ARTICLE Open Access

# Measurement and models accounting for cell death capture hidden variation in compound response

Song Yi Bae<sup>1</sup>, Ning Guan 6<sup>2</sup>, Rui Yan 6<sup>3</sup>, Katrina Warner 6<sup>4</sup>, Scott D. Taylor<sup>5</sup> and Aaron S. Meyer 6<sup>5,6,7,8</sup>

# Abstract

Cancer cell sensitivity or resistance is almost universally quantified through a direct or surrogate measure of cell number. However, compound responses can occur through many distinct phenotypic outcomes, including changes in cell growth, apoptosis, and non-apoptotic cell death. These outcomes have divergent effects on the tumor microenvironment, immune response, and resistance mechanisms. Here, we show that quantifying cell viability alone is insufficient to distinguish between these compound responses. Using an alternative assay and drug-response analysis amenable to high-throughput measurement, we find that compounds with identical viability outcomes can have very different effects on cell growth and death. Moreover, additive compound pairs with distinct growth/death effects can appear synergistic when only assessed by viability. Overall, these results demonstrate an approach to incorporating measurements of cell death when characterizing a pharmacologic response.

# Introduction

Quantifying cellular response to therapeutic compounds is essential to understanding their mechanisms of action and assessing therapeutic efficacy<sup>1–3</sup>. In the case of cancer treatments, and often with other diseases, drug activities are evaluated by quantifying the number of live cells after a short period using direct or surrogate measurements<sup>4,5</sup>. However, quantities beyond the number and viability of cells provide valuable information about the cellular response. Along with altering cell proliferation, promoting cell death is another important index of drug efficacy<sup>6,7</sup>. Incomplete eradication of drug-susceptible malignant cells allows the survival of drug-tolerant persister cell populations that can develop resistance by multiple routes<sup>8–10</sup>. Moreover, cell death can occur via a variety of mechanisms, including apoptosis and necroptosis, and selection among

these outcomes can potently modulate cancer immunogenicity<sup>11</sup>. Limited understanding of these underlying cellular responses further complicates the assessment of drug combinations. Combination treatments are typically evaluated for their ability to enact greater effects than either compound alone<sup>12</sup>, but typically only by quantifying viability.

Here, we show that directly measuring both cell growth and death can provide valuable information for interpreting the response of cells to single and combination treatments. We propose a framework for quantifying drug response that accounts for the compound-induced changes in rates of cell growth and death. This approach reveals extensive differences in cell response, otherwise hidden by simply quantifying cell number. Of course, trade-offs exist for the breadth versus depth of analysis that can be performed to characterize cell-compound response. We show that end-point analysis preserves much of the distinct outcomes we observe for kinetic measurements while allowing similarly high-throughput analysis to those of live-cell number surrogates. These results demonstrate the need for and an

© The Author(s) 2020

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/.



Correspondence: Aaron S. Meyer (a@asmlab.org)

<sup>&</sup>lt;sup>1</sup>Department of Pharmacology, University of Minnesota-Twin Cities, Minneapolis, MN, USA

<sup>&</sup>lt;sup>2</sup>Department of Biological Engineering, Massachusetts Institute of Technology, Cambridge, MA, USA

Full list of author information is available at the end of the article Edited by G. Melino

approach to more precisely quantify the nature of cell-compound response and interactions.

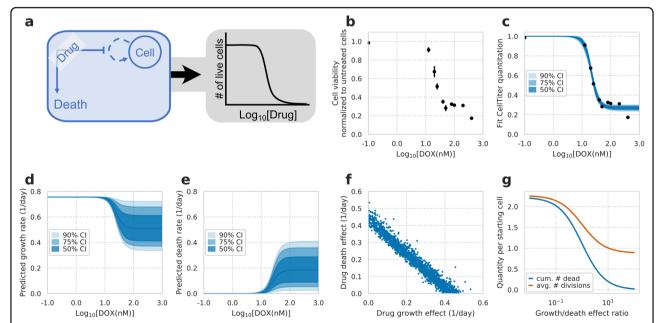
### Results

# Viability alone is insufficient to distinguish cell growth and death effects

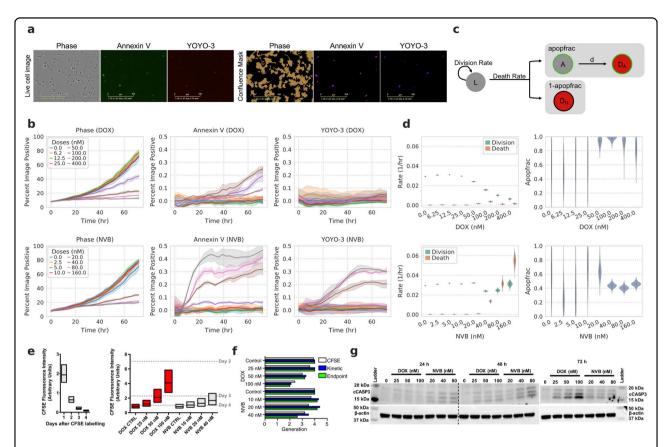
To test whether growth and death are confounded in livecell measurements, we first explored the uncertainty in a model using only these measurements (Fig. 1a). We fit typical dose-response measurements of H1299 non-smallcell lung cancer cells to the chemotherapy doxorubicin (Fig. 1b) to a model incorporating both cell growth and death. We assumed no cell death in the absence of drug to show the best-case scenario of assessing drug response. This model was identifiable for the live-cell number (Fig. 1c), and the  $IC_{50}$  and  $E_{max}$  of compound effect on cell viability were narrowly defined as  $21.4 \pm 1.0 \, \text{nM}$  and  $26.7 \pm 3.4\%$  (90% confidence interval, CI), respectively. In contrast, the model showed large uncertainty in inferred cellular growth or death rates (Fig. 1d, e). At the maximum dose, the predicted growth rate ranged 0.33-0.71 day<sup>-1</sup> and death rate 0.02-0.40 day<sup>-1</sup> (90% CI). The large uncertainty in outcome was due to a strong correlation in the fit values of drug effect on the growth and death rates (Fig. 1f). This shows that given only live-cell number one is incapable of distinguishing between reductions in cellular growth rate and increases in cell death. The number of divisions and cell deaths can vary largely while similarly fitting live-cell measurements. Moreover, the number of cell divisions and cumulative dead cells can differ drastically while resulting in the same cell viability (Fig. 1g).

# High-throughput measurements of cell death quantify compound response

To quantify pharmacologic response, we extended our experimental measurements to those of cell death. While quantifying the number of cells over time by phase, we used an Annexin V probe to measure phosphatidylserine exposure during apoptosis and a membrane-impermeable DNA dye, YOYO-3, to measure permeabilized apoptotic and dead cells (Fig. 2a)<sup>13,14</sup>. The areas occupied by cells, Annexin V, and YOYO-3 signal in each image were then analyzed to determine the total, apoptotic, and dead cells relative to the whole image area. We evaluated this realtime imaging method by measuring the response to doxorubicin (DOX) in H1299 cells. DOX strongly reduced the number of cells (Fig. 2b, top), as seen before (Fig. 1b). At the same time, we observed an increase in Annexin V while YOYO-3 increased minimally throughout the assay. Fitting these data to a model of cell growth and death (Fig. 2c), we



**Fig. 1 Confounding effects of cell growth and death on drug-response measurements. a** Schematic of drug response assessed by calculating relative changes in live-cell number after drug treatment (gray box), and cell growth and death rate that underlie the changes (blue box). Both growth and death rates were assumed to have a Hill curve relationship to drug concentration. **b** Cell viability of H1299 cells treated with doxorubicin (DOX) at varying concentrations for 72 h (N = 3). **c** Model fit to live-cell measurements. **d** Model fit and confidence intervals for the predicted growth rate of cells after fitting to measurements of live cell number. **e** Model fit and confidence intervals for the predicted death rate. **f** Model fit posterior samples of DOX's effect on growth versus its effect on cell death. **g** Model predictions of the cumulative number of dead cells and cell divisions throughout the experiment for a constant drug effect of reducing cell viability by 75%. *X*-axis indicates the relative ratio in magnitude of growth versus death drug effect. *Y*-axis indicates the varying quantity of predicted number of cell divisions or cumulative dead cells per starting cell.



**Fig. 2 High-throughput measurements of cell death accurately quantify compound response. a** Representative images from live-cell imaging and processing. Scale bar indicates 300 μm. **b** Experimental measurements of doxorubicin (DOX, top) and vinorelbine (NVB, bottom) response in H1299 cells over time. Phase indicates total cell confluence. Each line represents the mean of triplicate measurements for individual drug dose over time, and shaded areas show the ranges of measurements. **c** Schematic of the cell growth-death model. The live (L) cells grow at the rate of division (div), and die at the rate of death (deathRate). The model considers two fractions of cells in cell death: cells dying through apoptosis (apopfrac) and other modes (1–apopfrac). Cells in early apoptosis (A) proceed into late apoptosis (D<sub>A</sub>) by losing membrane integrity at rate d. **d** The model predicted div, deathRate, and apopfrac from the data represented in (**b**). Violins show posterior of model after fitting. **e** Cell division analysis of CFSE-labeled H1299 cells by flow cytometry. The distribution of cells according to CFSE fluorescence intensity on indicated days for untreated cells (left) or after indicated treatments (right) for 72 h are shown in boxplots with median, lower, and upper quartiles. Median CFSE intensities of 2–4 days post-CFSE labeling in left graph are marked as dashed lines on the right graph. **f** The number of generations for indicated conditions calculated by median fluorescence intensities from CFSE-based assay and model-predicted cell growth rates from kinetic and end-point measurements. **g** Western blot of cleaved-caspase-3 (cCASP3) after treating H1299 cells with indicated drug doses for 24, 48, and 72 h.

observed a strong decrease in the inferred growth rate (div), and only a modest increase in cell death rate (deathRate; Fig. 2d, top). We next compared these measurements to those with another chemotherapy, vinorelbine (NVB), again observing a dose-dependent decrease in the number of live cells (Fig. 2b, bottom). At the same time, we observed a large increase in both Annexin V and YOYO-3 signal. This was reflected in our subsequent analysis, inferring an increase in the death rate (Fig. 2d, bottom). The fraction of cells dying through apoptosis (apopfrac) was also inferred to be lower in NVB as compared with DOX treatment (Fig. 2d). While kinetic measurements provide a wealth of information, end-point measurement is more amenable to high-throughput experiments. We confirmed that our

analysis provided qualitatively similar results using only the first and last measurements in each experiment, demonstrating that cell growth and death can be quantified using either kinetic or endpoint measurements (Supplementary Fig. S1).

To independently verify these opposing outcomes upon DOX or NVB treatment, we used a carboxyfluorescein succinimidyl ester (CFSE)-based proliferation assay to verify the distinct growth rate effects inferred by our analysis. We measured CFSE intensity of untreated cells every 24 h starting from a day after cell labeling (Fig. 2e, left). The detected intensity over time was used to estimate the number of times each cell had divided. Consistent with the inferred growth rates of our model,

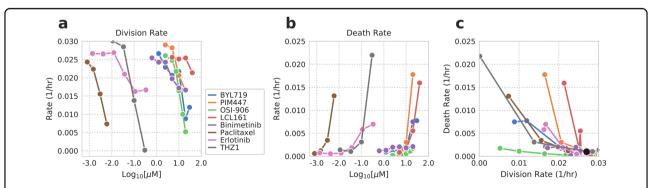
DOX-treated cells distributed more toward higher CFSE intensity in a dose-dependent manner, implying fewer cell divisions than with non-treated cells (Fig. 2e, right). In contrast, the distribution of NVB-treated cells remained more similar to non-treated cells. Our inferred cell growth rates and CFSE measurements were overall consistent (Fig. 2f).

To validate the inferred cell death rates, we measured the induction of cleaved caspase-3, an apoptotic marker, after treatment with DOX or NVB (Fig. 2g). After 24-h treatment, cleaved caspase-3 was detected in NVB-treated cells, but not DOX-treated cells. Both drugs induced the caspase-3 cleavage by 48-h treatment. These observations were consistent with the inferred cell death rates, indicating that NVB induces faster cell death than DOX. Interestingly, the level of cleaved caspase-3 after 72-h treatment was lower in NVB-treated cells compared with DOX-treated cells. This may be explained by the lower inferred apoptosis fraction in NVB treatment while the cell death rates increased (Fig. 2d) implying an increase in non-apoptotic cell death, such as caspase-independent cell death. Alternatively, a large fraction of NVB-treated cells are dead after 72 h. Phosphatidylserine (to which Annexin V binds) is irreversibly externalized to the cell surface by caspase-dependent scramblases during apoptosis, in contrast, it is not exposed by caspase-independent cell death<sup>15,16</sup>, and NVB may induce caspase-independent cell death as well as apoptosis. DOX and NVB are known operate through distinct mechanisms-inducing double-stranded breaks or preventing microtubule polymerization, respectively. As a result, each compound has a differing dependency on p53 status and leads to arrest in distinct cell cycle phases, supporting that each might engage distinct cell death programs 17-19. Collectively, measuring and analyzing phase, Annexin V and YOYO-3 signals can quantify both the growth and death rate effects of drugs on cells.

# Targeted compounds also display distinct phenotypic consequences

We next evaluated whether the growth-death model can dissect the response of cancer cells to targeted compounds as well. We treated a non-small cell lung cancer cell line PC9 with seven different targeted drugs whose effects on cell growth and death we expected to vary according to their mechanisms of action. We also used paclitaxel, a chemotherapeutic drug that is widely known to interfere with the cell cycle resulting in cell death and reduced proliferation, for comparison. The total, apoptotic and dead cell measurements from each compound treatment were diverse (Supplementary Fig. S2), and these differences were reflected in the inferred cell division and death rates (Fig. 3; Supplementary Fig. S3).

We were able to classify the tested compounds into three types: a compound that (1) both inhibits cell division and induces cell death, (2) only inhibits cell division, and (3) only induces cell death. As expected, paclitaxel fell into the first type by simultaneously exhibiting strong cell growth suppression and death. Similar to paclitaxel, the PI3Ka inhibitor BYL719, the pan-PIM kinase inhibitor PIM447, the EGFR tyrosine kinase inhibitor erlotinib, and CDK7/12 inhibitor THZ1 were grouped into the first type. In contrast, both OSI-906, a dual IGF-1R/INSR tyrosine kinase inhibitor, and binimetinib, a MEK1/2 inhibitor, mainly decreased cell division rates. OSI-906 inhibited the division rate as effective as paclitaxel within the range of tested doses. LCL161 is a small molecule SMAC mimetic that antagonizes multiple inhibitor of apoptosis (IAP) family proteins and augments apoptosis induction. Consistent with its mechanism of action, LCL161 showed a minimal effect on cell division while strongly enhancing cell death. Taken together, the growth-death model allows us to interpret the response of cancer cells to different targeted drugs in terms of cell division and death rates,



**Fig. 3 Comparing drug response for targeted compounds with distinct mechanisms of action.** Model predicted cell division (**a**) and death rates (**b**) for seven targeted and one chemotherapy compound. Measurements (phase, Annexin V, YOYO-3) are shown in Supplementary Fig. S2. The mean value of the model posterior for each dose is plotted. **c** Cell division and death rates are plotted together. See Supplementary Fig. S3 for the probability density of each compound's inferred div, deathRate and apopfrac. The black dot indicates the control rates.

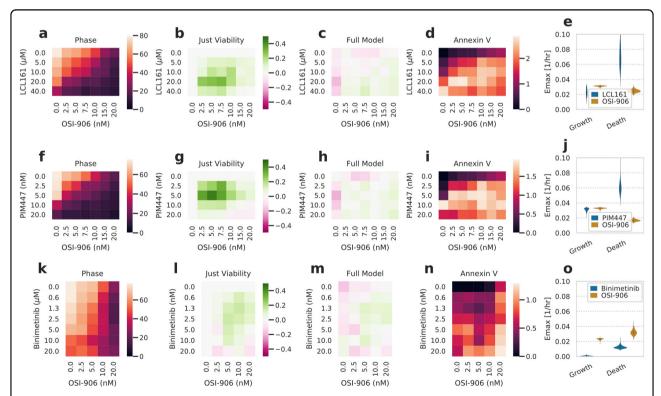
which otherwise would not be revealed from overall phenotypic changes without a panel of experiments.

# Compounds with disparate phenotypic outcomes can appear synergistic when only analyzed by viability

Based on the changes in rate parameters by targeted drugs in Fig. 3, we wondered how drugs with divergent phenotypic effects might influence cell behavior when combined. We selected one compound from each of the three groups identified from Fig. 3; PIM447 affected both cell division and death rates, OSI-906 affected only cell division, and LCL161 affected only cell death. Combination treatment between OSI-906 and LCL161 or PIM447 was quantified. Each compound's effect closely matched those we expected from the single-agent treatments (Figs. 3 and 4a, f; Supplementary Fig. S4a, b).

Intriguingly, the nature of each drug interaction was dependent upon whether we took cell death into account. A Bliss additivity model using just cell confluence indicated that both combinations led to a synergistic interaction (Fig. 4b, g). However, we also used our combination treatment data to fit a model in which both

growth and death rates respond based on a Hill curve dose-response relationship (Supplementary Fig. S4a, b). A model assuming Bliss additivity for the growth rate and an additive interaction for the death rate fit our measurements much more closely, despite also having to account for both phase and cell death measurements (Fig. 4c, h). Investigating the source of this discrepancy, we noted that the perceived synergy arose with an increase in cell death (Fig. 4b/g vs. d/i). We were surprised by the difference in outcomes between each model as the IC50 and Hill coefficient of the latter model (Fig. 4c, h) are assumed to be equivalent for both phenotypes. However, we expect that the difference arises from the very distinct  $E_{\text{max}}$ values for growth and death phenotypes (Fig. 4e, j). These terms do not simplify into one maximal effect term when entered into an exponential growth model accounting for both growth and death (methods), indicating a complex interaction between each phenotype's effects. Indeed, our model closely matched the results of just analyzing viability for co-treatment with binimetinib and OSI-906, which both preferentially inhibit growth (Fig. 4k-o). Therefore, we conclude that not only are measures of cell



**Fig. 4 Additive interactions can appear synergistic when only assessed by viability. a, f, k** Heatmap of percent confluence for combinations of LCL161 (**a**), PIM447 (**f**), or binimetinib (**k**) with OSI-906 at 72 h. Color scale indicates percent confluence. **b, g, l** Heatmap of deviation from Bliss additivity for measurements in **a, f, k**. Color scale indicates deviation from additive prediction on a scale of 0–1. **c, h, m** Heatmap of deviation from the additive model incorporating both growth and death processes for phase measurements. Color scale indicates deviation from additive prediction on a scale of 0–1, where positive and negative values indicate drug synergy and antagonism, respectively. **d, i, n** Heatmap of percent image area positive for Annexin V signal at 72 h. **e, j, o** Fit growth and death rate E<sub>max</sub> for each drug. Fits come from the additive interaction model incorporating both processes. The experimental measurements are shown in Supplementary Fig. S4.

death an important component of pharmacologic response but that ignoring cell death can lead to spurious conclusions of drug synergy.

# Discussion

More than half of preclinical agents entering clinical trials fail to be approved due to lack of efficacy 20,21, and the success rate for clinical approval of oncology drugs remains low as 13%<sup>22</sup>. These statistics reveal the limitation in current preclinical experimental models and analysis in predicting clinical outcomes. Recently, new strategies of evaluating drug response, such as patientderived xenografts and organoids, have been introduced to more closely match patient's tumors to effective compounds<sup>23–25</sup>. Two major phenotypic changes we can observe as cellular responses to drugs are cell growth and death. Each response differs according to the administered drug and contributes to the overall effect. In this paper, we demonstrate that cell viability measurements alone cannot distinguish these phenotypic outcomes (Fig. 1). These responses can strongly influence clinical outcome<sup>11</sup>. Thus, we propose a high-throughput assay that can be paired with analysis to quantify both the growth and death effects of drug response for better estimation of drug efficacy. Applying this analysis, we identify that compounds can have distinct outcomes by specifically driving growth, death, or mixed effects (Fig. 3). Cell death can also take distinct forms, which can easily be quantified as relatively more or less apoptotic (Fig. 2). Furthermore, additive drug interactions in terms of cell growth or death can appear synergistic when only assessed through cell viability (Fig. 4). Overall, these results show that cytotoxic drug response should be assessed by the distinct phenotypes of cell growth and death, and then demonstrate an approach to do so.

Separating these phenotypic outcomes provides future opportunities for cancer treatment. A more detailed view of phenotypic drug response should enable treatment optimization both in a population- and patient-specific manner<sup>26</sup>. For maximally effective cancer treatment, single or combination treatments should likely modulate both phenotypes. Purely cytostatic therapy leaves drugpersister cells able to undergo genetic or epigenetic changes giving rise to resistance<sup>9</sup>. On the other hand, cell death may not overcome the replenishment of tumor cells at toxicity-limiting doses. Among the targeted therapies we evaluated, an EGFR tyrosine kinase inhibitor erlotinib exhibited strong inhibition on cell division while modest induction on cell death. Consistent with our result, recent studies on mechanism of erlotinib resistance reported that drug-tolerant colonies maintain throughout drug exposure in quiescent state and expand over long period of time acquiring resistance<sup>8,10</sup>. A combination of erlotinib with a cell death-inducing agent may therefore prevent the survival of drug-tolerant subpopulation cells.

Our analysis results in drug combination also provided a plausible cause for frequent failure of preclinical regimens in clinical trials. The insulin-like growth factor-1 receptor (IGF-1R) signaling pathway is a wellcharacterized pathway involved in cancer cell survival and promoting drug resistance<sup>27</sup>. IGF-1R inhibitors have been validated for their efficacy in various cancer types as mono- and combined therapies in preclinical studies<sup>28–30</sup>. Despite promising preclinical evidence, only limited responses were observed in clinical trials<sup>31–34</sup>. From our drug combination experiments with the IGF-1R inhibitor OSI-906, we discovered that the analysis by cell viability alone found a strong synergistic interaction, while our analysis accounting cell growth and death identified an additive response. Thus, synergy observed in preclinical assays may have been due to the nature of the assay moreso than molecular synergy.

This more detailed analysis of phenotypic response can also help to optimize the microenvironment and host immune response to cancer. For example, host response is potently influenced by the route of cell death<sup>35,36</sup>. These methods can be applied not only in vitro but also in more complex, translationally relevant models, such as organoids and in vivo<sup>37</sup>. By doing so, treatments could be tailored to not just maximally reduce cell viability but optimize cytotoxicity and cell death programs to mount a host response<sup>38</sup>.

The approach here will benefit from improvements in the single-cell resolution of drug response and more exactly distinguishing cell death programs. Cell-to-cell variability impacts both drug response and the development of resistance<sup>39</sup>, and single-cell technologies have enabled in-depth molecular analyses of this heterogeneity<sup>40,41</sup>. Within a tumor population, drug treatment can dynamically shift the balance of variability present, and so dynamic information will likely be critical<sup>8,42</sup>. Automated cell tracking would enable drug-response quantification, with single-cell resolution, while preserving the lineage relationship of cells 43,44. Finally, while we distinguish relatively more or less apoptotic cell death here, various cell death programs exist<sup>45</sup>. Improved methods to visualize distinct forms of cell death in populations of cells will allow distinct forms of cell death to be separately quantified<sup>46</sup>. The diverse outcomes observed in our study suggest that deeper phenotypic characterization will uncover more differences between compounds in their elicited drug responses.

# **Materials and methods**

All analysis was implemented in Python, and can be found at https://github.com/meyer-lab/ps-growth-model, release 1.0.1 (https://doi.org/10.5281/zenodo.3742188).

# Compounds and cell culture

Doxorubicin, OSI-906, BYL719, binimetinib, and paclitaxel were purchased from LC Laboratories (Woburn, MA). PIM447 and LCL161 were obtained from Selleck Chemicals (Houston, TX). Vinorelbine was purchased from Sigma-Aldrich (St. Louis, MO). THZ1 was purchased from Cayman Chemical Company (Ann Arbor, MI). Human lung carcinoma PC9 cells were obtained from Sigma-Aldrich, and H1299 cells were provided from ATCC (Manassas, VA). All cell lines were grown in RMPI-1640 medium supplemented with 10% fetal bovine serum and 1% penicillin–streptomycin, at 37 °C and 5% CO<sub>2</sub>.

# End-point cell viability assay and time-lapse microscopy

For the end-point cell viability assay in Fig. 1, cells were seeded at  $1.5 \times 10^3$  cells per well in 96-well plates and cultured overnight. Then, cells were treated with doxorubicin. After 72 h, CellTiter Glo reagent (Promega, Madison, WI) was added to each well, and luminescence was detected according to the manufacturer's instructions.

Cells were seeded as indicated above for the time-lapse measurements. The next day, each indicated treatment was added, along with IncuCyte Annexin V Green Reagent (Essen BioScience, Ann Arbor, MI) and 300 nM YOYO-3 in media containing 1.25 mM CaCl<sub>2</sub>. Cells were then cultured and imaged within the IncuCyte Zoom or S3 (Essen BioScience) every 3 h. Four fields of view were taken per well. Fluorescence images were manually thresholded, and the fraction of image area with Annexin V and/or YOYO-3 signal was quantified using IncuCyte Zoom or S3 software (Essen BioScience). Finally, the fraction of area occupied by cells was analyzed by brightfield analysis.

# Hill curve identifiability model related to Fig. 1

A model of exponential growth along with death was fit to viability measurements assuming a Hill dose–response relationship. For comparing the model to the data, the fit residuals were assumed to be normally distributed. In the absence of drug, the growth rate was measured and experimentally set to be 0.0315 1/h, and cells were assumed to not undergo cell death. The minimum growth rate (at infinite concentration of drug) was fit using a uniform prior between 0.0 and the growth rate in the absence of drug. The maximal death rate (at infinite concentration of drug) was fit using a log-normal prior of  $-2.0\pm2.0\,1/h$  (log10 scale). The Hill slope was fit using a log-normal prior of  $0.0\pm1.0$  (log10 scale). Both the IC50 and Hill slope were assumed to be the same for growth and death rates.

# Growth model structure

Cell behavior was modeled using a series of kinetic equations incorporating cell growth and death. We

represent the overall state of a population of cells as  $v = [L, E, D_a, D_n]$ , respectively indicating the number of live cells, cells within early apoptosis, dead cells via apoptosis, and dead cells via a non-apoptotic process. Using such notation, the time derivative was defined as:

$$\dot{\nu} = \left[ (R_g - R_d)L, R_d \cdot f \cdot L - E, \tau \cdot E, R_d (1 - f)L \right]$$

where  $R_g$  (or div) is the rate of cell division,  $R_d$  (or deathRate) is the rate of cell death, f (or apopFrac) is the fraction of dying cells which go through apoptosis, and  $\tau$  (or d) determines the rate of conversion from early to late apoptosis.

If  $\gamma = R_g - R_d$ ,  $c = (R_d \cdot f)/(g + d)$ , and  $m = R_d(1 - f)$ , integrating these equations provides the solution:

$$u(t) = [e^{\gamma t}, c(L-e^{mt}), mc(L-1)/\gamma + c(e^{mt}-1),$$

$$m(L-1)/\gamma]$$

# Growth model inference

Predicted cell numbers were fit to experimental measurements using Markov chain Monte Carlo<sup>47</sup>. The percent area positive for cell confluence, Annexin V stain, or YOYO-3 stain was quantified and assumed to be proportional to the number of cells positive for each marker. Cell confluence was assumed to be the total of cells in all states. Apoptotic cells were assumed to be positive for Annexin V signal, then positive for both signals after late apoptosis. Non-apoptotic cells were assumed to just be positive for YOYO-3 signal after dying. Each rate parameter was fit to the corresponding measurements within a single drug condition over time. An entire experiment, corresponding to a set of different compounds and concentrations, was fit simultaneously, allowing for a background offset and conversion factor of each quantity to be fit across the experiment.

div was set to have a uniform prior of 0.0-0.35 1/h. deathRate, and d were set to have log-normal prior distributions of mean 0.01 1/h with standard deviation 0.5 (log<sub>10</sub> scale). By inspecting a calibration experiment and manually counting the cells within a field, we measured the conversion between number of cells and area of signal for the confluence, Annexin V, and YOYO-3 images. In addition, we quantified the ratio of positive area for each pair of signals when a single cell was positive for both. Each of these were set as log-normal prior distributions on the conversion values between number of cells and positive area. Finally, we observed appreciable background in the Annexin V and YOYO-3 signal, leading to signal in the absence of cells. Therefore, we set log-normal priors for the background levels with mean 0.1% of area and standard deviation of 0.1 (log<sub>10</sub> scale). Each data point was assumed to have independent, normally distributed error around the model prediction.

Sampling convergence was verified by checking that two independent runs generated insignificant differences, checking for ergodicity through the Geweke criterion comparing the first and second half of each run, and verifying an effective sample size of greater than 200. Sampling failures were solved by increasing the number of tuning samples.

## CFSE-based cell proliferation analysis

Cell division was measured using carboxyfluorescein diacetate succinimidyl ester (CFSE) dilution analysis. Cells were labeled with 5  $\mu M$  CFSE (Invitrogen, Carlsbad, CA) according to the manufacturer's protocol. The stained cells were seeded overnight in 60-mm dishes at a density of  $2\times10^5$  cells per dish, and then treated with indicated drugs next day. For 72 h at 24-h intervals, cells were collected and fixed in 4% paraformaldehyde prior to acquisition on a BD LSRFortessa flow cytometer (BD Biosciences, San Jose, CA). CFSE signal intensity of  $1\times10^4$  cells was recorded and analyzed to measure cell divisions. The same cell line was labeled the day of the analysis to determine initial labeling.

## Western blot analysis

Cells were seeded at a density of  $2 \times 10^5$  cells per 60-mm dish 24 h prior to drug treatment then treated with the indicated conditions for 24, 48, and 72 h. After incubation, cells were lysed in 10 mM Tris-HCl pH 8.0, 1 mM EDTA, 1% Triton-X 100, 0.1% Na deoxycholate, 0.1% SDS, and 140 mM NaCl, freshly supplemented with protease and phosphatase inhibitor (Boston Bio Products, Ashland, MA). Protein concentration was measured by a bicinchoninic acid assay. In total, 10 µg of protein from each cell lysate was subjected to SDS-PAGE, and then transferred to a polyvinylidene difluoride membrane. Each membrane was incubated overnight with antibody against cleaved caspase-3 (Cell Signaling Technology, Danvers, MA, #9664) or 1.5 h with HRP-conjugated β-actin antibody (Cell Signaling Technology, #12262). β-actin was used as a loading control for western blot analysis.

# **Drug interaction fitting**

Drug interaction was assumed to follow the Bliss independence model<sup>48</sup>. Where indicated, this was taken to be defined as a proportional decrease in the viability of cells. That is, cell viability was normalized to 1.0 for the control condition, and then the proportional decrease in cell viability was calculated by 1.0 minus cell viability. Synergy or antagonism was identified by a greater or lesser decrease in viability than predicted, respectively.

Alternatively, Bliss additivity was defined in conjunction with a model incorporating cell death. d and apopfrac were assumed to be constant across drug concentration or combination and fit using the same priors as before. The

growth rate in the absence of drug was fit using the lognormal prior of  $-1.5\pm0.1/h$  (log\_{10} scale) based on experimental growth measurement. Cells were assumed to undergo no cell death in the absence of drug. An  $E_{\rm max}$  of growth inhibition was fit using a Beta prior (a = 1.0,  $\beta=1.0$ ), where 1.0 indicates complete growth inhibition and 0.0 no growth inhibition. The  $E_{\rm max}$  of death effect was fit using a log-normal prior of  $-2.0\pm0.5/h$  (log\_{10} scale), where the value indicates the maximal death rate. The half-maximal concentration (EC\_{50} or IC\_{50}) and Hill coefficient of each compound were fit using the same priors as before for these quantities and assumed to be the same for both growth and death effects.

#### Acknowledgements

This work was supported by NIH U01-CA215709 to A.S.M. and in part by the UCLA Jonsson Comprehensive Cancer Center (JCCC) grant NIH P30-CA016042.

#### **Author details**

<sup>1</sup>Department of Pharmacology, University of Minnesota-Twin Cities, Minneapolis, MN, USA. <sup>2</sup>Department of Biological Engineering, Massachusetts Institute of Technology, Cambridge, MA, USA. <sup>3</sup>Institute for Computational and Mathematical Engineering, Stanford University, Stanford, CA, USA. <sup>4</sup>Biological and Biomedical Sciences Program, Harvard University, Cambridge, MA, USA. <sup>5</sup>Department of Bioengineering, University of California, Los Angeles, CA, USA. <sup>7</sup>Jonsson Comprehensive Cancer Center, University of California, Los Angeles, CA, USA. <sup>8</sup>Eli and Edythe Broad Center of Regenerative Medicine and Stem Cell Research, University of California, Los Angeles, CA, USA

## Conflict of interest

The authors declare that they have no conflict of interest.

## Publisher's note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

**Supplementary Information** accompanies this paper at (https://doi.org/10.1038/s41419-020-2462-8).

Received: 5 December 2019 Revised: 6 April 2020 Accepted: 7 April 2020 Published online: 20 April 2020

# References

- Heiser, L. M. et al. Subtype and pathway specific responses to anticancer compounds in breast cancer. Proc. Natl Acad. Sci. USA 109, 2724–2729 (2011).
- Sos, M. L. et al. Predicting drug susceptibility of non-small cell lung cancers based on genetic lesions. J. Clin. Invest. 119, 1727–1740 (2009).
- Garnett, M. J. et al. Systematic identification of genomic markers of drug sensitivity in cancer cells. Nature 483, 570–575 (2012).
- Tyson, D. R., Garbett, S. P., Frick, P. L. & Quaranta, V. Fractional proliferation: a method to deconvolve cell population dynamics from single-cell data. *Nat. Methods* 9, 923–928 (2012).
- Fallahi-Sichani, M., Honarnejad, S., Heiser, L. M., Gray, J. W. & Sorger, P. K. Metrics other than potency reveal systematic variation in responses to cancer drugs. *Nat. Chem. Biol.* 9, 708–714 (2013).
- Faber, A. C., Ebi, H., Costa, C. & Engelman, J. A. Apoptosis in targeted therapy responses: the role of BIM. Adv. Pharmacol. 65, 519–542 (2012).
- Long, J. S. & Ryan, K. M. New frontiers in promoting tumour cell death: targeting apoptosis, necroptosis and autophagy. *Oncogene* 31, 5045–5060 (2012).
- Sharma, S. V. et al. A chromatin-mediated reversible drug-tolerant state in cancer cell subpopulations. Cell 141, 69–80 (2010).

- Hata, A. N. et al. Tumor cells can follow distinct evolutionary paths to become resistant to epidermal growth factor receptor inhibition. *Nat. Med.* 22, 262–269 (2016).
- Ramirez, M. et al. Diverse drug-resistance mechanisms can emerge from drugtolerant cancer persister cells. Nat. Commun. 7, 10690 (2016).
- Gotwals, P. et al. Prospects for combining targeted and conventional cancer therapy with immunotherapy. Nat. Rev. Cancer 17, 286–301 (2017).
- Foucquier, J. & Guedj, M. Analysis of drug combinations: current methodological landscape. *Pharm. Res. Perspect.* 3, e00149 (2015).
- Kim, Y. E., Chen, J., Langen, R. & Chan, J. R. Monitoring apoptosis and neuronal degeneration by real-time detection of phosphatidylserine externalization using a polarity-sensitive indicator of viability and apoptosis. *Nat. Protoc.* 5, 1396–1405 (2010).
- Gelles, J. D. & Chipuk, J. E. Robust high-throughput kinetic analysis of apoptosis with real-time high-content live-cell imaging. Cell Death Dis. 7, e2493 (2016).
- Carter, B. Z. et al. Caspase-independent cell death in AML: caspase inhibition in vitro with pan-caspase inhibitors or in vivo by XIAP or Survivin does not affect cell survival or prognosis. *Blood* 102, 4179–4186 (2003).
- Segawa, K. & Nagata, S. An apoptotic "Eat Me" signal: phosphatidylserine exposure. Trends Cell Biol. 25, 639–650 (2015).
- Lukin, D. J., Carvajal, L. A., Liu, W.-j, Resnick-Silverman, L. & Manfredi, J. J. p53 promotes cell survival due to the reversibility of its cell-cycle checkpoints. *Mol. Cancer Res.* 13, 16–28 (2014).
- Ling, Y. H., el-Naggar, A. K., Priebe, W. & Perez-Soler, R. Cell cycle-dependent cytotoxicity, G2/M phase arrest, and disruption of p34cdc2/cyclin B1 activity induced by doxorubicin in synchronized P388 cells. *Mol. Pharmacol.* 49, 832–841 (1996).
- Klotz, D. M. et al. The microtubule poison vinorelbine kills cells independently of mitotic arrest and targets cells lacking the APC tumour suppressor more effectively. J. Cell. Sci. 125, 887–895 (2012).
- Arrowsmith, J. Phase II failures: 2008–2010. Nat. Rev. Drug Disco. 10, 328–329 (2011).
- 21. Arrowsmith, J. & Miller, P. Trial watch: phase II and phase III attrition rates 2011–2012. *Nat. Rev. Drug Disco.* **12**, 569 (2013).
- DiMasi, J. A., Reichert, J. M., Feldman, L. & Malins, A. Clinical approval success rates for investigational cancer drugs. Clin. Pharm. Ther. 94, 329–335 (2013).
- Gao, H. et al. High-throughput screening using patient-derived tumor xenografts to predict clinical trial drug response. Nat. Med. 21, 1318–1325 (2015).
- Hafner, M., Niepel, M., Chung, M. & Sorger, P. K. Growth rate inhibition metrics correct for confounders in measuring sensitivity to cancer drugs. *Nat. Methods* 13, 521–527 (2016).
- Meyer, C. T. et al. Quantifying drug combination synergy along potency and efficacy axes. Cell Syst. 8, 97–108.e16 (2019).
- Palmer, A. C. & Sorger, P. K. Combination cancer therapy can confer benefit via patient-to-patient variability without drug additivity or synergy. *Cell* 171, 1678–1691.e13 (2017).
- Scagliotti, G. V. & Novello, S. The role of the insulin-like growth factor signaling pathway in non-small cell lung cancer and other solid tumors. *Cancer Treat. Rev.* 38, 292–302 (2011).
- Ramcharan, R. et al. IGF-1R inhibition induces schedule-dependent sensitization of human melanoma to temozolomide. Oncotarget 6, 39877–39890 (2015).
- 29. de Lint, K. et al. Sensitizing triple-negative breast cancer to PI3K inhibition by cotargeting IGF1R. *Mol. Cancer Ther.* **15**, 1545–1556 (2016).

- Pivonello, C. et al. The dual targeting of insulin and insulin-like growth factor 1 receptor enhances the mTOR inhibitor-mediated antitumor efficacy in hepatocellular carcinoma. *Oncotarget* 7, 9718–9731 (2016).
- Ciuleanu, T.-E. et al. Randomised phase 2 study of maintenance linsitinib (OSI-906) in combination with erlotinib compared with placebo plus erlotinib after platinum-based chemotherapy in patients with advanced non-small cell lung cancer. Br. J. Cancer 117, 757–766 (2017).
- Oza, A. et al. Phase 2 study evaluating intermittent and continuous linsitinib and weekly paclitaxel in patients with recurrent platinum resistant ovarian epithelial cancer. *Gynecol. Oncol.* 149, 275–282 (2018).
- Leighl, N. B. et al. Phase 2 study of erlotinib in combination with linsitinib (OSI-906) or placebo in chemotherapy-naive patients with non-small-cell lung cancer and activating epidermal growth factor receptor mutations. *Clin. Lung Cancer* 18, 34–42.e2 (2016).
- Scagliotti, G. V. et al. Randomized, phase III trial of figitumumab in combination with erlotinib versus erlotinib alone in patients with nonadenocarcinoma nonsmall-cell lung cancer. *Ann. Oncol.* 26, 497–504 (2014).
- Snyder, A. G. et al. Intratumoral activation of the necroptotic pathway components RIPK1 and RIPK3 potentiates antitumor immunity. Sci. Immunol. 4, eaaw2004 (2019).
- Stanford, J. C. et al. Efferocytosis produces a prometastatic landscape during postpartum mammary gland involution. J. Clin. Invest. 124, 4737–4752 (2014).
- Phan, N. et al. A simple high-throughput approach identifies actionable drug sensitivities in patient-derived tumor organoids. Commun Biol. 2, 1–11 (2019).
- Zhou, Y. et al. Blockade of the phagocytic receptor MerTK on tumorassociated macrophages enhances P2X7R-dependent STING activation by tumor-derived cGAMP. *Immunity* 52, 357–373.e9 (2020).
- Meyer, A. S. & Heiser, L. M. Systems biology approaches to measure and model phenotypic heterogeneity in cancer. Curr. Opin. Syst. Biol. 17, 35–40 (2019)
- Tirosh, I. et al. Dissecting the multicellular ecosystem of metastatic melanoma by single-cell RNA-seq. Science 352, 189–196 (2016).
- Kimmerling, R. J. et al. Linking single-cell measurements of mass, growth rate, and gene expression. Genome Biol. 19, 1–13 (2018).
- 42. Gupta, P. B. et al. Stochastic state transitions give rise to phenotypic equilibrium in populations of cancer cells. *Cell* **146**, 633–644 (2011).
- Kuchen, E. E., Becker, N., Claudino, N. & Höfer, T. Long-range memory of growth and cycle progression correlates cell cycles in lineage trees. Preprint at https://doi.org/10.1101/373258 (2018).
- Comandante-Lou, N., Khaliq, M., Venkat, D., Manikkam, M. & Fallahi-Sichani, M. Phenotype-based probabilistic analysis of heterogeneous responses to cancer drugs and their combination efficacy. *PLoS Comput. Biol.* 16, e1007688 (2019)
- Galluzzi, L. et al. Molecular mechanisms of cell death: recommendations of the Nomenclature Committee on Cell Death 2018. Cell Death Differ. 25, 486–541 (2018).
- Plemel, J. R. et al. Unique spectral signatures of the nucleic acid dye acridine orange can distinguish cell death by apoptosis and necroptosis. J. Cell Biol. 216, 1163–1181 (2017).
- 47. Salvatier, J., Wiecki, T. V. & Fonnesbeck, C. Probabilistic programming in Python using PyMC3. *PeerJ Computer Sci.* **2**, e55 (2016).
- Greco, W. R., Bravo, G. & Parsons, J. C. The search for synergy: a critical review from a response surface perspective. *Pharmacol. Rev.* 47, 331–385 (1995).