Acute Effects of Drugs on Caenorhabditis elegans Movement Reveal Complex Responses and Plasticity

Mark Spensley,* Samantha Del Borrello,*† Djina Pajkic,* and Andrew G. Fraser*††
*Donnelly Centre for Cellular and Biomolecular Research, University of Toronto, Toronto, Ontario, M5S 3E1 Canada and
†Department of Molecular Genetics, University of Toronto, Toronto, Ontario, M5S 3E1 Canada

ORCID IDs: 0000-0001-6167-4461 (M.S.); 0000-0002-0117-0592 (S.D.B.); 0000-0001-9939-6014 (A.G.F.)

ABSTRACT Many drugs act very rapidly — they can turn on or off their targets within minutes in a whole animal. What are the acute effects of drug treatment and how does an animal respond to these? We developed a simple assay to measure the acute effects of drugs on C. elegans movement and examined the effects of a range of compounds including neuroactive drugs, toxins, environmental stresses and novel compounds on worm movement over a time period of 3 hr. We found a wide variety of acute responses. Many compounds cause rapid paralysis which may be permanent or followed by one or more recovery phases. The recoveries are not the result of some generic stress response but are specific to the drug e.g., recovery from paralysis due to a neuroactive drug requires neurotransmitter pathways whereas recovery from a metabolic inhibitor requires metabolic changes. Finally, we also find that acute responses can vary greatly across development and that there is extensive natural variation in acute responses. In summary, acute responses are sensitive probes of the ability of biological networks to respond to drug treatment and these responses can reveal the action of unexplored pathways.

Drugs are extremely powerful research tools. Addition of a drug can turn on or off a specific target protein and the effect on the organism can be followed over time. In model organisms, a well-characterized drug response can form the basis for genetic screens to identify the target protein and to find genes that modulate the effect of the drug. In C. elegans, for example, genetic screens for mutants with altered drug responses were key to finding the targets of several major anthelmintics (Lewis et al. 1980; Driscoll et al. 1989; Dent et al. 2000; Jones et al. 2005) as well as to identifying core components of conserved neuronal signaling pathways (Brenner 1974; Lewis et al. 1980; Nguyen et al. 1995; Miller et al. 1996).

Many genetic screens for C. elegans mutants with altered drug responses have used resistance to the toxic effects of chronic drug exposure as a target phenotype. In these screens, populations of mutant worms are typically exposed to drugs for long periods, ranging from hours to days (Rogalski et al. 1990; Kwok et al. 2006, 2008; Burns et al. 2006, 2015; Luciani et al. 2011). However, many drugs act very rapidly on worms yet currently relatively little is known about these acute responses of C. elegans to drugs. What do acute responses generally look like? Are acute responses the same at all developmental stages? Is there natural variation in acute responses as there is in mutant phenotypes or chronic drug responses? Do multiple drugs with the same long-term effect have the same acute responses or very different ones? What are the genes that affect the acute responses — are they the same or different genes as those that affect long-term drug effects?

Our goal in this study was to examine acute drug responses in C. elegans and address at least some of these questions. We developed an image-based method that accurately measures worm movement at high throughput. This allows us to gather rich information about the acute effects of drugs on worm movement, measuring both the effect of a range of doses of drugs and how the effects of any drug changes across time. We examined a wide range of drugs that were previously shown to be bioactive as well as a number of environmental stresses. We found that many drugs give responses with several distinct phases of paralysis.

KEYWORDS C. elegans Drug Assay Movement Acute Response

Diversity of Drug Responses in C. elegans

†Corresponding Author: Andrew G Fraser, Terrence Donnelly Centre for Cellular and Biomolecular Research, 160 College Street, Toronto, Ontario. M5S 3E1. Canada, andyfraser.utoronto@gmail.com, +1 416 978 2712
and recovery. We refer to these multi-phase responses throughout as “complex responses”. We show that the basis for these responses is drug-specific and that acute responses can differ greatly between different developmental stages. Finally, we compare acute responses in two isolates and find that there is substantial variation in acute responses and that this changes with dose, time and developmental stage. We thus find that measuring acute responses to drugs can reveal the action of unexplored pathways and that acute responses vary extensively and at many levels between different individuals. We suggest that ability to measure acute responses at high throughput may allow both drug screens and genetic screens to uncover the underlying molecular basis for these responses.

**MATERIALS AND METHODS**

**Worm Maintenance and Strains**

All worm stocks were maintained at 20°C on NGM agar plates seeded with *E.coli* strain OP50 as described elsewhere (Stiernagle 2006). In addition to the classical laboratory strain N2, we describe work using strains unc-38 (e264), lev-8 (x15), lev-1 (e211), unc-63 (x26), unc-29 (e1072), akr-8 (ok1240), akr-16 (ok789), gar-1 (ok755), gar-2 (ok520) and gar-3 (ok305). All strains were acquired from the Caenorhabditis Genetics Center. To prepare L1 worms, animals were washed from the agar plates with M9 buffer (Stiernagle 2006) and rinsed in one buffer change of M9. L1s were then isolated by filtration through 11 μm nylon mesh filter plates (Milipore Multiscreen). L1 larvae for immediate use in drug response assays were diluted to approximately 1.2 worms per microliter in either M9 buffer or modified NGM buffer (50 mM NaCl, 1 mM CaCl2, 1 mM MgSO4, 200 mM KH2PO4, 50 mM K2HPO4).

To assay drug response over different developmental stages, worm populations were first developmentally synchronized by filtration of L1s, as described above. Filtered L1s were returned to fresh NGM agar OP50 plates and incubated at 20°C until they had reached the required developmental stage (L1: no further incubation, L2: 18 hr, L3: 32 hr, L4: 42 hr, young adult: 54 hr). Worms were then washed from the plates and rinsed as above. To further synchronize developmental stages, worms were dispensed into assay wells using a COPAS Biosort machine (Union Biometrica), which can selectively dispense individual worms on the developmentally-correlated criteria of size and optical density (L1 worms: 100 / well, L2: 80 / well, L3: 50 / well, L4: 30 / well, young adult: 30 / well). Sorting parameters for each stage were determined empirically, as per the manufacturer’s instructions. The volume of liquid dispensed into the wells was determined and adjusted to 100 μl with buffer.

**Drug preparation**

Solutions of potassium cyanide (Sigma Aldrich 60178-25G), sodium chloride (BioShop SOD002.205), cycloheximide (Sigma Aldrich C7698-1G), abamectin (Sigma Aldrich S31732) and 2-deoxy-D-glucose (Sigma Aldrich D8375) were prepared fresh to twice the working concentration in M9 buffer, 1.6% DMSO. Stock solutions of aldicarb (Sigma Aldrich, 33886) were prepared to 1 M in DMSO. Solutions of levamisole hydrochloride (Acros Organics, 16595-80-5), oxotremorine M (Sigma Aldrich, O100) and mecamylamine hydrochloride (Sigma Aldrich, M9020) were prepared to 100 mM in water. These solutions were divided into single use aliquots and frozen. Stocks of arecoline hydrobromide (Acros Organics, 250130050) and atropine sulfate monohydrate (Sigma Aldrich, A0257) were prepared fresh on the day of use in M9 buffer. (-)-Nicotine (Sigma Aldrich, N3876) was stored frozen, undiluted in single use aliquots. Compound solutions described in Burns et al. (2015) were purchased as a custom library of 10 mM stock in DMSO (ChemBridge Corporation) and stored frozen.

**Preparation of drug response assays**

Assays were assembled in a final volume of 200 μl in flat-bottomed, polystyrene 96-well plates (Corning 3997) by combining 100 μl of worms in buffer with an equal volume of drug solution. To control for any confounding effects of drug solvent, all experiments were prepared to contain DMSO at 0.8% v/v, regardless of whether the drug solution was prepared in DMSO or water. All assays were assembled in M9 buffer apart from the screen for compounds that modify the response to aldicarb, which were assembled in modified NGM, which we found to cause fewer problems of drug precipitation. After assembling the assay, plates were sealed with transparent, self-adhesive films before imaging. The point at which worms and drugs were combined marked the start of the assay.

**Preparation of RNAi assays**

RNAi-by-feeding was performed as previously described (Lehner et al. 2006). After 4 days, RNAi-treated and mock treated worms were filtered to purify L1 worms and washed in two buffer changes of M9. Purified L1 samples were diluted to approximately 1.2 worms per microliter and 100 μl samples were transferred to microtiter plates for use in mobility assays.

**Image acquisition, analysis and calculation of fractional mobility score**

Images were acquired using a Nikon Ti Eclipse inverted microscope with 2x objective lens and DS-Qi1Mc camera. Image capture was automated in Nikon NIS-Elements AR software to capture two images of each well, separated by a 500 ms interval. This procedure was run in a 5 min loop for the 3 hr duration of the experiments. A sample NIS-elements script is available as supplementary information.

Since our microscope control software stores files in a proprietary image stack format (Nikon ND2), we began our image analysis with a file format conversion process to convert ND2 files to a collection of 8-bit TIFF images, with structured filenames identifying the timepoint and well coordinates of each image. The first step of this process, defined in the Python file *export_ND2.py*, converted the ND2 file to a series of TIFF files, using the Open Microscopy Environment Bio-Formats tool (Moore et al. 2015). This conversion yields a series of 12-bit TIFF files with names encoding the timepoint and order of acquisition of each image. The second step of the conversion process invoked the Image-Magick package to convert these files from 12-bit to 8-bit TIFF, modify filenames to include the well location, rather than order of acquisition and group files for each timepoint into separate subfolders. This file structure forms the input for the image analysis script.

Our image analysis pipeline was prepared with the Python programming language using the scikit-image library (van der Walt et al. 2014). All images were first processed with a Gaussian blur to reduce noise levels. Further image analysis consisted of two sets of processes to find moving and non-background, non-moving parts of the images.

To identify non-background, non-moving pixels, images were processed using a Sobel filter, which effectively produces the first derivative of the image, thus edges are highlighted and sharper edges appear brighter than shallow edges. An adaptive threshold was applied to the Sobel images to remove the background and yield a binary image. These binary images contained the outlines of the well and any objects, such as worms, within the well. Binary images were further processed with two iterations of binary closing to fill the pixels between sufficiently close edges. This process was hand-tuned to fill in the area occupied by worms but minimize overfilling into the areas between worms. A circular binary mask was applied to remove the edges of the wells from each
binary image and one of these images was designated as a reference image for each well (Figure 1A, Image B).

Pixels associated with movement were identified by calculating the absolute difference of the two consecutive images of a well (Figure 1A, Image A). A threshold was applied to the difference image to create a binary image (Figure 1A, Image B). This binary image contained the moving pixels from both parent images, showing double the number of movement-associated pixels. To correct for this, we multiplied the binary difference image by the reference image described above revealing only the non-background, movement associated pixels from a single frame (Figure 1A, Image C). Finally to identify pixels that did not move between wells, the reference image was multiplied by the complement of Image C (Figure 1A, Image D).

A fractional mobility score was calculated as the ratio of movement-associated pixels to total non-background pixels. This process is contained in the Python script `quantify_movement.py`. A second Python script, `quantify_movement_adult.py`, was used to analyze images of adult worms. This script contains an additional step to mask out objects below a certain number of pixels in size, to remove eggs from the analysis, since some chemicals stimulate egg-laying.

**Data analysis and curve-fitting**

All drug response data shown were normalized by dividing by the fractional mobility score of drug-free control wells for each time point. Fitted lines were determined by least squares minimization of either a three-parameter logistic function:

$$FMS_{\text{fractional}} = \frac{\text{movement-associated pixels}}{\text{total non-background pixels}}$$

**Figure 1 Outline of an image-based assay for worm mobility.** A: Worms are placed in a 96 well plate in M9 buffer and treated with drugs as required. Two images of each well (Im1 and Im2) are captured at an interval of 500 ms at each timepoint. Images are processed as described in Materials and Methods to identify portions of the worms that move between the two images (green) and parts that remain stationary (red). This allows the calculation of a raw fractional mobility score (FMS) as the fraction of total worm-associated pixels (red and green) that moved (green). This processing is illustrated in a magnified region of Im1 and Im2 (defined by blue box). B: The image-based assay produces a rich view of drug effects on worm movement. L4 worms were treated with a range of doses of aldicarb and movement measured at 5 min time points over a 3 hr time course. The heat map shows the FMS for each drug dose at each timepoint — white indicates full movement, red indicates lack of movement. Either dose response curves at a given time point (pink arrow) or kinetic responses to a single drug dose (red and blue arrows) can be extracted from these rich data. Data are means of four independent experiments. Error bars show standard error of the mean.
or an exponential function:
\[ f(x) = Me^{-x} + A \]

Where \( x \) denotes drug concentration and \( M, A \) and \( B \) denote parameters to be optimized. The choice of function was determined by the range of the dataset.

Proportional recovery scores were calculated using the formula:

\[ \frac{(\text{FMS after recovery} - \text{FMS at maximum paralysis})}{1 - \text{FMS at maximum paralysis}} \]

The value of FMS after recovery was calculated as the mean FMS over the period between 150 and 180 min. Maximum paralysis was determined as the minimum FMS value over the three hour course of the experiment.

Data availability

A full description of the library compounds we tested is available in Supplementary File S1 and response data for all compounds yielding acute movement effects are available in Supplementary File S1. All our Python scripts and an example dataset are available in our repository, https://github.com/fraser-lab-UofT/acute_assay. Supplementary material available at Figshare: https://doi.org/10.25387/g3.6670892.

RESULTS

Development of a high throughput image-based movement assay

We wanted to measure the acute effects of drugs on worms over the time period of minutes to hours. To sample a diverse range of compounds affecting movement in C. elegans, we focused on movement as a key phenotype — a very wide range of compounds affect movement including toxins, neuroactive compounds and environmental stresses. There are a wide variety of assays to measure the acute effects of drugs on movement, with different levels of throughput and resolution, and different requirements for specialized equipment such as worm trackers or microfluidic devices (Buckingham and Sattelle 2009; Swierczek et al. 2011; Yemini et al. 2011, 2013; Brown et al. 2013; Larsch et al. 2013). Here we focused on creating a simple assay for worm movement that we detect in our assay are specific to the action of elevated ACh on nAChRs. In the presence of 600 \( \mu \)M mecamylamine, the EC50 of aldicarb shifts from 0.76 mM to 1.6 mM, with a slight decrease in maximal response (Figure 2C). The effects of Ald on worm movement that we measure in our assay are thus specific to the action of elevated ACh on nAChRs.

We note that while we focus here on the use of this assay to measure the effects of drugs on worm movement, it can also be used to measure the effect of genetic perturbations on movement directly. To illustrate this, we used RNA-mediated interference (RNAi) to knock down 7 genes that are known to be required for wild-type movement and measured movement after RNAi. We found robust RNAi phenotypes for all 7 genes (Figure 2D) showing that our movement assay is applicable to both genetic and drug based screens. Thus, we expect our method to be applicable in a range of pharmacological, genetic, and RNAi experiments.

We have thus developed a simple method that allows us to measure the acute effects of drug treatments on worm movement at high throughput. We showed that our assay recapitulates the known effects of the well-characterized neuroactive drug Ald and that this is indeed due to its known mode of action through the activation of specific receptors. Crucially, our assay allows us to measure both dose responses and time-resolved responses to drugs.

Acute responses to drugs are often complex

We established a simple assay to measure acute responses of worms to drugs over time periods of minutes to hours. To sample a diverse range of acute responses, we exposed C. elegans L1 larvae to a wide range of
individual genes known to be required for normal movement. L1 progeny from these cultures were isolated by mean of 8 wells, containing 100 worms per well.

different compounds and treatments including compounds that inhibit core essential machineries such as the electron transport chain (inhibition by potassium cyanide, KCN, shown in Figure 3A) or the ribosome (effect of cycloheximide in Figure 3B), environmental stresses such as high salt conditions (NaCl shown in Figure 3C), known nematicides (abamectin shown in Figure 3D), compounds that affect movement via altered neurotransmission (e.g., aldicarb Figure 2A). For each of these chemically and mechanistically diverse treatments, we were able to elicit a paralysis response, demonstrating the broad applicability of our assay to different chemical perturbations. Unexpectedly, we found several of these compounds elicited responses with several distinct phases, where an initial phase of paralysis was followed by recovery of movement; we refer to these multi-phase responses as “complex responses”.

To estimate how widespread complex responses might be, we examined responses to 170 compounds that had been identified as affecting worm growth in chronic exposure, population growth assays (Burns et al. 2015; Supplementary File S1). Of the 53 compounds with acute effects on worm movement, more than half (31 of 53 with acute effects; 58%) showed complex time-resolved responses (Supplementary File S2), such as the examples illustrated in Figure 3E-G, including either sustained or transient recovery from initial drug effects, suggesting that complex acute responses may be relatively common in C. elegans.

Having looked at acute responses to across a range of different treatments, we found complex acute responses across every type of compound, including neuro-active compounds, environmental stresses, and toxins that target essential genes. These responses can be very complex — for example, the novel compound shown in Figure 3G has two distinct phases of recovery and several structurally related compounds have similarly complex responses (data not shown). We note that every one of these complex drug responses can be the basis either for a genetic screen for mutants with altered responses or for drug screens for compounds that modulate the response. This complexity of many acute drug responses underlines the need to be able to measure time-resolved acute effects of drugs on movement.

The complex effects of many compounds over time suggest that the worms are responding to many drug treatments in some way — for example, several drugs cause an initial rapid paralysis followed by a recovery phase where the worms begin to move again at almost normal rates. To gain some insights into mechanisms underlying such complex responses, we focused on the acute responses to two well-characterized compounds, aldicarb and cyanide.

**Agonists of acetylcholine signaling have complex effects on worm movement that change Across development**

As shown in Figure 2A treatment of adult animals with Ald results in a rapid reduction in movement and sustained paralysis. Interestingly, we found that L1 animals behave completely differently— Ald induces rapid paralysis of L1 worms but this is followed by a more gradual recovery (Figure 4A; Supplementary Figure S2). To further examine the Ald response across development, we tested all developmental stages and found the response is qualitatively very different across development (Figure 4A). L1 and L2 stage larvae show an initial phase where Ald treatment results in reduced movement, followed by a recovery phase during which paralysis is gradually relieved. As worms progress through the four larval stages, the initial paralysis remains similar but this recovery is reduced, becoming minimal by adulthood.

To explore the mechanisms underlying recovery of L1s from paralysis, we asked whether this is specific to Ald or whether other drugs that affect ACh signaling show similar acute responses. Ald affects ACh signaling by preventing the degradation of ACh causing this to accumulate in the synapse—it does not activate ACh receptors directly nor does its action have any specificity for any specific sub-type of ACh.
receptor. Nicotine (Nic) activates ACh signaling in a different way — it is a direct agonist of specific ACh receptors, the nicotinic ligand-gated ion channels. We compared their acute responses and found that both Ald and Nic show a rapid initial paralysis followed by a recovery of movement (Figure 4B). While these two responses have superficial similarities, there is a crucial difference in the recovery phase. For Nic, while L1 worms recover from the reduction of movement caused by low doses of Nic, increasing concentrations of Nic result in greater paralysis and less recovery until at high Nic doses there is no appreciable recovery (Figure 4D). For Ald, we see the precise opposite — there is no recovery at low doses of Ald, but recovery increases with increasing Ald (Figure 4G). This suggests that the recovery from Ald and Nic-induced paralysis is fundamentally different and that Ald is somehow driving recovery since the more Ald we add, the stronger the recovery. How could Ald be acting and how could this be different to Nic?

ACh is known to activate two completely different classes of receptor: ligand-gated ion channels (nAChRs) and muscarinic ACh receptors (mAChRs). nAChRs act rapidly and are stimulated directly by Nic and indirectly by Ald via its action to increase ACh. mAChRs have much more complex G-protein coupled signaling — crucially, they are not activated by Nic but are also activated following Ald treatment. We hypothesized that the Ald recovery phase might be mediated by mAChRs and test this below.
Recovery From rapid paralysis induced by Aldicarb occurs via muscarinic receptor signaling

To test whether recovery from Ald-induced paralysis might be due to the activation of mAChRs, we examined whether a mAChR antagonist, atropine (Atr), could block the recovery phase of the Ald response. This is indeed the case (Figure 5A) suggesting that recovery is the result of an activation of mAChR signaling following increased ACh levels after Ald treatment. The C. elegans genome encodes three muscarinic acetylcholine receptors, GAR-1, GAR-2 and GAR-3 (Hwang et al. 1999; Lee et al. 1999, 2000) — we obtained deletion mutants for each of these receptors and tested whether the effect Ald was altered in these mutant strains. The Ald response is clearly different in the gar-3(gk305) mutant strain: while the paralysis phase appears very similar to wild-type, the recovery phase is strongly suppressed in the gar-3(gk305) mutant, although not completely abolished (Figure 5B). We note that mutations in gar-1 and gar-2 also have effects on recovery but these are much weaker. Our data thus suggest that gar-3 is required for most of the recovery from nAChR-induced paralysis in L1 animals, although we note that we cannot exclude substantial redundancy between these genes.

We thus propose that Ald causes a complex acute response in L1 worms because the increased ACh levels have effects on two separate receptor pathways. The first phase of the acute response is a rapid reduction in movement due to activation of nAChRs. The second phase is a slower recovery due to activation of mAChRs which somehow relieve the nAChR-driven paralysis. To further test this, we examined whether we could drive recovery from Nic-induced paralysis with agonists of mAChRs. Nic can only stimulate nAChRs (Ishii and Kurachi 2006). Consistent with our model, we found that we could induce recovery from Nic-induced paralysis either by adding Ald (Figure 5C) or the mAChR agonists, Oxotremorine M (OxoM) or Arecoline (Are) (Figure 5D; Supplementary Figure S3). To confirm that the same mAChR requirement apply to OxoM-induced recovery as in Ald responses, we tested the ability of each gar mutant to recover from Nic-induced paralysis in the presence of OxoM (Figure 5E). We found that gar-3 is indeed the primary mAChR driving OxoM-induced recovery. Our data suggest that mAChR signaling can drive recovery from paralysis induced by sustained nAChR activation. Finally, we examined whether changes in mAChR signaling might underlie the difference in the ability of L1 and adults to recover from Ald paralysis. We find that stimulation of mAChR signaling by OxoM in adults cannot override the paralysis response (Figure 5F) and thus, changes in the mAChR signaling machinery might indeed account for the different responses across development.

Our data thus suggest that a model in which there is some kind of physiological cross talk between two types of ACh receptor, nAChRs and mAChRs. This model is summarized in Figure 6, and suggests that i) sustained activation of nAChRs alone results in long-term paralysis, ii) sustained activation of mAChRs alone has little effect on movement, but iii) dual activation of both nAChRs and the mAChR GAR-3, such as with Ald alone, or Nic + OxoM results in an initial phase of nAChR-driven paralysis followed by a slower mAChR-driven recovery. We conclude that the complex acute response of L1 worms to Ald is highly specific to the drug used and is not a generic stress or xenobiotic response. To examine whether this is true for other drugs, we next looked at the effects of a very different drug, KCN, on worm movement.
C. elegans has a complex response to treatment
With cyanide

Cyanide (KCN from here on) has a very well-characterized mode of action: it inhibits aerobic metabolism by blocking the mitochondrial electron transport chain (ETC) through binding to complex IV (reviewed in Hanstein 1976). KCN is toxic to C. elegans following prolonged exposure — here we examined the acute responses of worms to KCN exposure.

We immediately noticed that the dose response curves (DRCs) at early time points were unusual — while intermediate doses of KCN caused almost complete paralysis movement, high doses appeared to have little effect (Figure 7A). This was most pronounced in DRCs between 15-30 minutes — by ~60 min the DRCs appeared much more normal, with increased effects being seen with increased dose.

How can the effect of low concentrations of KCN be greater than the effect of high concentrations? We reasoned that this may be due to rewiring of metabolic pathways — worms might respond to high KCN concentrations by rapidly switching to use alternative metabolic pathways for generating ATP. An obvious candidate for such a pathway was anaerobic glycolysis which is the major pathway for ATP generation in anaerobic conditions in many animals — indeed we found that addition of 2-deoxy-D-glucose, an inhibitor of phosphoglucose isomerase, a critical enzyme in anaerobic glycolysis, caused a profound change in KCN DRCs. Worms can no longer continue moving in high doses of KCN when 2DG is also present — indeed we found that increased concentrations of KCN result in decreased movement at all time points (Figure 7A). This suggests that the unusual DRCs seen for KCN alone were due to a shift to utilization of anaerobic glycolysis at early time-points in high KCN (Figure 7B). In the absence of KCN, 2DG alone had no detectable effect in our assay over the same time period (Supplementary Figure S4).

This underscores the importance of measuring drug responses across time — if we had only measured KCN DRCs at 80 min, they would have looked completely unremarkable. It also confirms that the way that worms respond to acute drug treatment is highly specific to the drug itself — while the aldicarb response involves crosstalk between nAChRs and mAChRs, the KCN response involves metabolic switches. The rapid responses of worms to compounds thus do not appear to be generic stress or xenobiotic responses but are specific responses to the specific drug effects. They reveal how the organism can rewire and adapt to sudden inhibition of a key pathway.
Natural variation in acute drug response is complex

Drug responses are known to vary within natural populations. In C. elegans, natural variation in drug responses has been seen for several drugs including abamectin and etoposide (Ghosh et al. 2012; Zdraljevic et al. 2017). In a previous study we compared the effects of KCN on two isolates, the lab-adapted N2 strain and the natural isolate CB4856 (Vu et al. 2015). These differ by around 1SNP/800bp (Wicks et al. 2001), a similar degree of variation as that between two human genomes. In our previous study we simply examined one time point (90 min after drug treatment) and only looked at a single larval stage, the L1 stage — we wanted to expand this analysis to compare the effects of KCN on N2 and CB4856 across development. We decided to examine how a range of concentrations of KCN affect movement of all 4 larval stages of either N2 or CB4856 and in each case to examine how this response evolves over 3hr (Figure 8).

The results are complex and striking. In keeping with our previous study (Vu et al. 2015), we found that N2 animals are more severely affected by KCN at the L1 stage. However, we found the exact reverse at the L4 stage — CB4856 are substantially more sensitive here. At the L3 stage, the picture is even more complex — while KCN appears to affect CB4856 more rapidly (CB4856 L3 animals more severely affected at early timepoints), the same dose of KCN has a more severe effect on N2 at later timepoints. Taken together, these data show clearly that comparisons of drug sensitivity in natural populations are complex. It is impossible to make a blanket statement that isolate A is ‘more sensitive’ than isolate B — assessing the effect at a different dose, a different timepoint or a different developmental stage could give precisely the opposite conclusion.

DISCUSSION

Drugs have extremely rapid effects on their targets — they can switch on or off their target within minutes even in a whole animal. What is the effect on the animal and how does it respond? We looked at the effects of drugs on C. elegans movement to measure acute responses to drugs in a whole animal and find several unexpected results that have important implications for drug screens in animals.

First, we found that around half of all the bioactive compounds that we found to affect C. elegans movement show a complex response: the worms are rapidly affected, but after an early phase of near-paralysis, the animals recover movement — this recovery may be transient or longer lived. In some cases, like the responses to compound shown in Figure 3E-F, we see multiple phases of paralysis and recovery over time — studying these complex acute responses of worms over time will drive exciting new insights. That worms can recover from the rapid effects of many drugs indicates that the worms are responding in some way. We did not expect to see this kind of recovery so frequently and to such a wide range of drugs and treatments, or responses of such complexity — that surprise drove this study.

Second, at least in the examples we examined, the mechanisms underlying complex acute responses appear to be drug specific and not due to a generic stress or xenobiotic response. For example, recovery from paralysis due to signaling by one set of acetylcholine receptors (AChRs) is due to signaling from a different set of AChRs; recovery from paralysis due to inhibition of aerobic respiration is driven by the use of anaerobic pathways. We see complex acute responses to a wide range of drugs and thus worms can respond to many different acute perturbations of their core networks by rewiring and modulating the activity of specific pathways. This suggests a tremendous complexity of homeostatic mechanisms and sensors and this molecular biology can be revealed through studying acute drug responses. For example, worms recover from paralysis induced by activation of nAChRs through signaling from mAChRs — this cross talk is not evident in chronic drug assays but can be easily measured using acute responses. Acute responses are thus excellent reporters for the rewiring of genetic networks in response to the effects of a drug.

Finally, we find that many factors affect acute drug responses. The acute response to the same drug can be different at different developmental
stages, at different doses of drug, or in different individuals — measuring only a single dose, a single time point, or a single developmental stage can greatly affect conclusions. For example, we found that L1 larvae recover readily from Aldicarb-induced paralysis but adults cannot recover. The recovery of L1s from Ald-induced paralysis is highly dose-dependent — L1s do not recover at 2mM Ald but recover readily at the higher concentration of 6mM (Figure 4A). This kind of complexity means one cannot simply say ‘C. elegans can recover from Ald paralysis’ or ‘C. elegans cannot recover from Ald paralysis’. We see a similar complexity when we look at natural variation in acute responses. We compared the sensitivity to KCN of two natural isolates of C. elegans, N2 and CB4856 and find a complex picture — while N2 is more sensitive at the L1 stage, CB4856 is more sensitive at the L4 stage. Again, we cannot make a blanket statement of ‘N2 is more sensitive than CB4856’ — dose, time point, and developmental stage drive major differences in the drug responses and these factors are often neglected. For example a GWAS to identify QTL that affect ‘drug sensitivity’ would likely find different variants depending on drug dose and developmental stage assayed. Crucially, this means that to really understand acute responses, we need to measure them over a range of doses and for each developmental stage. The simple assay we report here allows us to gather this kind of rich data.

Taken together, our results show that worms have complex acute responses to a wide variety of drugs and that these complex responses are due to the specific rewiring of the pathways underlying worm biology. Acute responses are sensitive readouts of the pathways that are required after rewiring — some of these are largely unexplored. For example, activation of mACHR signaling alone only has subtle effects on worm biology (Steger and Avery 2004; Liu et al. 2007; Dittman and Kaplan 2008; Hendricks et al. 2012; Chan et al. 2013) but acute responses reveal a clear effect of mACHR signaling in driving recovery from nACHR-induced paralysis. Genetic screens to find mutants with altered acute drug responses can thus dissect otherwise intractable pathways and acute responses can also be the basis for sensitive drug screens. Understanding the genetics that underpin complex acute responses like those shown in Figure 3 will reveal novel insights into how genetic networks rewire and change following insult.

Importantly, we also show that acute drug responses vary in many ways — different drug doses, different developmental stages, and different genetic backgrounds can give different acute responses. Drawing broad conclusions about drug responses such as ‘C. elegans recovers from Aldicarb-induced paralysis’ or ‘the N2 isolate is more sensitive to KCN than the CB4856’ is thus very difficult — different doses or different developmental stages often give completely different results. Studies on drugs in whole animals need to be aware of this complexity and interpret results with care. We believe that high throughput quantitative assays for drug responses of the kind we describe here are key — they allow researchers to build up a rich view of a drug response rather than focus on an arbitrary dose, time point or developmental stage. Studying the full complexity of acute responses opens up a whole new set of whole animal phenotypes for drug and genetic screens and will reveal how animals rapidly rewire their genetic networks in response to drugs, toxins, and environmental stresses.

Figure 7 Dose sensitivity to KCN is complex. A: L1 worms were treated with a range of KCN doses and their movement measured over time. The Fractional Mobility Scores (FMS) are shown at various time points. At early time points the dose response curves are complex — intermediate doses cause more severe movement defects than high doses. B: Decreased sensitivity to KCN at higher doses is dependent on a metabolic transition to glycolysis. Inhibition of glycolysis with 2-dexoxy-D-glucose (61 mM) causes higher-doses of KCN to be more toxic than lower doses, in contrast to the dose-response relationship of KCN alone. This suggests that the relatively mild effects of higher KCN concentrations seen in panel A is at least partially attributable to KCN stimulating a shift in metabolic pathways from oxidative phosphorylation to glycolysis. Data shown are at the 20 min time point. A-B: Curves show mean of 3 repeats; error bars are standard error of the mean.

Figure 8 Variation in dose sensitivity to KCN changes across development. A: Worms of specific stages of either N2 or CB4856 isolates were purified using a worm sorter. Worms are sorted based on length and darkness — the exact windows chosen are indicated by colored boxes and examples of the sorted worms shown. B: N2 or CB4856 worms of specific developmental stages were exposed to a range of doses of KCN and their movement measured over a 3hr time course. The fractional mobility score (FMS) was calculated at each time point. The figure shows the difference in FMS scores for N2 and CB4856 — blue colors indicate that N2 has a lower score, red colors that CB4856 is moving less.
ACKNOWLEDGEMENTS

This work was supported by grant PJT-153036 to A.G.F from the Canadian Institutes for Health Research. Some strains were provided by the CGC, which is funded by NIH Office of Research Infrastructure Programs (P40 OD010440).

LITERATURE CITED

BouJnak, M., H. Ha, N. Maffey, and Y. Zhang, 2012 Compartmentalization
Jones, A. K., S. D. Buckingham, and D. B. Sattelle, 2005 Chemistry-to-gene
Kwok, T. C. Y., K. Hui, W. Kostelecki, N. Ricker, G. Selman et al., 2008 A
New Residues Required for DHP-Blockage of Mammalian Calcium Channels. PLoS Genet. 4: e1000067. https://doi.org/10.1371/journal.pgen.1000067
Kwok, T. C. Y., N. Ricker, R. Fraser, A. W. Chan, A. Burns et al., 2006 A small-molecule screen in C. elegans yields a new calcium channel antagonist. Nature 441: 91–95.
Larsch, J., D. Ventimiglia, C. I. Bargmann, and D. R. Albrecht, 2013 High-throughput imaging of neuronal activity in Caenorhabditis elegans. Proc. Natl. Acad. Sci. USA 110: E4266–E4273. https://doi.org/10.1073/pnas.1318325110
Lee, Y. S., Y. S. Park, D. J. Chang, J. M. Hwang, C. K. Min et al., 1999 Cloning and expression of a G protein-linked acetylcholine receptor from Caenorhabditis elegans. J. Neurochem. 72: 58–65.
Lee, Y. S., Y. S. Park, S. Nam, S. J. Suh, J. Lee et al., 2000 Characterization of GAR-2, a novel G protein-linked acetylcholine receptor from Caenorhabditis elegans. J. Neurochem. 75: 1800–1809.
Lehner, B., C. Crombie, J. Tischler, A. Fortunato, and A. G. Fraser, 2006 Systematic mapping of genetic interactions in Caenorhabditis elegans identifies common modifiers of diverse signaling pathways. Nat. Genet. 38: 896–903.
Lewis, J. A., C. H. Wu, H. Berg, and J. H. Levine, 1980 The genetics of levamisole resistance in the nematode Caenorhabditis elegans. Genetics 95: 905–928.
Liu, Y., B. LeBoeuf, and L. R. Garcia, 2007 G alpha(q)-coupled muscarinic acetylcholine receptors enhance nicotinic acetylcholine receptor signaling in Caenorhabditis elegans mating behavior. J. Neurosci. 27: 1411–1421.
Luciani, G. M., L. Magomedova, R. Puckrin, M. L. Urbanus, I. M. Wallace et al., 2011 Dafadine inhibits DAF-9 to promote dauer formation and longevity of Caenorhabditis elegans. Nat. Chem. Biol. 7: 891–893. https://doi.org/10.1038/nchembio.698
Miller, K. G., A. Alfonso, M. Nguyen, J. A. Crowell, C. D. Johnson et al., 1996 A genetic selection for Caenorhabditis elegans synaptic transmission mutants. Proc. Natl. Acad. Sci. USA 93: 12593–12598.
Moore, J., R. K. Linkert, Melissa Blackburn, Colin Carroll, Mark Ferguson, H. Flynn, K. Gilling, R. Leigh et al., 2015 OMERO and Bio-Formats: flexible access to large bioimaging datasets at scale. Proc. SPIE 9413, Med. Imaging 2015 Image Process. 941307.
Nguyen, M., A. Alfonso, C. D. Johnson, and J. B. Rand, 1995 Caenorhabditis elegans mutants resistant to inhibitors of acetylcholinoesterase. Genetics 140: 527–535.
Opperman, C. H., and S. Chang, 1991 Effects of Aldicarb and Fenamiphos on Acetylcholinesterase and Motility of Caenorhabditis elegans. J. Nematol. 23: 20–27.
Rand, J., 2007 Acetylcholine (The C. elegans Research Community, Ed.), Richmond, J. E., and E. M. Jorgensen, 1999 One GABA and two acetylcholine receptors function at the. Nat. Neurosci. 2: 791–798.
Rolagski, T. M., M. Golomb, and D. L. Riddle, 1990 Mutant Caenorhabditis elegans RNA polymerase II with a 20,000-fold reduced sensitivity to alpha-amanitin. Genetics 126: 889–898.
Steger, K. A., and L. Avery, 2004 The GAR-3 muscarinic receptor cooperates with calcium signals to regulate muscle contraction in the Caenorhabditis elegans pharynx. Genetics 167: 633–643.
Sternberg, T., 2006 Maintenance of C. elegans, in Wormbook, edited by The C. elegans Research Community.
Swierczek, N. A., A. C. Giles, C. H. Rankin, and R. A. Kerr, 2011 High-throughput behavioral analysis in C. elegans. Nat. Methods 8: 592–598. https://doi.org/10.1038/nmeth.1625
Towers, P. R., B. Edwards, J. E. Richmond, and D. B. Sattelle, 2005 The Caenorhabditis elegans lev-8 gene encodes a novel type of nicotinic acetylcholine receptor alpha subunit. J. Neurochem. 93: 1–9.
Vu, V., A. J. A. J. Verster, M. Schertzbarg, T. Chulunbaatar, M. Spensley et al., 2015 Natural Variation in Gene Expression Modulates the Severity of Mutant Phenotypes. Cell 162: 391–402. https://doi.org/10.1016/j.cell.2015.06.037
van der Walt, S., J. Schönberger, F. Nunez-Iglesias, J. Boulogne, J. Warner et al., 2014  scikit-image: Image Processing in Python. Peerj 2: e453. . https://doi.org/10.7717/peerj.453

Wicks, S. R., R. T. Yeh, W. R. Gish, R. H. Waterston, and R. H. Plasterk, 2001 Rapid gene mapping in Caenorhabditis elegans using a high density polymorphism map. Nat. Genet. 28: 160–164.

Yemini, E., T. Jucikas, L. J. Grundy, A. E. X. Brown, and W. R. Schafer, 2013 A database of Caenorhabditis elegans behavioral phenotypes. Nat. Methods 10: 877–879. https://doi.org/10.1038/nmeth.2560

Yemini, E., R. A. Kerr, and W. R. Schafer, 2011 Tracking Movement Behavior of Multiple Worms on Food. Cold Spring Harb. Protoc. 2011: pdb. prot067025.

Zdraljevic, S., C. Strand, H. S. Seidel, D. E. Cook, J. G. Doenck et al., 2017 Natural variation in a single amino acid substitution underlies physiological responses to topoisomerase II poisons. PLoS Genet. 13: e1006891. https://doi.org/10.1371/journal.pgen.1006891

Communicating editor: M. Boutros