

Cite this: *RSC Med. Chem.*, 2020, 11, 392

# Isoquinoline thiosemicarbazone displays potent anticancer activity with *in vivo* efficacy against aggressive leukemias†‡

Daniel L. Sun,<sup>§abc</sup> Soumya Poddar,<sup>§ab</sup> Roy D. Pan,<sup>id abc</sup> Ethan W. Rosser,<sup>id abc</sup> Evan R. Abt,<sup>ab</sup> Juno Van Valkenburgh,<sup>abc</sup> Thuc M. Le,<sup>ab</sup> Vincent Lok,<sup>a</sup> Selena P. Hernandez,<sup>c</sup> Janet Song,<sup>a</sup> Joanna Li,<sup>a</sup> Aneta Turlik,<sup>c</sup> Xiaohong Chen,<sup>c</sup> Chi-An Cheng,<sup>id cd</sup> Wei Chen,<sup>c</sup> Christine E. Mona,<sup>ab</sup> Andreea D. Stuparu,<sup>ab</sup> Laurent Vergnes,<sup>e</sup> Karen Reue,<sup>ef</sup> Robert Damoiseaux,<sup>g</sup> Jeffrey I. Zink,<sup>id c</sup> Johannes Czernin,<sup>ab</sup> Timothy R. Donahue,<sup>abh</sup> Kendall N. Houk,<sup>id c</sup> Michael E. Jung<sup>\*c</sup> and Caius G. Radu<sup>id \*ab</sup>

A potent class of isoquinoline-based  $\alpha$ -N-heterocyclic carboxaldehyde thiosemicarbazone (HCT) compounds has been rediscovered; based upon this scaffold, three series of antiproliferative agents were synthesized through iterative rounds of methylation and fluorination modifications, with anticancer activities being potentiated by physiologically relevant levels of copper. The lead compound, HCT-13, was highly potent against a panel of pancreatic, small cell lung carcinoma, prostate cancer, and leukemia models, with IC<sub>50</sub> values in the low-to-mid nanomolar range. Density functional theory (DFT) calculations showed that fluorination at the 6-position of HCT-13 was beneficial for ligand-copper complex formation, stability, and ease of metal-center reduction. Through a chemical genomics screen, we identify DNA damage response/replication stress response (DDR/RSR) pathways, specifically those mediated by ataxia-telangiectasia and Rad3-related protein kinase (ATR), as potential compensatory mechanism(s) of action following HCT-13 treatment. We further show that the cytotoxicity of HCT-13 is copper-dependent, that it promotes mitochondrial electron transport chain (mtETC) dysfunction, induces production of reactive oxygen species (ROS), and selectively depletes guanosine nucleotide pools. Lastly, we identify metabolic hallmarks for therapeutic target stratification and demonstrate the *in vivo* efficacy of HCT-13 against aggressive models of acute leukemias in mice.

Received 26th December 2019,  
Accepted 12th February 2020

DOI: 10.1039/c9md00594c

rsc.li/medchem

<sup>a</sup> Department of Molecular and Medical Pharmacology, University of California, Los Angeles, California 90095, USA. E-mail: cradu@mednet.ucla.edu

<sup>b</sup> Ahmanson Translational Imaging Division, University of California, Los Angeles, California 90095, USA

<sup>c</sup> Department of Chemistry and Biochemistry, University of California, Los Angeles, California 90095, USA. E-mail: jung@chem.ucla.edu

<sup>d</sup> Department of Bioengineering, University of California, Los Angeles, CA 90095, USA

<sup>e</sup> Department of Human Genetics, David Geffen School of Medicine, University of California, Los Angeles, California 90095, USA

<sup>f</sup> Molecular Biology Institute, University of California, Los Angeles, California 90095, USA

<sup>g</sup> UCLA Metabolomic Center, University of California, Los Angeles, Los Angeles, California 90095, USA

<sup>h</sup> Department of Surgery, University of California, Los Angeles, CA 90095, USA

† The authors declare the following competing financial interest(s): C. G. R., J. C., and M. E. J. are co-founders of Trethera Corporation. They and the University of California hold equity in Trethera Corporation. The University of California has patented additional intellectual property for thiosemicarbazone-based antiproliferative compounds.

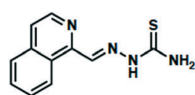
‡ Electronic supplementary information (ESI) available. See DOI: 10.1039/c9md00594c

§ These authors contributed equally. EWR, RDP, and CGR wrote the manuscript.

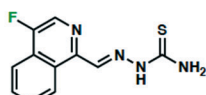
## Introduction

The diverse therapeutic potential of  $\alpha$ -N-heterocyclic carboxaldehyde thiosemicarbazones (HCTs) has been investigated since the 1940s, with tuberculostatic activity being first observed *in vivo* as early as 1946.<sup>1</sup> This class of compounds was subsequently shown to possess antitumor and antimicrobial activities, prompting decades of research and development.<sup>2–7</sup> In particular, isoquinoline-based HCTs such as 2-(isoquinolin-1-ylmethylene)hydrazine-1-carbothioamide (IQ-1, HCT-1)<sup>3</sup> were the subject of early interest due to evidence of preclinical anti-tumor efficacy (Fig. 1).<sup>5</sup> While the mechanisms of action of HCTs are multimodal and have not yet been fully defined,<sup>8–15</sup> their biological activities generally stem from the ability to chelate transition metals through their heterocyclic nitrogen, Schiff base nitrogen, and thiosemicarbazone sulfur. The resulting HCT-metal complexes undergo redox cycles and generate cytotoxic reactive oxygen species (ROS) through Fenton and/or Haber-

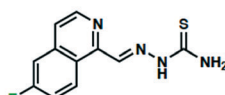
## 4' primary amines



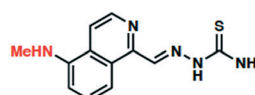
**HCT-1 (IQ-1)**  
 $2.21 \pm 0.12 \mu\text{M}$   
 ( $18.1 \pm 1.8 \mu\text{M}$ )



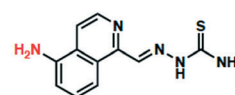
**HCT-2**  
 $2.06 \pm 0.08 \mu\text{M}$   
 ( $24.6 \pm 3.5 \mu\text{M}$ )



**HCT-3**  
 $2.04 \pm 0.34 \mu\text{M}$   
 ( $5.44 \pm 0.42 \mu\text{M}$ )

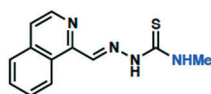


**HCT-4**  
 $5.33 \pm 1.57 \mu\text{M}$   
 ( $11.5 \pm 0.6 \mu\text{M}$ )

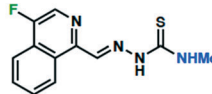


**HCT-5**  
 $71.1 \pm 0.1 \mu\text{M}$   
 ( $40.5 \pm 6.9 \mu\text{M}$ )

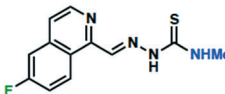
## 4' secondary amines



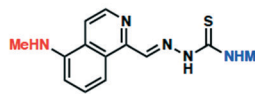
**HCT-6**  
 $0.331 \pm 0.002 \mu\text{M}$   
 ( $18.7 \pm 2.4 \mu\text{M}$ )



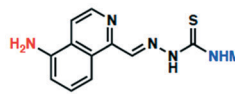
**HCT-7**  
 $0.114 \pm 0.007 \mu\text{M}$   
 ( $2.08 \pm 0.32 \mu\text{M}$ )



**HCT-8**  
 $0.233 \pm 0.054 \mu\text{M}$   
 ( $4.08 \pm 0.21 \mu\text{M}$ )

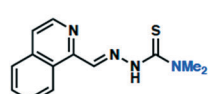


**HCT-9**  
 $0.607 \pm 0.004 \mu\text{M}$   
 ( $4.24 \pm 1.68 \mu\text{M}$ )

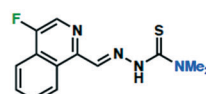


**HCT-10**  
 $7.08 \pm 0.63 \mu\text{M}$   
 ( $11.2 \pm 3.0 \mu\text{M}$ )

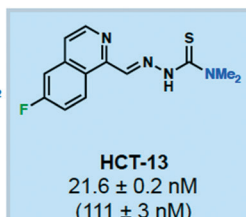
## 4' tertiary amines



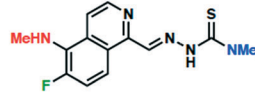
**HCT-11**  
 $73.7 \pm 0.8 \text{ nM}$   
 ( $29.6 \pm 7.4 \mu\text{M}$ )



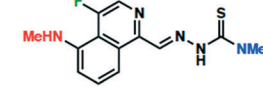
**HCT-12**  
 $26.6 \pm 3.2 \text{ nM}$   
 ( $274 \pm 6 \text{ nM}$ )



**HCT-13**  
 $21.6 \pm 0.2 \text{ nM}$   
 ( $111 \pm 3 \text{ nM}$ )



**HCT-14**  
 $38.4 \pm 1.4 \text{ nM}$   
 ( $272 \pm 30 \text{ nM}$ )



**HCT-15**  
 $42.3 \pm 13.8 \text{ nM}$   
 ( $327 \pm 39 \text{ nM}$ )

**Fig. 1** Cu(II) supplementation increases the potency of a subset of HCT compounds. MIAPACA2 cells were treated with HCT compound with 20  $\mu\text{M}$   $\text{CuCl}_2$  supplementation or with HCT compound alone (compound-only  $\text{IC}_{90}$  value shown in parentheses) for 72 h. Cell viability was then measured with CellTiter-Glo and  $\text{IC}_{90}$  values determined. Lead compound **HCT-13** is highlighted.

Weiss processes.<sup>16</sup> HCTs are particularly adept at binding copper,<sup>17</sup> which can be either beneficial or detrimental to the compound's biological activity. For instance, physiological concentrations of copper in human plasma (11–18  $\mu\text{M}$ )<sup>18,19</sup> interfere with the ribonucleotide reductase-inhibitory activity of 3-aminopyridine-2-carboxaldehyde thiosemicarbazone (3-AP),<sup>17</sup> while the cytotoxicities of Dp44mT<sup>20,21</sup> and NSC-319726<sup>22</sup> are potentiated by copper. Binding of this transition metal is intriguing from an anticancer therapy standpoint, as cancers rely upon higher intracellular levels of copper, relative to healthy cells, to promote angiogenesis, tumor growth, and metastasis.<sup>23,24</sup> Indeed, several therapeutic strategies have employed small molecules to disrupt copper homeostasis in cancers, either through chelation-mediated copper sequestration, or by increasing intracellular copper to cytotoxic levels through ionophoric modalities.<sup>22,25</sup>

The cytotoxic effects of HCTs coupled with their ability to bind copper make them a compelling scaffold from which to develop copper-mediated therapeutics. Our group has identified isoquinoline-based HCTs as a viable scaffold for this purpose, as previously reported analogs demonstrated both *in vitro* and *in vivo* efficacy and the chemical space and mechanism(s) of action remain insufficiently explored. Confounding factors have included lack of cancer tissue-specificity, difficulty stratifying susceptible cancer subtypes, poorly defined mechanisms of action, and potency and toxicity issues arising in part from aberrant metal chelation.<sup>12</sup>

Herein, the design and synthesis of a series of novel isoquinoline-based HCTs is reported, with identification of a promising lead compound – **HCT-13** – preceding mechanistic and pre-clinical studies. The lead compound potently inhibited proliferation of a panel of cancer models at nanomolar concentrations in the presence of physiologically relevant levels of copper, while non-cancerous cells were significantly less affected. A mechanism of action was delineated, with **HCT-13** being found to induce mitochondrial electron transport chain (mtETC) dysfunction and production of reactive oxygen species (ROS), deplete guanosine nucleotide pools, and engage DNA damage response/replication stress response pathways. The activity of **HCT-13** was potentiated by co-administration of the glycolysis inhibitor 2-deoxy-D-glucose (2-DG), indicating that cancers which rely more heavily upon oxidative phosphorylation, such as certain leukemias, will be more susceptible to **HCT-13** therapy. Lastly, we demonstrate the *in vivo* efficacy of a copper:**HCT-13** ( $\text{Cu}[\text{HCT-13}]$ ) complex in murine models of aggressive acute leukemias.

## Results

### Synthesis of isoquinoline-based HCT compounds

Among the previously reported isoquinoline HCTs, the 5-, 7-, and 8-fluoro analogs of **HCT-1** were of particular interest to our group. These compounds varied in terms of potency and

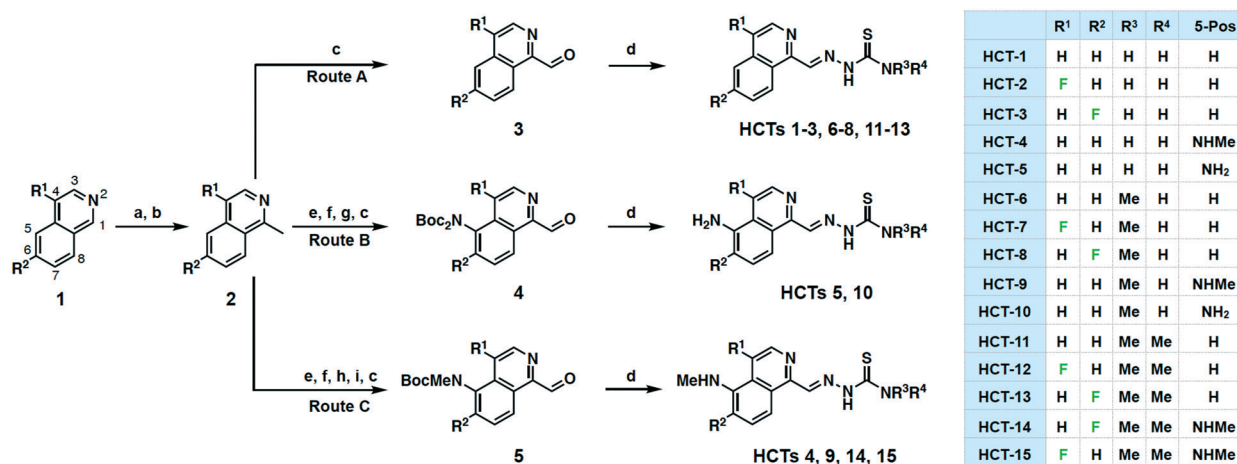
toxicity,<sup>6</sup> demonstrating that the effects of fluorination were dependent upon the position of isoquinoline substitution. The benefits that fluorination endows upon small-molecule drugs are well documented, and are also reflected in the marketplace, as approximately 20% of all pharmaceuticals are fluorinated.<sup>26–28</sup> Importantly, fluorination is associated with increased lipophilicity and may modulate redox activity upon metal complexation by virtue of its high electronegativity, a property which we wished to explore given our desire to leverage the copper-chelating potential of the scaffold.<sup>29</sup> With this in mind, we were curious as to how the previously unreported 4- and 6-fluoro analogs of **HCT-1** would perform in antiproliferative assays against cancer models, both with and without copper supplementation. Additionally, we were intrigued as to whether fluorination at these positions would complement 4' amine alkylation, as Kowol *et al.* discovered that 4' amine alkylation potentiated the cytotoxicity of 3-AP analogs.<sup>30</sup>

A total of 11 novel isoquinoline-based HCTs were synthesized, as well as four previously reported compounds (**HCT-1**, **HCT-4**, **HCT-5**, **HCT-6**) for comparison purposes. Our synthetic approach began with methylation of the appropriate isoquinoline **1** to generate **2** (Scheme 1). Depending upon the desired 5-position substituent, **2** was then subjected to either Route A (5-hydrido), Route B (5-amino), or Route C (5-methylamino). Syntheses of **HCTs 1–3**, **HCTs 6–8**, and **HCTs 11–13** were carried out by route A, wherein the methyl substituent of **2** was oxidized using selenium dioxide (SeO<sub>2</sub>) to furnish the carboxaldehyde **3**. Condensation with the appropriate thiosemicarbazide under acidic conditions yielded the desired HCT. **HCT-5** and **HCT-10** were synthesized *via* route B, which began with nitration of **2** followed by an iron-mediated reduction to the amine, which was subsequently Boc-protected and oxidized to produce carboxaldehyde **4**. This intermediate was then

simultaneously Boc-deprotected and condensed with the appropriate thiosemicarbazide under acidic conditions to furnish the target HCT. Syntheses of **HCT-4**, **HCT-9**, **HCT-14**, and **HCT-15** *via* route C proceeded from **2** with installation of a nitro group, subsequent conversion to the mono-Boc-methylamine, and SeO<sub>2</sub>-mediated oxidation to furnish **5**. Concurrent Boc-deprotection and thiosemicarbazide condensation were again achieved under acidic conditions to provide the desired HCT compound. While characterizing the HCTs, we occasionally observed the presence of a minor *Z*-isomeric product, particularly for **HCTs 11–15**, which was inseparable by HPLC from the *E*-isomer. When encountered, the isomeric mixtures were used for *in vitro* investigations, as previous studies have reported no significant difference in potency for these geometric isomers.<sup>31</sup>

### Fluorination and dimethylation display synergistic effects further potentiated by copper(II) supplementation

IC<sub>90</sub> values against MIA-PACA2 cells were first determined using conventional cell culture conditions (Fig. 1, compound-only IC<sub>90</sub> values reported in parentheses). Compounds were separated into three series – 4' primary amines, 4' secondary amines, and 4' tertiary amines – to reflect the relative degrees of 4' amine methylation. We first synthesized a series of non-methylated 4' primary amine compounds, with known compounds **HCT-1**, **HCT-4**, and **HCT-5** included to gauge whether fluorination of the isoquinoline proved beneficial for biological activity.<sup>32,33</sup> Within the 4' primary amine series, fluorination at the isoquinoline 4-position (**HCT-2**) did not show an increase in potency relative to unsubstituted analog **HCT-1**. However, fluorination at the 6-position (**HCT-3**) showed a 3-fold increase, demonstrating that the fluorine position impacts the activity of these compounds.



**Scheme 1** Synthesis of HCT compounds 1–15 from simple isoquinolines. (a) Allyl chloroformate, MeMgBr, THF; (b) Pd(PPh<sub>3</sub>)<sub>4</sub>, morpholine; DDQ, CH<sub>2</sub>Cl<sub>2</sub>; (c) SeO<sub>2</sub>, 1,4-dioxane, 60 °C; (d) appropriate thiosemicarbazide, HCl, EtOH, reflux or microwave 50 °C; (e) KNO<sub>3</sub>, H<sub>2</sub>SO<sub>4</sub>; (f) Fe, HCl, MeOH, reflux; (g) Boc<sub>2</sub>O, DMAP, TEA, THF; (h) Boc<sub>2</sub>O, DMAP, TEA, THF; NaHCO<sub>3</sub>, MeOH, reflux or K<sub>2</sub>CO<sub>3</sub>, MeOH, reflux; (i) NaH, THF; MeI. **HCTs 1–3**, **6–8**, and **11–13** synthesized through route A; **HCTs 5** and **10** synthesized through route B; **HCTs 4**, **9**, **14**, and **15** synthesized through route C. See Experimental section for full synthetic details.

In the 4' secondary amine series, we observed a synergistic effect when combining isoquinoline fluorination with 4' amine methylation. The 4-fluorine substituted **HCT-7** and the 6-fluorine substituted **HCT-8** were each significantly more potent than their corresponding non-fluorinated analog **HCT-6**, as well as more potent than their 4' primary amine counterparts (**HCT-2** and **HCT-3**, respectively). The trend of isoquinoline substitution and 4' secondary amine combining to enhance the potency of the drug held well for the 5-methylamino substituted **HCT-9** and 5-amino substituted **HCT-10** analogs. Together, these results suggested that combining isoquinoline substitution, particularly 4- or 6-fluorination, with 4' amine methylation produced greater-than-additive antiproliferative effects when compared with either modification alone.

The effects of fluorine substitution became significantly more pronounced for the dimethylated 4'-tertiary amine compounds **HCT-12** and **HCT-13**, whose  $IC_{90}$  values were in the nM range and were roughly 110- and 270-fold more potent, respectively, when compared to their non-fluorinated analog **HCT-11**. Fluorination at the 6-position (**HCT-13**) was found to be superior to fluorination at the 4-position (**HCT-12**), a trend which also held for the fluorine-substituted 5-methylamino compounds **HCT-14** and **HCT-15**.

HCT compounds are known to be copper chelators,<sup>34–36</sup> and a recent publication by Stockwell and coworkers demonstrated that the activity of a known 4' tertiary amine HCT (NSC-319726) was significantly potentiated by the addition of copper.<sup>22</sup> To test whether our compounds were similarly potentiated, we determined  $IC_{90}$  values against MIAPACA2 cells in media supplemented with physiologically relevant levels of copper (Fig. 1, ESI† Fig. S1a). While the activity of **HCT-5** was attenuated, all other compounds displayed significant increases in potency under copper-supplemented conditions, with +Cu  $IC_{90}$  values improving as the degree of 4' amine methylation increased. Fluorine substitution led to greater potency when compared with corresponding non-fluorinated analogs, and compounds bearing 4' tertiary amines were the most active, achieving +Cu  $IC_{90}$  values as low as 21.6 nM (**HCT-13**). Our results demonstrated that physiologically relevant levels of copper potentiated the activity of isoquinoline HCTs, and that 4'

amine methylation worked cooperatively with fluorine substitution. Due to its potency and straightforward synthesis, we chose **HCT-13** as our lead compound for in-depth mechanism of action studies.

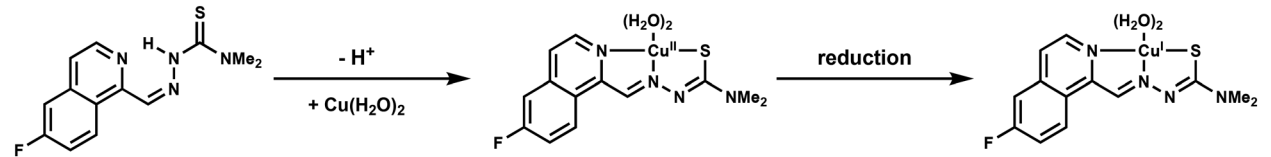
### Computational analysis shows fluorination increases the stability of copper:HCT complex

To elucidate potential contributing factors to **HCT-13**'s potency, density functional theory (DFT) calculations were performed and the relative energies of formation and Cu(II) reduction were determined for selected ligand–Cu(II) complexes (Table 1). All calculations were performed with Gaussian 16<sup>37</sup> using the B3LYP functional and the 6-31G(d) basis set. A correlation between  $IC_{90}$  value and energy of complex formation was found, with the latter consisting of ligand deprotonation and copper binding energies. **HCT-5**, which possesses an electron-donating substituent at the isoquinoline 6-position and was the least potent compound tested, bonded most weakly to Cu(II). Conversely, **HCT-12** and **HCT-13**, which bear electron-withdrawing fluorine substituents at the 4- and 6-positions, respectively, bonded more strongly. **HCT-11**, which possessed no isoquinoline substituents, had an energy of coordination to Cu(II) that was 4.0 kcal mol<sup>-1</sup> lower than that of **HCT-5**. These studies revealed that the ease of deprotonation in the thiosemicarbazone chain most strongly contributed to the trends observed in the energies of complex formation, where ligands with electron-withdrawing substituents had lower energies of deprotonation. It was also found that the reduction of Cu(II) to Cu(I) was most exergonic in the Cu:**HCT-13** complex, implying that this compound is most readily capable of participating in redox processes.

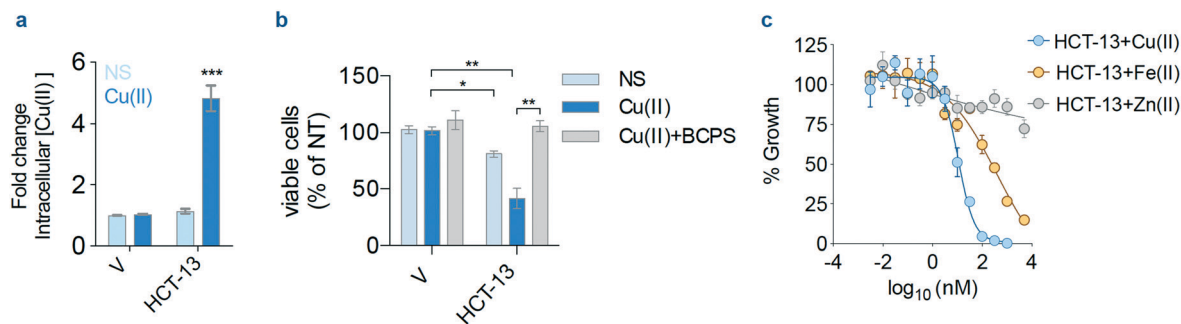
### HCT-13 induces copper-dependent toxicity and is highly potent against a panel of cancer models

To begin investigating the mechanism of action of **HCT-13**, its ionophoric capabilities were evaluated using inductively coupled plasma mass spectrometry (ICP-MS). Intracellular levels of copper increased when cells were treated with **HCT-13** in Cu(II)-supplemented media, but not with **HCT-13** or Cu(II) supplementation alone (Fig. 2a). To measure the effect of copper upon the lead compound's antiproliferative activity,

**Table 1** DFT calculations of relative deprotonation, binding, and complexation energies of selected Cu(II):HCT-13 complexes and Cu(II) to Cu(I) reduction energies



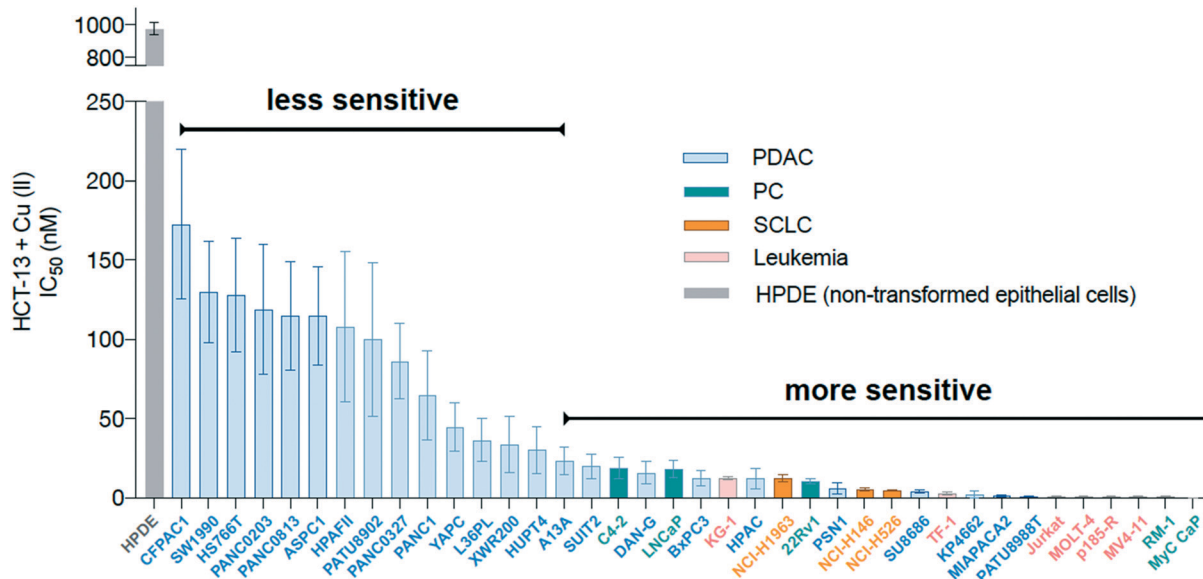
Compound	Rel. deprotonation energy (kcal mol <sup>-1</sup> )	Rel. binding energy (kcal mol <sup>-1</sup> )	Rel. complexation energy (kcal mol <sup>-1</sup> )	Rel. energy of reduction (kcal mol <sup>-1</sup> )
<b>HCT-5</b>	0	0	0	-0.9
<b>HCT-11</b>	-4.6	0.6	-4	0
<b>HCT-12</b>	-9.6	4.3	-5.5	-1.5
<b>HCT-13</b>	-10.5	4.4	-6.1	-1.5



**Fig. 2** Copper is required for HCT-13 activity. (a) Intracellular concentrations of copper measured by inductively coupled plasma mass spectrometry (ICP-MS) in MIAPACA2 PDAC cells treated with HCT-13 (25 nM) for 24 h  $\pm$  Cu(II) (20  $\mu$ M); \*\*\*  $P < 0.001$ ; V: vehicle. (b) Viability of MIAPACA2 cells treated with HCT-13 (25 nM) + Cu(II) (20  $\mu$ M) for 24 h  $\pm$  bathocuproine disulfonate (BCPS, 300  $\mu$ M) measured by trypan blue exclusion; \*  $P < 0.05$ ; \*\*  $P < 0.01$ ; V: vehicle, NS: no supplement. (c) Proliferation rate of MIAPACA2 PDAC cells measured by CellTiterGlo following HCT-13 treatment for 72 h  $\pm$  Cu(II), Fe(II) or Zn(II) (20  $\mu$ M). (mean  $\pm$  SD,  $n = 2$ , one-way ANOVA corrected for multiple comparisons by Bonferroni adjustment).

the viability of MIAPACA2 cells was assessed following treatment with HCT-13 in Cu(II)-supplemented media with and without bathocuproine disulfonate (BCPS), a membrane-impermeable copper chelator. The addition of BCPS completely abrogated the cytotoxicity of HCT-13, suggesting that the growth inhibitory effect of our lead compound is dependent upon copper availability (Fig. 2b). We next determined the effects of supplementation with selected metals, namely iron and zinc, upon the antiproliferative activity of HCT-13 and found its potency was highest in the presence of Cu(II), diminished but still active in the presence of Fe(II), and largely inactive in the presence of Zn(II) (Fig. 2c, 20  $\mu$ M supplementation). Collectively, these data suggest that HCT-13 is a Cu(II) ionophore which increases intracellular copper concentration and whose cytotoxicity is copper-dependent.

To determine whether HCT-13 displayed heterogeneous effects against different cell types, it was screened against a panel containing aggressive cancer models and a non-cancerous cell line. Pancreatic ductal adenocarcinoma (PDAC), small cell lung carcinoma (SCLC), and prostate cancer (PC) models were evaluated as these cancers rely upon elevated copper levels to sustain growth, and serum copper levels are elevated ( $>20 \mu$ M) in individuals with these types of tumors.<sup>24,38–43</sup> Aggressive leukemia models were also evaluated, as these may be more directly targeted relative to solid tumors and treatments for hematological malignancies such as acute myeloid leukemia (AML) remain an unmet need.<sup>44</sup> PDAC, SCLC, and PC cancer models were cultured in media supplemented with physiologically relevant levels of copper (20  $\mu$ M CuCl<sub>2</sub>) and treated with HCT-13, while the leukemia models were treated with pre-formed copper:HCT-



**Fig. 3** Activity of HCT-13 in a panel of cancer cell lines and non-transformed cells. IC<sub>50</sub> values in a panel of human and mouse prostate cancer (PC), small cell lung carcinoma (SCLC), pancreatic ductal adenocarcinoma (PDAC), leukemia, and non-transformed epithelial models treated with HCT-13 + Cu(II) (20  $\mu$ M) for 72 h measured by CellTiterGlo.

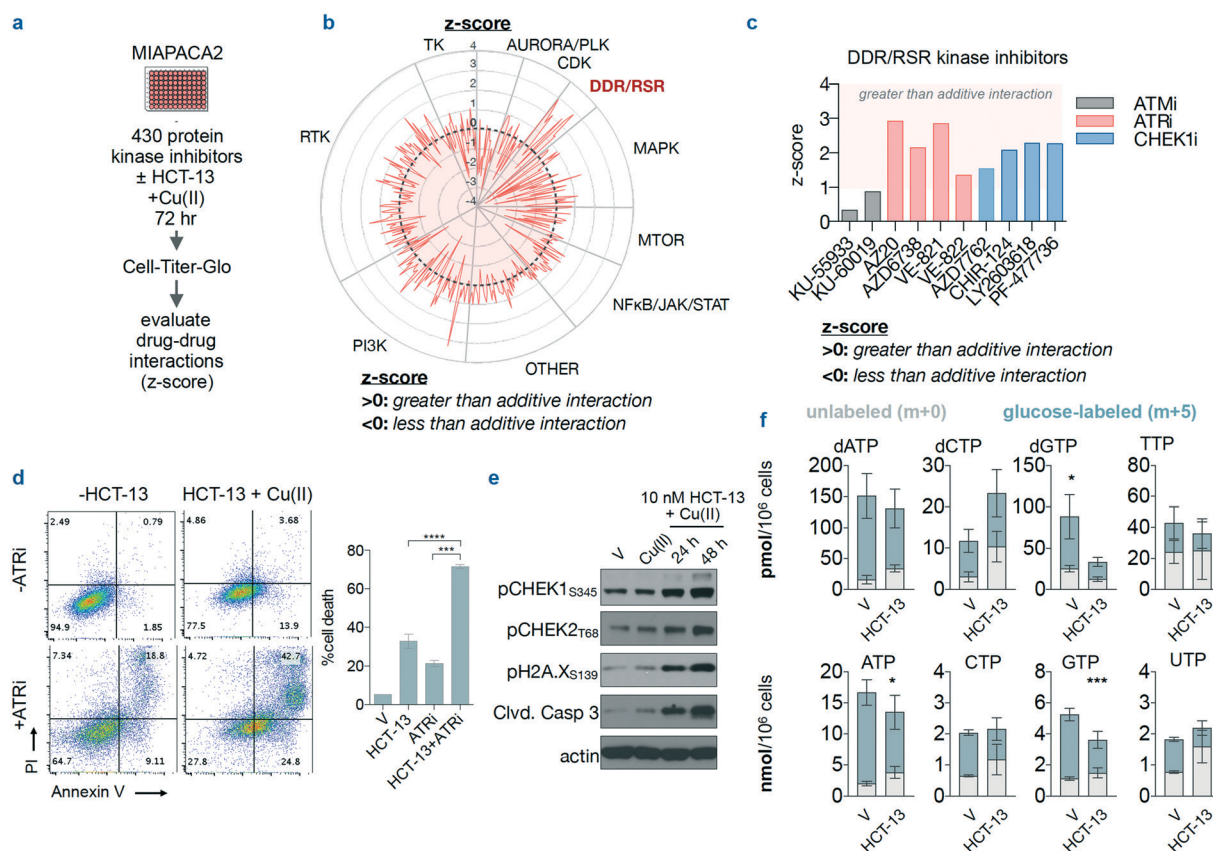
**13** (**Cu[HCT-13]**) complexes, as free copper is known to be toxic towards these cells (antiproliferative activity of **Cu[HCT-13]** was consistent with that of **HCT-13** + **Cu(II)**) (ESI† Fig. S1b). In this assay, **HCT-13** was shown to be a highly potent cancer cell growth inhibitor, with  $IC_{50}$  values ranging from 1 nM to 200 nM (Fig. 3). Importantly, the non-cancerous human epithelial cell line HPDE was markedly more resistant to treatment than the cancer models evaluated, while the cancer models displayed differential susceptibility. Together, these results indicate that **HCT-13** possesses a high degree of cancer-specific cytotoxicity, while its differential effects against the cancer models suggest its mechanism(s) of action is dependent upon specific cancer-subtypes/characteristics.

### Chemical genomics screen identifies DNA damage response and replication stress response as potential mechanisms of action

To begin identifying the mechanisms of action of **HCT-13**, we performed an unbiased pharmacological inhibition screen using a chemical genomics platform consisting of 430 kinase

inhibitors. MIAPACA2 cells were treated with the 430-member library, covering a 7-point concentration range spanning between 6.5 nM and 5  $\mu$ M, with and without 25 nM **HCT-13** in presence of 20  $\mu$ M  $CuCl_2$ . After 72 h, ATP content was measured using CellTiter-Glo (Fig. 4a, ESI† Fig. S2a). A composite synergy score was calculated for each combination, defined as the sum of the Bliss additivity score (% proliferation inhibition observed – % proliferation inhibition expected). A positive synergy score indicates greater-than-additive interaction, and a negative score indicates less-than-additive interaction *i.e.* antagonism.

The ten highest scoring compounds were kinase inhibitors contained in the DNA damage response/replication stress response (DDR/RSR) module, with the ataxia telangiectasia mutated serine/threonine kinase (ATM)/checkpoint kinase 2 (CHK2) and Rad3-related serine/threonine kinase (ATR)/CHK1 pathways featuring as the most prominent codependencies (Fig. 4b and c).<sup>45–47</sup> All eight ATR and CHEK1 inhibitors included in the library scored positively in the screen, implying that the DDR/RSR pathways are activated by **HCT-13**. The synergistic interaction of **HCT-13**



**Fig. 4** A chemical genomics screen identifies the replication stress response pathway as an actionable co-dependency of **HCT-13**-treated cells. (a) Experimental design of a synthetic lethality screen using a library of protein kinase inhibitors against **HCT-13**-treated cells in the presence of **Cu(II)** (20  $\mu$ M). (b) Radar plot of screen results. (c) Z-score values for kinase inhibitors within the DNA damage response/replication stress response (DDR/RSR) pathway module. (d) Annexin V/PI staining in MIAPACA2 cells to validate the synergistic interaction of **HCT-13** (25 nM) + **Cu(II)** (20  $\mu$ M) with ATRi (250 nM VE-822) treated for 72 h; \*\*\*\*  $P < 0.0001$ . (e) Representative immunoblot of DDR pathway and cell death biomarkers in MIAPACA2 PDAC cells treated with **HCT-13** (10 nM) + **Cu(II)** (20  $\mu$ M); V: vehicle. (f) Nucleotide pool measurements by LC-MS/MS-MRM in MIAPACA-2 cells treated with 25 nM **HCT-13** + **Cu(II)** (20  $\mu$ M) for 48 h (mean  $\pm$  SD,  $n = 3$ , comparison made between glucose-labeled pools using two-way ANOVA corrected for multiple comparisons by Bonferroni adjustment, \*  $P < 0.05$ ; \*\*\*  $P < 0.001$ ); V: vehicle.

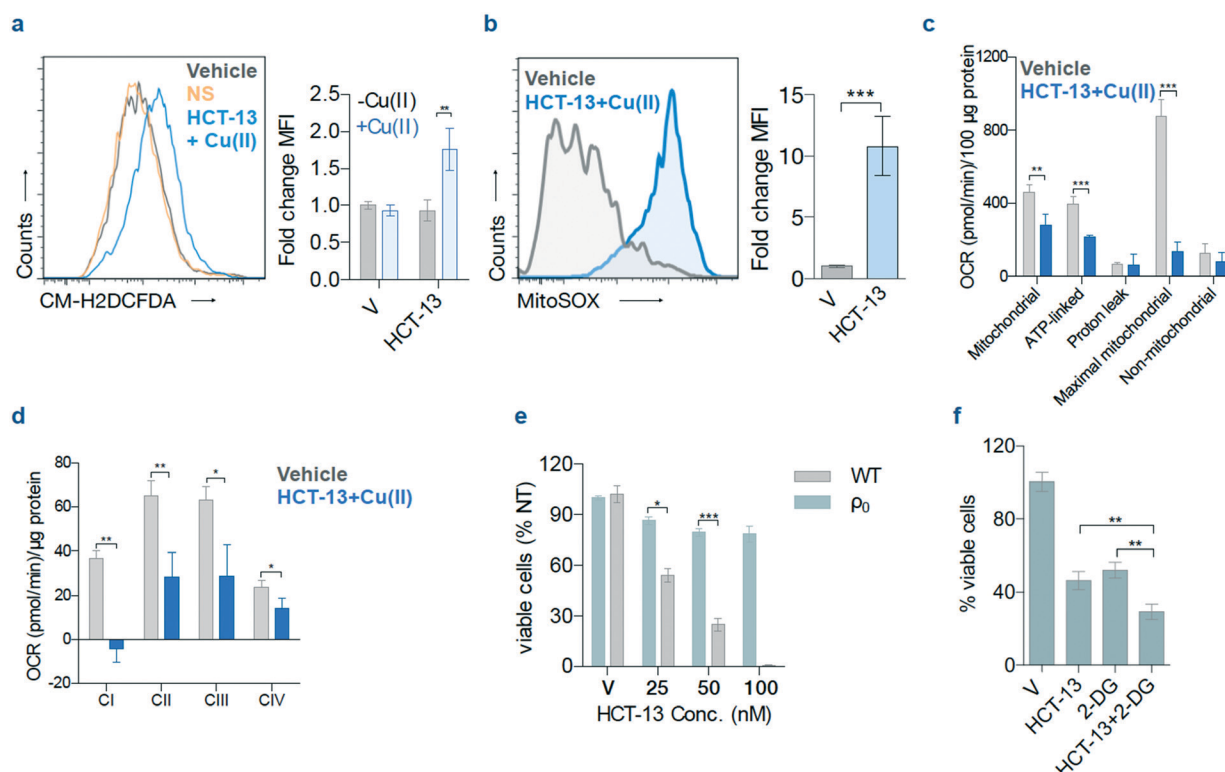
with ATR inhibition was further validated by annexin V/PI (apoptosis) and trypan blue viability staining in PDAC (MIAPACA2, CFPAC-1) and PC (C4-2) cell lines (Fig. 4d, ESI† Fig. S2b and c). Furthermore, phosphorylation of CHEK1 and CHEK2 kinases was consistently observed upon HCT-13 + Cu(II) treatment, as well as the induction of DNA damage marker pH2A.X and cleavage of the apoptotic marker caspase 3 (Fig. 4e). These observations are indicative of DDR/RSR pathway activation, which was hypothesized to arise from nucleotide insufficiency and/or, given the ability of copper to generate ROS, from ROS-mediated DNA damage.<sup>48–50</sup>

To test this hypothesis, nucleotide levels were measured in treated and non-treated cells using liquid chromatography-tandem mass spectrometry with multiple reaction monitoring (LC-MS/MS-MRM) as previously described.<sup>51</sup> Both dGTP and rGTP pools were decreased in HCT-13-treated cells, while levels of the other nucleotides either increased or the change was statistically insignificant (Fig. 4f). These findings indicate that HCT-13 preferentially decreases the dGTP and rGTP pools, which is potentially mediated by oxidative processes as guanine is the most readily oxidized nucleobase.<sup>52</sup> Together with the immunoblot results (Fig. 4e), these data suggest that dGTP pool

insufficiency in HCT-13-treated cells triggers the activation of the intra-S checkpoint, as measured by increased pChk1 levels, which in turn renders these cells dependent upon the activity of the replication stress response pathway.

### HCT-13 induces ROS production, displays mitochondrial-dependent toxicity and targets OXPPOS

Given that our dGTP and rGTP pool measurements pointed towards a guanosine-depleting mechanism of action, we set out to determine whether our lead compound was giving rise to ROS. We found that HCT-13 treatment induced ROS generation detectable by CM-H<sub>2</sub>DCFDA staining in MIAPACA2 cells (Fig. 5a).<sup>53</sup> ROS generation was also observed in the mitochondria, as measured by mitochondria-specific dye MitoSOX (Fig. 5b).<sup>53</sup> To further probe the ramifications of the HCT-13-generated ROS, we inquired as to whether oxidative phosphorylation (OXPHOS) in MIAPACA2 cells was altered following treatment. A Seahorse assay was performed to measure the alteration in overall OXPPOS, and an electron flow assay was performed in isolated mitochondria to assess which electron transport chain (ETC) complex is impacted (Fig. 5c and d, ESI† Fig. S3a and b).<sup>54–56</sup> These assays showed



**Fig. 5** HCT-13 + Cu(II) induces oxidative stress and has selective mitochondrial toxicity. (a) Reactive oxygen species (ROS) measurements using CM-H<sub>2</sub>DCFDA after HCT-13 (25 nM) ± Cu(II) (20 μM) treatment for 24 h; mean fluorescence intensity (MFI); mean ± SD, *n* = 2, Student *t*-test, \*\* *P* < 0.01. (b) Mitochondrial ROS detection using MitoSOX in MIAPACA2 PDAC cells treated with HCT-13 (25 nM) + Cu(II) (20 μM) for 24 h. mean ± SD, *n* = 2, Student *t*-test, \*\*\* *P* < 0.001. (c) Oxygen consumption rate (OCR) of MIAPACA2 PDAC cells treated with HCT-13 (25 nM) + Cu(II) (20 μM) for 24 h; \*\* *P* < 0.01; \*\*\* *P* < 0.001. (d) OCR of isolated mitochondria measured with or without HCT-13 (25 nM) + Cu(II) (20 μM); \* *P* < 0.05; \*\* *P* < 0.01. (e) Viability of 143 BTK parental (wild type, WT) and ρ<sub>0</sub> cells after 48 h of the indicated HCT-13 concentration + Cu(II) (20 μM) treatment, assessed with trypan blue staining; mean ± SD, *n* = 2, Student *t*-test, \* *P* < 0.05; \*\*\* *P* < 0.001; V: vehicle. (f) Cell viability by trypan blue staining in MIAPACA2 PDAC cells to interrogate the interaction of HCT-13 (25 nM) + Cu(II) (20 μM) with 2-DG (2 mM) for 48 h; mean ± SD, *n* = 2, Student *t*-test, \*\* *P* < 0.01; V: vehicle.

that cell respiratory capacity decreased significantly upon **HCT-13** treatment as measured by reduced oxygen consumption rate (OCR), that OXPHOS capacity was impaired, and that ETC complex 1 was most dramatically affected.

These findings suggested that **HCT-13** inhibited mtETC activity but did not indicate whether our lead compound's cytotoxicity stemmed from effects independent of mitochondrial function. Another HCT compound, Dp44mT, has been reported to induce ROS, though its cytotoxicity was not attributed to the functionality of the mitochondria.<sup>57</sup> To determine whether the cytotoxicity of **HCT-13** was ETC-dependent, we examined its effects upon 143 BTK  $\rho_0$ , a mitochondrial DNA (mtDNA)-deficient osteosarcoma cell line. Both 143 BTK  $\rho_0$  and parental (wild type, WT) cells were treated with **HCT-13** + 20  $\mu\text{M}$  Cu(II) for 48 h, after which cell viability was determined (Fig. 5e). Compared to WT, the  $\rho_0$  cells were significantly less sensitive to the treatment, with a concurrent decrease in levels of S-phase arrest (ESI $\ddagger$  Fig. S3c and d). Together, these results indicate that the cytotoxic effects of **HCT-13** are mitochondria-dependent and suggest that **HCT-13** may preferentially target tumor cells which rely more heavily upon OXPHOS than on glycolysis. To further explore this potentially stratifying metabolic hallmark, cells were treated with **HCT-13** with and without 2-deoxyglucose (2-DG), a molecule that competitively inhibits glycolysis, thereby forcing cells to rely upon OXPHOS for energy production (Fig. 5f).<sup>58</sup> Co-administration of 2-DG significantly potentiated the activity of **HCT-13**, conceivably by forcing the cancer cells to rely more heavily upon OXPHOS which was in turn impaired by **HCT-13**.

### **HCT-13 pre-complexed with copper is effective against aggressive B-ALL and AML models *in vivo***

With a mechanism of action outlined and a potentially stratifying metabolic hallmark identified, we next investigated the *in vivo* efficacy and tolerability of **HCT-13**. Two leukemia models were chosen – a primary murine BCR-ABL-expressing *Arf*-null pre-B (p185<sup>BCR-ABL</sup> *Arf*<sup>-/-</sup>) ALL model (p185) and a human systemic acute myeloid leukemia (AML) model (MV4-11) – as these leukemias possess aggressive phenotypes, have high intrinsic levels of OXPHOS, and there remains a persistent unmet need for effective therapeutic options, particularly in the case of AML. Both p185<sup>59-61</sup> and MV4-11<sup>62,63</sup> cell lines were engineered to express luciferase to monitor the systemic leukemic burden by bioluminescence imaging (BLI). To bypass the need for systemic copper supplementation, a one-to-one complex of copper and **HCT-13** (Cu[**HCT-13**]) was prepared for *in vivo* administration according to reported procedures for similar compounds<sup>64</sup> and characterized by UV-HPLC and HR-MS (ESI $\ddagger$  Fig. S4). The *in vitro* antiproliferative activity of Cu[**HCT-13**] was consistent with that of **HCT-13** + Cu(II) in all cell lines tested.

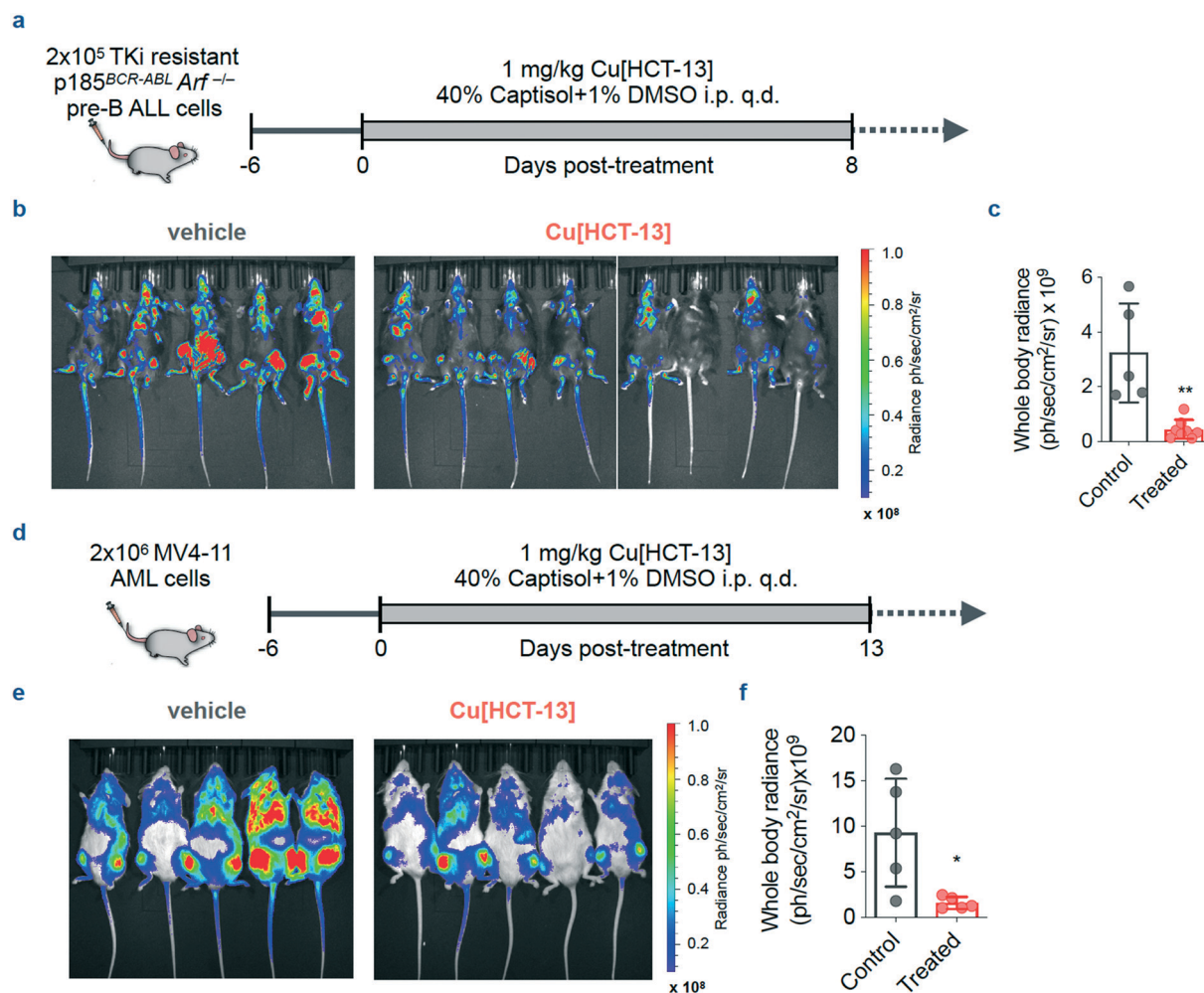
Treatment in both p185 and MV4-11 murine models was initiated on day six post-inoculation of cells, when all mice

showed evidence of systemic disease. Mice in treatment groups of the pre-B ALL and AML arms of the study were administered 1 mg kg<sup>-1</sup> Cu[**HCT-13**] q.d. for 8 and 13 days, respectively (Fig. 6a and d). The treatment was well tolerated as indicated by body weight measurements (ESI $\ddagger$  Fig. S5). 14 days after treatment initiation in the pre-B ALL arm, **HCT-13**-treated mice displayed significantly lower systemic disease burden than mice in the control group (Fig. 6b and c). Similarly, treatment group mice in the MV4-11 portion of the study had significantly lower disease burden on day 19 compared to the control group mice (Fig. 6e and f).

## Discussion

This study aimed to leverage the general ability of HCTs to bind copper and the anticancer activity of isoquinoline-based HCTs towards developing novel copper-binding analogs with a defined mechanism of action and *in vivo* efficacy against cancer models. Two potential modifications of the previously reported **HCT-1** (IQ-1) scaffold drew particular interest during our synthetic planning: fluorination of the isoquinoline ring, and sequential methylation of the 4' amine. With the goal of leveraging the copper chelating ability of the isoquinoline HCT scaffold against cancer cells, we sought to synthesize the novel 4- and 6-fluoro analogs of **HCT-1** and investigate what effects sequential 4' amine methylation would have upon antiproliferative activity. Following our synthetic campaign, analysis of the antiproliferative data revealed several trends. In conditions without copper supplementation, fluorination at either the 4- or 6-position of the isoquinoline ring led to an increase in potency for five out of six compounds, when compared with their corresponding non-fluorinated analogs (Fig. 1). In some cases, the change was dramatic – the IC<sub>90</sub> of **HCT-13** was nearly 270-fold lower than its non-fluorinated analog **HCT-11**. Secondly, 4' amine methylation in the absence of isoquinoline substitution or copper supplementation was detrimental to activity, as demonstrated by the decrease in potencies for **HCTs** 6 and 11 when compared with 4' primary amine **HCT-1**. However, combining 4' amine methylation and isoquinoline substitution in a single compound, as in **HCTs** 7–10, 12 and 13, produced greater-than-additive antiproliferative effects when compared with either their 4' primary amine or unsubstituted isoquinoline analogs; **HCT-13** again exemplified this trend. Interestingly, the fluorinated compounds **HCT-12** and **HCT-13** were qualitatively more stable than non-substituted **HCT-11**, the latter of which quickly began to decompose when solubilized in DMSO, whereas the former were stable for months when stored as a DMSO solution at room temperature, as monitored by <sup>1</sup>H NMR. This serendipitous discovery factored heavily into our decision to continue generating fluorinated analogs. The underlying mechanisms responsible for the observed synergy and stability remain under study by our group. Finally, the antiproliferative activities of all but one of our isoquinoline HCTs were potentiated by supplementation with





**Fig. 6** Cu[HCT-13] is effective in aggressive models of systemic B-ALL and AML. (a) Dose and schedule for the efficacy study in p185 pre-B ALL bearing mice. (b and c) Bioluminescence images and quantification of whole body radiance of mice treated with Cu[HCT-13] ( $n = 8$ ) or vehicle control ( $n = 5$ ). (d) Dose and schedule for the efficacy study in MV4-11 AML bearing mice. (e and f) Bioluminescence images and quantification of whole body radiance of mice treated with Cu[HCT-13] ( $n = 5$ ) or vehicle control ( $n = 5$ ) (mean  $\pm$  SD,  $n = 2$ , Student's paired  $t$ -test, \*  $P < 0.05$ ; \*\*  $P < 0.01$ ; \*\*\*  $P < 0.001$ ). q.d. once per day.

physiologically relevant levels of copper, with HCT-5 being the exception. DFT calculations were employed to gain further insight into the observed trends, revealing that presence of an electron withdrawing fluorine at the 4- and particularly the 6-position led to lower barriers to ligand deprotonation and greater complex stability. Additionally, the most exergonic reduction of Cu(II) to Cu(I) was observed in the Cu:HCT-13 complex, implying that this compound is most readily able to participate in redox processes. The observed trends culminated with identification of our lead compound as HCT-13, which was uncomplicated in its synthesis and highly potent.

The use of copper-chelating small molecules in anticancer therapy is an established strategy which is executed either through sequestration of copper from tumor tissue, or through increasing intracellular copper to cytotoxic levels.<sup>65</sup> HCT-13 behaves as an ionophore and increases intracellular levels of copper, a property that is essential for its cytotoxicity as sequestration of copper *via* BCPS-chelation negated our

lead compound's growth inhibitory effects (Fig. 2b). The potency of HCT-13 is highlighted by its nanomolar IC<sub>50</sub> values against a panel of PDAC, SCLC, PC, and leukemia cancer models (Fig. 3). Our chemical genomics screen pointed towards activation of DDR/RSR pathways following HCT-13 treatment. Upon further investigation, it was found that HCT-13 reduced dGTP and rGTP pools in treated cells, an effect consistent with oxidative damage. These results implicate a mechanism by which HCT-13 activates DDR/RSR pathways following oxidative depletion of guanosine nucleotide pools (Fig. 4f). Incorporation of oxidized nucleotides into newly synthesized DNA leads to DNA damage, while their removal from nucleotide pools may engender replication stress response due to insufficient nucleotide availability. Further probing of this mechanism led to confirmation of ROS presence in HCT-13-treated cells and the discovery of mitochondrial ROS and suppression of mitochondrial OXPHOS (Fig. 5a-d). Strikingly, the cytotoxicity of HCT-13 was greatly attenuated in cells lacking the

mitochondrial ETC (Fig. 5e). Conversely, 2-DG-mediated suppression of glycolysis sensitized MIA-PACA2 cells to treatment (Fig. 5f). Cumulatively, these data outline a mechanism in which in the presence of copper, **HCT-13** targets the mtETC and interferes with its function, generating ROS which in turn deplete GTP nucleotide pools and damage DNA either through nucleotide oxidation or through ROS-mediated ssDNA or dsDNA breaks.

The delineated mechanism of action illustrates a means for stratifying potential oncogenic targets – cancers which are more dependent upon OXPHOS and/or which possess defects in DDR are more likely to respond to **HCT-13** therapy than those without. Leukemias are reported to rely heavily upon OXPHOS in order to meet their energetic demands,<sup>66–72</sup> and its inhibition has been efficacious against preclinical models and is currently the subject of clinical investigation.<sup>73</sup> Additionally, preleukemic syndromes and myeloid malignancies rely heavily upon error-prone non-homologous end joining for the repair of double-strand breaks, a type of DNA damage commonly caused by ROS.<sup>74–76</sup> Coupled with preexisting oxidative stress, this reliance can lead to irreparable damage when challenged by treatment-induced ROS, a process which preferentially affects these malignant cells.<sup>77</sup> These characteristics informed our choice of pre-B ALL and AML models for preclinical studies, as leukemias fulfill the identified criteria for potential responders to **HCT-13** treatment. Gratifyingly, *in vivo* efficacy was realized, with a one-to-one copper:**HCT-13** complex (**Cu[HCT-13]**) being administered to avoid potential complications of systemic copper supplementation. Leukemic burden in both preclinical models as measured by BLI was significantly reduced in treatment group mice relative to control group mice, the latter of which had all succumbed to disease by the day of measurement (Fig. 6). In general, cancer cells exhibit higher levels of ROS and higher baseline oxidative stress than healthy cells, which may imbue our lead compound with selectivity towards cancerous tissue and explain the observed resistance of non-cancerous HPDE cells (Fig. 3).

While the data supporting our proposed mechanism of action is compelling, we cannot rule out additional mechanisms through which **HCT-13** may be effecting its anticancer activity. A recent report on the disulfiram metabolite diethyldithiocarbamate (DTC) details a mechanism by which DTC chelates copper *in vivo* and preferentially accumulates in tumor tissue before binding and aggregating NPL4, a protein which enables p97-mediated protein degradation.<sup>78</sup> Given the fact that NPL4 possesses two zinc finger domains which bind metal complexes, it is possible that **HCT-13** may derive some of its activity through mechanisms similar to the DTC-copper complex.<sup>79,80</sup> Additionally, the observed mechanisms of action suggest that **HCT-13** may synergize with radiotherapy, as therapeutic ionizing radiation increases ROS, thereby increasing oxidative stress and DNA damage in the targeted area(s). Therefore, **HCT-13** could also function as a radiosensitizer by further increasing the load of ROS, oxidative stress, and DNA damage

when administered in combination with radiation therapy. Further studies regarding mechanisms of action and potentials for combination therapy are ongoing in our group. Also of current interest is the use of alternative delivery strategies for administration of pre-complexed **Cu[HCT-13]**. Pre-complexation may prevent toxicity resulting from aberrant metal chelation following systemic administration when compared with **HCT-13** alone, and could promote preferential accumulation in cancer tissue, as has been previously observed.<sup>78</sup> Utilizing albumin-based formulations for delivery of this complex may bring further benefits for *in vivo* therapy and is currently under investigation.<sup>81</sup> Collectively, we think that identifying viable combination therapies and optimizing drug delivery will enhance the clinical efficacy of this already promising drug when combined with our elucidation of **HCT-13**'s mechanism(s) of action and identification of target-stratifying hallmarks.

## Conclusion

We have expanded upon a class of isoquinoline-based HCTs to produce a set of novel antiproliferative compounds. We demonstrated the synergistic effects of combining 4' amine methylation with isoquinoline substitution and identified **HCT-13** as a highly potent antiproliferative agent which is active against a panel of PDAC, SCLC, PC, and leukemia cancer models. We showed that presence of physiologically-relevant levels of Cu(II) greatly potentiated our lead compound's activity, and subsequent investigation into **HCT-13**'s mechanism of action revealed that it acts as a copper ionophore and requires copper to effect its cytotoxicity. Furthermore, **HCT-13** induces ROS production and mitochondrial dysfunction, decreases guanosine nucleotide pools, engages DDR/RSR pathways and synergizes with ATR inhibition, possesses mitochondrial-dependent cytotoxicity, and targets high-OXPHOS cells. Lastly, a one-to-one copper:**HCT-13** (**Cu[HCT-13]**) complex was demonstrated to be efficacious in preclinical models of aggressive leukemias. Taken together, the delineated mechanism of action of **HCT-13** and its demonstrated potency as a single agent therapeutic against aggressive tumor models *in vitro* and *in vivo* signal its promise as a therapeutic agent and warrant further investigations.

## Experimental section

### Chemistry

All chemicals, reagents and solvents were obtained from commercial sources and were used without further purification. Unless otherwise noted, reactions were carried out in oven-dried glassware under an atmosphere of argon using commercially available anhydrous solvents. Tetrahydrofuran (THF) was distilled from sodium under an argon atmosphere. Dichloromethane was distilled from calcium hydride. Solvents used for extractions and chromatography were not anhydrous. Analytical TLC was

carried out on precoated silica gel (Merck silica gel 60, F254) and visualized with UV light. Column chromatography was performed with silica (Fisher, 230–400 mesh).  $^1\text{H}$  NMR,  $^{13}\text{C}$  NMR, and  $^{19}\text{F}$  NMR spectra were measured in  $\text{CDCl}_3$  or  $\text{DMSO-}d_6$  on Bruker AV spectrometers at 400 or 500 MHz. Chemical shifts were reported in parts per million ( $\delta$ ) relative to residual solvent signals. The signals observed were described as follows: s (singlet), d (doublet), t (triplet), q (quartet), dd (doublet of doublets), dt (doublet of triplets), ddd (doublet of doublet of doublets), tt (triplet of triplets), tdd (triplet of doublet of doublets), m (multiplet), br s (broad singlet). Mass spectra were obtained on a Waters LCT Premier with ACQUITY UPLC mass spectrometer under electrospray ionization (ESI) or Thermo Fisher Scientific Exactive Plus with direct analysis in real time (DART) ionization. Purity of all compounds used in biological assays was determined to be >95% pure on a Hewlett Packard 1090 HPLC system using an Aquasil C18 column (250 mm  $\times$  2 mm, 5  $\mu\text{m}$ , Keystone Scientific) with an acetonitrile/water solvent system containing 0.1% TFA with detection performed at 254 nm (minute/% acetonitrile: 0/0, 8/0, 35/95, 43/95, 45/0, 55/0). All microwave-assisted reactions were carried out in a CEM Discover 908005 Microwave synthesizer system.

All final compounds (**HCT 1–15**) were synthesized according to the following synthetic procedure for **HCT13**, when incorporating the appropriate intermediates. The synthetic details and characterization information for all intermediates are shown in the ESI.†

**(E)-2-((6-Fluoroisoquinolin-1-yl)methylene)-N,N-dimethylhydrazine-1-carbothioamide** and **(Z)-2-((6-fluoroisoquinolin-1-yl)methylene)-N,N-dimethylhydrazine-1-carbothioamide (HCT13)**. To a solution of **S12** (8.6 mg, 0.0491 mmol) in EtOH (0.5 mL) was added 4,4-dimethyl-3-thiosemicarbazide (5.9 mg, 0.0491 mmol) and HCl (49  $\mu\text{L}$ , 0.294 mmol, 6 M in  $\text{H}_2\text{O}$ ). The mixture was stirred and refluxed for 1.5 hours and then cooled to room temperature (22  $^\circ\text{C}$ ). The hydrochloride salt that formed was then neutralized with saturated aqueous  $\text{NaHCO}_3$  solution (0.5 mL). The precipitate of the desired compound was collected by filtration, the filter cake sequentially washed with water and EtOH, and then dried to yield the isoquinoline **HCT13** as a pale-yellow solid containing a mixture of *E*- and *Z*-isomers (7.4 mg, 55%).  $^1\text{H}$  NMR (500 MHz,  $\text{DMSO-}d_6$ )  $\delta$  15.90 (s, 0.21H), 11.30 (s, 1H), 9.87 (dd,  $J$  = 9.5, 5.9 Hz, 1H), 8.91 (dd,  $J$  = 9.4, 5.4 Hz, 0.21H), 8.66 (m, 1.21H), 8.59 (s, 0.21H), 8.55 (d,  $J$  = 5.6 Hz, 1H), 7.97 (d,  $J$  = 5.6 Hz, 0.21H), 7.91 (dd,  $J$  = 9.6, 2.7 Hz, 0.21H), 7.79–7.82 (m, 2H), 7.73 (td,  $J$  = 9.1, 2.7 Hz, 0.21H), 7.62 (ddd,  $J$  = 9.6, 8.6, 2.8 Hz, 1H), 3.40 (s, 1.26H), 3.33 (s, 6H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{DMSO-}d_6$ )  $\delta$  180.78 (2C), 163.19 (d,  $^1\text{J}_{\text{C-F}}$  = 251.7 Hz), 162.66 (d,  $^1\text{J}_{\text{C-F}}$  = 250.6 Hz), 151.99 (d,  $^5\text{J}_{\text{C-F}}$  = 1.2 Hz), 150.63 (d,  $^5\text{J}_{\text{C-F}}$  = 0.9 Hz), 147.57, 143.42, 141.42, 138.83 (d,  $^3\text{J}_{\text{C-F}}$  = 15.5 Hz), 138.76 (d,  $^4\text{J}_{\text{C-F}}$  = 10.7 Hz), 132.08 (d,  $^3\text{J}_{\text{C-F}}$  = 9.3 Hz), 131.62, 128.64 (d,  $^3\text{J}_{\text{C-F}}$  = 9.9 Hz), 124.14, 123.10, 122.12 (d,  $^4\text{J}_{\text{C-F}}$  = 5.2 Hz), 121.56 (d,  $^4\text{J}_{\text{C-F}}$  = 5.1 Hz), 119.54 (d,  $^2\text{J}_{\text{C-F}}$  = 25.6 Hz), 119.14 (d,  $^2\text{J}_{\text{C-F}}$  = 24.4 Hz), 111.48 (d,  $^2\text{J}_{\text{C-F}}$  = 20.8 Hz), 110.90 (d,  $^2\text{J}_{\text{C-F}}$  = 20.7

Hz), 42.04 (4C).  $^{19}\text{F}$  NMR (376 MHz,  $\text{DMSO-}d_6$ )  $\delta$  -106.34, -107.95. DART-MS:  $m/z$  calcd. for  $\text{C}_{13}\text{H}_{14}\text{FN}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  277.09232, found 277.0905.

**(E)-2-((Isoquinolin-1-yl)methylene)hydrazine-1-carbothioamide (HCT1)**.  $^1\text{H}$  NMR (500 MHz,  $\text{DMSO-}d_6$ )  $\delta$  11.74 (s, 1H), 9.19 (d,  $J$  = 8.5 Hz, 1H), 8.60–8.54 (m, 2H), 8.49 (br s, 1H), 8.02 (d,  $J$  = 8.1 Hz, 1H), 7.86 (d,  $J$  = 5.6 Hz, 1H), 7.84–7.78 (m, 2H), 7.75 (ddd,  $J$  = 8.3, 6.8, 1.4 Hz, 1H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{DMSO-}d_6$ )  $\delta$  178.41, 150.78, 145.99, 142.13, 136.24, 130.47, 129.08, 127.22, 126.94, 125.58, 121.77. DART-MS:  $m/z$  calcd. for  $\text{C}_{11}\text{H}_{10}\text{N}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  231.06989, found 231.06938.

**(E)-2-((4-Fluoroisoquinolin-1-yl)methylene)hydrazine-1-carbothioamide (HCT2)**.  $^1\text{H}$  NMR (500 MHz,  $\text{DMSO-}d_6$ )  $\delta$  11.70 (s, 1H), 9.28 (d,  $J$  = 8.5 Hz, 1H), 8.56 (d,  $J$  = 1.5 Hz, 1H), 8.53 (s, 1H), 8.48 (s, 1H), 8.13 (d,  $J$  = 8.2 Hz, 1H), 7.94 (ddd,  $J$  = 8.2, 7.0, 0.9 Hz, 1H), 7.85 (m, 2H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{DMSO-}d_6$ )  $\delta$  178.84, 154.73 (d,  $^1\text{J}_{\text{C-F}}$  = 262.2 Hz), 148.03 (d,  $^3\text{J}_{\text{C-F}}$  = 5.2 Hz), 145.84, 131.75, 130.69, 128.10 (d,  $^2\text{J}_{\text{C-F}}$  = 23.3 Hz), 127.75, 127.35 (d,  $^4\text{J}_{\text{C-F}}$  = 3.0 Hz), 126.51 (d,  $^2\text{J}_{\text{C-F}}$  = 14.9 Hz), 119.79 (d,  $^3\text{J}_{\text{C-F}}$  = 4.6 Hz).  $^{19}\text{F}$  NMR (376 MHz,  $\text{DMSO-}d_6$ )  $\delta$  -137.31. DART-MS:  $m/z$  calcd. for  $\text{C}_{11}\text{H}_9\text{FN}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  249.06047, found 249.05042.

**(E)-2-((6-Fluoroisoquinolin-1-yl)methylene)hydrazine-1-carbothioamide (HCT3)**.  $^1\text{H}$  NMR (500 MHz,  $\text{DMSO-}d_6$ )  $\delta$  11.74 (s, 1H), 9.30 (dd,  $J$  = 9.4, 5.8 Hz, 1H), 8.55 (d,  $J$  = 5.6 Hz, 1H), 8.51 (s, 2H), 7.80–7.85 (m, 3H), 7.57 (td,  $J$  = 9.0, 2.8 Hz, 1H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{DMSO-}d_6$ )  $\delta$  178.88, 162.70 (d,  $^1\text{J}_{\text{C-F}}$  = 250.4 Hz), 151.35, 146.33, 143.50, 138.62 (d,  $^3\text{J}_{\text{C-F}}$  = 10.7 Hz), 131.52 (d,  $^3\text{J}_{\text{C-F}}$  = 9.5 Hz), 123.24, 121.79 (d,  $^4\text{J}_{\text{C-F}}$  = 5.0 Hz), 119.36 (d,  $^2\text{J}_{\text{C-F}}$  = 24.5 Hz), 110.86 (d,  $^2\text{J}_{\text{C-F}}$  = 20.7 Hz).  $^{19}\text{F}$  NMR (376 MHz,  $\text{DMSO-}d_6$ )  $\delta$  -107.79, extraneous peak found at -106.49. DART-MS:  $m/z$  calcd. for  $\text{C}_{11}\text{H}_{10}\text{FN}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  249.06047, found 249.05984.

**(E)-2-((5-(Methylamino)isoquinolin-1-yl)methylene)hydrazine-1-carbothioamide (HCT4)**.  $^1\text{H}$  NMR (500 MHz,  $\text{DMSO-}d_6$ )  $\delta$  12.32 (s, 1H), 9.07 (br s, 1H), 8.92 (s, 1H), 8.90 (s, 1H), 8.57 (d,  $J$  = 6.6 Hz, 1H), 8.52 (d,  $J$  = 6.7 Hz, 1H), 7.85 (t,  $J$  = 8.2 Hz, 1H), 7.56 (d,  $J$  = 8.3 Hz, 1H), 7.30 (br s, 1H), 7.01 (d,  $J$  = 8.0 Hz, 1H), 2.91 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{DMSO-}d_6$ )  $\delta$  179.40, 146.25, 146.00, 133.09, 130.09, 128.72, 126.75 (2C), 119.38, 111.17, 110.66, 30.39. DART-MS:  $m/z$  calcd. for  $\text{C}_{12}\text{H}_{14}\text{N}_5\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  260.09644, found 260.09501.

**(E)-2-((5-Aminoisoquinolin-1-yl)methylene)hydrazine-1-carbothioamide (HCT5)**.  $^1\text{H}$  NMR (500 MHz,  $\text{DMSO-}d_6$ )  $\delta$  11.66 (br s, 1H), 8.57 (s, 1H), 8.42 (d,  $J$  = 5.8 Hz, 1H), 8.31 (br s, 1H), 8.25 (d,  $J$  = 8.5 Hz, 1H), 7.98 (d,  $J$  = 5.8 Hz, 1H), 7.60 (br s, 1H), 7.42 (t,  $J$  = 8.1 Hz, 1H), 6.89 (d,  $J$  = 7.1 Hz, 1H), 6.02 (s, 2H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{DMSO-}d_6$ )  $\delta$  178.46, 150.36, 145.86, 144.62, 140.01, 129.74, 126.78, 125.83, 116.50, 113.12, 110.74. DART-MS:  $m/z$  calcd. for  $\text{C}_{11}\text{H}_{11}\text{N}_5\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  246.08079, found 246.08020.

**(E)-2-((Isoquinolin-1-yl)methylene)-N-methylhydrazine-1-carbothioamide (HCT6)**.  $^1\text{H}$  NMR (500 MHz,  $\text{DMSO-}d_6$ )  $\delta$  11.78 (br s, 1H), 9.11 (br s, 1H), 8.61 (s, 1H), 8.56 (d,  $J$  = 5.6 Hz, 1H), 8.31 (br s, 1H), 8.02 (d,  $J$  = 8.1 Hz, 1H), 7.89–7.79 (m,

2H), 7.76 (t,  $J = 7.7$  Hz, 1H), 3.07 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz, DMSO- $d_6$ )  $\delta$  178.36, 151.06, 144.76, 142.15, 136.25, 130.42, 128.86, 127.21, 126.82, 125.64, 121.48, 31.34. DART-MS:  $m/z$  calcd. for  $\text{C}_{12}\text{H}_{13}\text{N}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  245.08554, found 245.08505.

**(E)-2-((4-Fluoroisoquinolin-1-yl)methylene)-N-methylhydrazine-1-carbothioamide (HCT7).**  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ )  $\delta$  11.76 (s, 1H), 9.19 (d,  $J = 8.6$  Hz, 1H), 8.56 (d,  $J = 1.4$  Hz, 1H), 8.56 (s, 1H), 8.34 (d,  $J = 4.4$  Hz, 1H), 8.14 (d,  $J = 8.3$  Hz, 1H), 7.94–7.97 (m, 1H), 7.85–7.89 (m, 1H), 3.06 (d,  $J = 4.6$  Hz, 3H).  $^{13}\text{C}$  NMR (125 MHz, DMSO- $d_6$ )  $\delta$  178.84, 154.73 (d,  $^1\text{J}_{\text{C-F}} = 262.2$  Hz), 148.03 (d,  $^3\text{J}_{\text{C-F}} = 5.2$  Hz), 145.84, 131.75, 130.69, 128.10 (d,  $^2\text{J}_{\text{C-F}} = 23.3$  Hz), 127.75, 127.35 (d,  $^4\text{J}_{\text{C-F}} = 3.0$  Hz), 126.51 (d,  $^2\text{J}_{\text{C-F}} = 14.9$  Hz), 119.79 (d,  $^3\text{J}_{\text{C-F}} = 4.6$  Hz), 31.86.  $^{19}\text{F}$  NMR (376 MHz, DMSO- $d_6$ )  $\delta$  -137.53, extraneous peak found at -134.32. DART-MS:  $m/z$  calcd. for  $\text{C}_{12}\text{H}_{12}\text{FN}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  263.07612, found 263.07520.

**(E)-2-((6-Fluoroisoquinolin-1-yl)methylene)-N-methylhydrazine-1-carbothioamide (HCT8).**  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ )  $\delta$  11.80 (s, 1H), 9.20 (dd,  $J = 9.4, 5.7$  Hz, 1H), 8.55 (d,  $J = 5.6$  Hz, 1H), 8.54 (s, 1H), 8.35 (d,  $J = 4.7$  Hz, 1H), 7.83 (dd,  $J = 9.2, 3.9$  Hz, 2H), 7.60 (td,  $J = 9.0, 2.7$  Hz, 1H), 3.06 (d,  $J = 4.5$  Hz, 3H).  $^{13}\text{C}$  NMR (125 MHz, DMSO- $d_6$ )  $\delta$  178.56, 162.71 (d,  $^1\text{J}_{\text{C-F}} = 250.4$  Hz), 151.55, 145.22, 143.53, 138.62 (d,  $^3\text{J}_{\text{C-F}} = 10.6$  Hz), 131.28 (d,  $^3\text{J}_{\text{C-F}} = 9.5$  Hz), 123.29, 121.67 (d,  $^4\text{J}_{\text{C-F}} = 5.1$  Hz), 119.23 (d,  $^2\text{J}_{\text{C-F}} = 24.8$  Hz), 110.89 (d,  $^2\text{J}_{\text{C-F}} = 20.8$  Hz), 31.85.  $^{19}\text{F}$  NMR (376 MHz, DMSO- $d_6$ )  $\delta$  -106.55, extraneous peak found at -107.74. DART-MS:  $m/z$  calcd. for  $\text{C}_{12}\text{H}_{12}\text{FN}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  263.07612, found 263.07538.

**(E)-N-Methyl-2-((5-(methylamino)isoquinolin-1-yl)methylene)hydrazine-1-carbothioamide and (Z)-N-methyl-2-((5-(methylamino)isoquinolin-1-yl)methylene)hydrazine-1-carbothioamide (HCT9).**  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ )  $\delta$  14.74 (s, 0.15H), 12.22 (s, 1H), 9.39 (br s, 1H), 8.93 (q,  $J = 4.7$  Hz, 0.15H), 8.78 (s, 1H), 8.54 (d,  $J = 5.9$  Hz, 0.15H), 8.50 (d,  $J = 6.5$  Hz, 1H), 8.38 (s, 1H), 8.17 (s, 0.15H), 8.12 (d,  $J = 5.9$  Hz, 0.15H), 7.82 (d,  $J = 8.6$  Hz, 0.15H), 7.76 (t,  $J = 8.1$  Hz, 1H), 7.69 (s, 1H), 7.56 (t,  $J = 8.1$  Hz, 0.15H), 7.08 (br s, 1H), 6.92 (d,  $J = 7.7$  Hz, 1.15H), 6.72 (d,  $J = 7.8$  Hz, 0.15H), 3.07 (d,  $J = 4.6$  Hz, 3H), 3.02 (d,  $J = 4.6$  Hz, 0.45 H), 2.88 (s, 3H), 2.86 (s, 0.45H).  $^{13}\text{C}$  NMR (125 MHz, DMSO- $d_6$ )  $\delta$  178.84, 178.38, 150.11, 147.37, 145.83 (2C), 145.56, 138.94, 132.45, 130.58, 129.19, 128.23, 128.22, 126.87 (2C), 126.80, 118.47, 117.04, 111.57, 110.13, 109.63, 106.57, 31.59, 31.42, 30.42 (2C). DART-MS:  $m/z$  calcd. for  $\text{C}_{13}\text{H}_{16}\text{N}_5\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  274.11209, found 274.11104.

**(E)-2-((5-Aminoisoquinolin-1-yl)methylene)-N-methylhydrazine-1-carbothioamide and (Z)-2-((5-aminoisoquinolin-1-yl)methylene)-N-methylhydrazine-1-carbothioamide (HCT10).**  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ )  $\delta$  14.80 (s, 0.08H), 11.66 (br s, 1H), 8.95 (d,  $J = 4.9$  Hz, 0.08H), 8.62 (s, 1H), 8.52 (d,  $J = 5.9$  Hz, 0.8H), 8.42 (d,  $J = 5.8$  Hz, 1H), 8.25 (d,  $J = 3.3$  Hz, 1H), 8.18 (s, 0.08H), 8.11–8.15 (m, 1.08H), 7.99 (d,  $J = 5.9$  Hz, 1H), 7.83 (d,  $J = 8.4$  Hz, 0.08H), 7.48 (t,  $J = 7.9$  Hz, 0.08H), 7.43 (t,  $J = 8.0$  Hz, 1H), 6.97 (d,  $J = 7.6$  Hz, 0.08H), 6.91 (dd,  $J = 7.6, 0.9$  Hz, 1H), 6.21 (s, 0.16H), 6.04 (s, 2H), 3.05–3.07 (m, 3.24H).  $^{13}\text{C}$  NMR (125 MHz, DMSO- $d_6$ )  $\delta$

178.85, 178.52, 150.83, 150.19, 145.53, 145.20, 145.10, 140.49, 138.40, 130.28, 130.10, 129.16, 128.41, 127.36, 126.27, 126.16, 117.81, 116.96, 113.24, 111.66, 111.18, 110.61, 31.74, 31.59. DART-MS:  $m/z$  calcd. for  $\text{C}_{12}\text{H}_{14}\text{N}_5\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  260.09644, found 260.09563.

**(E)-2-(Isoquinolin-1-ylmethylene)-N,N-dimethylhydrazine-1-carbothioamide and (Z)-2-(isoquinolin-1-ylmethylene)-N,N-dimethylhydrazine-1-carbothioamide (HCT11).**  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ )  $\delta$  15.99 (s, 0.33H), 11.26 (br s, 1H), 9.77 (dd,  $J = 8.8, 5.1$  Hz, 1H), 8.81 (d,  $J = 8.6$  Hz, 0.33H), 8.70 (d,  $J = 1.7$  Hz, 1H), 8.69 (d,  $J = 1.2$  Hz, 0.33H), 8.63 (s, 0.33H), 8.55 (d,  $J = 5.5$  Hz, 1H), 8.12 (d,  $J = 8.2$  Hz, 0.33H), 8.01–7.96 (m, 1.33H), 7.92 (ddd,  $J = 8.1, 7.0, 1.1$  Hz, 0.33H), 7.88–7.76 (m, 2.33H), 7.72 (ddd,  $J = 8.4, 6.8, 1.4$  Hz, 1H), 3.43 (s, 1.98H), 3.35 (s, 6H).  $^{13}\text{C}$  NMR (125 MHz, DMSO- $d_6$ )  $\delta$  180.88, 180.81, 151.94, 150.58, 147.81, 142.50, 140.45, 136.86, 136.82, 131.80, 130.77, 129.48, 129.16, 128.21, 128.17 (2C), 127.68, 126.83, 125.87, 124.60, 122.53, 121.93, 42.08 (4C). DART-MS:  $m/z$  calcd. for  $\text{C}_{13}\text{H}_{15}\text{N}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  259.10119, found 259.10080.

**(E)-2-((4-Fluoroisoquinolin-1-yl)methylene)-N,N-dimethylhydrazine-1-carbothioamide and (Z)-2-((4-fluoroisoquinolin-1-yl)methylene)-N,N-dimethylhydrazine-1-carbothioamide (HCT12).**  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ )  $\delta$  15.52 (s, 0.15H), 11.28 (s, 1H), 9.87 (d,  $J = 8.7$  Hz, 1H), 8.87 (d,  $J = 9.0$  Hz, 0.15H), 8.77 (d,  $J = 1.9$  Hz, 0.15H), 8.69 (s, 1H), 8.57 (d,  $J = 1.6$  Hz, 1.15H), 8.23 (d,  $J = 8.2$  Hz, 0.15H), 8.15 (d,  $J = 8.2$  Hz, 1H), 8.02–8.05 (m, 0.15H), 7.92–7.97 (m, 1.15H), 7.86 (ddd,  $J = 8.3, 6.9, 1.3$  Hz, 1H), 3.42 (s, 0.90H), 3.36 (s, 6H).  $^{13}\text{C}$  NMR (125 MHz, DMSO- $d_6$ )  $\delta$  180.79, 180.73, 154.60 (d,  $^1\text{J}_{\text{C-F}} = 261.7$  Hz), 154.13 (d,  $^1\text{J}_{\text{C-F}} = 261.8$  Hz), 148.75 (d,  $^3\text{J}_{\text{C-F}} = 5.1$  Hz), 147.76 (d,  $^3\text{J}_{\text{C-F}} = 5.7$  Hz), 147.05, 132.52, 131.59 (d,  $^4\text{J}_{\text{C-F}} = 5.1$  Hz), 131.10, 130.49, 130.30, 128.60 (d,  $^4\text{J}_{\text{C-F}} = 3.3$  Hz), 128.41 (d,  $^4\text{J}_{\text{C-F}} = 1.0$  Hz), 127.90 (d,  $^2\text{J}_{\text{C-F}} = 23.3$  Hz), 127.19 (d,  $^4\text{J}_{\text{C-F}} = 2.6$  Hz), 126.88 (d,  $^2\text{J}_{\text{C-F}} = 14.8$  Hz), 126.71 (d,  $^2\text{J}_{\text{C-F}} = 14.7$  Hz), 126.35 (d,  $^2\text{J}_{\text{C-F}} = 25.2$  Hz), 124.98, 120.23 (d,  $^3\text{J}_{\text{C-F}} = 4.3$  Hz), 119.79 (d,  $^3\text{J}_{\text{C-F}} = 4.7$  Hz), 42.04 (4C).  $^{19}\text{F}$  NMR (376 MHz, DMSO- $d_6$ )  $\delta$  -134.93, -138.02. DART-MS:  $m/z$  calcd. for  $\text{C}_{13}\text{H}_{14}\text{FN}_4\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  277.09177, found 277.09096.

**(E)-2-((6-Fluoro-5-(methylamino)isoquinolin-1-yl)methylene)-N,N-dimethylhydrazine-1-carbothioamide and (Z)-2-((6-fluoro-5-(methylamino)isoquinolin-1-yl)methylene)-N,N-dimethylhydrazine-1-carbothioamide (HCT14).**  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ )  $\delta$  15.96 (s, 0.17H), 11.22 (Br s, 1H), 9.20 (s, 1H), 8.62–8.54 (m, 1.17H), 8.52 (s, 0.17H), 8.34 (d,  $J = 5.5$  Hz, 1H), 8.20 (d,  $J = 6.2$  Hz, 0.17H), 8.07 (dd,  $J = 9.3, 4.2$  Hz, 0.17H), 7.87 (br s, 1H), 7.56 (dd,  $J = 13.6, 9.2$  Hz, 0.17H), 7.33 (dd,  $J = 13.4, 9.5$  Hz, 1H), 6.10 (br s, 0.17H), 5.69 (br s, 1H), 3.41 (s, 1.02H), 3.27 (s, 6H), 3.10 (t,  $J = 5.5$  Hz, 0.51H), 3.05 (t,  $J = 5.2$  Hz, 3H). A  $^{13}\text{C}$  NMR was not obtained.  $^{19}\text{F}$  NMR (376 MHz, CDCl $_3$ )  $\delta$  -129.05, -129.53. DART-MS:  $m/z$  calcd. for  $\text{C}_{14}\text{H}_{17}\text{FN}_5\text{S}$  ( $\text{M} + \text{H}$ ) $^+$  306.11832, found 306.11719.

**(E)-2-((4-Fluoro-5-(methylamino)isoquinolin-1-yl)methylene)-N,N-dimethylhydrazine-1-carbothioamide and (Z)-2-((4-fluoro-5-(methylamino)isoquinolin-1-yl)methylene)-N,N-dimethylhydrazine-1-carbothioamide (HCT15).**  $^1\text{H}$  NMR (500

MHz, DMSO-*d*<sub>6</sub>)  $\delta$  15.46 (s, 0.33H), 11.13 (br s, 1H), 8.91 (dd,  $J = 8.4, 2.9$  Hz, 1H), 8.62 (s, 1H), 8.50 (d,  $J = 5.1$  Hz, 0.33H), 8.39 (s, 0.33H), 8.32 (d,  $J = 4.8$  Hz, 1H), 7.89 (dd,  $J = 8.5, 2.9$  Hz, 0.33H), 7.65 (t,  $J = 8.2$  Hz, 0.33H), 7.57 (t,  $J = 8.2$  Hz, 1H), 6.82 (d,  $J = 8.0$  Hz, 0.33H), 6.73 (d,  $J = 7.9$  Hz, 1H), 6.55 (dd,  $J = 11.9, 5.2$  Hz, 0.33H), 6.39 (dd,  $J = 12.4, 5.0$  Hz, 1H), 3.37 (s, 1.98H), 3.31 (s, 6H), 2.86–2.84 (m, 3.99H). <sup>13</sup>C NMR (125 MHz, DMSO-*d*<sub>6</sub>)  $\delta$  180.95, 180.72, 156.41 (d,  $J = 260.4$  Hz), 147.99 (d, <sup>4</sup>*J*C–F = 4.3 Hz), 147.41, 147.16, 144.92, 144.61 (d, <sup>4</sup>*J*C–F = 3.7 Hz), 131.90, 131.69, 130.83, 130.78, 129.29 (d, <sup>4</sup>*J*C–F = 2.4 Hz), 127.41 (d, <sup>2</sup>*J*C–F = 28.8 Hz), 125.43 (d, <sup>2</sup>*J*C–F = 30.5 Hz), 115.98 (d, <sup>2</sup>*J*C–F = 7.6 Hz), 113.89, 113.84, 110.26, 108.50, 107.78, 42.15 (4C), 30.95 (2C), one low-field carbon were either not observed or is overlapping with another low-field carbon. <sup>19</sup>F NMR (376 MHz, DMSO-*d*<sub>6</sub>)  $\delta$  –125.86, –129.02. DART-MS: *m/z* calcd. for C<sub>14</sub>H<sub>17</sub>FN<sub>5</sub>S (M + H)<sup>+</sup> 306.11832, found 306.11716.

**Cu[HCT13].** HR-MS (ESI<sup>+</sup>) data: *m/z* calculated for [C<sub>13</sub>H<sub>12</sub>CuFN<sub>4</sub>S]<sup>+</sup> = 338.0057; found 338.0038; *m/z* calculated for [C<sub>13</sub>H<sub>12</sub>CuFN<sub>4</sub>S + MeCN]<sup>+</sup> = 379.0323; found 379.0297 (Thermo LTQ-Orbitrap XL). Refer to Supplementary Information for HPLC trace and ESI data.

### Cell culture and culture conditions

Pancreatic adenocarcinoma cell lines: PATU8988T, MIAPACA2, SU8686, PSN1, HPAC, BXPC3, DANG, SUIT2, A13A, CAPAN2, T3M4, A2.1, HUPT4, XWR200, L36PL, YAPC, PANC0327, PANC1, PATU8902, HPAF11, ASPC1, PANC0813, PANC0203, HS766T, SW1990, and CFPAC1; prostate cancer cell lines: 22Rv1, LNCaP, RM1 and C4-2; and small cell lung carcinoma cell lines: NCI-H526, NCI-H146, and NCI-H1963 were obtained from American Type Culture Collection (ATCC). 143 BTK WT and 143 BTK  $\rho_0$ , BJ WT and BJ  $\rho_0$  cells were gifts from Prof. Michael Teitell in UCLA. Murine Prostate cancer cell lines MyC CaP was a kind gift from Prof. DLJ Thorek at WUSTL. Murine Pancreatic cancer cells KP4662 was kind gift from Prof. Robert Vonderheide at UPenn. Immortalized human pancreatic ductal epithelial (HPDE) cells were a kind gift from Dr. Ming-Sound Tsao at Ontario Cancer Institute (Invitrogen Cat#20). With a few exceptions, cell lines were cultured in DMEM (Corning) or RPMI (Corning) containing 10% fetal bovine serum (FBS, Omega Scientific) and were grown at 37 °C, 20% O<sub>2</sub> and 5% CO<sub>2</sub>. All cultured cells were incubated in antibiotic free media and were regularly tested for mycoplasma contamination using MycoAlert kit (Lonza) following the manufacturer's instructions, except that the reagents were diluted 1 : 4 from their recommended amount.

### HCT-13 stock solution

HCT-13 was solubilized up to a concentration of 20 mg mL<sup>-1</sup> in an aqueous solution of 40% Captisol with 1% DMSO with the aid of heating at 50 °C and sonication for 15 minutes.

### Proliferation assay

Cells were plated in 384-well plates (500 cells per well for adherent cell lines in 30  $\mu$ l volume). Drugs were serially diluted to the desired concentrations and an equivalent volume of DMSO was added to vehicle control. Following 72 h incubation, ATP content was measured using CellTiter-Glo reagent according to manufacturer's instructions (Promega, CellTiter-Glo Luminescent Cell Viability Assay), and analyzed by SpectraMax luminometer (Molecular Devices). IC<sub>50</sub> and IC<sub>90</sub> values, concentrations required to inhibit proliferation by 50% and 90% respectively compared to DMSO treated cells, were calculated using Prism 6.0 h (Graphpad Software). The 430-member protein kinase inhibitor library used for the chemical genomics screen was purchased from Selleckchem, catalog no. L1200.

### Western blot

Cells were lysed using RIPA buffer supplemented with protease (ThermoFisher, 78430) and phosphatase (ThermoFisher, 78420) inhibitors, scraped, sonicated, and centrifuged (20 000g at 4 °C). Protein concentrations in the supernatant were determined using the Micro BCA Protein Assay kit (Thermo), and equal amounts of protein were resolved on pre-made bis-tris polyacrylamide gels (Life Technologies). Primary antibodies: pAMPK<sub>T172</sub> (cell signaling, #2535, 1:1000), HO-1 (cell signaling, #5061S, 1:1000), pS345 CHEK1 (cell signaling, #2348L, 1:1000), pT68 CHEK2 (cell signaling, #2197S, 1:1000), pS139 H2A.X (Millipore, 05-636, 1:1000), clvd. Casp3 (cell signaling, #9662, 1:1000), and anti-actin (Cell Signaling Technology, 9470, 1:10 000). Primary antibodies were stored in 5% BSA (Sigma-Aldrich) and 0.1% NaN<sub>3</sub> in TBST solution. Anti-rabbit IgG HRP-linked (Cell Signaling Technology, 7074, 1:2500) and anti-mouse IgG HRP-linked (Cell Signaling Technology, 7076, 1:2500) were used as secondary antibodies. Chemiluminescent substrates (ThermoFisher Scientific, 34077 and 34095) and autoradiography film (Denville) were used for detection.

### Viability/apoptosis assay

Viable cells were measured by trypan blue staining using a Vi-Cell counter (Beckman Coulter, CA, USA). Apoptosis and cell death were assayed using annexin V-FITC and PI according to manufacturer's instructions (FITC annexin V apoptosis detection kit, BD Sciences, #556570).

### Cell cycle

Cell cycle was assessed using propidium iodide staining at indicated timepoints. Cells were pulsed with EdU 1 h before collection at different time points. Cells were fixed 4% paraformaldehyde, permeabilized with perm/wash reagent (Invitrogen), stained with Azide-AF647 (using click-chemistry, Invitrogen; Click-iT EdU Flow cytometry kit, #C10634) and FxCycle-Violet (Invitrogen), and then analyzed by flow cytometry.

## ROS measurements

Cellular ROS measurement was assayed with CM-H<sub>2</sub>DCFDA staining after treatment according to manufacturer's instructions (reactive oxygen species (ROS) detection reagents, Invitrogen, #D399). The cells were then incubated with 5 μM of CM-H<sub>2</sub>DCFDA for 30 min, spun down at 450g for 4 min, and the supernatant was replaced with fresh media containing lethal compounds and/or Cu(II). Then, the cells were incubated for 30 min, spun down, and the supernatant was replaced with PBS. The samples were analyzed using flow cytometry.

Mitochondrial ROS was measured using MitoSOX staining according to manufacturer's instructions (MitoSOX, Invitrogen, #M36008). Cells were treated with HCT-13, washed and treated with MitoSOX. Cells were then incubated for 30 minutes at 37 °C. After incubation, media is aspirated and cells are washed with PBS and analyzed by flow cytometry.

## Mito stress test and electron flow assay

All oxygen consumption rate (OCR) was measured using a XF24 Analyzer (Agilent) and normalized per μg protein. For cellular OCR, cells were incubated in unbuffered DMEM containing 25 mM glucose, 1 mM pyruvate and 2 mM glutamine. OCR was measured before (total respiration) and after the sequential injection of 1 μM oligomycin (complex V inhibitor), 0.75 μM FCCP (uncoupler), and 1 μM of rotenone and myxothiazol (complex I and III inhibitors, respectively), as described previously.<sup>54</sup> Mitochondrial respiration was calculated by subtracting the non-mitochondrial respiration left after rotenone and myxothiazol injection. Oligomycin-sensitive respiration represents ATP-linked respiration (coupled respiration).

To measure electron transport chain complex activity from cells, cells were incubated in MAS buffer with 10 mM pyruvate (complex I substrate), 2 mM malate, 4 μM FCCP, 4 mM ADP, and 1 nM of XF plasma membrane permeabilizer (PMP) reagent (Agilent). OCR was measured before and after the sequential injection of 2 μM rotenone, 10 mM succinate (complex II substrate), 4 μM antimycin A (complex III inhibitor), and a mix of 10 mM ascorbate and 100 μM TMPD (complex IV substrates), as described previously.<sup>55</sup> Antimycin A-sensitive respiration represents the complex III respiration.

To measure OCR directly from mitochondria, mitochondria were isolated from fresh mouse liver by dual centrifugation at 800g and 8000g and seeded by centrifugation.<sup>55</sup> Mitochondria were incubated with 1 mM pyruvate (complex I), 2 mM malate, 4 μM FCCP in MAS buffer, as well as the "corresponding drugs" for 30 min at 37 °C. OCR was measured before and after the sequential injections described in the previous paragraph.

## Intracellular Cu(II) measurement

Cells were plated in 6-well plates and cultured for one day. Vehicle of HCT-13 were added to the cells the following day

and incubated for 24 hours. The plates were then washed 2 times with PBS containing 1 mM EDTA and 2 times with PBS alone. The concentration of Cu(II) was measured using inductive coupled plasma mass spectrometry (ICP-MS) using standard procedure.

## FACS analyses

All flow cytometry data were acquired on a five-laser LSRII cytometer (BD), and analyzed using the FlowJo software (Tree Star).

## Isotopic labeling in cell culture

Cells were transferred into DMEM without glucose and supplemented with 10% dialyzed FBS (Gibco) containing [U-13C<sub>6</sub>]glucose (Sigma-Aldrich, 389374) at 11 mM. The cells were incubated for 48 h before sample collection and processing as previously described.<sup>51</sup>

## Animal studies

Mice were housed under specific pathogen-free conditions and were treated in accordance with UCLA Animal Research Committee protocol guidelines. All C57BL/6 female mice were purchased from the UCLA Radiation Oncology breeding colony. All NCG female mice were purchased from the Jackson Labs (JAX).

## In vivo leukemia models and treatment regimens

All animal studies were approved by the UCLA Animal Research Committee (ARC). For development of systemic murine B ALL model, C57BL/6 female mice were injected intravenously with 200 000 firefly luciferase expressing p185BCR-ABL Arf<sup>-/-</sup> pre-B-ALL cells (kindly gifted by Dr. Nidal Boulous and the CERN Foundation).<sup>59,82</sup> For development of the systemic human AML model, NCG female mice (from Jackson Labs) were injected intravenously with 5 × 10<sup>6</sup> firefly luciferase expressing MV4-11 cells.<sup>83</sup> The leukemic burden was monitored using bioluminescence imaging. All Cu[HCT-13] treatments were performed using a formulation consisting of 40% Captisol and 1% DMSO. The treatments were performed by intra-peritoneal (i.p.) injections using 100 μL volume daily.

## Abbreviations used

HCTs	α-N-Heterocyclic carboxaldehyde thiosemicarbazones
3-AP	3-Aminopyridine-2-carboxaldehyde thiosemicarbazone
RNR	Ribonucleotide reductase
ROS	Reactive oxygen species
PDAC	Pancreatic ductal adenocarcinoma
SCLC	Small cell lung carcinoma
PC	Prostate cancer
BCPS	Bathocuproine disulfonate
mtETC	Mitochondrial electron transport chain
DDR/RSR	DNA damage response/replication stress response

ATR	Ataxia-telangiectasia mutated (ATM) and Rad3-related protein kinase
dGTP	Deoxyguanosine triphosphate
OXPPOS	Oxidative phosphorylation
2-DG	2-Deoxy-D-glucose
BLI	Bioluminescence imaging
OCR	Oxygen consumption rate

## Conflicts of interest

The authors declare the following competing financial interest(s): C. G. R., J. C., and M. E. J. are co-founders of Trethera Corporation. They and the University of California hold Equity in Trethera Corporation. The University of California has patented additional intellectual property for thiosemicarbazone-based antiproliferative compounds.

## Acknowledgements

We thank Dr. Nagichettiar Satyamurthy for his expert chemistry advice, and Dr. Kym Faull at the Pasarow Mass Spectrometry Laboratory for continued tutelage regarding mass spectrometry. We additionally thank Joseph R. Capri, Anthony E. Cabebe, and Wesley R. Armstrong for their excellent technical assistance. This work was supported by the National Cancer Institute [Grant R01CA187678-01] administered by the National Institutes of Health, Trethera Corporation [Contract Number 20162965]. This material is additionally based upon work supported by the National Science Foundation Graduate Research Fellowship Program [Grant No. DGE-1650504 to E. W. R.].

## References

- 1 R. W. Brockman, J. R. Thomson, M. J. Bell and H. E. Skipper, Observations on the Antileukemic Activity of Pyridine-2-Carboxaldehyde Thiosemicarbazone and Thiocarbohydrazone, *Cancer Res.*, 1956, **16**(2), 167–170.
- 2 H. Beraldo and D. Gambino, The Wide Pharmacological Versatility of Semicarbazones, Thiosemicarbazones and Their Metal Complexes, *Mini-Rev. Med. Chem.*, 2004, **4**(1), 31–39.
- 3 A. C. Sartorelli, Effect of Chelating Agents Upon the Synthesis of Nucleic Acids and Protein: Inhibition of DNA Synthesis by 1-Formylisoquinoline Thiosemicarbazone, *Biochem. Biophys. Res. Commun.*, 1967, **27**(1), 26–32.
- 4 M. C. Liu, T. C. Lin and A. C. Sartorelli, Synthesis and Antitumor Activity of Amino Derivatives of Pyridine-2-Carboxaldehyde Thiosemicarbazone, *J. Med. Chem.*, 1992, **35**(20), 3672–3677.
- 5 K. C. Agrawal, P. D. Mooney and A. C. Sartorelli, Potential Antitumor Agents. 13. 4-Methyl-5-Amino-1-Formylisoquinoline Thiosemicarbazone, *J. Med. Chem.*, 1976, **19**(7), 970–972.
- 6 F. A. French, E. J. Blanz, J. R. DoAmaral and D. A. French, Carcinostatic Activity of Thiosemicarbazones of Formyl Heteroaromatic Compounds. VI. 1-Formylisoquinoline Derivatives Bearing Additional Ring Substituents, with Notes on Mechanism of Action, *J. Med. Chem.*, 1970, **13**(6), 1117–1124.
- 7 F. A. French and E. J. Blanz, Jr, The Carcinostatic Activity of  $\alpha$ -(N) Heterocyclic Carboxaldehyde Thiosemicarbazones I. Isoquinoline-1-Carboxaldehyde Thiosemicarbazone, *Cancer Res.*, 1965, **29**(9), 1454–1458.
- 8 R. A. Finch, M. C. Liu, A. H. Cory, J. G. Cory and A. C. Sartorelli, Triapine (3-Aminopyridine-2-Carboxaldehyde Thiosemicarbazone; 3-AP): an Inhibitor of Ribonucleotide Reductase with Antineoplastic Activity, *Adv. Enzyme Regul.*, 1999, **39**, 3–12.
- 9 Y. Yu, J. Wong, D. B. Lovejoy, D. S. Kalinowski and D. R. Richardson, Chelators at the Cancer Coalface: Desferrioxamine to Triapine and Beyond, *Clin. Cancer Res.*, 2006, **12**(23), 6876–6883.
- 10 A. B. Alvero, W. Chen, A. C. Sartorelli, P. Schwartz, T. Rutherford and G. Mor, Triapine (3-Aminopyridine-2-Carboxaldehyde Thiosemicarbazone) Induces Apoptosis in Ovarian Cancer Cells, *J. Soc. Gynecol. Invest.*, 2006, **13**(2), 145–152.
- 11 Y. Yu, E. Gutierrez, Z. Kovacevic, F. Saletta, P. Obeidy, Y. Suryo Rahmanto and D. R. Richardson, Iron Chelators for the Treatment of Cancer, *Curr. Med. Chem.*, 2012, **19**(17), 2689–2702.
- 12 A. M. Merlot, D. S. Kalinowski and D. R. Richardson, Novel Chelators for Cancer Treatment: Where Are We Now?, *Antioxid. Redox Signaling*, 2013, **18**(8), 973–1006.
- 13 T. B. Chaston, D. R. Richardson, D. B. Lovejoy and R. N. Watts, Examination of the Antiproliferative Activity of Iron Chelators: Multiple Cellular Targets and the Different Mechanism of Action of Triapine Compared with Desferrioxamine and the Potent Pyridoxal Isonicotinoyl Hydrazone Analogue 311, *Clin. Cancer Res.*, 2003, **9**, 402–414.
- 14 J. G. Cory, A. H. Cory, G. Rappa, A. Lorico, L. Mao-Chin, L. Tai-Shun and A. C. Sartorelli, Inhibitors of Ribonucleotide Reductase: Comparative Effects of Amino- and Hydroxy-Substituted Pyridine-2-Carboxaldehydethiosemicarbazones, *Biochem. Pharmacol.*, 1994, **48**(2), 335–344.
- 15 D. R. Richardson, D. S. Kalinowski, V. Richardson, P. C. Sharpe, D. B. Lovejoy, M. Islam and P. V. Bernhardt, 2-Acetylpyridine Thiosemicarbazones Are Potent Iron Chelators and Antiproliferative Agents: Redox Activity, Iron Complexation and Characterization of Their Antitumor Activity, *J. Med. Chem.*, 2009, **52**(5), 1459–1470.
- 16 A. Popović-Bijelić, C. R. Kowol, M. E. S. Lind, J. Luo, F. Himo, É. A. Enyedy, V. B. Arion and A. Gräslund, Ribonucleotide Reductase Inhibition by Metal Complexes of Triapine (3-Aminopyridine-2-Carboxaldehyde Thiosemicarbazone): a Combined Experimental and Theoretical Study, *J. Inorg. Biochem.*, 2011, **105**(11), 1422–1431.
- 17 K. Ishiguro, Z. P. Lin, P. G. Penketh, K. Shyam, R. Zhu, R. P. Baumann, Y.-L. Zhu, A. C. Sartorelli, T. J. Rutherford and E. S. Ratner, Distinct Mechanisms of Cell-Kill by Triapine and Its Terminally Dimethylated Derivative Dp44mT Due to a Loss or Gain of Activity of Their Copper(II) Complexes, *Biochem. Pharmacol.*, 2014, **91**(3), 312–322.
- 18 D. Denoyer, S. A. S. Clatworthy, S. Masaldan, P. M. Meggyesy and M. A. Cater, Heterogeneous Copper Concentrations in Cancerous Human Prostate Tissues, *Prostate*, 2015, **75**(14), 1510–1517.

- 19 E. Madsen and J. D. Gitlin, Copper and Iron Disorders of the Brain, *Annu. Rev. Neurosci.*, 2007, **30**(1), 317–337.
- 20 P. J. Jansson, T. Yamagishi, A. Arvind, N. Seebacher, E. Gutierrez, A. Stacy, S. Maleki, D. Sharp, S. Sahni and D. R. Richardson, Di-2-Pyridylketone 4,4-Dimethyl-3-Thiosemicarbazone (Dp44mT) Overcomes Multidrug Resistance by a Novel Mechanism Involving the Hijacking of Lysosomal P-Glycoprotein (Pgp), *J. Biol. Chem.*, 2015, **290**(15), 9588–9603.
- 21 M. Whitnall, J. Howard, P. Ponka and D. R. Richardson, A Class of Iron Chelators with a Wide Spectrum of Potent Antitumor Activity That Overcomes Resistance to Chemotherapeutics, *Proc. Natl. Acad. Sci. U. S. A.*, 2006, **103**(40), 14901–14906.
- 22 K. Shimada, E. Reznik, M. E. Stokes, L. Krishnamoorthy, P. H. Bos, Y. Song, C. E. Quartararo, N. C. Pagano, D. R. Carpizo, A. C. deCarvalho, D. C. Lo and B. R. Stockwell, Copper-Binding Small Molecule Induces Oxidative Stress and Cell-Cycle Arrest in Glioblastoma- Patient-Derived Cells, *Cell Chem. Biol.*, 2018, **25**(5), 585–594.
- 23 D. Chen, Q. C. Cui, H. Yang and Q. P. Dou, Disulfiram, a Clinically Used Anti-Alcoholism Drug and Copper-Binding Agent, Induces Apoptotic Cell Death in Breast Cancer Cultures and Xenografts via Inhibition of the Proteasome Activity, *Cancer Res.*, 2006, **66**(21), 10425–10433.
- 24 G. J. Brewer, R. D. Dick, D. K. Grover, V. LeClaire, M. Tseng, M. Wicha, K. Pienta, B. G. Redman, T. Jahan, V. K. Sondak, M. Strawderman, G. LeCarpentier and S. D. Merajver, Treatment of Metastatic Cancer with Tetrathiomolybdate, an Anticopper, Antiangiogenic Agent: Phase I Study, *Clin. Cancer Res.*, 2000, **6**(1), 1–10.
- 25 D. Cen, R. I. Gonzalez, J. A. Buckmeier, R. S. Kahlon, N. B. Tohidian and F. L. Meyskens, Disulfiram Induces Apoptosis in Human Melanoma Cells: a Redox-Related Process, *Mol. Cancer Ther.*, 2002, **1**(3), 197–204.
- 26 K. Müller, C. Faeh and F. Diederich, Fluorine in Pharmaceuticals: Looking Beyond Intuition, *Science*, 2007, **317**(5846), 1881–1886.
- 27 S. Purser, P. R. Moore, S. Swallow and V. Gouverneur, Fluorine in Medicinal Chemistry, *Chem. Soc. Rev.*, 2008, **37**(2), 320–330.
- 28 E. P. Gillis, K. J. Eastman, M. D. Hill, D. J. Donnelly and N. A. Meanwell, Applications of Fluorine in Medicinal Chemistry, *J. Med. Chem.*, 2015, **58**(21), 8315–8359.
- 29 C. Belle, C. Beguin, I. Gautier-Luneau, S. Hamman, C. Philouze, J. L. Pierre, F. Thomas, S. Torelli, E. Saint-Aman and M. Bonin, Dicopper(II) Complexes of H-BPMP-Type Ligands: pH-Induced Changes of Redox, Spectroscopic (<sup>19</sup>F NMR Studies of Fluorinated Complexes), Structural Properties, and Catecholase Activities, *Inorg. Chem.*, 2002, **41**(3), 479–491.
- 30 C. R. Kowol, R. Trondl, P. Heffeter, V. B. Arion, M. A. Jakupec, A. Roller, M. Galanski, W. Berger and B. K. Keppler, Impact of Metal Coordination on Cytotoxicity of 3-Aminopyridine-2-Carboxaldehyde Thiosemicarbazone (Triapine) and Novel Insights Into Terminal Dimethylation, *J. Med. Chem.*, 2009, **52**(16), 5032–5043.
- 31 C. R. Kowol, W. Miklos, S. Pfaff, S. Hager, S. Kallus, K. Pelivan, M. Kubanik, É. A. Enyedy, W. Berger, P. Heffeter and B. K. Keppler, Impact of Stepwise NH<sub>2</sub>-Methylation of Triapine on the Physicochemical Properties, Anticancer Activity, and Resistance Circumvention, *J. Med. Chem.*, 2016, **59**(14), 6739–6752.
- 32 K. C. Agrawal, B. A. Booth and A. C. Sartorelli, Potential Antitumor Agents. I. a Series of 5-Substituted 1-Formylisoquinoline Thiosemicarbazones, *J. Med. Chem.*, 1968, **11**(4), 700–703.
- 33 P. D. Mooney, B. A. Booth, E. C. Moore, K. C. Agrawal and A. C. Sartorelli, Potential Antitumor Agents. 10. Synthesis and Biochemical Properties of 5-N-Alkylamino-,N,N-Dialkylamino-, and N-Alkylacetamido-1-Formylisoquinoline Thiosemicarbazones, *J. Med. Chem.*, 1974, **17**(11), 1145–1150.
- 34 M. F. Zaltariov, M. Hammerstad, H. J. Arabshahi, K. Jovanović, K. W. Richter, M. Cazacu, S. Shova, M. Balan, N. H. Andersen, S. Radulović, J. Reynisson, K. K. Andersson and V. B. Arion, New Iminodiacetate-Thiosemicarbazone Hybrids and Their Copper(II) Complexes Are Potential Ribonucleotide Reductase R2 Inhibitors with High Antiproliferative Activity, *Inorg. Chem.*, 2017, **56**(6), 3532–3549.
- 35 K. Malarz, A. Mrozek-Wilczkiewicz, M. Serda, M. Rejmund, J. Polanski and R. Musiol, The Role of Oxidative Stress in Activity of Anticancer Thiosemicarbazones, *Oncotarget*, 2018, **9**(25), 17689–17710.
- 36 K. C. Park, L. Fouani, P. J. Jansson, D. Wooi, S. Sahni, D. J. R. Lane, D. Palanimuthu, H. C. Lok, Z. Kovacevic, M. L. H. Huang, D. S. Kalinowski and D. R. Richardson, Copper and Conquer: Copper Complexes of Di-2-Pyridylketone Thiosemicarbazones as Novel Anti-Cancer Therapeutics, *Metallomics*, 2016, **8**(9), 874–886.
- 37 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman and D. J. Fox, *Gaussian 16, Revision A.03*, Gaussian, Inc., Wallingford CT, 2016.
- 38 D. Denoyer, H. B. Pearson, S. A. S. Clatworthy, Z. M. Smith, P. S. Francis, R. M. Llanos, I. Volitakis, W. A. Phillips, P. M. Meggyesy, S. Masaldan and M. A. Cater, Copper as a Target for Prostate Cancer Therapeutics: Copper-Ionophore Pharmacology and Altering Systemic Copper Distribution, *Oncotarget*, 2016, **7**(24), 37064–37080.



- 39 M. A. Cater, H. B. Pearson, K. Wolyniec, P. Klaver, M. Bilandzic, B. M. Paterson, A. I. Bush, P. O. Humbert, S. La Fontaine, P. S. Donnelly and Y. Haupt, Increasing Intracellular Bioavailable Copper Selectively Targets Prostate Cancer Cells, *ACS Chem. Biol.*, 2013, **8**(7), 1621–1631.
- 40 R. Safi, E. R. Nelson, S. K. Chitneni, K. J. Franz, D. J. George, M. R. Zalutsky and D. P. McDonnell, Copper Signaling Axis as a Target for Prostate Cancer Therapeutics, *Cancer Res.*, 2014, **74**(20), 5819–5831.
- 41 J. Kodydkova, L. Vavrova, B. Stankova, J. Macasek, T. Krechler and A. Zak, Antioxidant Status and Oxidative Stress Markers in Pancreatic Cancer and Chronic Pancreatitis, *Pancreas*, 2013, **42**(4), 614–621.
- 42 M. R. Lener, R. J. Scott, A. Wiechowska-Kozłowska, P. Serrano-Fernández, P. Baszuk, K. Jaworska-Bieniek, G. Sukiennicki, W. Marciniak, M. Muszyńska, J. Kładny, T. Gromowski, K. Kaczmarek, A. Jakubowska and J. Lubiński, Serum Concentrations of Selenium and Copper in Patients Diagnosed with Pancreatic Cancer, *Cancer Res. Treat.*, 2016, **48**(3), 1056–1064.
- 43 X. Zhang and Q. Yang, Association Between Serum Copper Levels and Lung Cancer Risk: a Meta-Analysis, *J. Int. Med. Res.*, 2018, **46**(12), 4863–4873.
- 44 P. H. Wiernik, Inching Toward Cure of Acute Myeloid Leukemia: a Summary of the Progress Made in the Last 50 Years, *Med. Oncol.*, 2014, **31**(8), 136.
- 45 A. Maréchal and L. Zou, DNA Damage Sensing by the ATM and ATR Kinases, *Cold Spring Harbor Perspect. Biol.*, 2013, **5**(9), a012716.
- 46 A. M. Weber and A. J. Ryan, ATM and ATR as Therapeutic Targets in Cancer, *Pharmacol. Ther.*, 2015, **149**, 124–138.
- 47 J. Yang, Y. Yu, H. E. Hamrick and J. Duerksen-Hughes, ATM ATR and DNA-PK: Initiators of the Cellular Genotoxic Stress Responses, *Carcinogenesis*, 2003, **24**(10), 1571–1580.
- 48 B. Halliwell and J. M. Gutteridge, Role of Free Radicals and Catalytic Metal Ions in Human Disease: an Overview, *Methods Enzymol.*, 1990, **186**, 1–85.
- 49 C. Duncan and A. R. White, Copper Complexes as Therapeutic Agents, *Metallomics*, 2012, **4**(2), 127–138.
- 50 J. Willis, Y. Patel, B. L. Lentz and S. Yan, APE2 is Required for ATR-Chk1 Checkpoint Activation in Response to Oxidative Stress, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**(26), 10592–10597.
- 51 T. M. Le, S. Poddar, J. R. Capri, E. R. Abt, W. Kim, L. Wei, N. T. Uong, C. M. Cheng, D. Braas, M. Nikanjam, P. Rix, D. Merkurjev, J. Zaretsky, H. I. Kornblum, A. Ribas, H. R. Herschman, J. Whitelegge, K. F. Faull, T. R. Donahue, J. Czernin and C. G. Radu, ATR Inhibition Facilitates Targeting of Leukemia Dependence on Convergent Nucleotide Biosynthetic Pathways, *Nat. Commun.*, 2017, **8**(1), 1–14.
- 52 K. Kino, M. Hirao-Suzuki, M. Morikawa, A. Sakaga and H. Miyazawa, Generation, Repair and Replication of Guanine Oxidation Products, *Genes Environ.*, 2017, **39**, 21.
- 53 S. I. Dikalov and D. G. Harrison, Methods for Detection of Mitochondrial and Cellular Reactive Oxygen Species, *Antioxid. Redox Signaling*, 2014, **20**(2), 372–382.
- 54 M. Wu, A. Neilson, A. L. Swift, R. Moran, J. Tamagnine, D. Parslow, S. Armistead, K. Lemire, J. Orrell, J. Teich, S. Chomicz and D. A. Ferrick, Multiparameter Metabolic Analysis Reveals a Close Link Between Attenuated Mitochondrial Bioenergetic Function and Enhanced Glycolysis Dependency in Human Tumor Cells, *Am. J. Physiol.*, 2007, **292**(1), C125–C136.
- 55 L. Vergnes, G. R. Davies, J. Y. Lin, M. W. Yeh, M. J. Livhits, A. Harari, M. E. Symonds, H. S. Sacks and K. Reue, Adipocyte Browning and Higher Mitochondrial Function in Periadrenal But Not SC Fat in Pheochromocytoma, *J. Clin. Endocrinol. Metab.*, 2016, **101**(11), 4440–4448.
- 56 M. D. Brand and D. G. Nicholls, Assessing Mitochondrial Dysfunction in Cells, *Biochem. J.*, 2011, **435**(2), 297–312.
- 57 S. Krishan, D. R. Richardson and S. Sahni, The Anticancer Agent, Di-2-Pyridylketone 4,4-Dimethyl-3-Thiosemicarbazone (Dp44mT), Up-Regulates the AMPK-Dependent Energy Homeostasis Pathway in Cancer Cells, *Biochim. Biophys. Acta, Mol. Cell Res.*, 2016, **1863**(12), 2916–2933.
- 58 S. Dar, J. Chhina, I. Mert, D. Chitale, T. Buekers, H. Kaur, S. Giri, A. Munkarah and R. Rattan, Bioenergetic Adaptations in Chemoresistant Ovarian Cancer Cells, *Sci. Rep.*, 2017, **7**(1), 8760.
- 59 N. Boulous, H. L. Mulder, C. R. Calabrese, J. B. Morrison, J. E. Rehg, M. V. Relling, C. J. Sherr and R. T. Williams, Chemotherapeutic Agents Circumvent Emergence of Dasatinib-Resistant BCR-ABL Kinase Mutations in a Precise Mouse Model of Philadelphia Chromosome-Positive Acute Lymphoblastic Leukemia, *Blood*, 2011, **117**(13), 3585–3595.
- 60 T. P. Hughes, G. Saglio, A. Quintás-Cardama, M. J. Mauro, D. W. Kim, J. H. Lipton, M. B. Bradley-Garelak, J. Ukropec and A. Hochhaus, BCR-ABL1 Mutation Development During First-Line Treatment with Dasatinib or Imatinib for Chronic Myeloid Leukemia in Chronic Phase, *Leukemia*, 2015, **29**(9), 1832–1838.
- 61 R. T. Williams, M. F. Roussel and C. J. Sherr, Arf Gene Loss Enhances Oncogenicity and Limits Imatinib Response in Mouse Models of Bcr-Abl-Induced Acute Lymphoblastic Leukemia, *Proc. Natl. Acad. Sci. U. S. A.*, 2006, **103**(17), 6688–6693.
- 62 M. Rahmani, J. Nkwocha, E. Hawkins, X. Pei, R. E. Parker, M. Kmiecik, J. D. Levenson, D. Sampath, A. Ferreira-Gonzalez and S. Grant, Cotargeting BCL-2 and PI3K Induces BAX-Dependent Mitochondrial Apoptosis in AML Cells, *Cancer Res.*, 2018, **78**(11), 3075–3086.
- 63 T. Yamaura, T. Nakatani, K. Uda, H. Ogura, W. Shin, N. Kurokawa, K. Saito, N. Fujikawa, T. Date, M. Takasaki, D. Terada, A. Hirai, A. Akashi, F. Chen, Y. Adachi, Y. Ishikawa, F. Hayakawa, S. Hagiwara, T. Naoe and H. Kiyoi, A Novel Irreversible FLT3 Inhibitor, FF-10101, Shows Excellent Efficacy Against AML Cells with FLT3 Mutations, *Blood*, 2018, **131**(4), 426–438.
- 64 P. J. Jansson, P. C. Sharpe, P. V. Bernhardt and D. R. Richardson, Novel Thiosemicarbazones of the ApT and DpT Series and Their Copper Complexes: Identification of Pronounced Redox Activity and Characterization of Their Antitumor Activity, *J. Med. Chem.*, 2010, **53**(15), 5759–5769.

- 65 B. Englinger, C. Pirker, P. Heffeter, A. Terenzi, C. R. Kowol, B. K. Keppler and W. Berger, Metal Drugs and the Anticancer Immune Response, *Chem. Rev.*, 2018, **119**(2), 1519–1624.
- 66 M. Goto, H. Miwa, K. Suganuma, N. Tsunekawa-Imai, M. Shikami, M. Mizutani, S. Mizuno, I. Hanamura and M. Nitta, Adaptation of Leukemia Cells to Hypoxic Condition Through Switching the Energy Metabolism or Avoiding the Oxidative Stress, *BMC Cancer*, 2014, **14**(1), 76–79.
- 67 M. Goto, H. Miwa, M. Shikami, N. Tsunekawa-Imai, K. Suganuma, S. Mizuno, M. Takahashi, M. Mizutani, I. Hanamura and M. Nitta, Importance of Glutamine Metabolism in Leukemia Cells by Energy Production Through TCA Cycle and by Redox Homeostasis, *Cancer Invest.*, 2014, **32**(6), 241–247.
- 68 T. Simsek, F. Kocabas, J. Zheng, R. J. DeBerardinis, A. I. Mahmoud, E. N. Olson, J. W. Schneider, C. C. Zhang and H. A. Sadek, The Distinct Metabolic Profile of Hematopoietic Stem Cells Reflects Their Location in a Hypoxic Niche, *Cell Stem Cell*, 2010, **7**(3), 380–390.
- 69 M. Škrčić, S. Sriskanthadevan, B. Jhas, M. Gebbia, X. Wang, Z. Wang, R. Hurren, Y. Jitkova, M. Gronda, N. Maclean, C. K. Lai, Y. Eberhard, J. Bartoszko, P. Spagnuolo, A. C. Rutledge, A. Datti, T. Ketela, J. Moffat, B. H. Robinson, J. H. Cameron, J. Wrana, C. J. Eaves, M. D. Minden, J. C. Y. Wang, J. E. Dick, K. Humphries, C. Nislow, G. Giaever and A. D. Schimmer, Inhibition of Mitochondrial Translation as a Therapeutic Strategy for Human Acute Myeloid Leukemia, *Cancer Cell*, 2011, **20**(5), 674–688.
- 70 S. Sriskanthadevan, D. V. Jeyaraju, T. E. Chung, S. Prabha, W. Xu, M. Škrčić, B. Jhas, R. Hurren, M. Gronda, X. Wang, Y. Jitkova, M. A. Sukhai, F.-H. Lin, N. Maclean, R. Laister, C. A. Goard, P. J. Mullen, S. Xie, L. Z. Penn, I. M. Rogers, J. E. Dick, M. D. Minden and A. D. Schimmer, AML Cells Have Low Spare Reserve Capacity in Their Respiratory Chain That Renders Them Susceptible to Oxidative Metabolic Stress, *Blood*, 2015, **125**(13), 2120–2130.
- 71 E. D. Lagadinou, A. Sach, K. Callahan, R. M. Rossi, S. J. Neering, M. Minhajuddin, J. M. Ashton, S. Pei, V. Grose, K. M. O'Dwyer, J. L. Liesveld, P. S. Brookes, M. W. Becker and C. T. Jordan, BCL-2 Inhibition Targets Oxidative Phosphorylation and Selectively Eradicates Quiescent Human Leukemia Stem Cells, *Cell Stem Cell*, 2013, **12**(3), 329–341.
- 72 I. Samudio, R. Harmancey, M. Fiegl, H. Kantarjian, M. Konopleva, B. Korchin, K. Kaluarachchi, W. Bornmann, S. Duvvuri, H. Taegtmeier and M. Andreeff, Pharmacologic Inhibition of Fatty Acid Oxidation Sensitizes Human Leukemia Cells to Apoptosis Induction, *J. Clin. Invest.*, 2010, **120**(1), 142–156.
- 73 J. R. Molina, Y. Sun, M. Protopopova, S. Gera, M. Bandi, C. Bristow, T. McAfoos, P. Morlacchi, J. Ackroyd, A.-N. A. Agip, G. Al-Atrash, J. Asara, J. Bardenhagen, C. C. Carrillo, C. Carroll, E. Chang, S. Ciurea, J. B. Cross, B. Czako, A. Deem, N. Daver, J. F. de Groot, J.-W. Dong, N. Feng, G. Gao, J. Gay, M. G. Do, J. Greer, V. Giuliani, J. Han, L. Han, V. K. Henry, J. Hirst, S. Huang, Y. Jiang, Z. Kang, T. Khor, S. Konoplev, Y.-H. Lin, G. Liu, A. Lodi, T. Lofton, H. Ma, M. Mahendra, P. Matre, R. Mullinax, M. Peoples, A. Petrocchi, J. Rodriguez-Canale, R. Serreli, T. Shi, M. Smith, Y. Tabe, J. Theroff, S. Tiziani, Q. Xu, Q. Zhang, F. Muller, R. A. DePinho, C. Toniatti, G. F. Draetta, T. P. Heffernan, M. Konopleva, P. Jones, M. E. Di Francesco and J. R. Marszalek, An Inhibitor of Oxidative Phosphorylation Exploits Cancer Vulnerability, *Nat. Med.*, 2018, **24**(7), 1036–1046.
- 74 F. V. Rassool, T. J. Gaymes, N. Omidvar, N. Brady, S. Beurlet, M. Pla, M. Reboul, N. Lea, C. Chomienne, N. S. B. Thomas, G. J. Mufti and R. A. Padua, Reactive Oxygen Species, DNA Damage, and Error-Prone Repair: a Model for Genomic Instability with Progression in Myeloid Leukemia?, *Cancer Res.*, 2007, **67**(18), 8762–8771.
- 75 K. D. Mills, D. O. Ferguson and F. W. Alt, The Role of DNA Breaks in Genomic Instability and Tumorigenesis, *Immunol. Rev.*, 2003, **194**(1), 77–95.
- 76 J. Fan, L. Li, D. Small and F. Rassool, Cells Expressing FLT3/ITD Mutations Exhibit Elevated Repair Errors Generated Through Alternative NHEJ Pathways: Implications for Genomic Instability and Therapy, *Blood*, 2010, **116**(24), 5298–5305.
- 77 P. S. Hole, R. L. Darley and A. Tonks, Do Reactive Oxygen Species Play a Role in Myeloid Leukemias?, *Blood*, 2011, **117**(22), 5816–5826.
- 78 Z. Skrott, M. Mistrik, K. K. Andersen, S. Friis, D. Majera, J. Gursky, T. Ozdian, J. Bartkova, Z. Turi, P. Moudry, M. Kraus, M. Michalova, J. Vaclavkova, P. Dzubak, I. Vrobel, P. Pouckova, J. Sedlacek, A. Miklovicova, A. Kutt, J. Li, J. Mattova, C. Driessen, Q. P. Dou, J. Olsen, M. Hajduch, B. Cvek, R. J. Deshaies and J. Bartek, Alcohol-Abuse Drug Disulfiram Targets Cancer via P97 Segregase Adaptor NPL4, *Nature*, 2017, **552**(7684), 194–199.
- 79 A. Lass, E. McConnell, K. Fleck, A. Palamarchuk and C. Wójcik, Analysis of Npl4 Deletion Mutants in Mammalian Cells Unravels New Ufd1-Interacting Motifs and Suggests a Regulatory Role of Npl4 in ERAD, *Exp. Cell Res.*, 2008, **314**(14), 2715–2723.
- 80 I. Voráčková, Š. Suchanová, P. Ulbrich, W. E. Diehl and T. Ruml, Purification of Proteins Containing Zinc Finger Domains Using Immobilized Metal Ion Affinity Chromatography, *Protein Expression Purif.*, 2011, **79**(1), 88–95.
- 81 M. T. Larsen, M. Kuhlmann, M. L. Hvam and K. A. Howard, Albumin-Based Drug Delivery: Harnessing Nature to Cure Disease, *Mol. Cell. Ther.*, 2016, **4**(1), 3.
- 82 D. A. Nathanson, A. L. Armijo, M. Tom, Z. Li, E. Dimitrova, W. R. Austin, J. Nomme, D. O. Campbell, L. Ta, T. M. Le, J. T. Lee, R. Darvish, A. Gordin, L. Wei, H.-I. Liao, M. Wilks, C. Martin, S. Sadeghi, J. M. Murphy, N. Boulos, M. E. Phelps, K. F. Faull, H. R. Herschman, M. E. Jung, J. Czernin, A. Lavie and C. G. Radu, Co-Targeting of Convergent Nucleotide Biosynthetic Pathways for Leukemia Eradication, *J. Exp. Med.*, 2014, **211**(3), 473–486.
- 83 N. Jacque, A. M. Ronchetti, C. Larrue, G. Meunier, R. Birsén, L. Willems, E. Saland, J. Decroocq, T. T. Maciel, M. Lambert,

L. Poulain, M. A. Hospital, P. Sujobert, L. Joseph, N. Chapuis, C. Lacombe, I. C. Moura, S. Demo, J. E. Sarry, C. Recher, P. Mayeux, J. Tamburini and D. Bouscary, Targeting

Glutaminolysis Has Antileukemic Activity in Acute Myeloid Leukemia and Synergizes with BCL-2 Inhibition, *Blood*, 2015, **126**(11), 1346–1356.