and **30** in 50–60% yields (Scheme 2b).



Scheme 1 Synthesis of nucleophilic substitution products. Reagents and conditions: (a) corresponding amine, Et₃N or K₂CO₃, EtOH or Et₂O, r.t., 18–72 h.

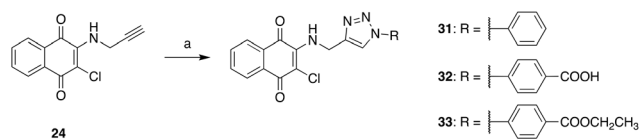




Scheme 2 Synthesis of salt products. Reagents and conditions: (a) corresponding amine, Et₃N, EtOH, r.t., 8 h; (b) 1 : 3 TFA/DCM, r.t., 3 h.

Since the triazole ring is a common heterocycle found in drugs, it is of great interest to explore the effect of introducing the triazole ring into the analogues. Through click chemistry, the alkyne group on **24** was cyclised with appropriate phenyl azides in the presence of CuSO₄·5H₂O and sodium ascorbate as catalysts in THF/H₂O (5 : 1) at 90 °C for 10 h to give triazole analogues **31–33** in 16–89% yield (Scheme 3).

BH10 was previously found to be unstable in the biological body and can be easily metabolised.^{7,8} It was postulated that this was due to the presence of a good Michael addition acceptor in the scaffold of **BH10** as well as the chlorine atom at the 3-position of the molecule acting as a good leaving group. A possible solution to mitigate this limitation could be the protection of the Michael addition acceptor and the conversion of the labile chlorine into a stable amide group. To achieve this, the amide intermediates were first synthesized, following a literature method described by Ho *et al.*, where the chloramine starting material **34** was reacted with excess alkyl anhydride in the presence of concentrated H₂SO₄ at room temperature for 10 h (Scheme 4).¹⁵ The desired amide compounds **35–38** were isolated in moderate to high (50–90%) yields. To expand the diversity of the analogue library, a benzyl-substituted amide was synthesized by reacting the chloramine compound **34** with phenylacetyl chloride in the presence of the Lewis acid boron trifluoride diethyl etherate in toluene at 90 °C for 1–2 h to afford **49** in moderate yield of 65% (Scheme 4).¹⁶ The synthesized chloramide intermediates **35–38** and **49** were



Scheme 3 Synthesis of triazole products: (a) corresponding azido compounds, CuSO₄·5H₂O, sodium ascorbate, THF/H₂O, 90 °C, 10 h.

then reacted with aminopropylpiperidine following a modified literature method by Brun *et al.*, giving the desired products **39–43** and **50** in 30–80% yield.¹⁷

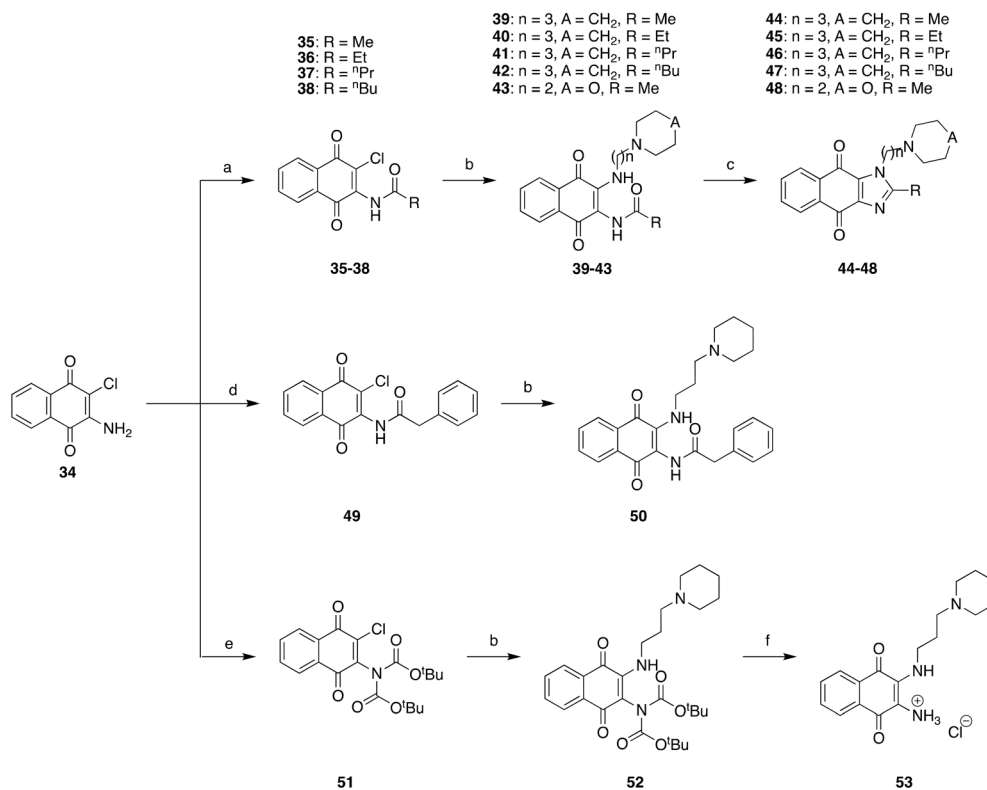
Increasing the rigidity of a molecule could increase both its selectivity and potency, as the compound might bind to fewer targets and the entropic energy cost in binding could be reduced.¹⁸ Therefore, an extra fused ring was introduced into the scaffold to investigate the effect of rigidity on anticancer activity of the analogues. The synthesis of these analogues was achieved by dehydration of amide compounds **39–43** following a literature method described by Ho *et al.*¹⁵ Compounds **39–43** were heated under basic conditions in EtOH for 1–2 h to produce the ring-closed imidazole compounds **44–48** in moderate to high yields of 47–100% (Scheme 4). The effect of replacing the chlorine atom in the 1,4-naphthoquinone with an amino group on the anticancer activity of the compound was also explored and the ammonium salt **53** was synthesized. The primary amino group in compound **34** was first protected using di-*tert*-butyl dicarbonate (Boc₂O) to give compound **51**, which was then reacted with aminopropylpiperidine to afford compound **52**. The deprotection of the Boc-protecting group in compound **52** gave final compound **53** as a primary ammonium chloride salt.

3. *In vitro* cell cytotoxicity and structure–activity relationship study

To evaluate the anticancer potency and selectivity of the synthesized compounds, they were screened for cytotoxicity against human endometrial cancer (HEC1A) cells and noncancerous human endometrial stromal (MAD11) cells using the 3-(4,5-dimethylthiazole-2-yl)-2,5-diphenyltetrazolium bromide (MTT) reduction assay (Table 1). The hit compound **BH10** was used as a control molecule.

Upon testing of compounds **5–11**, it was found that biological activity was abolished once the halogen was





Scheme 4 Synthesis of amide compounds and naphthoquinoneimidazole compounds. Reagents and conditions: (a) corresponding anhydrides, concentrated H₂SO₄, 50 °C, 1–2 h; (b) corresponding amines, Et₃N, THF, r.t., 18 h; (c) 2 M NaOH, EtOH, 50 °C, 1–2 h; (d) phenylacetyl chloride, BF₃·Et₂O, toluene, 90 °C, 7 h; (e) Boc₂O, pyridine, THF, r.t., 24 h; (f) HCl, Et₂O, r.t., 18 h.

removed from the C3-position of the naphthoquinone ring. The unsubstituted compounds 5–6 and the 2-methyl substituted compound 11 were unable to reduce the viability of either cell line by at least 50% at the maximum tested dose of 40 μM while the IC₅₀ of 7 also increased to 17.21 μM. In contrast, 2-bromosubstituted compounds 8–10 have better cytotoxicity profiles, with IC₅₀ of 9.55, 4.16 and 1.24 μM, respectively, against HEC1A compared to that of **BH10** which possessed IC₅₀ of 10.22 μM against HEC1A. Compounds 8–10 also have similar selectivity ratio of 2.15, 2.90 and 2.64, respectively, compared to that of 2.51 possessed by **BH10** (Table 1). These results suggest that an electron-withdrawing group such as a halogen at the C3 position of the naphthoquinone ring promotes cytotoxicity.

Interestingly, modifying the pendent amino group also had great influence on cytotoxicity (Table 1). Removal of the morpholine ring was found to have varied effects on cytotoxicity. Compound 23 with a terminal butyl chain, compound 24 with a terminal alkyne group and compound 25 with a terminal ester saw a drastic decrease in cytotoxicity, with IC₅₀ of 32.00, 21.92 and 20.00 μM, respectively, compared to **BH10** with IC₅₀ of 10.22 μM. Moreover, the addition of an amino acid moiety in compounds 26 and 27 also resulted in a loss of potency (IC₅₀ > 40 μM) against tumour cells. In contrast, compound 16 with a terminal 1-chloroethyl group possessed IC₅₀ of 5.65 μM and was more potent than **BH10** although the selectivity ratio of 2.14

possessed by compound 16 is slightly lower than that of 2.51 possessed by **BH10**. Furthermore, compounds 12 and 22 bearing a dimethylamino group showed IC₅₀ of 1.57 and 2.36 μM, respectively, and were at least four times more potent than that of **BH10**. However, their selectivity of 1.83 and 2.23, respectively, is again lower than that of **BH10**. This suggested that a terminal amino group provides the analogues superior cytotoxicity potency while the selectivity of the analogues is reduced as a trade-off.

Similarly, when the terminal morpholine ring system of **BH10** was replaced by other cyclic amine ring systems as in compounds 13–15, the potency was significantly increased while selectivity was decreased with respect to **BH10** (Table 1). Amongst compounds 13–15 with different terminal cyclic amine ring size, compound 14 with a terminal six-membered piperidine ring was the most potent compound in this study, showing an IC₅₀ of 1.07 μM and being almost 10-times more potent than **BH10**. On the other hand, the pyrrolidinyl derivative 13 and azapenyl derivative 15 share the same IC₅₀ of 1.29 μM and were almost 8-times more potent than that of **BH10** although they are slightly less potent than compound 14. However, all three compounds possessed lower selectivity than **BH10**. Given that compound 14 exhibited the highest potency amongst compounds 13–15 with different ring sizes, an additional CH₂ was introduced into the terminal alkyl chain (compound 21) to assess its impact. The results showed that the IC₅₀ of compound 21 slightly decreased to 1.83 μM compared to



Table 1 IC₅₀ values^a of **BH10** analogues against human endometrial cancer (HEC1A) cells and noncancerous human endometrial stromal (MAD11) cells, and their corresponding selectivity ratio and clog *P* value

Compound	HEC1A IC ₅₀ (μM)	MAD11 IC ₅₀ (μM)	Selectivity ratio	clog <i>P</i>
BH10	10.22	25.61	2.51	2.96
5	>40	>40	—	6.85
6	>40	>40	—	2.47
7	17.21	>40	>2.32	3.68
8	9.55	20.51	2.15	6.70
9	4.16	12.06	2.90	3.33
10	1.24	3.27	2.64	4.54
11	>40	>40	—	6.86
12	1.57	2.87	1.83	2.94
13	1.29	2.38	1.84	3.61
14	1.07	2.33	2.18	4.17
15	1.29	2.38	1.84	4.73
16	5.65	12.09	2.14	3.38
17	15.83	39.56	2.50	4.85
18	2.91	3.40	1.17	5.78
19	18.60	40.34	2.24	4.60
20	6.41	21.92	3.42	3.24
21	1.83	4.79	2.62	4.45
22	2.36	5.27	2.23	3.28
23	32.0	>40	>1.25	4.37
24	21.92	22.39	1.02	3.11
25	20.0	20.1	1.00	2.73
26	>40	>40	—	4.20
27	>40	>40	—	4.20
29	4.68	11.47	2.45	3.26
30	38.01	>40	—	2.50
31	19.51	34.02	1.74	3.85
32	>40	>40	—	3.59
33	11.13	21.32	1.92	4.35
39	26.04	>40	—	2.51
40	14.11	38.20	2.70	3.04
41	15	>40	—	3.52
42	15	>40	—	4.03
43	>40	>40	—	1.02
44	6.40	23.00	3.59	3.54
45	12.0	35.0	2.92	4.01
46	19	>40	—	4.52
47	14	24	1.80	5.13
48	>40	>40	—	1.91
50	8.8	39.0	4.43	4.20
53	27	24	0.90	2.34

^a IC₅₀ values were obtained from MTT cell viability assays (*n* = 3 biological repeats in HEC1A and MAD11 cell lines).

1.07 μM for compound **14**, suggesting that the extra CH₂ in the alkyl chain is not beneficial to potency. These results highlight that the removal of the ether functionality from the morpholine ring is beneficial for potency, with the six-membered piperidine ring and ethylene linker length to be the optimum ring size and linker length, respectively, for anticancer activity. This suggested that hydrophobic terminal groups might elicit a better effect in this position of the molecule.

The incorporation of an additional nitrogen in the pendent amine ring yielded varied effects in potency. The Boc-protected intermediate **17** and indole compound **19** with a terminal indole ring showed IC₅₀ of 15.83 and 18.60 μM, respectively, and were less potent than **BH10**. In contrast, the piperazinyl salt **29** with an IC₅₀ of 4.68 μM and selectivity ratio of 2.45 was 2.2 times more potent than **BH10** and have similar selectivity as **BH10**. However, another salt analogue

30 showed lower potency with IC₅₀ of 38.01 μM respectively. On the other hand, the bulky analogue **18** possessed IC₅₀ of 2.91 μM and was 3.5-times more potent than **BH10**, however, it was less selective with selectivity ratio of only 1.17. Therefore, large terminal groups attached to the pendent amine ring are not suitable for structural modifications aimed at simultaneously enhancing potency and selectivity.

Naphthoquinone compounds **31–33** with a 1,2,3-triazole linker and a terminal phenyl ring at the *N*¹ position were also synthesized and tested as the triazole moiety is commonly found in bioactive compounds. However, compounds **31–33** showed moderate to low anticancer activity, with none of their potency and selectivity being better than **BH10**. This demonstrated that the introduction of a triazole ring into naphthoquinone is not a good structural modification strategy.



Analogues with a nitrogen-containing group substituted at the C3 position were synthesized to examine their impact on potency and specificity. This series of compounds demonstrated that replacing the halogen with an amide functionality improves specificity while retaining potency. The propionamide analogue **40**, butyramide analogue **41**, pentanamide analogue **42** and phenylacetamide analogue **50** showed IC₅₀ of 14, 15, 15 and 8.8 μM, respectively, with similar potency as **BH10**, but the newly synthesized compounds possess higher selectivity of 2.7, 3.7, 3.9 and 4.4, respectively.

When the amide-substituted compounds were cyclised into an imidazole functionality, potency generally improved for some compounds while the specificity was retained from the amide-substituted compounds. The most potent imidazole analogue was the 2-methyl substituted analogue **44** with an IC₅₀ of 6.4 μM, followed by the 2-ethyl substituted analogue **45** with an IC₅₀ of 12 μM, 2-butyl substituted analogue **47** with an IC₅₀ of 14 μM and the 2-propyl substituted analogue **46** with an IC₅₀ of 19 μM. Upon comparing the uncyclised amide compounds **39–42** with their corresponding cyclised imidazole counterparts **44–47**, the cyclised imidazole compound **44** showed a sharp increase in potency with IC₅₀ of 6.4 μM compared to its corresponding uncyclised compound **39**, which possesses IC₅₀ of 26 μM. Cyclised imidazole compounds **45** and **47** also showed slight increases of potency with IC₅₀ decreased to 12 and 14 μM, respectively, compared to their corresponding uncyclised amide compounds **40** and **42**, which possess IC₅₀ of 14 and 15 μM, respectively. However, when the uncyclised amide compound **41** was cyclised into the imidazole compound **46**, the potency dropped slightly, with the IC₅₀ increased from 15 μM to 19 μM. On the other hand, the good selectivity observed for the imidazole compounds could possibly be attributed to the fact that the imidazole ring is relatively stable and rigid. The above results show that the cyclisation of amide analogues into imidazole compounds generally improves the potency of compounds, while retaining specificity.

Based on the biotesting results and analysis, several structure–activity relationships (SAR) were identified (Fig. 3). Firstly, incorporating an imidazole ring into the naphthoquinones enhances both potency and selectivity. Secondly, terminal cyclic amine groups generally exhibit higher activity. Additionally, introducing an amide group at the C3 position can improve selectivity while maintaining potency. Lastly, the presence of halogen atoms is crucial for maintaining potency.

To further assess the therapeutic potential and antiproliferative activity of selected potent analogues **21** and **44**, additional *in vitro* cytotoxicity assays were performed against two human cancer cell lines (CALU-1 (lung squamous cell carcinoma) and Mia-Pa-Ca-2 (pancreatic cancer)). Among the tested compounds, compounds **21** and **44** exhibited better potency compared to **BH10**, especially compound **21** demonstrating significantly higher efficacy against the selected cancer cell lines (Table 2).

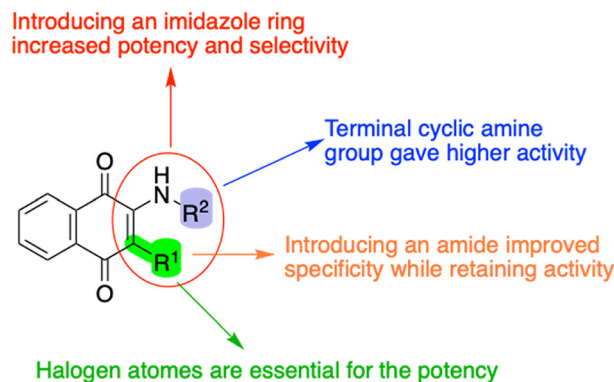


Fig. 3 SAR conclusions of the synthesized **BH10** derivatives. The incorporation of an imidazole ring enhances both potency and selectivity. Terminal cyclic amine groups generally exhibit higher activity. Introducing an amide group at the C3 position enhances selectivity while preserving potency. The presence of halogen atoms is critical for maintaining potency.

4. Investigating the potential mechanism of action (MOA) of compounds **44** and **21**

We wanted to find compounds that were more potent and had greater selectivity (cytotoxicity towards cancer cells than non-cancerous cells) than hit compound **BH10**. The imidazole **44** (IC₅₀ = 6.40 μM; selectivity ratio = 3.6) and **21** (IC₅₀ = 1.83 μM; selectivity ratio = 2.62) were selected for investigating whether their mechanism of action was similar to **BH10** (Fig. 4a).

4.1. Thiol rescue experiment

A thiol rescue experiment was performed to investigate the effect of thiol-containing compounds (antioxidants) on the cytotoxicity of **BH10** analogues against cancer cells in a cellular environment. This allowed the exploration of the mechanism of action or selectivity of the newly synthesized analogues. The thiol rescue experiment followed the same protocol as the MTT assay with an additional step. HEC1A cancer cells were seeded on day 1 as normal. On day 2, the cells were pre-treated with thiol-containing compounds *N*-acetyl cysteine (NAC) or glutathione (GSH) (at 2 mM) for 40 min (Fig. 4b). After that, the test compounds were added to the plate and cells were incubated for 48 h, as per previous

Table 2 IC₅₀ values of **BH10**, **21** and **44** against other human cancer cell lines^a

Compound	Mean IC ₅₀ (μM)	
	CALU-1	Mia-Pa-Ca-2
BH10	16.67	19.68
21	4.70	4.75
44	15.38	14.72

^a Values were obtained from MTT cell viability assays (*n* = 3 biological repeats in all cell lines).



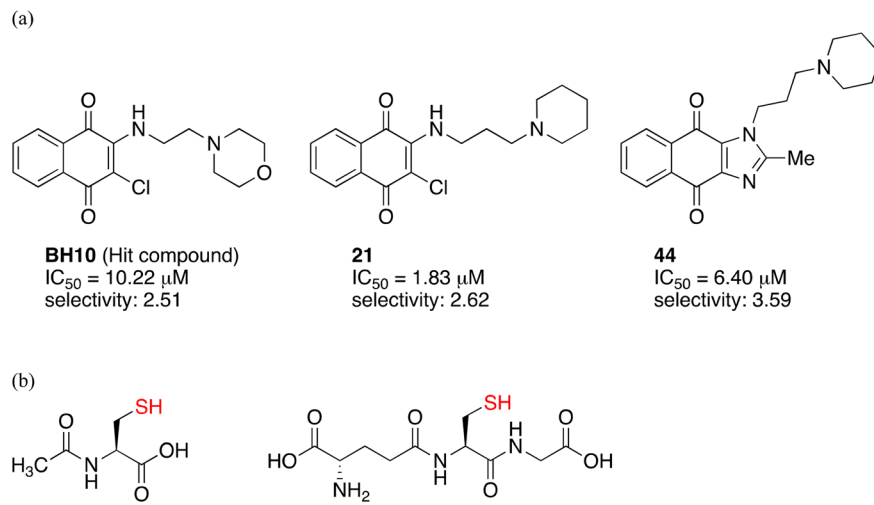


Fig. 4 (a) Chemical structures of the hit compound **BH10** and its most promising analogues, **21** and **44**. (b) Chemical structures of thiol-containing compounds, *N*-acetyl cysteine (NAC, left) and glutathione (GSH, right), with thiol functional groups highlighted in red. The thiol rescue experiment was conducted following the standard MTT assay protocol with an additional step, where HEC1A cancer cells were seeded as per the standard protocol on day 1. Cells were pre-treated with NAC or GSH (2 mM) for 40 minutes, followed by the addition of test compounds on day 2. The cells were then incubated for 48 hours according to the standard MTT assay protocol.

MTT protocol.⁸ An increase in IC₅₀ for cells treated with thiol-containing compounds compared to the control group indicates rescue from cell death by the thiols (Table 3).

In general, HEC1A cancer cells treated with compounds containing the C3 halogen (**BH10**, **21**) were rescued by the thiol-containing compounds NAC and GSH. **BH10** lost its cytotoxicity as demonstrated by the increased IC₅₀ value of 58.00 μM and greater than 80.00 μM with NAC and GSH pre-treatment, respectively. The bioactivity also decreased 4.6 times with NAC pre-treatment. Similarly, the cytotoxicity of **21** was also partially rescued by both NAC and GSH because the IC₅₀ values increased to 13.00 and 16.00 μM, and potency decreased by 7.1 and 8.7 times, respectively. These results suggested that compound **21** likely shares the same mechanism as **BH10**. In contrast, pretreating cancer cells with either NAC or GSH did not rescue the cytotoxicity of compound **44**, with no significant change in IC₅₀ values from the control group observed. An example of cellular viability curves for **BH10** and **44** alone or in the presence of NAC or GSH is shown in ESI† S1.

It was hypothesized that the thiol compounds may act as scavenger molecules, reacting with the halogen-containing compounds, leading to their deactivation by preventing binding to other cellular targets. Therefore, the presence of the halogen at C3 appeared to be essential for thiol-mediated cell-rescue. Halogens are very good leaving groups, making

Michael addition-type nucleophilic attack on the naphthoquinone scaffold possible. When the halogen is replaced with much poorer leaving groups, such as amides and amines, Michael addition becomes less likely. In the imidazole analogues, nucleophilic attack is very unlikely as the enone functionality is in between a fused ring system, making it no longer available to react as any reactions that occur at that position would disrupt the aromaticity of the system and hence such reactions are energetically unfavourable.

4.2. Measurements of oxygen consumption

Our initial study identified **BH10** as a 'hit' compound based on a phenotypic screen that investigated compounds that increased cellular oxygen consumption rate (OCR).⁷ Therefore, oxygen consumption measurements were performed with the newly synthesized analogues using a Seahorse Extracellular Flux Analyser, which measures the OCR of cultured cells over time. High OCR can indicate a high rate of nutrient oxidation in the mitochondria caused by oxidative phosphorylation, or it can also be influenced by external mechanisms such as redox cycling.¹⁹ Although measurements of OCR do not pinpoint the location of oxygen consumption, it is a good initial indicator to compare the mechanism of action of new analogues with that of **BH10**. It

Table 3 IC₅₀ values for **BH10** analogues alone, or pre-treated with NAC or GSH, in HEC1A endometrial cancer cells

Compound	Alone IC ₅₀ (μM)	+NAC IC ₅₀ (μM)	ΔIC ₅₀ ^a (μM)	+GSH IC ₅₀ ± SEM (μM)	ΔIC ₅₀ ^a (μM)
BH10	10.22	58.00	+47.78	>80.00	>+69.78
21	1.83	13.00	+11.17	16.00	+14.17
44	6.40	6.80	+0.40	7.40	+1.00

^a ΔIC₅₀ is IC₅₀ with thiol compound minus IC₅₀ of compound alone.



was hypothesized that compounds without the C3 halogen and were unaffected by thiols in the cell rescue experiment might act *via* a different mechanism to **BH10**, and hence give different results in OCR.

In this assay, HEC1A cells were treated with compounds **BH10**, **21** and **44** at various concentrations, and OCR was measured over 120 minutes. The results demonstrated that **BH10** increased OCR in cells by up to 200% (Table 4 and ESI† S2), with the increase correlating with drug concentration, though 10 μM appeared to approach the maximum effect. Compound **21**, another C3 halogen-containing compound, showed a similar trend to **BH10**, with a maximum OCR increase of approximately 280% at 20 μM . In contrast, compound **44**, which lacks the C3 halogen group, induced a maximum OCR increase of 130% at doses equivalent to that of **BH10** and **21**, compared to the vehicle (DMSO) control. These OCR results suggest that the new analogue **44**, lacking the C3 halogen, likely acts *via* a different mechanism compared to **BH10**.

5. Pharmacokinetic profiling of compound **44** *in vivo*

One of the major limitations of **BH10** as an anticancer compound is its short half-life *in vivo*, with a half-life of 21.6 min when administered at 10 mg kg⁻¹ *via* oral gavage in C57BL/6 mice.⁸ This means that the drug is rapidly metabolised, so larger doses would be required for *in vivo* efficacy, which in turn increases the risk of toxicity. By replacing the halogen at C3 in the hit compound **BH10** to more metabolically stable functional groups such as amides and imidazole rings, it was hypothesized that the half-life would increase. To confirm this, pharmacokinetic profiling of the most potent analogue **44** was performed in mice. In such experiments, mice were administered the compound *via* oral gavage, and blood was drawn from their tail at regular intervals. Blood was then centrifuged, and the plasma collected was mixed with 10% MeOH in acetonitrile to precipitate protein. Mass spectrometry analyses of each sample and drug standards (drugs spiked into blood) enabled quantitation of each drug in the plasma over time to generate a pharmacokinetic profile. From this, maximum plasma concentration (C_{max}) and half-life ($t_{1/2}$) were determined (Table 5). The novel analogue **44**, lacking the halogen at the C3 position, demonstrated a half-life of 26.8 min *via* oral administration (see ESI† S3), which is 5.2 min longer than

BH10 ($t_{1/2}$ = 21.6 min).⁸ Although this represents a slight improvement, further optimization is necessary to enhance the compound's half-life.

6. Molecular docking study

A molecular docking simulation was conducted to examine the interaction between **BH10/44** and the Keap1 protein (PDB: 4XMB), aiming to elucidate their potential binding modes. The results indicated that both compounds occupy the same hydrophobic pocket within Keap1 but engage in distinct interactions. For **BH10**, the carbonyl group forms hydrogen bonds with key residues GLY-367, VAL-465, and VAL-606 (Fig. 5a). Additionally, the amine hydrogen forms hydrogen bonds with the carbonyl groups of GLY-367 and VAL-606, and the nitrogen in the morpholine ring forms another hydrogen bond with THR-560. In contrast, compound **44** exhibited a simpler interaction profile, with only the carbonyl group forming hydrogen bonds with residues VAL-465 and VAL-606 (Fig. 5b). Notably, compound **44** had a docking score of -7.36, closely aligned with the fitness score of -8.73 for **BH10**, indicating a comparable binding affinity of compound **44** for the target protein.

7. Conclusion

In conclusion, a series of 40 analogues based on the **BH10** scaffold were successfully synthesized, accompanied by a detailed structure-activity relationship study. Several compounds exhibited promising anticancer potency with IC₅₀ of 1–2 μM but with decreased selectivity. Among the synthesized analogues, the imidazole compound **44** demonstrated the most favourable balance, with an IC₅₀ of 6.4 μM and a selectivity factor of 3.6. *In vitro* investigations into the mechanism revealed compound **44** was unaffected by thiol compounds in the cytotoxicity assay, and did not increase OCR significantly which indicates that **44** likely acts *via* a different mechanism to the hit compound **BH10**. This could indicate differences in the metabolic profiles of **44** and **BH10**. Pharmacokinetic profiling *in vivo* showed only a slight improvement in the half-life of compound **44** compared to **BH10**, indicating the need for further optimization to improve its pharmacokinetic properties.

8. Experimental

8.1. Chemical synthesis

8.1.1. General information. Commercially available reagents were purchased from Acros Organics, Alfa Aesar, Fluka, Combi-Blocks, Oakwood, Lancaster and Sigma Aldrich, and were used without further purification. All reactions requiring anhydrous conditions were performed under nitrogen atmosphere and anhydrous solvents were obtained using a PureSolv MD solvent purification system. Melting points were measured using a Mel-Temp melting point apparatus and are uncorrected. ¹H and ¹³C NMR spectra were

Table 4 Effect of **BH10**, compound **21** and compound **44** on percent OCR (basal OCR is defined as 100%) over 120 minutes in HEC1A cancer cells

Compound	OCR (%)			
	DMSO	5 μM	10 μM	20 μM
BH10	100	125	200	200
21	100	175	260	280
44	100	100	130	130



Table 5 Pharmacokinetic profile data^a

Compound	Treatment type	Dose (mg kg ⁻¹)	Mouse strain	C _{max} (μM)	t _{1/2} (min)
44	Oral	20	C57BL/6	0.70	26.8

^a Values are the average of three runs. C_{max}, maximum concentration; t_{1/2}, half-life.

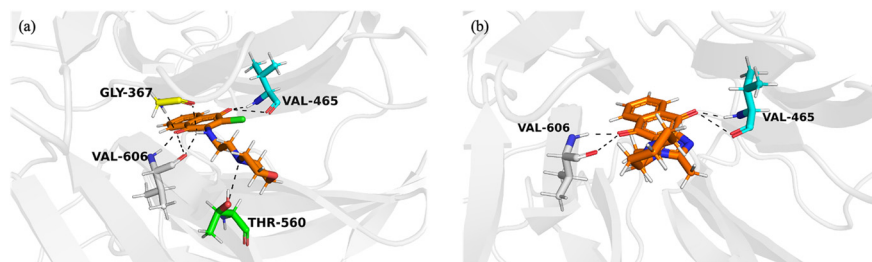


Fig. 5 Predicted binding model of BH10 (a) and 44 (b) with Keap1 (PDB code: 4XMB). Hydrogen bonding interactions are indicated by black dashed lines. (a) The carbonyl group of BH10 forms hydrogen bonds with residues GLY-367, VAL-465, and VAL-606 while the amine hydrogen forms hydrogen bonds with the carbonyl groups of GLY-367 and VAL-606. (b) The carbonyl group of compound 44 forms hydrogen bonds with residues VAL-465 and VAL-606.

obtained in the designated solvents on a Bruker Avance III HD (300 or 400 MHz) spectrometer. ¹H NMR spectral data are reported as follows: chemical shift measured in parts per million (ppm) downfield from TMS (δ); multiplicity; observed coupling constant (J) in Hertz (Hz) and proton count. Multiplicities are recorded as singlet (s), broad singlet (br s), doublet (d), triplet (t), quartet (q), multiplet (m) and combinations of these. ¹³C NMR chemical shifts are reported in ppm downfield from TMS (δ), and identifiable carbons are given. Infrared spectra were recorded on a Perkin Elmer Spotlight 400 FTIR microscope. High-resolution mass spectrometry was performed by the bioanalytical mass spectrometry unit, UNSW. Ajax Finechem Silica 200–325 mesh was used for column chromatography and Merck silica gel 60H was used for flash chromatography. Reactions were monitored using thin layer chromatography, performed on Merck DC aluminium foil coated with silica gel (60GF₂₅₄). Compounds were detected using short or long wavelength ultraviolet light.

General synthetic method a for 5–27, 39–43 and 50. This method was based on a modification of the procedure reported by Brun *et al.*¹⁷ The appropriate naphthoquinone (1 equivalent) was dissolved in EtOH or Et₂O (20 mL) and to the resulting mixture was added 1.0–1.2 equivalent of the appropriate amine and 1.0–1.2 equivalent of Et₃N or K₂CO₃. The resulting mixture was stirred at r.t. for 18–72 h. The reaction mixture was then concentrated *in vacuo*, diluted in EtOAc and washed with water, and the organic layer was dried over anhydrous Na₂SO₄. The solvent was evaporated, and the resulting residue was purified *via* flash column chromatography (when required) in the appropriate solvent system to afford the desired product.

General synthetic method B for 29 and 30. TFA (1 mL) was added to a solution of corresponding Boc-compound (0.30 g, 0.67 mmol) in DCM (10 mL) under nitrogen on ice. The reaction mixture was stirred at r.t. for 5 h. The solvent removed *in vacuo* yielding corresponding products.

General synthetic method C for 31–33. Under an argon atmosphere, alkyne 24 and the corresponding azido moieties were reacted in THF: H₂O (5:1), sodium ascorbate 20 mol%, CuSO₄·5H₂O 10 mol% at 50 °C for 20 h. The reaction was then quenched in H₂O (20 mL), and the organic layer extracted by EtOAc (2 × 30 mL) with a single brine wash (1 × 10 mL). The combined extracts were dried over Na₂SO₄, and the volatiles evaporated under reduced pressure. The crude product was purified using column chromatography on silica eluting with a Hexane: EtOAc solvent system to afford the target products.

General synthetic method D for amide intermediates 35–38. The appropriate excessive anhydride was added to compound 34. Concentrated sulfuric acid (2 drops) was added. The reaction mixture was stirred at r.t. for 2 h. Excess anhydride was removed by an ether wash. Column purification was performed when required.

General synthetic method E for imidazole analogues 44–48. Aqueous NaOH (2 M) was added to a solution of appropriate amide analogue 39–43 in EtOH (2–10 mL). The reaction mixture was heated at 50 °C for 1–2 h. The solvent was removed *in vacuo* and the residue was purified *via* flash column chromatography (when required) in the appropriate solvent system to afford the product.

8.1.2. Synthesis of intermediates

Methyl N⁶-(*tert*-butoxycarbonyl)-N²-(3-chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)-L-lysinate (28). Following general synthetic procedure A, *N*-Boc-L-lysine methyl ester hydrochloride (0.66 g, 2.2 mmol) and K₂CO₃ (0.10 g, 0.73 mmol) were added to a solution of 2,3-dichloro-[1,4]-naphthoquinone (0.51 g, 2.20 mmol) in 10% THF in EtOH (10 mL). The product eluted from hexane: EtOAc (5:1) as an orange solid (0.34 g, 34%). ¹H NMR (400 MHz, CDCl₃) δ 8.17 (dd, J = 7.6, 1.3 Hz, 1H), 8.08 (dd, J = 7.7, 1.3 Hz, 1H), 7.76 (td, J = 7.6, 1.4 Hz, 1H), 7.67 (td, J = 7.6, 1.4 Hz, 1H), 6.34 (s, 1H), 5.25 (m, 1H), 4.55 (s, 1H), 3.82 (s,



3H), 3.13 (t, $J = 6.6$ Hz, 2H), 2.05–1.84 (m, 2H), 1.60 (m, 2H), 1.58–1.52 (m, 2H), 1.45 (s, 9H); ^{13}C NMR (101 MHz, CDCl_3) δ 180.10, 176.84, 172.37, 155.95, 143.47, 134.90, 132.73, 132.31, 129.88, 126.96, 126.84, 79.25, 56.22, 52.78, 33.71, 29.79, 28.40, 22.38; IR (cm^{-1}) 3371, 3311, 2965, 2846, 1735, 1671, 1605; HRMS (+ESI): found m/z 473.1446; $[\text{M} + \text{Na}]^+$; $\text{C}_{22}\text{H}_{27}\text{ClN}_2\text{O}_6\text{Na}$ required 473.1450.

N-(3-Chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)acetamide (35).¹⁵ Following general synthetic method D, acetic anhydride (5 mL) was added to **34** (0.21 g, 1.0 mol). Concentrated sulfuric acid (2 drops) was added. The reaction mixture was stirred at r. t. for 2 h. The resulting yellow precipitate was washed with ether and collected (1.30 g, 52%).¹⁵ ^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 7.98 (d, $J = 7.7$ Hz, 1H), 7.84–7.66 (m, 2H), 7.60 (t, $J = 7.6$ Hz, 1H), 2.60 (s, 3H); ^{13}C NMR (101 MHz, $\text{DMSO}-d_6$) δ 179.47, 172.64, 167.78, 163.39, 158.06, 135.47, 131.16, 130.33, 130.03, 125.98, 122.87, 14.17; MS (+ESI): found m/z 250.05; $[\text{M} + \text{H}]^+$; $\text{C}_{12}\text{H}_9\text{ClNO}_3$ required 250.03.

N-(3-Chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)propionamide (36).²⁰ Following general synthetic method D, propionic anhydride (5 mL) was added to **34** (0.21 g, 1.0 mmol). Concentrated sulphuric acid (1 drop) was added. The reaction mixture was stirred well at r.t. for 2 h. The resulting yellow precipitate was washed with ether and collected (0.24 g, 86%).^{15,20} ^1H NMR (400 MHz, CDCl_3) δ 8.25–8.06 (m, 2H), 7.86–7.73 (m, 2H), 7.73–7.63 (m, 1H), 2.57 (q, $J = 7.5$ Hz, 2H), 1.30 (t, $J = 7.5$ Hz, 3H); ^{13}C NMR (101 MHz, CDCl_3) δ 179.98, 177.68, 170.38, 139.13, 134.81, 134.14, 131.58, 131.10, 130.27, 127.54, 127.07, 30.52, 9.32; MS (+ESI): Found m/z 264.05; $[\text{M} + \text{H}]^+$; $\text{C}_{13}\text{H}_{11}\text{ClNO}_3$ required 264.04.

N-(3-Chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)butyramide (37).²⁰ Following general synthetic method D, butyric anhydride (5 mL) was added to **34** (0.21 g, 1.0 mmol). Concentrated sulphuric acid (3 drops) was added. The reaction mixture was stirred at r.t. for 2 h. The resulting yellow precipitate was washed with ether and collected (0.18 g). The product eluted from hexane: EtOAc (5:1) column as a yellow solid (0.15 g, 53%).²⁰ ^1H NMR (400 MHz, CDCl_3) δ 8.25–8.15 (m, 1H), 8.16–8.08 (m, 1H), 7.79 (dd, $J = 6.7, 5.8$ Hz, 2H), 7.73–7.66 (m, 1H), 2.52 (t, $J = 7.4$ Hz, 2H), 1.82 (q, $J = 7.4$ Hz, 2H), 1.07 (t, $J = 7.4$ Hz, 3H); ^{13}C NMR (101 MHz, CDCl_3) δ 179.99, 177.68, 169.64, 139.11, 134.81, 134.14, 133.19, 131.59, 130.27, 127.54, 127.06, 39.19, 18.85, 13.62; MS (+ESI): found m/z 278.05; $[\text{M} + \text{H}]^+$; $\text{C}_{14}\text{H}_{13}\text{ClNO}_3$ required 278.06.

N-(3-Chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)pentanamide (38).²⁰ Following general synthetic method D, valeric anhydride (5 mL) was added to **34** (0.20 g, 0.98 mmol). Concentrated sulphuric acid (2 drops) was added. The reaction mixture was stirred well at r.t. for 2 h. The resulting yellow precipitate was washed with ether and collected (0.25 g). Product eluted from hexane: EtOAc column (5:1) as a yellow solid (0.19 g, 65%).²⁰ ^1H NMR (400 MHz, CDCl_3) δ 8.26–8.05 (m, 1H), 7.87–7.65 (m, 2H), 2.53 (t, $J = 7.5$ Hz, 1H), 1.88–1.70 (m, 1H), 1.55–1.37 (m, 1H), 0.99 (t, $J = 7.3$ Hz, 2H); ^{13}C NMR (101 MHz, CDCl_3) δ 179.98, 177.67, 169.81, 139.13, 134.80, 134.13, 133.18,

131.58, 130.27, 127.53, 127.05, 37.09, 27.39, 22.25, 13.79; MS (+ESI): found m/z 292.05; $[\text{M} + \text{H}]^+$; $\text{C}_{15}\text{H}_{15}\text{ClNO}_3$ required 292.07.

N-(3-Chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)-2-phenylacetamide (49).¹⁶ A solution of aminochloro compound **34** (0.90 g, 4.3 mmol) in toluene (9 mL) was stirred at 90 °C. A solution of borontrifluoride diethyl etherate (600 μL , 3.2 mmol) and phenylacetyl chloride (3.0 mL, 16 mmol) in toluene (4 mL) was added dropwise over 30 min. The reaction mixture was then left stirring at 90 °C for a further 7 h. The reaction mixture was cooled to r.t. and the solvent evaporated. The resulting pale-yellow green precipitate was washed with ether and collected (0.78 g, 55%).¹⁶ ^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 10.38 (s, 1H), 8.12–7.98 (m, 2H), 7.95–7.86 (m, 2H), 7.39–7.30 (m, 4H), 7.30–7.22 (m, 1H), 3.81 (s, 2H); ^{13}C NMR (101 MHz, $\text{DMSO}-d_6$) δ 179.23, 178.06, 169.26, 141.61, 135.75, 135.31, 135.04, 135.03, 131.44, 131.30, 129.69, 128.78, 127.17, 127.12, 42.51. MS (+ESI): found m/z 326.06; $[\text{M} + \text{H}]^+$; $\text{C}_{18}\text{H}_{13}\text{ClNO}_3$ required 326.05.

8.1.3. Synthesis of final products

2-((2-Morpholinoethyl)amino)naphthalene-1,4-dione (5).²¹ Following general synthetic procedure A, 1,4-naphthoquinone (0.35 g, 2.20 mmol), EtOH (20 mL), 4-(2-aminoethyl) morpholine (0.30 mL, 2.20 mmol) and Et_3N (0.31 mL, 2.20 mmol) were reacted. Reaction stirred for 18 h. Product eluted with EtOAc/hexane in flash column chromatography to afford a dark orange powder (0.49 g, 79%): mp: 147–148 °C; ^1H NMR (300 MHz, CDCl_3) δ 8.21–8.11 (m, 1H), 8.11–8.01 (m, 1H), 7.59–7.79 (m, 2H), 7.00 (s, 1H), 4.06–3.91 (m, 2H), 3.85–3.71 (m, 4H), 2.69 (t, $J = 6.0$ Hz, 2H), 2.64–2.46 (m, 4H). ^{13}C NMR (75 MHz, CDCl_3) δ 180.33, 147.00, 134.71, 132.42, 132.32, 130.05, 126.96, 126.75, 67.05, 56.65, 52.94, 41.17. IR (cm^{-1}) 3342, 2969, 2832, 1671, 1595, 1567, 1517, 1113, 865, 724; MS (+ESI): found m/z 287.25, $[\text{M} + \text{H}]^+$; $\text{C}_{16}\text{H}_{19}\text{N}_2\text{O}_3$ required 287.13.

2-((3-Morpholinopropyl)amino)-[1,4]-naphthoquinone (6). Following general synthetic procedure A, 1,4-naphthoquinone (0.35 g, 2.20 mmol), EtOH (20 mL), 3-morpholinopropylamine (0.32 mL, 2.20 mmol) and Et_3N (0.31 mL, 2.20 mmol) were reacted. Reaction stirred for 18 h. Product eluted with EtOAc/hexane in flash column chromatography to afford a bright red crystals (0.42 g, 64%), mp: 129–131 °C; ^1H NMR (300 MHz, CDCl_3) δ 8.09 (m, 2H), 7.79–7.57 (m, 3H), 5.72 (s, 1H), 3.91 (t, $J = 4.7$ Hz, 4H), 3.32 (q, $J = 5.8$ Hz, 2H), 2.58 (s, 6H), 1.92 (s, 2H); ^{13}C NMR (75 MHz, CDCl_3) δ 182.93, 181.94, 148.64, 134.59, 133.80, 131.84, 130.69, 126.22, 126.12, 100.36, 66.64, 57.65, 53.74, 42.81, 23.37; IR (cm^{-1}) 3258, 2963, 2852, 1677, 1592, 1562, 1519, 1111, 726; HRMS (+ESI): found m/z 301.1547, $[\text{M} + \text{H}]^+$; $\text{C}_{17}\text{H}_{21}\text{N}_2\text{O}_3$ required 301.1545.

2-((3-Piperidin-1-yl)propyl)amino)-[1,4]-naphthoquinone (7). Following general synthetic procedure A, 1,4-naphthoquinone (0.35 g, 2.20 mmol), EtOH (20 mL), 3-aminopropylpiperidine (0.35 mL, 2.20 mmol) and Et_3N (0.31 mL, 2.20 mmol) were reacted. Product eluted at EtOAc/hexane = 1:1 in flash column chromatography to afford a dark red solid (0.06 g, 10%): mp:



108–110 °C; ^1H NMR (400 MHz, CDCl_3): δ 8.11 (m, 2H), 7.92 (br s, 1H), 7.73 (t, $J = 7.5$ Hz, 1H), 7.61 (t, $J = 7.6$ Hz, 1H), 5.70 (s, 1H), 3.29 (q, $J = 17.2$ Hz, 2H), 2.53–2.45 (m, 6H), 1.90–1.70 (m, 6H), 1.53 (t, $J = 4.1$ Hz, 2H); ^{13}C NMR (CDCl_3): δ 182.90, 181.94, 148.88, 134.46, 133.88, 131.69, 130.79, 126.19, 126.06, 100.16, 58.10, 43.08, 30.93, 25.68, 24.47, 23.75; IR (cm^{-1}) 3259, 3076, 2918, 2784, 1679, 1605, 1518, 725; HRMS (+ESI): found m/z 2.991754, $[\text{M} + \text{H}]^+$; $\text{C}_{18}\text{H}_{23}\text{N}_2\text{O}_2$ required 299.1756.

2-Bromo-3-((2-(4-morpholinyl)ethyl)amino)-[1,4]-naphthoquinone (8). Following general synthetic procedure A, 2,3-dibromo-[1,4]-naphthoquinone (0.70 g, 2.20 mmol), Et_2O (20 mL), 4-(2-aminoethyl)morpholine (0.30 mL, 2.20 mmol), Et_3N (0.30 mL, 2.20 mmol). Reaction stirred for 72 h. Product eluted at EtOAc/hexane = 1:4 in flash column chromatography to afford a red powder (0.58 g, 72%): mp: 120–122 °C; ^1H NMR (300 MHz, CDCl_3): δ 8.14 (d, $J = 7.6$ Hz, 1H), 8.04 (d, $J = 8.7$ Hz, 1H), 7.74 (t, $J = 8.9$ Hz, 1H), 7.69 (t, $J = 7.9$ Hz, 1H), 6.98 (br s, 1H), 4.00 (q, $J = 16.8$ Hz, 2H), 3.77 (t, $J = 9.2$ Hz, 4H), 2.69 (t, $J = 11.9$ Hz, 2H), 2.54 (t, $J = 8.9$ Hz, 4H); ^{13}C NMR (CDCl_3): δ 180.33, 134.70, 132.31, 130.52, 126.96, 126.74, 67.05, 56.65, 52.94, 41.18; IR (cm^{-1}) 3210, 2948, 2831, 1676, 1563, 1450, 1285, 718; HRMS (+ESI): found m/z 365.0495, $[\text{M} + \text{H}]^+$; $\text{C}_{16}\text{H}_{17}\text{BrN}_2\text{O}_3$ required 365.0490.

2-Bromo-3-((3-morpholinopropyl)amino)-[1,4]-naphthoquinone (9). Following general synthetic procedure A, 2,3-dibromo-[1,4]-naphthoquinone (0.70 g, 2.20 mmol), Et_2O (20 mL), 3-morpholinopropylamine (0.32 mL, 2.20 mmol), Et_3N (0.31 mL, 2.20 mmol). Reaction stirred for 19 h. Product eluted at EtOAc/hexane = 1:1 in flash column chromatography to afford an orange powder (0.18 g, 22%): mp: 144–146 °C; ^1H NMR (400 MHz, CDCl_3): δ 8.16 (d, $J = 8.6$ Hz, 1H), 8.04 (d, $J = 8.6$ Hz, 1H), 7.73 (t, $J = 16.5$ Hz, 1H), 7.64 (t, $J = 8.9$ Hz, 1H), 4.02–4.11 (m, 2H), 3.80–3.92 (m, 4H), 2.40–2.70 (m, 6H), 1.89 (s, 2H); ^{13}C NMR (CDCl_3): δ 180.34, 147.06, 134.73, 132.51, 132.25, 129.96, 127.03, 126.81, 66.57, 57.54, 53.92, 45.50, 25.63; IR (cm^{-1}) 3082, 2948, 2782, 1678, 1561, 1291, 1110, 721; HRMS (+ESI): found m/z 379.0652, $[\text{M} + \text{H}]^+$; $\text{C}_{17}\text{H}_{19}\text{BrN}_2\text{O}_3$ required 379.0649.

2-Bromo-3-((3-(piperidin-1-yl)propyl)amino)-[1,4]-naphthoquinone (10). Following general synthetic procedure A, 2,3-dibromo-[1,4]-naphthoquinone (0.70 g, 2.20 mmol), Et_2O (20 mL), *N*-(3-aminopropyl)piperidine (0.35 mL, 2.20 mmol), Et_3N (0.31 mL, 2.20 mmol). Reaction stirred for 17 h. Product eluted at EtOAc/hexane = 1:1 in flash column chromatography to afford a dark red solid (0.21 g, 25%): mp: 102–104 °C; ^1H NMR (300 MHz, CDCl_3): δ 8.19–8.16 (m, 1H), 8.07–8.03 (m, 1H), 7.72 (td, $J = 7.5$, 1.4 Hz, 1H), 7.63 (td, $J = 7.5$, 1.4 Hz, 1H), 7.56 (s, 1H), 4.04 (q, $J = 6.0$ Hz, 2H), 2.53 (d, $J = 8.1$ Hz, 6H), 1.98–1.83 (m, 2H), 1.73 (q, $J = 5.6$ Hz, 4H), 1.52 (d, $J = 7.1$ Hz, 2H); ^{13}C NMR (76 MHz, CDCl_3): δ 180.38, 161.96, 134.63, 132.54, 132.16, 131.22, 126.96, 126.79, 57.65, 54.90, 45.63, 26.02, 25.38, 24.31; IR (cm^{-1}) 3065, 2935, 2765, 1679, 1562, 1502; HRMS (+ESI): found m/z 377.0859, $[\text{M} + \text{H}]^+$; $\text{C}_{18}\text{H}_{22}\text{BrN}_2\text{O}_2$ required 377.0858.

2-Methyl-3-((2-(4-morpholinyl)ethyl)amino)-[1,4]-naphthoquinone (11). Following general synthetic procedure

A, Menadione (0.38 g, 2.20 mmol), EtOH (20 mL), 4-(2-aminoethyl)morpholine (0.31 mL, 2.20 mmol). Reaction stirred for 21 h. Product eluted at EtOAc/hexane = 1:4 in flash column chromatography to afford an orange oil that was recrystallized from 70% EtOH to afford bright red crystals. The crystals were filtered and the filtrate was further evaporated *in vacuo* to afford a red powder (66 mg, 9%): mp: 70–72 °C; ^1H NMR (300 MHz, CDCl_3): δ 7.92–7.88 (m, 1H), 7.83–7.78 (m, 1H), 7.53 (td, $J = 7.5$, 1.4 Hz, 1H), 7.43 (td, $J = 7.5$, 1.4 Hz, 1H), 6.25 (s, 1H), 3.68–3.59 (m, 4H), 3.54 (q, $J = 5.6$ Hz, 2H), 2.57–2.47 (m, 2H), 2.47–2.37 (m, 4H), 2.07 (s, 3H). ^{13}C NMR (75 MHz, CDCl_3): δ 183.30, 182.39, 146.52, 134.12, 133.37, 131.74, 130.41, 126.02, 125.81, 112.46, 66.95, 57.33, 53.06, 41.35, 10.96. IR (cm^{-1}) 2800, 1671, 1609, 1560, 1500, 1451, 1337, 1294, 1105, 715; HRMS (+ESI): found m/z 301.1547, $[\text{M} + \text{H}]^+$; $\text{C}_{17}\text{H}_{21}\text{N}_2\text{O}_3$ required 301.1546.

2-Chloro-3-((2-(dimethylamino)ethyl)amino)-[1,4]-naphthoquinone (12). Following general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (0.51 g, 2.20 mmol), EtOH (20 mL), 3-(dimethylamino)-1-ethylamine (0.24 mL, 2.20 mmol) and Et_3N (0.31 mL, 2.20 mmol) were reacted. Reaction stirred for 18 h. Product eluted at EtOAc/hexane = 1:1 in flash column chromatography to afford a dark orange oil. The oil was recrystallized from 70% EtOH to afford a red powder 16 (0.41 g, 72%): mp: 80–82 °C; ^1H NMR (300 MHz, CDCl_3): δ 8.16 (d, $J = 8.6$ Hz, 1H), 8.00 (d, $J = 9.0$ Hz, 1H), 7.71 (t, $J = 16.5$ Hz, 1H), 7.61 (t, $J = 16.4$ Hz, 1H), 6.80 (br s, 1H), 3.97 (q, $J = 17.2$ Hz, 2H), 2.63 (t, $J = 11.9$ Hz, 2H), 2.32 (s, 6H); ^{13}C NMR (CDCl_3): δ 180.67, 176.71, 134.90, 132.89, 132.45, 130.12, 126.84, 77.57, 76.73, 58.17, 45.06, 41.98; IR (cm^{-1}) 3222, 2940, 2821, 2773, 1677, 1607, 1573, 1488, 1281, 718, 677; HRMS (+ESI): found m/z 301.0714, $[\text{M} + \text{Na}]^+$; $\text{C}_{14}\text{H}_{15}\text{ClN}_2\text{O}_2$ required 301.0716.

2-Chloro-3-((2-(1-pyrrolidinyl)ethyl)amino)-[1,4]-naphthoquinone (13). Following general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (0.51 g, 2.20 mmol), EtOH (20 mL), 1-(2-aminoethyl)pyrrolidine (0.28 mL, 2.20 mmol) and Et_3N (0.31 mL, 2.20 mmol) were reacted. Reaction stirred for 18 h. Product eluted with EtOAc/hexane in flash column chromatography to afford an orange oil (0.44 g, 63%): mp: 63–64 °C; ^1H NMR (300 MHz, CDCl_3): δ 8.15 (d, $J = 9.0$ Hz, 1H), 8.04 (d, $J = 9.0$ Hz, 1H), 7.64 (t, $J = 16.5$ Hz, 1H), 7.53 (t, $J = 16.4$ Hz, 1H), 6.76 (br s, 1H), 3.89 (q, $J = 17.5$ Hz, 2H), 2.73 (t, $J = 12.2$ Hz, 2H), 2.52–2.59 (m, 4H), 1.62–1.75 (m, 4H); ^{13}C NMR (75 MHz, CDCl_3): δ 180.14, 176.30, 144.60, 134.63, 132.47, 132.20, 129.74, 126.52, 126.42, 54.83, 53.63, 42.92, 23.55. IR (cm^{-1}) 3246, 2959, 2822, 1600, 1567, 1288, 1267, 717; MS (+ESI): found m/z 305.20, $[\text{M} + \text{H}]^+$; $\text{C}_{16}\text{H}_{18}\text{ClN}_2\text{O}_2$ required 305.11.

2-Chloro-3-((2-(1-piperidinyl)ethyl)amino)-[1,4]-naphthoquinone (14). Following general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (0.25 g, 1.10 mmol), EtOH (10 mL), 1-(2-aminoethyl)piperidine (0.16 mL, 1.20 mmol) and Et_3N (0.31 mL, 2.20 mmol) were reacted. Reaction stirred for 18 h. Product eluted at EtOAc/hexane = 1:4 in flash column chromatography to afford a dark red powder



(0.20 g, 57%): mp: 108–110 °C; ^1H NMR (300 MHz, CDCl_3): δ 8.15 (d, $J = 9.0$ Hz, 1H), 8.03 (d, $J = 9.5$ Hz, 1H), 7.73 (t, $J = 9.1$ Hz, 1H), 7.63 (t, $J = 9.0$ Hz, 1H), 7.06 (br s, 1H), 3.97 (q, $J = 17.3$ Hz, 2H), 2.65 (t, $J = 12.2$ Hz, 2H), 2.40 (br 4H), 1.62–1.67 (m, 4H), 1.47–1.51 (m, 2H); ^{13}C NMR (CDCl_3): δ 180.67, 176.0, 144.4, 134.7, 132.3, 131.8, 130.0, 126.7, 56.6, 54.0, 41.1, 26.0, 24.3. IR (cm^{-1}) 3208, 2923, 2847, 1674, 1599, 1568, 1267, 717; MS (+ESI): Found m/z 319.25, $[\text{M} + \text{H}]^+$; $\text{C}_{17}\text{H}_{20}\text{ClN}_2\text{O}_2$ required 319.12.

2-((2-(Azepan-1-yl)ethyl)amino)-3-chloro-[1,4]-naphthoquinone (15). Following general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (0.51 g, 2.20 mmol), hexamethyleneimine (0.50 mL, 4.40 mmol) and Et_3N (0.31 mL, 2.20 mmol) were reacted. Reaction refluxed for 18 h. Product eluted at EtOAc/hexane = 1:4 in flash column chromatography to afford a dark orange oil. The oil was recrystallised from 70% EtOH to afford an orange powder 24 (0.17 g, 24%): mp: 87–89 °C; ^1H NMR (300 MHz, CDCl_3): δ 8.05 (d, $J = 7.7$ Hz, 1H), 7.97 (d, $J = 8.1$ Hz, 1H), 7.69 (t, $J = 8.9$ Hz, 1H), 7.58 (t, $J = 7.5$ Hz, 1H), 7.14 (br s, 1H), 3.87 (q, $J = 16.8$ Hz, 2H), 2.76 (t, $J = 11.9$ Hz, 2H), 2.70–2.63 (m, 4H), 1.72–1.58 (m, 8H); ^{13}C NMR (CDCl_3): δ 180.48, 176.41, 144.86, 134.62, 132.77, 130.00, 126.58, 56.30, 54.77, 42.00, 31.56, 28.69, 26.93, 22.63; IR (cm^{-1}) 3207, 2920, 2846, 1674, 1562, 1262, 715; HRMS (+ESI): found m/z 333.1364, $[\text{M} + \text{H}]^+$; $\text{C}_{18}\text{H}_{22}\text{ClN}_2\text{O}_2$ required 333.1367.

2-Chloro-3-((2-chloroethyl)amino)-[1,4]-naphthoquinone (16). To a suspension of 2,3-dichloro-[1,4]-naphthoquinone (1.00 g, 4.40 mmol) in MeCN (40 mL) were added 2-chloroethylamine hydrochloride (0.52 g, 4.40 mmol) and diisopropylethylamine (1.53 mL, 8.80 mmol) and the resulting light orange mixture was stirred at r.t. for 72 h. The resulting mixture was concentrated *in vacuo*, diluted in EtOAc and washed with water, and the organic layer was dried over anhydrous Na_2SO_4 . The solvent was evaporated, and the resulting residue was purified *via* flash column chromatography (DCM/hexane = 1:4) to afford a red powder (0.73 g, 61%): mp: 166–168 °C; ^1H NMR (300 MHz, CDCl_3): δ 8.14 (d, $J = 7.1$ Hz, 1H), 8.05 (d, $J = 7.7$ Hz, 1H), 7.77 (t, $J = 8.9$ Hz, 1H), 7.67 (t, $J = 8.1$ Hz, 1H), 6.33 (br s, 1H), 4.23 (q, $J = 18.2$ Hz, 2H), 3.79 (t, $J = 11.2$ Hz, 2H); ^{13}C NMR (75 MHz, CDCl_3) δ 180.20, 176.89, 143.81, 135.01, 132.74, 132.42, 129.81, 126.96, 126.94, 45.93, 44.16; IR (cm^{-1}) 3284, 1679, 1564, 1511, 721, 678; HRMS (+ESI): found m/z 291.9903, $[\text{M} + \text{Na}]^+$; $\text{C}_{12}\text{H}_9\text{Cl}_2\text{NO}_2\text{Na}^+$ required 291.9902.

2-Chloro-3-((2-(1-bocpiperazinyl)ethyl)amino)-[1,4]-naphthoquinone (17). 2-chloro-3-((2-chloroethyl)amino)-[1,4]-naphthoquinone (0.59 g, 2.20 mmol), Boc-piperazine (1.32 g, 6.60 mmol), K_2CO_3 (0.33 g, 2.42 mmol), NaI (0.03 g, 0.22 mmol). Reaction heated at reflux for 22 h. The resulting mixture was concentrated *in vacuo*, diluted in EtOAc and washed with water, and the organic layer was dried over anhydrous Na_2SO_4 . The solvent was evaporated, and the resulting residue was purified *via* flash column chromatography (EtOAc/hexane = 1:4) to afford an orange solid (0.13 g, 14%): mp: 73–75 °C; ^1H NMR (300 MHz, CDCl_3)

δ 8.15 (dd, $J = 7.7, 1.4$ Hz, 1H), 8.02 (dd, $J = 7.6, 1.4$ Hz, 1H), 7.73 (td, $J = 7.6, 1.4$ Hz, 1H), 7.63 (td, $J = 7.5, 1.4$ Hz, 1H), 6.89 (s, 1H), 4.03–3.91 (m, 2H), 3.53–3.42 (m, 4H), 2.69 (t, $J = 6.0$ Hz, 2H), 2.48 (t, $J = 5.0$ Hz, 4H), 1.48 (s, 9H); ^{13}C NMR (75 MHz, CDCl_3) δ 180.58, 176.64, 171.12, 154.71, 144.63, 134.81, 132.72, 132.36, 129.93, 126.73, 126.68, 79.76, 56.41, 52.36, 40.93, 28.41, 22.64. IR (cm^{-1}) 3286, 2970, 2810, 1679, 1609, 1420, 1236, 1129, 1005, 720, 678; HRMS (+ESI): found m/z 442.1504, $[\text{M} + \text{Na}]^+$; $\text{C}_{21}\text{H}_{25}\text{ClN}_3\text{NaO}_4^+$ required 442.1502.

2-Chloro-3-(4-(2-((3-chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)amino)ethyl)piperazin-1-yl)-[1,4]-naphthoquinone (18). Following general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (1.14 g, 5.00 mmol), Et_2O (40 mL), 1-(2-aminoethyl)piperazine (0.66 mL, 5.00 mmol), Et_3N (0.70 mL, 5.00 mmol). Reaction stirred for 63 h. Product eluted at EtOAc/hexane = 1:9 in flash column chromatography to afford a dark brown solid (0.88 g, 29%): mp: 166–168 °C; ^1H NMR (300 MHz, CDCl_3): δ 8.17–8.17 (m, 2H), 8.15–8.11 (m, 2H), 7.76–7.61 (m, 4H), 7.26 (s, 1H), 4.04 (q, $J = 8.0$ Hz, 2H), 3.70 (t, $J = 9.5$ Hz, 4H), 2.77 (m, 6H); ^{13}C NMR (75 MHz, CDCl_3) δ 180.33, 174.06, 147.03, 134.71, 132.43, 132.32, 130.06, 126.97, 126.75, 67.06, 56.65, 52.94, 41.18; IR (cm^{-1}) 3326, 2890, 1733, 1667, 1557, 1246, 993, 718; HRMS (+ESI): found m/z 510.0982, $[\text{M} + \text{H}]^+$; $\text{C}_{26}\text{H}_{22}\text{O}_4\text{N}_3\text{Cl}_2$ required 510.0979.

2-((2-(1H-Indol-3-yl)ethyl)amino)-3-chloronaphthalene-1,4-dione (19).²² Following general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (1.14 g, 5.00 mmol), Et_2O (40 mL), tryptamine (0.9 g, 5.00 mmol), Et_3N (0.70 mL, 5.00 mmol). Reaction stirred for 12 h. Product eluted at EtOAc/hexane = 1:9 in flash column chromatography to afford a dark brown solid (1.19 g, 68%): mp: 189–190 °C; ^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 10.86 (s, 1H), 7.97 (d, $J = 7.6$ Hz, 1H), 7.91 (d, $J = 7.6$ Hz, 1H), 7.82 (td, $J = 7.5, 1.4$ Hz, 1H), 7.72 (td, $J = 7.5, 1.3$ Hz, 1H), 7.61 (d, $J = 7.7$ Hz, 1H), 7.50 (s, 1H), 7.31 (d, $J = 7.8$ Hz, 1H), 7.19 (d, $J = 2.3$ Hz, 1H), 7.08–7.03 (m, 1H), 7.01–6.96 (m, 1H), 4.03 (dt, $J = 8.2, 6.7$ Hz, 2H), 3.08–2.97 (m, 2H). ^{13}C NMR (101 MHz, $\text{DMSO}-d_6$) δ 180.66, 136.74, 135.30, 133.04, 127.56, 126.92, 126.20, 123.62, 121.48, 118.82, 111.86, 111.27, 46.11, 45.15. IR (cm^{-1}) 3281, 1679, 1596, 1565, 1516, 1338, 749; MS (+ESI): found m/z 351.10, $[\text{M} + \text{H}]^+$; $\text{C}_{20}\text{H}_{16}\text{ClN}_2\text{O}_2$ required 351.09.

2-Chloro-3-((3-morpholinopropyl)amino)naphthalene-1,4-dione (20).²³ Follow general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (0.50 g, 2.20 mmol), Et_2O (20 mL), 3-morpholinopropylamine (0.30 mL, 2.20 mmol), Et_3N (0.31 mL, 2.20 mmol). Reaction stirred for 17 h. Upon evaporating solvent following work-up a red solid was afforded (0.34 g, 47%): mp: 148–150 °C; ^1H NMR (300 MHz, CDCl_3) δ 8.19–8.14 (m, 1H), 8.14–8.01 (m, 1H), 7.74 (td, $J = 7.6, 1.4$ Hz, 1H), 7.68–7.61 (m, 1H), 7.55 (s, 1H), 4.05 (q, $J = 6.0$ Hz, 2H), 3.87 (s, 4H), 2.55 (s, 6H), 1.89 (s, 2H). ^{13}C NMR (75 MHz, $\text{DMSO}-d_6$) δ 180.70, 135.37, 133.07, 132.55, 126.97, 126.26, 66.46, 56.80, 53.86, 44.14, 26.81. IR (cm^{-1}) 2952, 2785, 1680, 1601, 1565, 1512, 1456, 1331, 1297, 1264, 1111, 722; MS (+ESI): found m/z 335.20, $[\text{M} + \text{H}]^+$; $\text{C}_{17}\text{H}_{20}\text{ClN}_2\text{O}_3$ required 335.11.



2-Chloro-3-((3-(piperidin-1-yl)propyl)amino)-[1,4]-naphthoquinone (21). Follow general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (0.50 g, 2.20 mmol), Et₂O (20 mL), *N*-(3-aminopropyl)piperidine (0.35 mL, 2.20 mmol), Et₃N (0.31 mL, 2.20 mmol). Reaction stirred for 17 h. Product eluted at EtOAc/hexane = 1:1 in flash column chromatography to afford a dark orange solid 27 (0.55 g, 75%): mp: 105–107 °C; ¹H NMR (300 MHz, CDCl₃) δ 8.18–8.13 (m, 1H), 8.05–8.01 (m, 1H), 7.72 (td, *J* = 7.5, 1.4 Hz, 2H), 7.61 (td, *J* = 7.6, 1.4 Hz, 1H), 4.10–3.92 (m, 2H), 2.52–2.40 (m, 6H), 1.90–1.76 (m, 2H), 1.69 (q, *J* = 5.6 Hz, 4H), 1.59–1.43 (m, 2H); ¹³C NMR (75 MHz, CDCl₃) δ 180.74, 134.64, 132.86, 132.11, 126.66, 126.63, 58.11, 55.02, 45.81, 25.98, 25.63, 24.51; IR (cm⁻¹) 3065, 2921, 2817, 2765, 1679; HRMS (+ESI): found *m/z* 333.1364, [M + H]⁺; C₁₈H₂₂ClN₂O₂ required 333.1365.

2-Chloro-3-((3-(dimethylamino)propyl)amino)naphthalene-1,4-dione (22). Follow general synthetic procedure A, 2,3-dichloro-1,4-naphthoquinone (0.51 g, 2.20 mmol), Et₂O (20 mL), 3-(dimethylamino)-1-propylamine (0.28 mL, 2.20 mmol), Et₃N (0.31 mL, 2.20 mmol). Reaction stirred for 19 h. Upon evaporating the solvent following the workup, the resulting orange oil was recrystallized from 70% EtOH to afford dark red crystals (0.18 g, 29%): mp: 88–92 °C; ¹H NMR (400 MHz, DMSO-*d*₆) δ 7.83–7.99 (m, 3H), 7.78 (td, *J* = 7.5, 1.4 Hz, 1H), 7.69 (td, *J* = 7.5, 1.4 Hz, 1H), 3.78 (q, *J* = 6.5 Hz, 2H), 2.38 (t, *J* = 6.5 Hz, 2H), 2.17 (s, 6H), 1.74 (p, *J* = 6.6 Hz, 2H). ¹³C NMR (101 MHz, DMSO-*d*₆) δ 180.53, 135.20, 132.94, 132.43, 126.82, 126.13, 57.23, 45.15, 44.02, 27.53; IR (cm⁻¹) 3076, 2928, 2816, 2763, 1675, 1556, 1264; HRMS (+ESI): found *m/z* 315.0871, [M + Na]⁺; C₁₅H₁₇O₂N₂ClNa⁺ required 315.0872.

2-(Butylamino)-3-chloro-[1,4]-naphthoquinone (23). Following general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (0.51 g, 2.20 mmol), EtOH (20 mL), butylamine (0.26 mL, 2.60 mmol), Et₃N (0.31 mL, 2.20 mmol). Reaction stirred for 18 h. Product eluted at DCM/hexane = 1:1 in flash column chromatography to afford red crystals (0.41 g, 71%): mp: 112–114 °C; ¹H NMR (300 MHz, CDCl₃): δ 8.12 (d, *J* = 1.3 Hz, 1H), 8.02 (d, *J* = 8.5 Hz, 1H), 7.70 (t, *J* = 8.9 Hz, 1H), 7.62 (t, *J* = 8.9 Hz, 1H), 6.04 (br s, 1H), 3.83 (t, *J* = 8.0 Hz, 2H), 1.75–1.66 (m, 2H), 1.50–1.42 (m, 2H), 0.99 (t, *J* = 12.0 Hz, 3H); ¹³C NMR (CDCl₃): δ 180.70, 176.80, 134.94, 132.84, 132.36, 129.75, 126.85, 126.80, 44.71, 33.04, 19.83, 13.74. IR (cm⁻¹) 3308, 2934, 2870, 1672, 1598, 1565, 1512, 1441, 1330, 1293, 719; MS (+ESI): found *m/z* 264.20, [M + H]⁺; C₁₄H₁₅ClNO₂ required 264.08.

2-Chloro-3-(prop-2-yn-1-ylamino)naphthalene-1,4-dione (24).²⁴ Following general synthetic procedure A, 2,3-dichloro-[1,4]-naphthoquinone (0.51 g, 2.20 mmol), EtOH (20 mL), propargylamine (0.17 mL, 2.60 mmol), Et₃N (0.31 mL, 2.20 mmol). Reaction stirred for 18 h. Product eluted at EtOAc/hexane = 1:1 in flash column chromatography to afford red solid (0.35 g, 65%): mp: 171–172 °C; ¹H NMR (400 MHz, CDCl₃) δ 8.17 (dd, *J* = 7.6, 1.3 Hz, 1H), 8.08 (dd, *J* = 7.7, 1.3 Hz, 1H), 7.76 (td, *J* = 7.6, 1.4 Hz, 1H), 7.67 (td, *J* = 7.6, 1.3 Hz, 1H), 6.05 (s, 1H), 4.66 (dd, *J* = 6.0, 2.5 Hz, 2H), 2.42 (t, *J* = 2.5 Hz, 1H); ¹³C

NMR (101 MHz, CDCl₃) δ 180.12, 177.00, 143.57, 134.96, 134.69, 132.79, 132.34, 129.82, 127.86, 126.93, 79.24, 73.43, 35.02. IR (cm⁻¹) 3319, 3246, 1676, 1596, 1568, 1509, 1297, 1139, 1070, 713; MS (+ESI): found *m/z* 246.10, [M + H]⁺; C₁₃H₉ClNO₂ required 246.03.

Ethyl (3-chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)glycinate (25).²⁵ Following general synthetic procedure A, L-glycine ethyl ester HCl (0.161 g, 1.1 mmol) and K₂CO₃ (35 mg, 0.26 mmol) were added to a stirred solution of 2,3-dichloro-[1,4]-naphthoquinone (0.249 g, 1.1 mmol) in 10% THF in EtOH (10 mL). Product eluted with DCM: hexane (1:1) in flash column chromatography as an orange solid (43 mg, 13%); mp: 128–129 °C; ¹H NMR (400 MHz, CDCl₃) δ 8.21–8.15 (m, 1H), 8.11–8.05 (m, 1H), 7.76 (m, 1H), 7.68 (m, 1H), 6.52 (s, 1H), 4.62 (d, *J* = 5.5 Hz, 2H), 4.32 (q, *J* = 7.1 Hz, 2H), 1.35 (t, *J* = 7.2 Hz, 3H); ¹³C NMR (101 MHz, CDCl₃) δ 180.10, 176.88, 169.53, 143.89, 134.89, 132.71, 132.34, 129.89, 126.89, 126.83, 62.07, 46.28, 14.18. IR (cm⁻¹) 3306, 2925, 2853, 1734, 1603, 1572, 1502, 1287, 1211, 725; MS (+ESI): found *m/z* 294.20, [M + H]⁺; C₁₄H₁₃ClNO₄ required 294.05.

Methyl (3-chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)-L-tryptophanate (26). Following general synthetic procedure A, L-tryptophan methyl ester hydrochloride (0.28 g, 1.1 mmol) and K₂CO₃ (37 mg, 0.27 mmol) were added to a solution of 2,3-dichloro-[1,4]-naphthoquinone (0.25 g, 1.1 mmol) in 10% THF in EtOH (5 mL). Product eluted from hexane: EtOAc (3:2) in flash column chromatography as a red solid (0.29 g, 65%): mp: 81.8–82.0 °C; ¹H NMR (300 MHz, CDCl₃) δ 8.19–8.13 (m, 1H), 8.14–8.09 (m, 1H), 7.94 (d, *J* = 7.7 Hz, 1H), 7.72 (td, *J* = 7.5, 1.4 Hz, 1H), 7.65–7.55 (m, 2H), 7.39–7.29 (m, 1H), 7.19 (m, 1H), 7.16–7.07 (m, 2H), 6.36 (s, 1H), 5.56 (dt, *J* = 8.2, 5.9 Hz, 1H), 3.76 (s, 3H), 3.52–3.38 (m, 2H); ¹³C NMR (101 MHz, CDCl₃) δ 172.41, 171.42, 136.51, 134.94, 132.82, 127.35, 127.04, 126.91, 123.52, 122.82, 120.18, 118.78, 111.58, 109.35, 60.66, 56.81, 53.01, 30.03, 21.32, 14.46; IR (cm⁻¹) 3302, 2919, 2850, 1734, 1672, 1592, 1563; HRMS (+ESI): found *m/z* 409.0951, [M + H]⁺; C₂₂H₁₈ClN₂O₄ required 409.0950.

Methyl (3-chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)-L-phenylalaninate (27). Following general synthetic procedure A, L-phenylalanine methyl ester hydrochloride (0.24 g, 1.1 mmol) and K₂CO₃ (26 mg, 0.19 mmol) were added to a solution of 2,3-dichloro-[1,4]-naphthoquinone (0.25 g, 1.1 mmol) in 10% THF in EtOH (5 mL). Product eluted with hexane: EtOAc (5:1) in flash column chromatography as an orange solid (20 mg, 5%); mp: 158.5–158.7 °C; ¹H NMR (300 MHz, CDCl₃) δ 8.17–8.11 (m, 1H), 8.06–7.98 (m, 1H), 7.74 (td, *J* = 7.5, 1.5 Hz, 1H), 7.68–7.61 (m, 1H), 7.34–7.26 (m, 3H), 7.21–7.15 (m, 2H), 6.27 (s, 1H), 5.58–5.49 (m, 1H), 3.79 (s, 3H), 3.35–3.17 (m, 2H). ¹³C NMR (101 MHz, CDCl₃) δ 180.07, 176.74, 171.71, 134.93, 134.83, 132.71, 129.30, 128.89, 127.58, 126.91, 126.79, 57.29, 52.69, 40.02; IR (cm⁻¹) 3312, 2950, 1738, 1593, 1567; HRMS (+ESI): found *m/z* 370.0842, [M + H]⁺; C₂₀H₁₇ClNO₄ required 370.0841.

2-Chloro-3-((2-(1-piperazinyl)ethyl)amino)-[1,4]-naphthoquinone salt (29). 2-Chloro-3-((2-(1-bocpiperazinyl)



ethyl)amino)-[1,4]-naphthoquinone **17** (0.50 g, 1.20 mmol) was dissolved in DCM (20 mL) and to this suspension was added TFA (3 mL). The resulting orange mixture was stirred at r.t. for 3 h. The mixture was evaporated *in vacuo* to afford an oil which was then recrystallized from Et₂O to give an orange salt (234.0 mg, 60%); mp: 198–200 °C; ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.87 (s, 2H), 7.96–8.00 (m, 2H), 7.85 (td, *J* = 7.5, 1.5 Hz, 1H), 7.77 (td, *J* = 7.5, 1.4 Hz, 1H), 7.37 (s, 1H), 3.94 (d, *J* = 6.3 Hz, 2H), 3.21 (s, 4H), 2.97 (s, 6H). ¹³C NMR (75 MHz, DMSO-*d*₆) δ 180.57, 175.91, 158.56, 135.42, 133.31, 132.31, 126.99, 126.31, 57.02, 49.31, 42.44; IR (cm⁻¹) 3323, 3015, 2479, 1675, 1130, 721; HRMS (+ESI): found *m/z* 320.1160, [M + H]⁺; C₁₆H₁₉ClN₃O₂ required 320.1164.

Methyl (3-chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)-l-lysinate (30). TFA (10 mL) was added to a solution of Boc-compound **16e** (0.30 g, 0.67 mmol) in DCM (10 mL) under nitrogen on ice. The reaction mixture was stirred at r.t. for 5 h. The solvent removed *in vacuo* yielding analogue **30** as a sticky solid (0.16 g, 50%). ¹H NMR (400 MHz, DMSO-*d*₆) δ 8.04–7.96 (m, 3H), 7.86 (td, *J* = 7.5, 1.4 Hz, 1H), 7.79 (td, *J* = 7.5, 1.3 Hz, 1H), 6.93 (d, *J* = 8.3 Hz, 1H), 5.08–5.02 (m, 1H), 3.71 (s, 3H), 3.57 (s, 1H), 2.78 (m, 2H), 2.06–1.86 (m, 2H), 1.56 (m, 2H), 1.43 (q, *J* = 7.7 Hz, 2H); ¹³C NMR (101 MHz, DMSO-*d*₆) δ 180.43, 172.61, 158.61, 135.51, 133.58, 131.95, 130.44, 127.13, 126.37, 66.81, 56.94, 52.88, 39.02, 32.18, 27.06, 22.63; IR (cm⁻¹) 2946, 1672, 1580, 1568; HRMS (+ESI): found *m/z* 351.1105, [M]⁺; C₁₇H₂₀ClN₂O₄ required 351.1106.

2-Chloro-3-(((1-phenyl-1H-1,2,3-triazol-4-yl)methyl)amino)naphthalene-1,4-dione (31). **24** (0.2 g, 0.8 mmol) was dissolved in 10 mL of THF:H₂O (5:1) under a nitrogen atmosphere, and azido benzene (0.19 g, 1.2 mmol), sodium ascorbate 20 mol%, and CuSO₄·5H₂O, 10 mol%, were added. The reaction mixture was stirred at 50 °C for 12 h and then quenched by addition of 15 mL ice water and extracted with ethyl acetate (15 mL × 3). The combined organic extracts were washed with brine, dried over anhydrous Na₂SO₄, and evaporated under reduced pressure. Purification was completed with flash column chromatography on silica using a DCM: MeOH solvent system to yield a red crystal (0.047 g, 16%); mp: 204.2 °C. ¹H NMR (600 MHz, DMSO-*d*₆) δ 8.62 (s, 1H), 8.01 (dt, *J* = 7.6, 1.5 Hz, 2H), 7.88–7.85 (m, 2H), 7.83 (td, *J* = 7.5, 1.4 Hz, 1H), 7.75 (td, *J* = 7.6, 1.3 Hz, 1H), 7.60–7.54 (m, 3H), 7.49–7.45 (m, 1H), 5.16 (d, *J* = 6.6 Hz, 2H). ¹³C NMR (151 MHz, DMSO-*d*₆) δ 180.61, 176.14, 146.77, 145.79, 137.27, 135.22, 133.19, 132.50, 130.66, 130.22, 129.00, 126.94, 126.30, 121.44, 120.62, 110.97, 40.45. HRMS (+ESI): found *m/z* 365.08011, [M + H]⁺; C₁₉H₁₄ClN₄O₂ requires 365.08053.

4-(4-(((3-Chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)amino)methyl)-1H-1,2,3-triazol-1-yl)benzoic acid (32). **24** (0.2 g, 0.8 mmol) was dissolved in 10 mL of THF:H₂O (5:1) under a nitrogen atmosphere, and 4-azido benzoic acid (0.3 g, 1.8 mmol), sodium ascorbate 20 mol%, and CuSO₄·5H₂O, 10 mol%, were added. The reaction mixture was stirred at 50 °C for 12 h and then quenched by addition of 15 mL ice water and extracted with ethyl acetate (15 mL × 3). The combined organic extracts were washed with brine, dried

over anhydrous Na₂SO₄, and evaporated under reduced pressure. Purification was completed with flash column chromatography on silica using a DCM: MeOH solvent system to yield as an orange powder (0.2 g, 89%); mp: 187.2 °C. ¹H NMR (600 MHz, DMSO-*d*₆) δ 8.73 (s, 1H), 8.14–8.10 (m, 2H), 8.03–7.99 (m, 4H), 7.83 (td, *J* = 7.5, 1.3 Hz, 1H), 7.76 (td, *J* = 7.5, 1.2 Hz, 1H), 7.60 (t, *J* = 6.7 Hz, 1H), 5.17 (d, *J* = 6.6 Hz, 2H). ¹³C NMR (151 MHz, DMSO-*d*₆) δ 180.61, 176.15, 166.72, 147.24, 145.77, 140.09, 135.23, 133.21, 132.49, 131.42, 131.30, 130.67, 126.96, 126.31, 121.57, 120.30, 111.01, 40.33. HRMS (+ESI): found *m/z* 409.06981, [M + H]⁺; C₂₀H₁₄ClN₄O₄ requires 409.07036.

Ethyl 4-(4-(((3-chloro-1,4-dioxo-1,4-dihydronaphthalen-2-yl)amino)methyl)-1H-1,2,3-triazol-1-yl)benzoate (33). **24** (0.2 g, 0.8 mmol) was dissolved in 10 mL of THF:H₂O (5:1) under a nitrogen atmosphere, and ethyl-4-azidobenzoate (0.33 g, 1.8 mmol), sodium ascorbate 20 mol%, and CuSO₄·5H₂O, 10 mol%, were added. The reaction mixture was stirred at 50 °C for 12 h and then quenched by addition of 15 mL ice water and extracted with ethyl acetate (15 mL × 3). The combined organic extracts were washed with brine, dried over anhydrous Na₂SO₄, and evaporated under reduced pressure. Purification was completed with flash column chromatography on silica using a DCM: MeOH solvent system to yield as a dark-orange crystals (0.03 g, 8%); mp: 194.0 °C. ¹H NMR (400 MHz, DMSO-*d*₆) δ 8.84 (s, 1H), 8.15–8.10 (m, 2H), 8.09–8.04 (m, 2H), 8.03–7.94 (m, 3H), 7.84 (td, *J* = 7.5, 1.4 Hz, 1H), 7.77 (td, *J* = 7.5, 1.4 Hz, 1H), 5.12 (d, *J* = 6.8 Hz, 2H), 4.34 (q, *J* = 7.1 Hz, 2H), 1.34 (t, *J* = 7.1 Hz, 3H). ¹³C NMR (101 MHz, DMSO-*d*₆) δ 180.52, 176.14, 165.30, 147.93, 147.01, 140.21, 135.32, 133.29, 132.34, 131.38, 130.61, 128.80, 129.97, 127.02, 126.29, 121.49, 120.16, 61.56, 49.07 (MeOH), 14.60. HRMS (+ESI): found *m/z* 459.0830, [M + Na]⁺; C₂₂HClN₄O₄Na requires 459.0836.

N-(1,4-Dioxo-3-(((3-(piperidin-1-yl)propyl)amino)-1,4-dihydronaphthalen-2-yl)acetamide (39). Following general synthetic method A, aminopropylpiperidine (0.35 mL, 2.2 mmol) and Et₃N (0.31 mL, 2.2 mmol) were added to a solution of amide **35** (0.55 g, 2.2 mmol) in diethyl ether (20 mL). The reaction mixture was refluxed for 18 h. The product purified by flash column chromatography using gradient elution (0–15% MeOH in DCM) to give a red solid (0.16 g, 21%); mp: 285–296 °C; ¹H NMR (400 MHz, CDCl₃) δ 8.09–7.99 (m, 2H), 7.67 (t, *J* = 7.6 Hz, 1H), 7.57 (t, *J* = 7.6 Hz, 1H), 7.33 (s, 1H), 3.58 (s, 2H), 2.43 (s, 6H), 2.22 (s, 3H), 1.78 (s, 2H), 1.70 (s, 4H), 1.50 (s, 2H); ¹³C NMR (101 MHz, CDCl₃) δ 182.12, 179.35, 170.83, 143.37, 134.49, 132.57, 132.08, 130.51, 126.40, 126.10, 111.49, 57.44, 54.75, 50.65, 43.22, 25.80, 25.35, 24.30, 23.41; IR (cm⁻¹) 3337, 2928, 1662, 1605, 1600; HRMS (+ESI): found *m/z* 378.1784, [M + Na]⁺; C₂₀H₂₅N₃O₃Na requires 378.1788.

N-(1,4-Dioxo-3-(((3-(piperidin-1-yl)propyl)amino)-1,4-dihydronaphthalen-2-yl)propionamide (40). Following general synthetic method A, aminopropylpiperidine (0.14 mL, 0.9 mmol) and Et₃N (0.12 mL, 0.9 mmol) were added to amide **36** (0.24 g, 0.9 mmol) in THF (8 mL) and ether (3 mL). The



reaction mixture was refluxed for 18 h. The organic layer was collected, dried with sodium sulfate and the solvent evaporated yielding product as a red solid (0.37 g, 100%); mp: 101.1–101.8 °C; ^1H NMR (400 MHz, CDCl_3) δ 8.06–7.98 (m, 2H), 7.67 (t, $J = 7.5$ Hz, 1H), 7.60–7.54 (m, 1H), 7.49 (s, 1H), 7.35–7.30 (m, 1H), 3.58–3.51 (m, 2H), 2.48 (t, $J = 7.6$ Hz, 2H), 2.43–2.34 (m, 6H), 2.16 (s, 3H), 1.75 (q, $J = 6.3$ Hz, 2H), 1.67 (t, $J = 5.7$ Hz, 4H), 1.48 (q, $J = 5.8$ Hz, 2H), 1.25 (td, $J = 7.3, 4.3$ Hz, 4H); ^{13}C NMR (101 MHz, CDCl_3) δ 206.99, 182.18, 179.39, 174.40, 143.05, 134.48, 132.56, 132.09, 130.56, 126.42, 126.08, 111.54, 60.40, 57.68, 54.87, 43.37, 30.94, 30.32, 29.93, 26.02, 25.61, 24.48, 21.06, 14.20, 9.92; IR (cm^{-1}) 3339, 2928, 1669, 1605, 1572; HRMS (+ESI): found m/z 370.2127, $[\text{M} + \text{H}]^+$; $\text{C}_{21}\text{H}_{28}\text{N}_3\text{O}_3$ required 370.2125.

N-(1,4-Dioxo-3-((3-(piperidin-1-yl)propyl)amino)-1,4-dihydronaphthalen-2-yl)butyramide (**41**). Following general synthetic method A, aminopropylpiperidine (0.07 mL, 0.36 mmol) and Et_3N (0.06 mL, 0.36 mmol) were added to a solution of amide **37** (0.10 g, 0.36 mmol) in THF (10 mL) and Et_2O (10 mL). The reaction mixture was refluxed for 24 h. The product eluted from gradient column (0–20% MeOH in DCM) as a dark red solid (61 mg, 44%). ^1H NMR (400 MHz, CDCl_3) δ 8.11–7.99 (m, 2H), 7.69 (td, $J = 7.6, 1.3$ Hz, 1H), 7.60 (td, $J = 7.6, 1.4$ Hz, 1H), 7.43 (s, 1H), 7.36 (s, 1H), 3.60 (q, $J = 5.3$ Hz, 2H), 2.59–2.48 (m, 4H), 2.47 (d, $J = 7.4$ Hz, 3H), 1.82 (dt, $J = 6.5, 2.5$ Hz, 3H), 1.73 (q, $J = 5.6$ Hz, 4H), 1.52 (s, 2H), 1.05 (t, $J = 7.4$ Hz, 3H); ^{13}C NMR (101 MHz, CDCl_3) δ 182.26, 179.29, 173.73, 142.87, 134.48, 132.45, 132.12, 130.59, 126.98, 126.40, 126.08, 111.98, 57.13, 54.63, 54.47, 42.95, 38.79, 28.70, 25.82, 25.11, 24.12, 21.24, 19.20, 13.93; IR (cm^{-1}) 2929, 1662, 1604, 1590; HRMS (+ESI): found m/z 384.2279, $[\text{M} + \text{H}]^+$; $\text{C}_{22}\text{H}_{30}\text{N}_3\text{O}_3$ required 384.2282.

N-(1,4-Dioxo-3-((3-(piperidin-1-yl)propyl)amino)-1,4-dihydronaphthalen-2-yl)pentanamide (**42**). Following general synthetic method A, aminopropylpiperidine (57 μL , 0.37 mmol) and Et_3N (53 μL , 0.37 mmol) were added to a solution of amide **38** (0.15 g, 0.37 mmol) in THF (6 mL) in Et_2O (6 mL). The reaction mixture was refluxed for 18 h. The product eluted from gradient column (0–15% MeOH in DCM) as a dark red solid (0.10 g, 51%). ^1H NMR (400 MHz, CDCl_3) δ 8.08–7.97 (m, 2H), 7.70–7.63 (m, 1H), 7.57 (td, $J = 7.6, 1.4$ Hz, 1H), 7.31 (s, 1H), 3.58 (d, $J = 5.8$ Hz, 2H), 2.45 (t, $J = 7.7$ Hz, 4H), 1.80 (s, 2H), 1.72 (m, 6H), 1.47 (s, 2H), 1.41 (p, $J = 7.4$ Hz, 2H), 0.94 (t, $J = 7.4$ Hz, 3H); ^{13}C NMR (101 MHz, CDCl_3) δ 182.22, 173.74, 134.41, 132.38, 132.05, 130.56, 126.33, 126.01, 54.63, 42.95, 36.60, 27.74, 25.81, 24.13, 22.45, 13.79; IR (cm^{-1}) 3233, 2929, 2855, 1671, 1605, 1565; HRMS (+ESI): found m/z 398.2433, $[\text{M} + \text{H}]^+$; $\text{C}_{23}\text{H}_{32}\text{N}_3\text{O}_3$ required 398.2438.

N-(3-((2-Morpholinoethyl)amino)-1,4-dioxo-1,4-dihydronaphthalen-2-yl)acetamide (**43**). Following general synthetic method A, ethylaminomorpholine (0.30 mL, 2.2 mmol) and Et_3N (0.31 mL, 2.2 mmol) were added to amide **35** (0.55 g, 2.2 mmol) in diethyl ether (20 mL). The reaction mixture was refluxed for 18 h. Product eluted from gradient column (0–15% MeOH in DCM) as a red solid (0.25 g, 34%); mp: 147.0–147.1 °C; ^1H NMR (400 MHz, CDCl_3) δ 8.12–8.01

(m, 2H), 7.72 (t, $J = 7.5$ Hz, 1H), 7.66–7.60 (m, 1H), 7.24 (s, 1H), 6.86 (s, 1H), 3.77 (t, $J = 4.7$ Hz, 4H), 3.60 (d, $J = 7.8$ Hz, 2H), 2.63 (s, 2H), 2.52 (s, 4H), 2.27 (s, 3H); ^{13}C NMR (101 MHz, CDCl_3) δ 182.23, 170.67, 142.37, 134.63, 132.38, 130.55, 126.46, 126.14, 66.95, 56.87, 53.08, 39.11, 23.57; IR (cm^{-1}) 3463, 3166, 2821, 1658, 1604, 1569; HRMS (+ESI): found m/z 344.1603, $[\text{M} + \text{H}]^+$; $\text{C}_{18}\text{H}_{22}\text{N}_3\text{O}_4$ required 344.1605.

2-Methyl-1-(3-(piperidin-1-yl)propyl)-1H-naphtho[2,3-d]imidazole-4,9-dione (**44**). Following general procedure E, aqueous NaOH (2 M, 2 drops) was added to a solution of amide **39** (100 mg, 0.3 mmol) in EtOH (2 mL). The reaction mixture was heated at 50 °C for 1 h. The solvent was removed *in vacuo* yielding a yellowy-green solid (50 mg, 53%); mp: 125.6–126.0 °C; ^1H NMR (400 MHz, CDCl_3) δ 8.27–8.10 (m, 2H), 7.78–7.66 (m, 2H), 4.44 (t, $J = 7.1$ Hz, 2H), 2.63 (s, 3H), 2.35 (t, $J = 6.5$ Hz, 6H), 2.08–2.00 (m, 2H), 1.56 (q, $J = 5.6$ Hz, 4H), 1.45 (d, $J = 5.7$ Hz, 2H); ^{13}C NMR (101 MHz, CDCl_3) δ 179.06, 176.14, 153.48, 143.41, 133.65, 133.37, 133.20, 132.83, 132.21, 126.98, 126.44, 55.41, 54.49, 44.05, 27.22, 25.92, 24.40, 13.30; IR (cm^{-1}) 2934, 2765, 1656, 1583; HRMS (+ESI): found m/z 338.1860, $[\text{M} + \text{H}]^+$; $\text{C}_{20}\text{H}_{24}\text{N}_3\text{O}_2$ required 338.1863.

2-Ethyl-1-(3-(piperidin-1-yl)propyl)-1H-naphtho[2,3-d]imidazole-4,9-dione (**45**). Following general synthetic method E, aqueous NaOH (2 M, 6 drops) was added to a solution of amide **40** (130 mg, 0.4 mmol) in EtOH (12 mL). The reaction mixture was heated at 50 °C for 2 h. The solvent was removed *in vacuo* yielding a yellowy-green solid (67 mg, 54%). ^1H NMR (300 MHz, CDCl_3) δ 8.31–8.19 (m, 1H), 8.19–8.08 (m, 1H), 7.79–7.65 (m, 2H), 4.50–4.36 (m, 2H), 2.93 (q, $J = 7.5$ Hz, 2H), 2.36 (t, $J = 6.6$ Hz, 6H), 2.02 (p, $J = 6.9$ Hz, 3H), 1.56 (q, $J = 5.5$ Hz, 4H), 1.47 (t, $J = 7.5$ Hz, 5H), 1.34–1.20 (m, 1H); ^{13}C NMR (101 MHz, CDCl_3) δ 179.10, 176.21, 157.84, 143.46, 133.59, 133.31, 133.18, 132.83, 132.15, 126.95, 126.37, 55.48, 54.51, 43.72, 27.54, 25.95, 24.40, 20.24, 11.83; IR (cm^{-1}) 2935, 2768, 1653, 1584; HRMS (+ESI): found m/z 352.2016, $[\text{M} + \text{H}]^+$; $\text{C}_{21}\text{H}_{26}\text{N}_3\text{O}_2$ required 352.2020.

1-(3-(Piperidin-1-yl)propyl)-2-propyl-1H-naphtho[2,3-d]imidazole-4,9-dione (**46**). Following general synthetic method E, aqueous NaOH (2 M, 4 drops) was added to a solution of amide **41** (31 mg, 0.08 mmol) in EtOH (5 mL). The reaction mixture was heated at 50 °C for 2 h. The solvent was removed *in vacuo* yielding a dark green solid (39 mg, 100%). ^1H NMR (400 MHz, CDCl_3) δ 8.28–8.17 (m, 1H), 8.17–8.05 (m, 1H), 7.77–7.67 (m, 2H), 4.48–4.40 (m, 1H), 3.62 (q, $J = 7.1$ Hz, 1H), 2.92–2.86 (m, 1H), 2.38 (d, $J = 5.5$ Hz, 4H), 2.07–1.96 (m, 2H), 1.89–1.80 (m, 2H), 1.55 (s, 3H), 1.55–1.40 (m, 4H), 1.03–0.90 (m, 3H); ^{13}C NMR (101 MHz, CDCl_3) δ 176.21, 157.11, 133.61, 133.35, 133.20, 132.84, 126.97, 126.40, 77.24, 58.23, 55.46, 54.49, 43.75, 29.87, 27.54, 26.56, 25.85, 24.34, 22.65, 20.18, 18.34, 13.86, 13.77; IR (cm^{-1}) 2829, 2869, 1655, 1587; HRMS (+ESI): found m/z 366.2173, $[\text{M} + \text{H}]^+$; $\text{C}_{22}\text{H}_{28}\text{N}_3\text{O}_2$ required 366.2176.

2-Butyl-1-(3-(piperidin-1-yl)propyl)-1H-naphtho[2,3-d]imidazole-4,9-dione (**47**). Following general synthetic method E, aqueous NaOH (2 M, 4 drops) was added to a solution of amide **42** (64 mg, 0.16 mmol) in EtOH (5 mL).



The reaction mixture was heated at 50 °C for 2 h. The product was purified by flash column chromatography using gradient elution (0–5% MeOH in DCM) to give a yellow sticky solid (29 mg, 47%). ¹H NMR (400 MHz, CDCl₃) δ 8.22 (m, 1H), 8.15–8.07 (m, 1H), 7.75–7.65 (m, 2H), 4.43 (t, *J* = 7.4 Hz, 2H), 2.87 (m, 2H), 2.46 (s, 4H), 2.08–2.00 (m, 2H), 1.84 (td, *J* = 8.1, 1.7 Hz, 2H), 1.62 (dd, *J* = 8.2, 3.3 Hz, 4H), 1.51–1.39 (m, 4H), 0.97 (td, *J* = 7.4, 1.0 Hz, 3H); ¹³C NMR (101 MHz, CDCl₃) δ 206.98, 179.09, 176.25, 157.14, 143.59, 133.63, 133.33, 133.19, 132.86, 131.99, 127.00, 126.38, 55.27, 54.35, 53.43, 50.83, 43.61, 30.93, 29.87, 27.27, 26.57, 25.49, 24.09, 22.65, 13.79; IR (cm⁻¹) 2928, 2853, 1620, 1589; HRMS (+ESI): found *m/z* 380.2331, [M + H]⁺; C₂₃H₃₀N₃O₂ required 380.2333.

2-Methyl-1-(2-morpholinoethyl)-1H-naphtho[2,3-*d*]imidazole-4,9-dione (48). Following general synthetic method E, aqueous NaOH (2 M, 3 drops) was added to a solution of amide **43** (65 mg, 0.2 mmol) in EtOH (4 mL). The reaction mixture was heated at 50 °C for 1 h. The solvent was removed *in vacuo* yielding a black-green solid (68 mg, 100%); mp: 133.3–133.8 °C; ¹H NMR (400 MHz, CDCl₃) δ 8.30–8.21 (m, 1H), 8.18–8.09 (m, 1H), 7.79–7.68 (m, 2H), 4.50 (t, *J* = 6.4 Hz, 2H), 3.68–3.64 (m, 4H), 2.76 (t, *J* = 6.4 Hz, 2H), 2.63 (s, 3H), 2.58–2.51 (m, 4H); ¹³C NMR (101 MHz, CDCl₃) δ 178.99, 176.33, 153.55, 143.24, 133.77, 133.45, 133.10, 132.81, 132.15, 127.04, 126.42, 66.94, 58.40, 54.03, 43.40, 13.52; IR (cm⁻¹) 2948, 2931, 1656, 1589; HRMS (+ESI): found *m/z* 348.1313, [M + H]⁺; C₁₈H₁₉N₃O₃Na required 348.1319.

***N*-(1,4-Dioxo-3-((3-(piperidin-1-yl)propyl)amino)-1,4-dihydronaphthalen-2-yl)-2-phenylacetamide (50).** Following general synthetic method A, aminopropylpiperidine (0.36 mL, 2.4 mmol) and Et₃N (0.34 mL, 2.4 mmol) were added to a solution of amide **49** (0.78 g, 2.4 mmol) in THF (20 mL). The reaction mixture was refluxed for 18 h. The product was purified by flash column chromatography using gradient elution (0–10% MeOH in DCM) to give a red solid (0.71 g, 69%). ¹H NMR (400 MHz, CDCl₃) δ 8.08–7.99 (m, 2H), 7.69 (td, *J* = 7.5, 1.4 Hz, 1H), 7.59 (td, *J* = 7.6, 1.3 Hz, 1H), 7.48–7.38 (m, 4H), 7.36–7.30 (m, 1H), 7.22 (s, 1H), 3.83 (s, 2H), 3.42 (s, 2H), 2.49 (s, 6H), 1.76 (s, 6H), 1.49 (s, 2H); ¹³C NMR (101 MHz, CDCl₃) δ 182.32, 179.32, 171.58, 143.23, 134.93, 134.72, 132.64, 132.31, 130.75, 129.84, 129.23, 127.65, 126.61, 126.28, 54.73, 44.33, 43.10, 25.66, 24.24; IR (cm⁻¹) 3239, 2931, 2771, 1604, 1568; HRMS (+ESI): found *m/z* 432.2280, [M + H]⁺; C₂₆H₃₀N₃O₃ required 432.2282.

1,4-Dioxo-3-((3-(piperidin-1-yl)propyl)amino)-1,4-dihydronaphthalen-2-ammonium chloride (53). HCl-dioxane (4 M, 1.1 mL, 4.4 mmol) was added to a solution of **52** (0.42 g, 0.81 mmol) in diethyl ether (10 mL) under nitrogen on ice. The reaction mixture was stirred at r.t. for 18 h. The resulting brown precipitate was collected (0.30 g, 100%); mp: 228.4–228.7 °C; ¹H NMR (400 MHz, DMSO-*d*₆) δ 7.85 (dd, *J* = 7.1, 1.7 Hz, 2H), 7.73–7.63 (m, 2H), 3.38 (d, *J* = 12.2 Hz, 2H), 3.24 (t, *J* = 6.9 Hz, 2H), 3.05 (dd, *J* = 9.4, 6.6 Hz, 2H), 2.81 (t, *J* = 12.3 Hz, 2H), 1.82 (m, 4H), 1.60 (m, 4H), 1.38 (m, 2H); ¹³C NMR (101 MHz, DMSO-*d*₆) δ 180.14, 178.49, 135.21, 133.78, 131.49, 130.46, 126.42, 126.10, 83.49, 79.25, 78.92, 53.90, 52.83,

52.72, 44.18, 31.18, 31.03, 27.82, 22.87, 22.34, 21.44; IR (cm⁻¹) 3369, 3080, 2936, 2482, 1622, 1561; HRMS (+ESI): found *m/z* 314.1860, [M]⁺; C₁₈H₂₄N₃O₂ required 314.1863.

8.2. Cytotoxicity assays

Cell lines. HEC-1-A endometrial cancer and Mia-Pa-Ca-2 pancreatic cancer cell lines were sourced from the American Type Culture Collection (ATCC). The CALU-1 lung squamous cell carcinoma cell line was provided by Dr. Joshua McCarroll from the Children's Cancer Institute, Australia. The MAD11 endometrial stromal cell line, derived from non-cancerous patients and immortalized using human telomerase reverse transcriptase (hTERT), was obtained from Dr. Hui Li at the University of Virginia.

Reagents and Equipment. Phosphate buffered saline (Sigma Aldrich), trypsin-EDTA Solution 10× (Sigma Aldrich), Dulbecco's modified Eagle's medium with 10% foetal bovine serum (Sigma Aldrich), minimum essential medium (ThermoFisher Scientific), MTT Reagent (5 mg mL⁻¹ MTT in PBS), MTT Solvent (4 mM HCl, 0.1% Nondet P-40, in isopropanol), dimethyl sulfoxide (DMSO; Sigma Aldrich), *N*-acetylcysteine (NAC, Sigma Aldrich) and reduced glutathione (GSH, Sigma Aldrich).

Methods. Cytotoxicity assays were performed as previously described.⁷ Briefly, HEC1A (5000/well) and MAD11 (2500/well) cells were seeded in 50 μL of media in 96-well tissue culture plates and incubated at 37 °C overnight prior to drug treatment. All compounds, except NAC and GSH, were reconstituted in dimethyl sulfoxide (DMSO), serially diluted in cell culture medium (to final concentrations of 160 μM, 80 μM, 40 μM, 20 μM, 10 μM, 5 μM, 2.5 μM, 1.25 μM) and 50 μL of each was added to wells in triplicate. For the thiol rescue experiments, NAC and GSH were prepared in cell culture media (neutralised to pH 7.4) and the cells were treated with 2 mM of each for 40 minutes prior to the addition of the compounds (BH10, 21, and 44). In order to eliminate variations caused by the presence of DMSO (vehicle), the drug dilutions and control wells were treated with the same final concentration of DMSO. The maximum concentration of DMSO added to the cells did not exceed 0.5% v/v. Cell viability was then determined after 48 h by the addition of 20 μL per well of 5 mg mL⁻¹ thiazolyl blue tetrazolium bromide (MTT) reagent (3-hour incubation). Formazan crystals were solubilized in solvent (4 mM HCl, 0.1% NP-40 in isopropanol) and absorbance read at 590/620 nm using an EnSight Multimode plate reader (PerkinElmer). Viability of drug-treated cells is displayed as a percentage of control cells *i.e.* cells with equivalent concentrations of the appropriate vehicle (DMSO). The same experimental protocol used for HEC-1-A cells was applied to the CALU-1 and Mia-Pa-Ca-2 cancer cell lines, with the following modifications: CALU-1 cells were cultured in minimum essential medium (MEM) instead of Dulbecco's modified Eagle medium (DMEM).

Calculations of IC₅₀ values, selectivity and clogP. The half maximal inhibitory concentration (IC₅₀), that is, the amount



of compound required to inhibit cell viability by 50% relative to vehicle controls, was calculated using GraphPad Prism software. The selectivity ratio is the IC_{50} value of the non-cancerous cells (MAD11) divided by the IC_{50} value of the cancer cells (HEC1A), for each compound.

8.3. Oxygen consumption rate assay

Reagents and Equipment. Seahorse XF96 analyser calibrant solution, Seahorse XF media pH 7.4 (DMEM (8.3 g L^{-1}), sodium pyruvate (1 mM), L-glutamine (4 mM), glucose (25 mM) Sigma-Aldrich).

Method. The methods used were adapted from those described by Byrne *et al.*, 2014.²⁶ HEC1A cells were seeded in Seahorse 96-well tissue culture plates at a density of 15 000 cells per well in complete media (DMEM supplemented with 10% (v/v) FBS, and 0.1% penicillin (10 000 IU mL^{-1})/streptomycin (10 mg mL^{-1}). Cells were then allowed to adhere for 24 hours in a 37 °C humidified incubator with 5% CO_2 . Prior to the commencement of the assay, cell culture media was replaced with Seahorse media (unbuffered phenol-red free DMEM supplemented with 25 mM glucose, 4 mM L-glutamine, and 1 mM sodium pyruvate, pH 7.4) and cells were incubated at 37 °C in a non- CO_2 incubator for 40 minutes to allow for de-gassing. The cartridge plate was then equilibrated in the Seahorse XF analyser for 20 min, prior to adding the cell culture plate. Baseline measurements were recorded 3 times prior to the injection of the compounds. Oxygen Consumption Rate (OCR) was measured subsequently for 11 cycles, each consisting of a 3-minute waiting, 3-minute mixing, and 2-minute measurement period. Data from Seahorse assays was acquired using the Agilent Seahorse Wave Desktop (V2.6) software. The OCR was normalised to the third baseline reading.

8.4. Pharmacokinetic profiling *in vivo*

Reagents and Equipment. Lithium heparinised tubes (Sarstedt), centrifuge (4 °C, 2000 × *g*, 10 min), 10% MeOH in ACN (Sigma-Aldrich), dimethyl sulphoxide (DMSO, Sigma-Aldrich), Tween-80 (2%) (Sigma-Aldrich), captisol (20%) (ligand technology). Mass spectrometry was performed using UHPLC-MS/MS (Thermo Fischer Scientific). Data processing of chromatograms (integration) was performed using Xcalibur software package version 2.0.7 (Thermo Fisher Scientific) and exported to Excel. Mice were obtained from Australian BioResources Ltd. All animal procedures were performed in accordance with the Guidelines for Care and Use of Laboratory Animals of University of New South Wales (UNSW) and approved by the UNSW Animal Ethics Committee (ethics numbers 20/43A and 18/76A).

Method. Compounds were delivered (oral gavage) and blood was extracted (14 μL) at various time points up to 3 h. Blood was also collected before treatment from separate mice to prepare 'spiked' drug concentrations (0, 0.1, 1, 10 ng mL^{-1}) in order to generate a standard curve. All samples were then spun down using a centrifuge (4 °C, 2000 × *g*, 10 min)

and serum (7 μL) was extracted. Serum was then added to a solution of 10% MeOH in ACN (100 μL) to precipitate plasma proteins. Following another spin (4 °C, 2000 × *g*, 10 min), supernatant was collected (60 μL) and MS was performed. MS gave an area per time point. The equation of the line from standard curve of samples with known concentrations converted value of unknown sample to concentration. Concentration was then plotted against time show the concentration of drug in plasma over time.

8.5. *In silico* molecular docking study

The Keap1 protein (PDB code: 4XMB) was obtained from the Protein Data Bank. Molecular docking simulations were performed using Maestro software (Schrödinger, New York, NY, USA) in standard precision mode. All workflows, including protein preparation, LigPrep, and receptor grid generation, were executed with default parameter settings. The predicted protein-ligand complexes were further analyzed in Maestro and visualized using PyMOL (Schrödinger, New York, NY, USA).

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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