Title: Uncovering active modulators of native macroautophagy through novel high-content screens

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Abstract

Autophagy is an evolutionarily conserved pathway mediating the breakdown of cellular proteins and organelles. Emphasizing its pivotal nature, autophagy dysfunction contributes to many diseases; nevertheless, development of effective autophagy modulating drugs is hampered by fundamental deficiencies in available methods for measuring autophagic activity, or flux. To overcome these limitations, we introduced the photoconvertible protein Dendra2 into the *MAP1LC3B* locus of human cells via CRISPR/Cas9 genome editing, enabling accurate and sensitive assessments of autophagy in living cells by optical pulse labeling. High-content screening of 1,500 tool compounds provided construct validity for the assay and uncovered many new autophagy modulators. In an expanded screen of 24,000 diverse compounds, we identified additional hits with profound effects on autophagy. Further, the autophagy activator NVP-BEZ235 exhibited significant neuroprotective properties in a neurodegenerative disease model. These studies confirm the utility of the Dendra2-LC3 assay, while simultaneously highlighting new autophagy-modulating compounds that display promising therapeutic effects.

Introduction

Macroautophagy (hereafter referred to as autophagy) is an essential pathway for protein homeostasis whereby cytoplasmic proteins and organelles are delivered to lysosomes for degradation¹. Through the coordinated action of a series of autophagy-related (ATG) proteins and cargo receptors including p62/SQSTM1, NBR1, and optineurin², substrates are sequestered within double membrane vesicles called autophagosomes. Autophagosomes mature as they traffic along microtubules and eventually fuse with lysosomes to form autolysosomes, wherein hydrolases degrade autophagic cargo. The protein LC3 (ATG8) is an obligate component of autophagosome membranes and is itself degraded within autolysosomes. For these reasons, it often

serves as both a marker of autophagosomes and as a representative autophagy substrate³.

Underscoring the critical requirement of autophagy in cellular homeostasis, deletion of core autophagy genes in mice results in embryonic lethality^{4,5,6}. Accordingly, dysfunctional autophagy is linked to a wide spectrum of human disease including neurodegeneration, cancer, metabolic disorders, infectious and cardiovascular diseases⁷. Often these conditions involve deficiencies in one or more steps of autophagy, resulting in impaired clearance of potentially toxic cellular components, and/or a failure to replenish amino acids required for anabolic processes. In these instances, enhancing the rate of autophagic cargo clearance, commonly referred to as flux, would be beneficial. Conversely, autophagy can promote tumor progression and resistance to chemotherapy for some cancers^{8,9,10,11}. Here, autophagy inhibition may represent a more apt therapeutic strategy⁷.

Autophagy is of particular importance in the central nervous system (CNS). Deletion of essential autophagy genes within the CNS of mice leads to progressive neurodegeneration marked by accumulation of protein aggregates^{12,13,14,15,16}. Defective autophagy is a common feature of many neurodegenerative diseases including Alzheimer's disease^{17,18}, Parkinson's disease^{19,20,21,22,23}, polyglutamine disorders^{24,25,26}, amyotrophic lateral sclerosis (ALS)²⁷ and frontotemporal dementia (FTD)^{28,29,30,31}. Moreover, mutations in several autophagy related proteins including p62/SQSTM1³², optineurin³³, C9ORF72^{34,35}, TBK1³⁶, and UBQLN2³⁷ results in familial ALS and FTD.

Due to its broad therapeutic potential, autophagy modulation has received considerable attention as a target for drug development⁷. Nevertheless, these efforts have thus far failed to translate into effective therapies for patients. This is in part due to the intrinsic difficulties in measuring autophagic flux, and consequent inability of many conventional and widely used autophagy assays to accurately estimate flux³. One prominent limitation of these assays is an implicit reliance on the steady-state abundance of pathway intermediates such as LC3-II, the lipidated isoform of LC3. Due to the dynamic nature of autophagy, changes in such intermediates may equally reflect increased autophagy induction or late-stage inhibition of autophagosome clearance; although discriminating among these mechanisms is crucial for drug development, many assays are effectively unable to do so. While lysosomal inhibitors such as bafilomycin-A1 can be used to isolate autophagy induction from inhibition, this approach obscures estimates of substrate clearance, perhaps the most relevant measure of autophagic flux.

Bafilomycin-A1 and related compounds are also inherently toxic, further confounding flux measurements^{38,39}. Yet another common shortcoming is an inherent reliance on static “snapshots” of separate cellular populations that cannot be followed prospectively or longitudinally due to the need for cell lysis and measurement of pathway intermediates.

We previously developed a technique called optical pulse labeling (OPL), enabling non-invasive measurements of autophagic flux in living cells without the need for lysosomal inhibition⁴⁰. In this technique, LC3 is labeled with the photoconvertible protein Dendra2⁴¹. Upon exposure to blue (405nm) light, Dendra2 fluorescence irreversibly shifts from green to red. Since the generation of red-fluorescent Dendra2-LC3 is limited by blue light, LC3 turnover can be determined independent of new protein synthesis by tracking the time-dependent reduction in red fluorescence following a brief pulse of blue light (Fig. 1A). LC3 is an autophagy substrate, and therefore its degradation kinetics serves as a faithful proxy for estimates of autophagic flux. While OPL offered several advantages over conventional assays, it was nonetheless limited by its reliance on protein overexpression; in effect, Dendra2-LC3 overexpression floods the pathway under investigation with an obligate substrate. Burdening the cell with non-physiological concentrations of substrate might artificially prolong flux estimates, or conversely enhance flux via regulatory feedback mechanisms. Moreover, because autophagy regulation is intricately tied to amino acid availability^{42,43} and the ubiquitin proteasome system⁴⁴, any perturbations to these pathways brought on by protein overexpression may further confound measurements of flux.

To overcome these drawbacks, we labeled *native* LC3 with Dendra2 using CRISPR/Cas9 editing, producing a novel autophagy reporter cell line capable of assaying flux in living cells without the need for drug treatment, protein overexpression or measurements of pathway intermediates, thus establishing a faithful reporter of endogenous autophagy activity unadulterated by exogenous manipulations. Leveraging this cell line for its unique perspective on autophagy and the opportunities it presents, we adapted Dendra2-LC3 cells for conducting high-content screens and identified several new and active autophagy modulators with promising therapeutic properties.

Results

Creation of a novel reporter of autophagic flux

We employed CRISPR/Cas9 genome editing to label native LC3 by introducing Dendra2 into the *MAP1LC3B* locus (encoding LC3) of human embryonic kidney (HEK)

cells⁴⁵ (Fig. 1B). To minimize the risk of undesired insertions/deletions via non-homologous end-joining we used a dual-nickase strategy⁴⁶, in which Cas9(D10A) was expressed along with two single-guide RNAs (sgRNAs) targeting sequences immediately upstream and downstream of the *MAP1LC3B* start codon. Unlike wild-type Cas9, Cas9(D10A) induces single-stranded nicks rather than double-stranded breaks in the DNA, limiting recombination to the region marked by the sgRNAs. In addition, a vector containing the Dendra2 open reading frame (ORF) flanked by 400bp of homologous sequence 5' and 3' to the *MAP1LC3B* start codon was introduced to facilitate homology directed repair (HDR), thereby creating a sequence encoding Dendra2 fused to the N-terminus of LC3. Positive cells were selected based on Dendra2 fluorescence and enriched by sequential passaging until a homogenous population was achieved. Western blotting confirmed the successful insertion of the Dendra2 ORF into the desired locus (Fig. 1C). Transfection with siRNA targeting LC3 substantially reduced both Dendra-LC3 protein levels and native GFP fluorescence, providing further validation of successful on-target CRISPR editing (Fig. 1C,D).

In untreated cells, Dendra2-LC3 fluorescence was diffusely distributed, matching the predicted localization of the non-lipidated, cytosolic LC3-I isoform (Fig. 1E, Supplemental movie 1). Treatment with the potent autophagy inducer Torin1⁴⁷ elicited the formation of visible fluorescent puncta and reduced the intensity of diffuse Dendra2-LC3 (Fig. 1E, Supplemental movie 2), reflecting the incorporation of Dendra2-LC3 into autophagosome membranes. In agreement with previous studies of autophagosome dynamics^{26,48}, live cell imaging revealed that a subset of Dendra2-LC3 puncta were highly mobile (Supplemental movie 2). As expected, inhibiting the clearance of autophagosomes via treatment with the lysosomal V-ATPase inhibitor bafilomycin-A1 lead to the accumulation of large bright puncta without an accompanying decrease in diffuse Dendra2-LC3 fluorescence (Supplemental movie 3). Together these data confirm that Dendra2-tagged version of LC3 behaves as expected in modified HEK293T cells⁴⁰, and that these cells can be used to visualize autophagy modulation by a variety of stimuli.

Development and validation of an autophagic flux assay

In these cells, endogenous Dendra2-LC3 could be efficiently photoconverted with minimal toxicity using 4s pulses of 405nm light, producing a strong red signal (Fig. 2A) concurrent with a substantial reduction in green fluorescence. Using time-lapsed

microscopy, we measured fluorescence intensity in both the red (TRITC) and green (GFP) channels at regular intervals over the span of 13.5h. In vehicle-treated cells red fluorescence decayed with a half-life of approximately 7.5h. Treatment with Torin1 significantly accelerated this decay, reducing Dendra2-LC3 half-life ~3-fold to 2.5h. In contrast, bafilomycin-A1 completely stabilized Dendra2-LC3 and blocked the Torin1-induced reduction in Dendra2-LC3 half-life (Fig. 2A, B). Thus, endogenous Dendra2-LC3 flux measured by OPL responds appropriately to bidirectional modulation of autophagy.

To confirm autophagy-dependent degradation of Dendra2-LC3 in modified HEK cells, we asked whether genetic inhibition of autophagy extended Dendra-LC3 half-life. HEK cells were transfected with siRNA targeting the autophagy gene ATG5⁴⁹, achieving a marked reduction in ATG5 levels (Fig. S1). ATG5 knockdown attenuated Torin1's effects on Dendra2-LC3 half-life but had no discernible impact on Dendra2-LC3 turnover in vehicle treated cells (Fig. 2C). These data show that Dendra2-LC3 clearance in response to Torin1 is mediated by autophagy, and also suggest that ATG5 expression levels may be rate limiting only upon autophagy induction.

Consistent with effective photoconversion of Dendra2-LC3, GFP intensity dropped by approximately 40% in pulsed cells, returning to pre-conversion levels within 13h (Fig. 2D). This return to steady-state GFP intensity likely reflects an equilibrium point at which new Dendra-LC3 production is balanced with its turnover. Treatment with Torin1 shifted this balance, not only preventing the return in GFP signal, but also further reducing GFP intensity over time. Application of bafilomycin-A1 (Fig. 2D) or ATG5 knockdown (Fig. 2E) both prevented Torin1-induced reductions in Dendra2-LC3 GFP intensity. Conversely, bafilomycin-A1 treatment led to a supra-physiological increase in GFP signal (Fig. 2D). Thus, while the decay of photoconverted (red) Dendra2-LC3 can be used to accurately measure autophagic flux because it decouples protein turnover and synthesis, time-dependent changes in native (green) Dendra2-LC3 fluorescence mirror those observed in the red channel, and provide complementary estimates of flux.

To confirm that the metabolism of endogenous Dendra2-LC3 reflects autophagic flux, while simultaneously validating the use Dendra2-LC3 cells for identifying new autophagy-modulating strategies, we used the cells to screen an Enzo tool compound library that includes several drugs with purported effects on autophagy (Fig. 3A, Figure 3-source data). These experiments helped gauge the generalizability of the assay beyond the effects of strong autophagy modulators such as Torin1 and bafilomycin-A1,

and also helped determine its ability to identify drugs that impact autophagy through a variety of mechanisms.

Cells were imaged once prior to photoconversion to measure background RFP intensity. As in previous experiments, a 4s pulse of 405nm light was used for photoconversion. Imaging occurred again immediately following photoconversion to determine the maximum RFP signal. All subsequent RFP intensities were normalized to this value after subtraction of the measured background RFP signal. Library drugs were added at 10 μ M via a robotic liquid handler, cells were imaged every 1.5h for 16h, and time-dependent changes in RFP intensity were measured for each condition. Using DMSO and Torin1 as negative and positive controls, respectively, we observed a Z' =.79 at 9h after drug application, demonstrating high sensitivity and reproducibility of the assay.

Many canonical autophagy-modulating drugs demonstrated clear effects on autophagic flux (Fig. 3A), establishing construct validity for the assay. Among compounds that significantly enhanced autophagic flux were the allosteric mTOR inhibitor rapamycin and the dual PI3K/mTOR inhibitors NVP-BEZ235 and PI-103. Torin1 and PI-103 exerted particularly strong effects in line with their action as ATP competitive antagonists^{47,50}. 10-NCP, an AKT inhibitor that we previously identified as a neuronal autophagy-inducing compound^{40,51}, also increased autophagic flux in Dendra2-LC3 HEK cells. Rottlerin, a compound that demonstrated autophagy-enhancing effects via inhibition of mTOR as well as protein kinase C (PKC) δ , greatly accelerated the degradation of Dendra2-LC3^{52,53}.

Bafilomycin-A1, also present in the compound library, registered as a strong inhibitor of Dendra2-LC3 flux (Fig. 3A), providing internal consistency with regards to our initial investigations. Rather than decrease over time, the intensity of photoconverted (red) Dendra2-LC3 in cells treated with bafilomycin-A1 progressively increased, eventually exceeding levels immediately following photoconversion (Fig. 3B). This phenomenon reflects a peculiar imaging property unique to Dendra2-linked proteins that accumulate within large puncta, in the process sequestering diffuse, low-intensity signal within relatively small regions (Fig. S2). Thus, the time-dependent degradation Dendra2-LC3 is inhibited by bafilomycin-A1, and the observed increase in red fluorescence intensity is due to the accumulation of Dendra2-LC3 within large clusters of perinuclear autophagosomes.

The protein translation inhibitor cyclohexamide stabilized Dendra2-LC3 turnover (Fig. 3A,B), in keeping with autophagy inhibition downstream of amino acid accumulation and mTORC1 activation⁵⁴. This is in contrast to what is observed in the green channel, where inhibiting the synthesis of Dendra2-LC3 results in a decrease in GFP fluorescence as expected (Fig. 5-source data). These results highlight the pivotal ability of the assay to decouple autophagy inhibition from new protein synthesis.

Dendra2-LC3 RFP fluorescence was also stabilized by the proteasome inhibitor MG132 and the protease/proteasome inhibitor ALLN (Fig. 3A; Fig. 3-source data), albeit to a far lesser degree than with bafilomycin-A1 or other strong inhibitors. This suggests that while Dendra2-LC3 serves as valid reporter of autophagic flux, it is not degraded exclusively via autophagy. Therefore, to confirm that hits arising from this assay were indeed capable of affecting Dendra2-LC3 turnover via their actions on autophagy, we tested their effects in ATG5-deficient cells. As seen with Torin1 (Fig. 2), ATG5 knockdown attenuated the autophagy-inducing effects of NVP-BEZ235 and rapamycin (Fig. 3C,D), verifying that these drugs stimulate Dendra2-LC3 clearance by enhancing autophagic flux.

Because of the nature of the screen, compounds exhibiting intrinsic fluorescence could result in an artificially high RFP signal, leading to their subsequent misclassification as autophagy inhibitors. To address this possibility we rescreened all hits in unmodified HEK293 cells that do not express Dendra2. We found that 4 out of the 35 tested drugs, including 3 out of the 10 drugs that were identified as inhibitors, exhibited intrinsic fluorescence in the RFP channel (Fig. 3-source data, Fig. S3). For instance, curcumin, a purported autophagy modulator⁵⁵ with known autofluorescent properties⁵⁶ produced a substantial increase in background RFP signal that precluded any estimations of its effects on autophagy in this assay. In contrast, the PKC inhibitor bim-1 had little effect on background fluorescence, but instead accumulated within perinuclear autofluorescent puncta that resembled those observed in cells treated with bafilomycin-A1. These results underscore the importance of counter-screening to exclude intrinsically fluorescent drug properties that can confound or obscure results.

Establishing a high-content screening platform for autophagy modulators

Using Torin1 and bafilomycin-A1, we next evaluated the sensitivity of Dendra2-LC3 cells for detecting small changes in autophagic flux in a dose-dependent manner. We tested the effects of 10 serial dilutions of each drug in both the GFP and RFP

channels. As in previous experiments, baseline GFP and background RFP measurements were acquired prior to photoconversion. GFP measurements were normalized to this baseline value while RFP was normalized to the background-subtracted postconversion intensity. Following drug treatment, we imaged every 30m for 13.5h and analyzed time- and concentration-dependent effects in each channel. We observed a tunable and proportional response to increasing drug concentrations for both Torin1 and bafilomycin-A1 (Fig. 4A). This was perhaps most evident for bafilomycin-A1, where the assay had sufficient resolution to discriminate 2nM changes in concentration (Fig. 4C, E). Notably, the GFP channel was nearly as sensitive as the RFP channel for detecting differences in autophagic flux (Fig. 4D, E). While the GFP fluorescence gradually returned to equilibrium in vehicle treated cells over a 12h span, it continued to drop with Torin1 treatment in a dose-dependent manner (Fig. 4D). Conversely, the GFP intensity quickly surpassed pre-photoconversion levels in cells treated with bafilomycin-A1, and the rate of increase was proportional to the drug dose (Fig. 4E). For Torin1, bafilomycin-A1, rapamycin and NVP-BE235, the dose response relationships for each drug were strikingly similar between channels, producing nearly identical half maximal effective concentration (EC50) and half maximal inhibitory concentration (IC50) values for each compound (Fig. 4F).

These data indicate that both the GFP and RFP channels provide accurate information regarding autophagic flux upon drug addition. Since imaging in the GFP channel does not require photoconversion, experiments take only a fraction of the time that would otherwise be needed to track Dendra-LC3 turnover in the RFP channel. We took advantage of this fact in developing a high-throughput and high-content screening platform in Dendra2-LC3 HEK cells. Cells were imaged in the GFP channel immediately prior to drug addition (GFP_{0H}) and then again 15h later (GFP_{15H}). Autophagy enhancers were defined as those drugs that reduce the GFP_{15H}/GFP_{0H} ratio by more than 3 standard deviations from the mean of vehicle (DMSO), and inhibitors as drugs that increase the GFP_{15H}/GFP_{0H} ratio by more than 3 standard deviations. In 3 replicate 90/10 studies where 90% of wells were treated with DMSO and 10% were treated with Torin1, we observed a 1% false positive rate, no false negatives, and a mean Z'=.52, thus validating this method as a reliable primary screening assay (Fig. 5A, Table 1).

TABLE 1: 90/10 experiments evaluating primary high-throughput screening assay.

	Experiment 1(n=96)	Experiment 2	Experiment 3
Z'	0.51	0.49	0.56
false positives	1	3	4
false negatives	0	0	0
mean DMSO-mean Torin1	0.75	1.46	1.14
DMSO SD/DMSO mean	6.40%	11%	9%

We then devised a layered screening scheme where hits from the primary assay were filtered based on toxicity, then subjected to a counter-screen where Dendra2-LC3 half-life is determined following photoconversion and imaging in the red channel. This organization combines the added throughput of imaging in the green channel with the ability to selectively monitor Dendra2-LC3 degradation in the red channel. Custom scripts were used to exclude toxic compounds based on their effects on cell number—this was particularly important since drugs that cause cells to die might significantly lower GFP intensity and could therefore be misconstrued as false-positives.

We first applied this screening strategy to the Prestwick drug library, a collection of 1,280 off-patent small molecules, 95% of which have gained regulatory approval by the FDA, EMA and other agencies. Eighteen compounds were filtered out due to toxicity; 17 of which would have otherwise been identified as autophagy enhancers due to their ability to significantly reduce GFP intensity. Among the remaining compounds, 129 exhibited significant effects on Dendra2-LC3 levels, with 88 significantly reducing the GFP_{15H}/GFP_{0H} ratio (i.e. enhancing autophagy) and 41 increasing the ratio (i.e. inhibiting autophagy) (Fig. 5B, Fig. 5-source data).

We also assessed the frequency of Dendra2-LC3 puncta—corresponding to autophagosomes—in treated cells, since a change in the number or size of puncta could indicate either autophagy induction or late-stage autophagy inhibition³. In fact, 2 previous high-throughput screens utilized changes in LC3 puncta number to identify autophagy modulators^{57,58}. In native Dendra2-LC3 cells, we observed an increase in Dendra2-LC3 puncta number in response to 38 drugs, but only 13 of these compounds significantly affected autophagic flux in the primary screen. Among these 13, 11 reduced the GFP_{15H}/GFP_{0H} ratio and were counted as enhancers, while 2 compounds acted as inhibitors and increased the GFP_{15H}/GFP_{0H} ratio (Fig. 5B, Fig. 5-source data).

All 129 hits, along with 2 drugs that narrowly missed significance in the primary assay, were then evaluated for their ability to affect the degradation of photoconverted Dendra2-LC3. 17 compounds (13%) significantly modulated autophagic flux; of these, 11 enhanced flux and 6 inhibited it. Counter-screening in unmodified HEK cells uncovered two drugs, diacerin and mitoxanthrone, with intrinsic red fluorescence independent of their effects on autophagy Fig. 5-source data, Fig. S3).

Among notable enhancers were the antiarrhythmic drugs digoxigenin, a metabolite of digoxin, and clofilium tosylate. Ciclopirox olamine, an off-patent anti-fungal agent found in both the Prestwick and Enzo libraries, potently enhanced autophagy.

Similarly, the anthelmintic niclosamide inhibited autophagy to a comparable extent as bafilomycin-A1 in the Enzo compound screen, and was the strongest inhibitor tested in the Prestwick library, demonstrating significant consistency of the assay across different libraries. As seen previously, cyclohexamide modestly stabilized photoconverted (red) Dendra2-LC3, despite lowering green Dendra2-LC3 levels in the primary screen. This ability to exclude compounds that lower LC3 levels due to translation inhibition demonstrates the value of screening in both the GFP and RFP channels. These data, along with the observation that 87% of primary screen “enhancers” failed to stimulate autophagy in counter-screening, suggest that the primary screen is a sensitive but not specific method to identify autophagy enhancers, but the added selectivity of the secondary screen provides a means to successfully filter out false enhancers.

An expanded screen identifies novel autophagy inhibitors

Our data indicate that Dendra2-LC3 cells provide a robust, accurate and precise measure of autophagic flux that can be readily adapted for compound screening. To identify new autophagy modulating drugs, we took advantage of the screening assay we developed to investigate a library of 24,000 drugs spanning considerable chemical diversity curated from the Maybridge library. In our screen of the Prestwick library a large percentage of hits from the primary screen (time-dependent changes in the GFP intensity) failed to show an effect in the secondary assay (time-dependent changes in RFP intensity post-photoconversion). Since we aimed to test ~20 times as many compounds as before, we chose to filter out false positives by repeating the primary assay and retesting hits after exclusion of toxic compounds, before progressing to secondary screening involving clearance of photoconverted Dendra2-LC3. We also confirmed each hit by repeating the secondary screen prior to further filtering based on the solubility and permeability of each hit. Finally, the intrinsic fluorescence of each candidate compound was assessed in unmodified HEK239T cells, and all remaining hits were confirmed using orthogonal flux assays (Fig. 6A).

Similar to the 10% hit rate observed with the Prestwick library, we identified 2160 compounds (9%) as potential autophagy modulators from the primary screen. Of these, 1958 registered as autophagy enhancers, and 202 as inhibitors. Upon repeating the screen after excluding toxic compounds, 232 candidates (1%) remained as hits, underscoring the need to repeat the primary screen. The retest reduced the number of enhancers more than 10-fold while decreasing the number of inhibitors by a factor of 2.4,

demonstrating a predilection towards false-enhancers in the primary screen. Following the secondary screen and retest, 23 compounds remained. Notably, all enhancers identified in the primary screen failed to pass the secondary screens (Fig. 6B, Table 2).

Of the 23 candidate autophagy inhibitors, 7 were excluded based on limited solubility and permeability, and 11 more were discarded upon acquisition of fresh powder from commercial sources. Among the 5 remaining autophagy inhibitors (Fig 6C, Table 2), the top 3 candidates consistently tested among the most potent inhibitors in all assays. Additionally, each compound elicited a dramatic perinuclear accumulation of Dendra2-LC3 in modified HEK293T cells (Fig. 6C), suggestive of a late-stage block in autophagosome maturation.

We next sought to validate the ability of these compounds to inhibit autophagy using an alternative flux assay. To do so we utilized a HeLa cell line stably overexpressing the tandem RFP-GFP-LC3 reporter⁵⁹. In this system, LC3 is fused to an acid-sensitive GFP as well as an acid-insensitive RFP. Upon progression from autophagosome to autolysosome, GFP fluorescence is quenched as the sensor enters an acidic environment. Application of autophagy inhibitors such as bafilomycin-A1 that inhibit lysosomal acidification result in the appearance of GFP(+)/RFP(+) (yellow) autophagosomes (Fig. S4A). All 5 of the newly-identified compounds significantly increased yellow puncta accumulation in RFP-GFP-LC3 HeLa cells, indicative of effective autophagy inhibition. To assess this effect in an automated and unbiased manner, we developed an image analysis pipeline that identifies cytoplasmic puncta and reports the fraction of GFP(+)/RFP(+) puncta (Fig. S4B). Using this pipeline we observed dose-dependent effects for each compound across similar concentration ranges as those observed in Dendra2-LC3 HEK293T cells (Fig. S4C; Fig. S5). After 12h, all compounds save #2 had reached their maximal response (Fig. S4D). Compounds 1,4, and 5 exerted more than half their maximum effect immediately after drug addition. Quinacrine, a previously reported autophagy inhibitor⁶⁰, showed similar kinetics (Fig. S4E), while the response of compound 3 more closely matched the kinetics of bafilomycin-A1.

We next asked whether these compounds exhibited intrinsic fluorescence that might interfere with our assessment of autophagic flux. Compound 4 did not produce red fluorescence at any concentration tested, but we observed dose-dependent red fluorescence for all other compounds (Fig. S5A). Even so, each compound inhibited autophagy in both flux assays at concentrations where intrinsic fluorescence is minimal

(Fig. S5B). In the green channel, all drugs produced significant green fluorescence with the exception of compound 4. Application of compound 3 resulted in the appearance of bright perinuclear puncta that could confound estimates of GFP(+)/RFP(+) puncta in HeLa cells. Indeed, in this assay a lower concentration of compound 3 was needed to exert an effect than in the Dendra2-LC3 flux assay, which should be unaffected by changes in green fluorescence (Fig. S5B).

To exclude any possible contribution of fluorescence in the measurement of flux, we assessed autophagy inhibition using an orthogonal assay that does not rely on fluorescent readouts. Treatment with compounds 1 and 2 resulted in a significant and reproducible accumulation of LC3-II by western blotting, indicative of impaired autophagosome clearance (Fig. 6D,E). Together these data demonstrate that both compounds are indeed autophagy inhibitors; however, the intrinsic fluorescence of compound 2, and to a lesser extent compound 1, can complicate readouts of flux in fluorescence-based autophagy assays.

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393 **TABLE 2:** Summary of Maybridge library screen and orthogonal autophagic flux assays.

Phase	# compounds	%	Enhancer	Inhibitor
Maybridge Library	24000	100	NA	NA
Primary	2160	9	1958	202
Primary confirmation	232	0.97	148	84
Secondary	41	0.17	3	39
Secondary confirmation	23	0.1	0	23
passed med chem filters	16	0.07	0	16
retest reordered drugs	5	0.02	0	5
GFP-RFP-LC3 HeLa	5	0.02	0	5
LC3II western blot	2	0.01	0	2

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Enhancing autophagic flux suppresses toxicity in a primary neuron model of ALS

We previously demonstrated that autophagy induction using a family of small molecules protected against TDP43-mediated toxicity in models of ALS/FTD⁴⁰. Here we sought to test whether any enhancers identified in our compound screens conferred similar neuroprotection. To accomplish this we turned to automated fluorescence microscopy (AFM), a technology capable of individually tracking thousands of cells prospectively over time^{61,62}. Primary rodent spinal and cortical neurons were transfected with Dendra2-LC3, photoconverted with a brief pulse of blue light and imaged by AFM (Fig. 7A). Because the platform measures single-cell intensity within the TRITC channel over time, we are able to calculate a half-life for Dendra2-LC3, corresponding to autophagic flux, within each neuron. Cortical neurons exhibited slightly higher basal rates of autophagy than spinal neurons, with a mean single-cell Dendra2-LC3 half-life of 33.2h compared to 37.1h seen in spinal neurons (Fig. S6A,B, $p=7.1 \times 10^{-4}$, Welch two sample t-test).

We also determined the lifespan of each neuron using custom scripts that assign a time of death for each cell^{62,63}. To assess the relationship between basal rates of autophagy in neurons and their survival, we incorporated Dendra2-LC3 half-life as a continuous variable into a Cox proportional hazards model of neuronal survival⁶⁴ (Fig. 7B). For both cortical and spinal neurons, rapid turnover of Dendra2-LC3 was associated with extended lifespan (Fig. 7C, cortical: $p=3.4 \times 10^{-9}$, spinal $p=1.1 \times 10^{-6}$, Cox hazards analysis). These results indicate that higher rates of basal autophagic flux are associated with prolonged neuronal survival *in vitro*, providing further rationale for the development of autophagy activators for neurodegenerative diseases.

NVP-BEZ235 (dactolisib) was among the most potent autophagy enhancers identified using Dendra2-LC3 HEK cells (Figs. 3,4). In light of our prior data attesting to the benefit of autophagy induction in ALS/FTD models and the proportional relationship between autophagic flux and neuronal survival (Fig. 7C), we predicted that NVP-BEZ235 could prevent neurodegeneration in disease models. We therefore tested this compound in neurons overexpressing TDP43, an RNA binding protein whose accumulation is integrally connected with both ALS and FTD^{65,66}. In prior studies, TDP43 overexpression reproduced characteristic features of disease, including the formation of ubiquitin- and TDP43-positive neuronal inclusions, cytoplasmic TDP43 mislocalization, and neurodegeneration^{40,67,68}. As expected, TDP43 overexpression resulted in an increase in the risk of death in comparison to neurons transfected with EGFP, a control protein.

429 Treatment with NVP-BEZ235 significantly prolonged the survival of neurons
430 overexpressing TDP43 ($p < 0.05$, HR=0.89, Cox hazards analysis), without adversely
431 affecting EGFP-expressing neurons (Fig. 7D, Table 3). These data demonstrate the
432 therapeutic potential of NVP-BEZ235 as an autophagy inducer capable of extending
433 neuronal survival in ALS/FTD models.
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435 **TABLE 3:** Summary of Cox proportional hazards analysis

Comparison	Hazard Ratio	p-value
GFP/DMSO vs. TDP43-GFP/DMSO	1.74	1.02E-18
GFP/DMSO vs. GFP/25nM NVP-BEZ235	1.02	0.67
WT-TDP43-GFP DMSO vs. WT-TDP43-GFP/ 25nM NVPBEZ235	0.89	0.03

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Discussion

In this study we developed a unique human reporter cell line enabling the non-invasive measure of autophagic flux in living cells, without interference from pathway intermediates or potentially toxic pathway inhibitors. Building on previous efforts to isolate autophagic clearance from induction^{69,40,59}, here we created a system with the pivotal capacity to assess *native* autophagic flux, thereby avoiding several basic confounds associated with overexpression of autophagy reporters^{70,71}. This is of particular relevance considering the intersection between autophagy and nutrient/energy sensing⁴², the role of microtubule associated transport in autophagosome maturation⁴⁸, and the crosstalk between autophagy and the ubiquitin proteasome system⁴⁴. Increasing protein dosage can also induce aberrant aggregation of misfolded proteins, and influence the likelihood of protein phase separation^{70,72}. Beyond the potential toxicity associated with these outcomes, overexpression of LC3 and LC3-based reporters is sufficient to produce visible puncta that could be mistaken for autophagosomes.

Through the targeted insertion of Dendra2 into the *MAP1LC3B* locus, we generated reporter cell lines in which Dendra2-LC3 expression is driven by the endogenous *MAP1LC3B* promoter. As such, the baseline fluorescence intensity of non-photoconverted (green) Dendra2-LC3 reflects the steady-state balance between LC3 synthesis and degradation. We took advantage of this relationship to quickly and accurately gauge the effects of compounds that enhance or inhibit autophagy, without the need for photoconversion of Dendra2-LC3. Such an approach can be problematic in overexpression-based systems, but it provided a robust means of estimating autophagic flux on a high-throughput basis in Dendra2-LC3 HEK cells. Future studies employing a full bleach or photoconversion of Dendra2-LC3 could be useful for investigating autophagy regulation at the level of transcription or protein synthesis, and for identifying genetic and/or pharmacologic approaches that induce autophagy at an early stage.

Previous high-throughput screens for autophagy modulators utilized the formation of LC3-positive puncta as the major criterion for autophagy induction^{58,57}. As late stage autophagy inhibition is equally effective as autophagy induction for triggering the accumulation of LC3 puncta, such analyses are unable to discriminate whether a drug truly enhances or blocks flux. For example, the drugs loperamide and astemizole, two drugs found here to enhance flux, were also identified as potential autophagy inducers based on their ability to increase LC3 puncta⁵⁸. However, niclosamide was also labeled as an inducer because of its effect on LC3 puncta, but was in fact a strong

inhibitor of flux in Dendra2-LC3 cells. Further supporting the apparent disconnect between puncta number and autophagic activity, we identified several compounds that increased Dendra2-LC3 puncta without markedly impacting Dendra2-LC3 levels (Fig. 5A). As such, reporters that judge the clearance of autophagy substrates are intrinsically more suited to gauging autophagic flux than are those that focus on pathway intermediates such as autophagosome number. Because autophagy is a dynamic pathway that requires coordinated regulation of several critical steps, increasing autophagy *induction* without an accompanying downstream increase in substrate clearance is likely to be of little therapeutic benefit and may even be maladaptive^{73,74}. Therefore, we placed particular emphasis on enhancing *productive* autophagy and the measurement of autophagic flux through the use of non-invasive reporters.

In comparison to alternative methods for measuring autophagic flux, Dendra2-LC3 cells offer unique advantages. Not only do these cells afford the only means of estimating *native* autophagic flux in living cells, but they also preclude the need for overexpression of LC3 analogues, thereby avoiding many of the pitfalls that plague other approaches. For instance, while the GFP-LC3-RFP-LC3ΔG reporter⁶⁹ is likewise capable of discriminating LC3 synthesis from degradation, measurements of autophagic flux using this probe require relating the degradation rates of two proteins, one of which is an autophagy marker and substrate (GFP-LC3) and another that is untethered within the cytoplasm (RFP-LC3ΔG). Conditions that stabilize RFP-LC3ΔG without accelerating GFP-LC3 clearance could be misinterpreted as increasing flux. Supporting this notion, in a screen using the GFP-LC3-RFP-LC3ΔG reporter, MG132 was identified as an autophagy enhancer⁶⁹, while we found that MG132 instead stabilized Dendra2-LC3. Conversely, compounds such as loperamide that enhance proteolytic clearance via the ubiquitin-proteasome pathway⁷⁵ may be mislabeled as autophagy inhibitors because of their preferential effects on RFP-LC3ΔG.

Several drugs purported to stimulate autophagy—including trehalose^{76,77}, metformin^{78,79,80,81}, and carbamazepine^{82,83,84}—failed to do so in our Dendra2-LC3 cell line. Despite promising results in mouse models of Huntington's disease⁷⁶, ALS⁸⁵, and Parkinson's disease⁸⁶, evidence that trehalose induces autophagy is variable, with some studies claiming that the drug actually inhibits flux^{87,88,69}. Likewise, the ability of carbamazepine to enhance autophagic flux and prevent neurodegeneration was based upon changes in steady-state levels of autophagy intermediates^{83,84}. Our data suggest that the protective effects of these drugs may not be a result of autophagy stimulation.

However, the discrepancy in findings may be due to the high concentration of these drugs used in previous studies^{81,79,80,89,77,82} relative to the 10uM dose used in our screens, as well as species and cell type specific differences.

Our screen of the Maybridge library suggests that uncovering novel autophagy enhancers may be considerably more challenging than inhibitors. Testing larger libraries or incorporating iterative chemical synthesis guided by structure-activity relationships⁵⁰ may be required to effectively identify new autophagy-inducing compounds. Even so, compounds 1 and 2 were repeatedly found to inhibit autophagy in Dendra2-LC3 cells, RFP-GFP-HeLa cells, and via immunoblotting. While these drugs hold potential for the treatment of neoplasms that rely on autophagy for survival, their potency, activity, and bioavailability could be improved through similar means.

Despite the broad therapeutic potential of autophagy modulation, there are no clinically-approved drugs that have been specifically developed to target autophagy⁷. Our work with NVP-BEZ235 presents a potential template to evaluate the efficacy of putative autophagy modulators. We validated the autophagy-enhancing effects of the compound in Dendra2-LC3 HEK293T cells, and further utilized these cells to establish a dose-response relationship that informed subsequent studies demonstrating the drug's therapeutic effects in a model of ALS and FTD. Given previous data indicating the ability of NVP-BEZ235 to suppress amyloid- β induced neurotoxicity⁹⁰, and our results showing its potential in a model of ALS and FTD, this compound may be broadly effective in stimulating autophagy and preventing neuron loss arising from the accumulation of misfolded proteins in Alzheimer's disease as well as ALS, FTD, and related conditions.

Autophagy has alternatively been reported to play protective or detrimental roles in ALS^{91,92}. In our previous work, pharmacological activation of neuronal autophagy suppressed TDP43 mediated toxicity⁴⁰. This is in accordance with the protective effects of rapamycin in a TDP43 mouse model⁸⁴. In contrast, rapamycin administration in SOD1 mice exacerbated muscle degeneration and shortened lifespan⁹³. One study in which autophagy was genetically ablated from motor neurons in SOD1 (G93A) mice provided further insight into the conflicting roles of autophagy in ALS⁷³. Loss of autophagy in motor neurons accelerated disease onset but also prolonged survival. This study suggested that early in disease autophagy plays a critical role in the maintenance of neuromuscular junctions. However, at later stages it promotes disease progression in a non-cell-autonomous manner. Our work further supports a protective role of autophagy enhancement in ameliorating toxicity associated with the accumulation of TDP43. Future

studies may determine whether NVP-BEZ235 confers similar beneficial effects in other disease models involving SOD1 or RNA binding proteins related to TDP43. Even so, neurons respond poorly to most autophagy-inducing stimuli, making them a particularly challenging cell type to target for therapy development^{51,94,95}. Additional investigations involving the introduction of Dendra2 into the *MAP1LC3B* locus of induced pluripotent stem cells may be critical for defining the cell type-specific mechanisms governing autophagy within neurons and other cell types.

While this technology represents a considerable advancement, it is not without limitations. Chief among these is the reliance on reductions in Dendra2-LC3 signal intensity to indicate enhanced flux. For this reason, we developed automated programs to selectively remove drugs with toxic effects that might otherwise be misclassified as false-positives. Since HEK293T cells are actively dividing, compounds that merely inhibit growth rate may also be falsely identified as enhancers when measuring GFP fluorescence, necessitating the use of counter-screens involving the measurement of photoconverted (red) Dendra2-LC3 half-life to eliminate these false-positives from the final pool of candidate compounds. We imaged multiple frames/well and multiple wells to account for autofluorescence artifacts that are common in the red channel, but the use of brighter fluorophores or photoconvertible fluorescent proteins that emit in the far-red wavelengths may avoid these complications⁹⁶.

Additionally, due to the assay's reliance on measuring fluorescent intensity, drugs that exhibit intrinsic fluorescence can confound flux measurements. In unlabeled HEK293T cells, compound 3 from the Maybridge library screen and the drug Bim-1 accumulated within bright fluorescent perinuclear puncta with striking resemblance to puncta observed in bafilomycin-A1 treated Dendra2-LC3 HEK293T cells (Fig. S3). We therefore counter-screened all candidates in unlabeled HEK293T cells and eliminated those exhibiting intrinsic fluorescence at the concentrations tested. We also confirmed autophagy inhibition using a non-imaging based flux assay. Despite the fact that all 5 Maybridge library hits demonstrated autophagy-inhibiting properties in RFP-GFP-LC3 expressing HeLa cells, immunoblotting revealed that only compounds 1 and 2 produced a significant increase in LC3-II levels. Therefore, due to its reliance on measuring fluorescence intensity, the RFP-GFP-LC3 flux assay suffers from the same problem in misidentifying intrinsically fluorescent drugs as inhibitors. Any fluorescence-based autophagy assays are likely to be similarly impacted by intrinsic fluorescence, emphasizing the need to account for these effects in screening efforts.

Ideally, future studies will evaluate autophagic flux using complimentary reporters that provide valuable information on mechanism of action, in addition to magnitude and potency. Such an approach, coupled with a shift toward analyzing the productive autophagic clearance of substrates expressed at endogenous levels, promises to accelerate and facilitate the discovery of novel autophagy-modulating compounds with wide-ranging therapeutic potential.

Methods

Cell Culture

HEK293T Dendra2-LC3 and HeLa RFP-GFP-LC3 cells were cultured in HEK complete media consisting of Dulbecco's Modified Eagle Medium (DMEM) (GIBCO, Waltham, MA, cat #: 11995-065) supplemented with 20% fetal bovine serum, 1% GlutaMAX (GIBCO, cat #: 35050-061) and penicillin-streptomycin. For imaging experiments cells were placed in Neumo media (Cell Guidance Systems, Cambridge, UK, cat #: M07-500) with SOS supplement (Cell Guidance Systems, cat #: M09-50).

CRISPR editing

Single guide RNA pairs (sgRNAs; Table 4) were selected using the CRISPR design tool available at <http://crispr.mit.edu/>. Sense and antisense oligonucleotides encoding each sgRNA (Table 4) were annealed and inserted into the BbsI site of the pX335 vector (Addgene, 42335), according to protocols available on the Addgene website.

596 **TABLE 4:** Sense and antisense oligonucleotides cloned into pX335 plasmid

oligo	sequence
LC3bD2_Forward_Sense	CAC CGT TCG GTG AGT GTC GCC GCG A
LC3bD2_Forward_Antisense	AAA CTC GCG GCG ACA CTC ACC GAA C
LC3bD2_Reverse_Sense	CAC CGT TCT CCG ACG GCA TGG TGC A
LC3bD2_Reverse_Antisense	AAA CTG CAC CAT GCC GTC GGA GAA C

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The homology directed repair (HDR) donor vector was synthesized in the pUCminusMCS backbone by Blue Heron Biotechnology (Bothell, WA). The donor sequence consisted of the Dendra2 ORF flanked by 400bp of homologous sequence upstream and downstream of *MAP1LC3B* start codon. 1.25ug of pX335-Forward-sgRNA, 1.25ug of pX335-Reverse-sgRNA, and 2.5ug of the homology donor were transfected into HEK-293T cells using lipofectamine 2000 (Invitrogen, Waltham, MA), according to the manufacturers suggested protocol. Cells were selected based on fluorescence and passaged to homogeneity.

Western Blotting and antibodies

HEK293T cells were lysed in RIPA buffer (Thermo, Waltham, MA, cat #89900) containing protease inhibitors (Roche, Basel, Switzerland, cat #11836170001).

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612 **TABLE 5:** Antibodies used in this study

Antibody	dilution	manufacturer	Cat #
Rabbit anti-LC3	1:1000	Cell Signaling	2775S
Rabbit anti-ATG5	1:1000	Cell Signaling	129945
Mouse anti-GAPDH	1:1000	Millipore	MAB374
Donkey anti-Mouse iRDye 680RD	1:10,000	LICOR	926-68072
Donkey anti-Rabbit iRDye 800CW	1:10,000	LICOR	926-32213

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siRNA knockdown

HEK293T cells were plated at 60% confluency, then transfected the next day using DharmaFECT 1 Transfection reagent (Dharmacon, LaFayette, CO, T-2001-02) and the following siRNAs from Dharmacon: ON-TARGETplus ATG5 Smartpool siRNA (L-004374-00-0005), ON-TARGETplus MAP1LC3B Smartpool siRNA (L-012846-00-0005) or non-targeting siRNA (D-001810-01-05), at a concentration of either 20nM and 40nM. Cells were imaged and lysates were collected 2d after siRNA transfection.

Primary screen

HEK293T Dendra2-LC3 cells were plated in HEK complete media at 1.1×10^5 cells/mL on ViewPlate 384w plates (Perkin Elmer, Waltham, MA, cat #: 6007460) using a Multidrop Combi (Thermo Scientific, cat #: 5840300), adding 50uL to each well. Approximately 48h later, and immediately prior to imaging, HEK complete media was exchanged with Neumo+SOS media using a Biomek FX^P laboratory automation workstation (Beckman Coulter, Brea, CA). To avoid dislodging cells during the media exchange, 35uL of the HEK media was removed and replaced with 45uL Neumo+SOS. Another 45uL was removed and replaced with Neumo+SOS, effectively diluting out the concentration of HEK complete media to 6.25%. Cells were imaged with an ImageXpress Micro (Molecular Devices, San Jose, CA) equipped with a 20x objective lens. After imaging in the GFP channel (Semrock, Rochester, NY, FITC-3540B-NTE-ZERO filter) to measure baseline fluorescence intensity, compounds were added using a BioMek FX pintool (Beckman Coulter) at a concentration of 10μM, with the exception of the positive control Torin1, which was added at 1μM. Plates were imaged again 15h after drug addition. One image was acquired per well. Images were background subtracted using FIJI (<https://fiji.sc/>) with a rolling ball radius of 150. Mean GFP fluorescence intensity was calculated on a whole well basis. Enhancers are defined as: $[\text{Sample}_{15\text{H GFP/0H GFP}}] < [\text{DMSO}_{15\text{H GFP/0H GFP}} - 3\text{SD}_{\text{DMSO}}]$ and inhibitors as: $[\text{Sample}_{15\text{H GFP/0H GFP}}] > [\text{DMSO}_{15\text{H GFP/0H GFP}} + 3\text{SD}_{\text{DMSO}}]$. Using a custom FIJI script that measured the area of occupied by cells, toxic compounds were identified as those that elicited a reduction in cellular area $\geq 3\text{SD}$, verified by eye.

Secondary screen

Plating of HEK293T Dendra2-LC3 cells and media exchange were performed as in the primary screen, described above. Two sites per well were imaged in the Texas Red

(RFP) (Semrock TxRed-4040C-NTE-ZERO filter) channel prior to photoconversion to establish background fluorescence levels. Photoconversion was accomplished using a 4s DAPI (Semrock Brightline DAPI-5060-NTE-ZERO filter) exposure, and afterwards the cells were immediately imaged once more in the RFP channel. The plate was then removed from the ImageXpress stage and compounds were added to 3 replicate wells via the BioMekFX automation station at a working concentration of 10 μ M. The plate was then returned to the ImageXpress and imaged every 1.5h in a recurring loop for 16h, while maintaining 5% CO₂, humidity and a temperature of 37C. Initial optimization studies demonstrated a maximal Z' = 0.79 within 9h of drug addition, and therefore this time was selected for measuring autophagic flux. We defined enhancers and inhibitors by the following criteria: enhancer, $[\text{Sample}_{9\text{H RFP}}/(\text{Postconversion RFP} - \text{background RFP})] < [\text{DMSO}_{9\text{H RFP}}/(\text{Postconversion RFP} - \text{background RFP}) - 3\text{SD}_{\text{DMSO}}]$; inhibitor, $[\text{Sample}_{9\text{H RFP}}/(\text{Postconversion RFP} - \text{background RFP})] > [\text{DMSO}_{9\text{H RFP}}/(\text{Postconversion RFP} - \text{background RFP}) + 3\text{SD}_{\text{DMSO}}]$. Images with autofluorescent artifacts were excluded and the remaining images were averaged on a per compound basis.

RFP-GFP-LC3 assay

HeLa RFP-GFP-LC3 cells⁵⁹ were plated at 80% confluency in HEK complete media. Prior to imaging, cells were switched to Neumo+SOS media. Cells were imaged at baseline as well as 0, 4, 8, and 12h after drug addition. Images were background subtracted using the rolling ball background subtraction plugin in FIJI, with a radius = 150. LC3 puncta were identified and quantified using CellProfiler 3.0 (<https://cellprofiler.org/>) utilizing a customized pipeline (<https://github.com/BarmadaLab/LC3-puncta>). Briefly, a series of image processing operations are performed to segment a cell into nuclear and cytoplasmic compartments. The contrast between puncta and background is further enhanced to emphasize LC3 puncta in both GFP and RFP images. Object-oriented colocalization then records the number of cytoplasmic red puncta that are also green, representing autophagosomes.

Primary neuron survival assay

Primary neurons were dissected from the cortex of embryonic day 20 rat pups and plated at a density of 1x10⁵ cells/well on a laminin/poly-D-lysine coated 96 well plate⁶⁸⁶³⁹⁷ in Neumo complete media (Cell Guidance Systems M07-500). Transfection and longitudinal microscopy were performed as previously described⁶². Briefly, DIV4

neurons were transfected using lipofectamine 2000 (Invitrogen). Neurons were incubated with Lipofectamine-DNA mixtures for 20 minutes followed by washes with Neurobasal media containing 1 mM kynurenic acid to remove residual lipofectamine. Neurons were then placed in half conditioned media and half fresh Neumo+SOS media. A Eclipse Ti inverted microscope (Nikon, Tokyo, Japan) equipped with PerfectFocus, Semrock GFP and TRITC filters, Lambda XL lamp (Sutter Instruments, Novato, CA) and an Andor Zyla 4.2(+) sCMOS camera (Oxford Instruments, Abingdon, UK) was used to acquire images at 20x. To maintain a temperature of 37C and 5% CO₂ levels, the microscope was encased in a custom-built plexiglass environmental chamber. Automated stage movements, filter turret rotation, and image acquisition were controlled via µManager with original code written in BeanShell. In-house software was used to assign a barcode for each neuron, measure its fluorescent intensity, and register time of death as described previously^{61,62,63,98,99}. For optical pulse labeling experiments a 1.5s pulse of 405nm light was used for photoconversion. For studies relating the rate of Dendra2-LC3 turnover to neuronal survival only cells that lived the entire duration of OPL imaging were included in the survival analysis.

Figure Legends

Figure 1: Creation of a stable cell line serving as a reporter for autophagic flux.

(A) Illustration depicting the use of optical pulse labeling (OPL) to measure autophagic flux. Dendra2 is a photoconvertible protein that upon exposure to 405nm light irreversibly shifts its fluorescence from green to red. Photoconverted Dendra2-LC3 is degraded in lysosomes, resulting in a drop in red fluorescence intensity. The time dependent decay of red signal serves as an estimate of autophagic flux independent of new (green) LC3-Dendra2 synthesis. (B) Schematic for tagging native LC3 using CRISPR/Cas9 genome editing. In HEK293T cells, the Dendra2 ORF was introduced into the *MAP1LC3B* locus upstream of exon 1 creating an N-terminal fusion protein upon translation. (C) Western blot confirming the successful labeling of LC3 with Dendra2. Dendra2-LC3 HEK293T cells were treated with 20nM siRNA targeting LC3 or scrambled siRNA. Lysates were collected after 48h and immunoblotted with an LC3 antibody, demonstrating the Dendra2-LC3 fusion protein running at the expected MW of 43kDa that disappears upon siRNA-mediated knockdown of LC3. GAPDH serves as a loading control. (D) Dendra2-LC3 reporter line imaged in GFP channel 48h after application of siRNA. Scale bar =

100 μ m (E) Dendra2-LC3 cells imaged 6 hours after treatment with vehicle, 1 μ M Torin1, and 20nM Bafilomycin-A1. Scale bar =10 μ m

Figure 2: Time dependent decay of Dendra2-LC3 serves as an accurate measure of autophagic flux.

(A) Dendra2-LC3 HEK293T cells were imaged prior to photoconversion to measure background RFP intensity. Immediately following photoconversion, cells were treated with DMSO, 1 μ M Torin1, or 10nM Bafilomycin-A1 and imaged at the indicated times. Images are pseudocolored to better highlight intensity differences. Scale bar = 50 μ m (B-E) Time dependent changes in photoconverted Dendra2-LC3 fluorescence in the RFP (B, C) and GFP (D, E) channels. Intensity measurements were obtained prior to (dark grey) and following photoconversion (light grey) and normalized. For RFP measurements, the background intensity prior to conversion was set to 0 and the post-conversion value to 1. GFP values are scaled to the pre-conversion intensity. Error bars represent SEM from 3 replicate experiments. (B) Treatment with 1 μ M Torin1 accelerates Dendra2-LC3 decay, reflecting enhanced autophagic degradation of the reporter, while treatment with bafilomycin-A1 stabilizes reporter half-life. (D) Photoconversion results in a 40% drop in GFP intensity. As new Dendra2-LC3 is synthesized, GFP levels return to pre-photoconversion levels over the span of 13.5h. Torin1 blocks the observed return in GFP fluorescence by accelerating flux. Genetic inhibition of autophagy via siRNA-mediated knockdown of ATG5 2d prior attenuates Torin1's effects in both the RFP (C) and GFP (E) channels. For plots B-D * indicates significant difference ($p < 0.05$) using DMSO as reference group with Tukey's multiple comparisons test. # indicates $p < 0.05$ with the scramble control for each drug treatment as the reference group (i.e Scramble siRNA 1 μ M Torin1 vs. ATG5 siRNA 1 μ M Torin1). Superscript number indicates the first time point when significance was achieved.

Figure 3: Small-scale screen in Dendra2-LC3 HEK293T cells confirm assay validity and identify new autophagy modulators.

(A) An unbiased screen of the Enzo autophagy compound library identified several known autophagy-modulating compounds, including enhancers (rapamycin, NVP-BEZ235, AKT inhibitor X) and inhibitors (bafilomycin-A1). All drugs were added at a final concentration of 10 μ M via liquid handler, and autophagic flux estimated by their effects upon clearance of photoconverted (red) Dendra2-LC3 9h after drug addition. (B)

Representative images of red Dendra2-LC3 immediately post-conversion and 9h after drug addition, demonstrating relatively rapid clearance with application of enhancers and marked accumulation of Dendra2-LC3 with inhibitors. Scale bar = 100µm (C) The effect of autophagy enhancers on Dendra2-LC3 half-life is attenuated by siRNA-mediated knockdown of ATG5 2d prior to drug application. Two-Way ANOVA indicated a significant interaction between time and treatment $F(126,300)=15.22$, $p<0.0001$ and significant effects for treatment $F(9,300)=3427$, $p<0.0001$ and time $F(14,300)=957.3$, $p<0.0001$ (D) Similar effects are observed in the green channel. Two-Way ANOVA found a significant interaction between time and treatment $F(126,300)=20.71$, $p<0.0001$. as well as significant main effects for treatment $F(9,300)=721$, $p<0.0001$ and time $F(14,300)=320.8$, $p<0.0001$. For (C) and (D) Error bars represent SEM from 3 replicate experiments. # indicates $p<0.05$ using Tukey's multiple comparisons test with the scramble control for each drug treatment as the reference group (i.e Scramble siRNA Torin1 vs. ATG5 siRNA Torin1). Superscript number indicates the first time point when significance was achieved.

Figure 4: Proportional and bidirectional effects of autophagy modulators highlight assay sensitivity.

(A) Dendra2-LC3 HEK293T cells were treated with increasing concentrations of Torin1 and bafilomycin-A1. Representative images in the GFP (top) and RFP (bottom) channels, pseudocolored to accentuate intensity variations. Scale bar = 100µm (B) Dendra2-LC3 clearance increased in a dose-dependent manner with Torin1 (B), while bafilomycin-A1 resulted in dose-dependent prolongation of Dendra2-LC3 half-life (C). These changes are even more apparent in the GFP channel for both Torin1 (D) and bafilomycin-A1 (E). Error bars represent SEM from 8 replicate wells. For plots B-E * indicates significant difference ($p<0.05$) using DMSO as reference group with Dunnett's multiple comparisons test. Superscript number indicates the first time point when significance was achieved. (F) Dose-response curves for Torin1, NVP-BEZ235, rapamycin, and bafilomycin-A1. For autophagy enhancers, the minimal RFP intensity 7h after drug treatment relative to DMSO was set to 1, and the maximal value set to 0. For inhibitors, the maximum effect represents the maximal RFP intensity within 7h after drug treatment. Dose-response was determined similarly for the GFP channel, utilizing values 14h after drug treatment. Concentration is plotted in nM on a log(2) scale, with ≥ 3

replicate wells for each channel shown as colored dots. EC50 and IC50 values are reported along the x-axis for both RFP and GFP.

Figure 5: Screening the Prestwick library reveals previously unrecognized autophagy-modulating drugs.

(A) “90/10” experiment validating the measurement of Dendra-LC3 in the GFP channels as a primary assay. 320 wells of a 384w plate were treated with DMSO and 32 were treated with 1 μ M Torin1. Plates were imaged in GFP immediately before, and 15h after drug treatment. $Z' = .52 \pm 0.04$ in 3 replicates. (B) Primary screen of the Prestwick drug library. Z-score is calculated as number of SD_{DMSO} greater or less than mean DMSO_{15H/0H}. Significant changes in flux, toxicity, or puncta for each drug are indicated by the color and size of representative dots, according to the key. The magnitude of effect is represented as both % bafilomycin-A1 (left y-axis) and % Torin1 (right y-axis) (C) Time-dependent decay in red (photoconverted) Dendra2-LC3 was used as a secondary screen of the non-toxic candidates emerging from the primary screen. Error bars represent the SEM from 6 images (2 images/well of 3 replicate wells).

Figure 6: High throughput screening identifies novel autophagy inhibitors.

(A) Schematic depicting the screening hierarchy used. (B) The primary screen was performed using the Maybridge 24K library, consisting of 24,000 chemically diverse compounds. For enhancers, changes in Dendra2-LC3 GFP intensity were normalized to Torin1’s effects. Non-toxic compounds that passed the primary screen were filtered by re-testing in the GFP channel, then evaluated in a secondary screen involving calculation of Dendra2-LC3 half-life in the RFP channel. Hits were re-tested in the secondary screen, followed by repeat evaluation in the RFP channel using fresh compound from a different distributor. The color and size of each dot denotes the stage at which individual compounds were eliminated, in accordance with the key. Candidates that passed all filters are shown in black. (C) Representative images of the primary screen, secondary screen, and repeat secondary screen with fresh drug for the 5 putative autophagy inhibitors, ranked based on the magnitude of inhibition measured in the initial secondary screen. Z-scores are reported for each screening phase. Scale bar = 100 μ m. (D) Unmodified HEK293T cells were treated with vehicle or each compound at either 10 μ M or 100 μ M. 9H after treatment lysates were collected and immunoblotted with an LC3 antibody. (E) Quantification of three replicate experiments demonstrating

that at 100 μ M of compounds 1 and 2 significantly inhibit autophagic flux. For each group LC3-II was normalized to the loading control GAPDH and then scaled to 100nM bafilomycin-A1. Error bars represent standard error of the mean. One-way ANOVA showed significant differences between groups ($F=24.28$, $P<0.0001$). * $p<0.01$ compared to DMSO, Dunnet's multiple comparison test.

Figure 7: Autophagic flux predicts neuronal survival and mitigates toxicity in an ALS/FTD disease model.

(A) Mixed rat spinal neurons were transfected on DIV 4 with Dendra2-LC3, imaged 24h later (DAY 1 pre), then pulsed with 405nm light to photoconvert Dendra2-LC3 before imaging repeatedly and longitudinally over several days to track the time-dependent loss of red fluorescence and neuronal survival. Scale bars = 100 μ m in each panel. (B) Experimental outline for determining the relationship between Dendra2-LC3 half-life and neuronal survival. After calculating Dendra2-LC3 half-life for individual neurons (Stage 1), each cell is prospectively tracked using automated fluorescence microscopy to determine its time of death (Stage 2; red number and corresponding arrow). (C) Penalized spline Cox proportional hazards model depicting Dendra2-LC3 half-life (x-axis) vs. relative risk of death (y-axis) for primary rat cortical (black) and spinal (red) neurons, demonstrating a strong linear relationship for both populations (cortical: $p=3.4\times10^{-9}$; spinal $p=1.1\times10^{-6}$, linear Cox proportional hazards). Cortical and spinal neurons with shorter Dendra2-LC3 half-lives, and therefore higher rates of basal autophagy, lived longer. Each hash mark represents an individual neuron, collected from 3 biological and 8 technical replicates each. Grey dotted lines mark 95% confidence intervals. (D) NVP-BEZ235 (25nM NVP) treatment suppresses toxicity in primary rat cortical neurons overexpressing WT-TDP43-GFP. Table 3 summarizes the hazard ratio and statistical significance of each comparison as determined by Cox proportional hazards analysis. N for each group represents total neurons pooled from three replicate experiments. * $p<0.05$, cox proportional hazards analysis.

Figure S1: siRNA knockdown of ATG5

HEK293T cells were transfected with 20nM and 40nM ATG5 siRNA, or 40nM non-targeting siRNA. Lysates were collected two days later and blotted with ATG5 and GAPDH antibodies. Relative to scramble siRNA, 20nM and 40nM ATG5 siRNA

produced a 58% and 64% knockdown of ATG5 protein, respectively. ATG5 levels were normalized to GAPDH.

Figure S2: Aggregation of Dendra2 tagged proteins causes a supraphysiological increase in signal.

Primary Cortical Neurons were co-transfected on DIV4 with GFP (blue) and a Dendra2-tagged fragment of mutant huntingtin (HTT) carrying a pathologic expansion of 74 glutamine residues (Dendra2-HTT-exon1-Q74)¹⁰⁰. On the first day following transfection Dendra2-HTT-exon1-Q74 was photoconverted with a 1s pulse of 405nm light and subsequently imaged via automated fluorescence microscopy. The decay in TRITC (red) intensity was measured every hour as a metric for protein degradation. The TRITC intensity spikes upon aggregate formation, preventing accurate measurements of protein half-life. Lines in the plot correspond to the cell labels in white in the image panels. Red cell labels indicate the time point when a cell has formed an aggregate. Scale bar = 50µm

Figure S3: Drugs identified as autophagy modulators display intrinsic fluorescence.

Unmodified HEK293T cells imaged in the Texas Red channel 9H after drug treatment. Scale bar = 50µm

Figure S4: Novel inhibitors block autophagic flux when assessed using independent measures of autophagy.

(A) HeLa cells expressing a tandem LC3 reporter (RFP-GFP-LC3) were treated with the indicated compounds and imaged 12h later in the RFP (middle panels) and GFP (bottom panels) channels. Composite images are displayed on the top row. Concentrations for each compound (3nM bafilomycin-A1; 1µM Quinacrine; 50µM 245536; and 100µM of 254522, 45808, 234794, and 237373) correspond to the lowest dose resulting in the maximum degree of colocalization between GFP and RFP puncta as calculated in (C). Scale bar: 50µm (B) The percentage of RFP(+)/GFP(+) puncta was determined using CellProfiler¹⁰¹. Images from the GFP (1a) and RFP (1b) channels are uploaded into CellProfiler, nuclei are identified using the GFP channel (2a-green outlines) and a nuclear mask (2b) is generated. Nuclei that do not pass size or intensity thresholds, or are on the edges of an image, are excluded (2a-purple outlines). Cellular boundaries are

identified (3a-purple outlines), and the nuclear mask is subtracted from the newly created cellular mask to produce a cytoplasmic mask (3b). The intensity of cytoplasmic GFP (4a) and RFP (5a) puncta are enhanced, allowing the generation of masks corresponding to puncta in both channels (4b, 5b) and the identification of GFP(+)/RFP(+) autophagosomes (6a, b). Scale bar: 50 μ m. (C) Dose response curves showing the degree of autophagy inhibition with increasing drug concentrations, plotted on a log₂ scale. The y-axis represents the proportion of RFP(+) puncta that are GFP(+), with the maximum and minimum set to 1.0 and 0, respectively. Dots represent individual technical replicates. (D) RFP-GFP-LC3 HeLa cells were imaged, treated with the lowest dose that produced the maximum response as calculated in (C), then imaged again 0, 4, 8 and 12h after drug treatment. (E) Normalized data from (D), depicting the time to maximal effect for each compound. Data in (D, E) represent mean $\pm$ SEM from 3 biological replicates, 8 technical replicates each.

Figure S5: Intrinsic fluorescence confounds flux estimates in a subset of Maybridge library hits

Unmodified HEK293T cells were treated with each compound that registered as an autophagy inhibitor in the screen of the Maybridge 24K library (Fig. 6) Images were acquired in the GFP and TRITC channels 9h after treatment of each drug at either 10 μ M or 80 μ M. Images are pseudocolored to accentuate intensity differences. For compounds 1 and 5, 80 μ M images are presented at higher brightness settings to emphasize low levels of intrinsic fluorescence. All other images are presented at equivalent brightness and contrast. Scale Bar=50 μ m (B) Dose response relationships for each drug in the Dendra-LC3 HEK293T (black), HeLa RFP-GFP-LC3 (blue) autophagic flux assays, as well as intrinsic fluorescence in the GFP (green) and TRITC (red) channels in unmodified HEK cells. In Dendra2-LC3 HEK293T cells the maximum effect represents the greatest increase in RFP intensity 14h after drug treatment. In RFP-GFP-LC3 HeLa cells the effect represents the proportion RFP(+) puncta that are GFP(+), with the maximum and minimum set to 1.0 and 0, respectively. For unlabeled HEK293T cells the maximum fluorescence intensity observed 9h after drug treatment was set to 1 and the lowest to 0. Concentration is plotted in nM on a log(2) scale, with ≥ 3 replicate wells for each concentration shown as colored dots. Dotted vertical lines correspond to 10 μ M, which is the concentration the compounds were initially screened at.

Figure S6: Primary cortical and spinal neurons display subtly different rates of autophagic flux.

(A) Histogram depicting single cell Dendra2-LC3 half-lives in cortical and spinal neurons. Relative to spinal neurons (n=1043), a leftward shift was observed in the Dendra2-LC3 half-life distribution of cortical neurons (n=4058) indicating a greater frequency of cortical neurons exhibiting high rates of autophagic flux compared to spinal neurons ($p=8.2 \times 10^{-4}$, two-sample Kolmogorov-Smirnov (KS) test). **(B)** Consistent with this, the mean single-cell Dendra2-LC3 half-life was slightly reduced in cortical neurons (33.2h) compared to spinal neurons (37.1h) ($p=7.1 \times 10^{-4}$, Welch two sample t-test), indicating higher rates of basal autophagy in cortical neurons.

Supplemental Movie 1: Dendra2-LC3 HEK cells treated with DMSO.

Dendra2-LC3 HEK cells imaged 6 hours after treatment with DMSO. Presented at 3x actual speed. Scale bar = 10 μ m

Supplemental Movie 2: Dendra2-LC3 HEK cells treated with Torin1.

Dendra2-LC3 HEK cells imaged 6 hours after treatment with 1 μ M Torin1. Presented at 3x actual speed. Scale bar = 10 μ m

Supplemental Movie 3: Dendra2-LC3 HEK cells treated with Bafilomycin-A1.

Dendra2-LC3 HEK cells imaged 6 hours after treatment with 20nM Bafilomycin-A1. Presented at 3x actual speed. Scale bar = 10 μ m

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- 1270

Fig. 1

Fig. 2

Fig. 3

Fig. 4

Fig. 5

Fig. 6

A

B

C

COMPOUND	GFP		RFP		Reordered		Z-SCORE				
	0H	15H	0H	6H	0H	6H	PRIMARY		SECONDARY		
							INIT	CONF	INIT	CONF	FRESH
1							3.9	5.4	36.3	28.9	9.5
2							3.3	6.4	19.7	15.2	10.3
3							4.7	15.8	10.3	19.2	20.6
4							3.6	3.9	3.9	3.7	4.5
5							18.1	23.2	3.4	3.6	3.6

D

E

Fig. S1

Fig. S2

Fig. S3

Curcumin

U74389G

BIM-1

Mitoxanthrone

Fig. S4

Fig. S5

Fig. S6

Figure 3-source data

Row Labels	RFP T9/T0	DMSO normalized	Torin1 equivalent	Z-score	status (T9/T0)	Half-Life	status (half-life)	9H RFP unlabelled	intrinsic fluorescence
Torin1	0.18451368	0.250268516	1	16.99580141	enhancer	7.67058648	enhancer	#N/A	#N/A
Rotlerin	0.31034468	0.42094171	0.772354239	13.12677926	enhancer	9.60509818	enhancer	0.797956473	NO
PI-103	0.323565	0.438873329	0.74843685	12.72028408	enhancer	8.05049221	enhancer	0.716262471	NO
DISULFIRAM	0.48471352	0.657450091	0.456896791	7.765327117	enhancer	10.6878055	enhancer	0.752091873	NO
GERI-BP002A	0.52314922	0.709583059	0.387361272	6.583515251	enhancer	11.4711285	non-significant	0.677914507	NO
Penitrem A	0.52607752	0.713554907	0.382063577	6.493476682	enhancer	10.5485005	enhancer	0.718935518	NO
CICLOPIROX OLAMINE	0.5327825	0.722649326	0.369933342	6.287313614	enhancer	9.7168813	enhancer	0.709758632	NO
Forskolin	0.55416044	0.751645695	0.33125767	5.62989858	enhancer	13.2529446	non-significant	0.748086874	NO
BIX01294 (hydrochloride hydrate)	0.55768925	0.756432066	0.324873557	5.521486456	enhancer	10.7263467	enhancer	1.171951492	YES
NVP-BE2235	0.55997579	0.759533446	0.320736902	5.451180696	enhancer	10.5383637	enhancer	0.697428956	NO
Akt Inhibitor X	0.57102512	0.774520418	0.300747116	5.111438266	enhancer	11.0935641	enhancer	0.695030009	NO
Rapamycin	0.57405931	0.778635893	0.295257851	5.018143808	enhancer	11.2950243	enhancer	0.701176479	NO
LOPERAMIDE HYDROCHLORIDE	0.5814055	0.788600034	0.281967571	4.792264848	enhancer	10.9771341	enhancer	0.708235618	NO
3-AMINOBENZAMIDE	0.58168351	0.78897711	0.281464624	4.783716855	enhancer	11.4498773	non-significant	0.760392802	NO
CAPTOPRIL	0.58606178	0.794915658	0.273543723	4.6490948	enhancer	13.4935297	non-significant	0.736081605	NO
Salmeterol	0.59690097	0.809617597	0.253934118	4.315813847	enhancer	12.9062832	non-significant	0.829920965	NO
Etoposide	0.59823237	0.811423461	0.251525437	4.274876374	enhancer	11.3927922	non-significant	0.822720117	NO
Rho Kinase Inhibitor III, Rockout	0.60285122	0.817688328	0.243169289	4.132856948	enhancer	12.4600793	non-significant	0.719856483	NO
Tenovin-6	0.6149372	0.834081369	0.221304073	3.76124008	enhancer	10.9745764	enhancer	0.739717126	NO
SP 600125	0.61560171	0.83498269	0.220101881	3.740807856	enhancer	13.9317686	non-significant	0.648998264	NO
Cumene hydroperoxide	0.61677262	0.836570869	0.217983551	3.704805148	enhancer	11.2409634	enhancer	0.749216947	NO
NSC 56817	0.62234912	0.844134667	0.207894784	3.533340001	enhancer	11.8493846	non-significant	0.741542122	NO
Piceatannol	0.63003189	0.854555326	0.193995686	3.297112151	enhancer	12.5709004	non-significant	0.777472288	NO
Apicidin	0.63325373	0.858925335	0.188166922	3.198047633	enhancer	15.0575098	non-significant	0.772912482	NO
DE09	0.63514326	0.861488237	0.184748495	3.139948726	enhancer	12.7414421	non-significant	0.869500013	NO
MCCSL-JR-1-012	0.64088565	0.869277029	0.17435972	2.963383168	non-significant	14.0174064	non-significant	#N/A	#N/A
PromethazineA?HCl	0.64276142	0.871821268	0.170966185	2.905707332	non-significant	13.7611322	non-significant	#N/A	#N/A
RESVERATROL	0.64341426	0.872706748	0.169785123	2.885634229	non-significant	12.9221536	non-significant	#N/A	#N/A
TEMPO	0.64413424	0.873683309	0.168482574	2.863496377	non-significant	13.614936	non-significant	#N/A	#N/A
Tunicamycin	0.6456152	0.875688382	0.165808187	2.818043023	non-significant	12.0358666	non-significant	#N/A	#N/A
Carvedilol	0.64691527	0.87745541	0.163451305	2.777985928	non-significant	13.237755	non-significant	#N/A	#N/A
HC Toxin	0.64870611	0.879884452	0.160211423	2.722921527	non-significant	13.8631547	non-significant	#N/A	#N/A
BAICALEIN	0.65130539	0.883410037	0.155508959	2.642999383	non-significant	12.4019022	non-significant	#N/A	#N/A
ABC294640?HCl	0.65332468	0.886148939	0.151855782	2.580910707	non-significant	12.909977	non-significant	#N/A	#N/A
CAY10433	0.65389278	0.88691949	0.150828013	2.563442952	non-significant	12.938385	non-significant	#N/A	#N/A
UNC0224	0.65591371	0.889660605	0.147171884	2.501304107	non-significant	14.5576992	non-significant	#N/A	#N/A
UNC0638	0.65621399	0.890067899	0.14662863	2.492071084	non-significant	12.827412	non-significant	#N/A	#N/A
ESCULETIN	0.65813824	0.892677894	0.14314739	2.43290462	non-significant	14.9047114	non-significant	#N/A	#N/A
3230-2939	0.66092334	0.89645552	0.138108752	2.347268926	non-significant	13.1212868	non-significant	#N/A	#N/A
6164173	0.66255535	0.89866912	0.135156228	2.297088417	non-significant	12.8188547	non-significant	#N/A	#N/A
Ibuprofen	0.66472477	0.901611654	0.131231445	2.230383584	non-significant	12.7006296	non-significant	#N/A	#N/A
17-AAG	0.66519393	0.90224801	0.130382666	2.215957899	non-significant	16.4447973	non-significant	#N/A	#N/A
PROBUCOL	0.66521051	0.902270495	0.130352675	2.21544818	non-significant	13.5699457	non-significant	#N/A	#N/A
NSC 326231	0.66724978	0.905036501	0.126663346	2.152745082	non-significant	12.5508278	non-significant	#N/A	#N/A
2-PCPA (hydrochloride)	0.66751795	0.905400239	0.12617819	2.144499451	non-significant	13.0812726	non-significant	#N/A	#N/A
Garcinol	0.66770357	0.905652009	0.125842376	2.138792029	non-significant	12.5616332	non-significant	#N/A	#N/A
ROTENONE	0.66787783	0.905888368	0.125527117	2.133433947	non-significant	18.4946894	inhibitor	#N/A	#N/A
PD-98059	0.66906334	0.907496353	0.123382369	2.09698225	non-significant	14.2859897	non-significant	#N/A	#N/A
Selenomethionine	0.66909226	0.907535581	0.123300447	2.096092987	non-significant	12.9149126	non-significant	#N/A	#N/A
Rosmarinic acid	0.66920591	0.907689738	0.123124431	2.092598378	non-significant	14.6654539	non-significant	#N/A	#N/A
Sorafenib tosylate	0.66960661	0.908233228	0.122399517	2.080277891	non-significant	21.4134279	inhibitor	#N/A	#N/A
Imiquimod	0.67094631	0.910050353	0.119975817	2.039085163	non-significant	12.9047114	non-significant	#N/A	#N/A
NSC 407286	0.67099712	0.910119266	0.1198839	2.037522958	non-significant	13.288539	non-significant	#N/A	#N/A
TAMOXIFEN CITRATE	0.67325743	0.913185079	0.115794685	1.968023466	non-significant	12.2535968	non-significant	#N/A	#N/A
C2-dihydroceramide	0.67364093	0.913705257	0.115100866	1.956231453	non-significant	13.3055546	non-significant	#N/A	#N/A
PAEONOL	0.67390289	0.914060563	0.114626954	1.94817695	non-significant	12.672234	non-significant	#N/A	#N/A
1,2-Dithiole-3- thione	0.67462114	0.915034783	0.113327529	1.926092182	non-significant	13.2754268	non-significant	#N/A	#N/A
UB3836EA-2?HCl	0.67472586	0.915176821	0.113138078	1.922872309	non-significant	13.8966008	non-significant	#N/A	#N/A
4112-3315	0.67722351	0.918564556	0.108619481	1.846075122	non-significant	13.2941244	non-significant	#N/A	#N/A
FK-866	0.67891105	0.92085347	0.105566501	1.794187291	non-significant	13.8154129	non-significant	#N/A	#N/A
AMIODARONE HYDROCHLORIDE	0.67954766	0.921716957	0.104414773	1.774612748	non-significant	13.3344818	non-significant	#N/A	#N/A
TRIFLUOPERAZINE HYDROCHLORIDE	0.68246093	0.92566842	0.099144268	1.685036298	non-significant	12.3804936	non-significant	#N/A	#N/A
DCHA	0.68295555	0.926339302	0.098249439	1.669827962	non-significant	14.9835837	non-significant	#N/A	#N/A
7-Ketocholesterol	0.68360324	0.927217812	0.097077674	1.649912861	non-significant	13.527055	non-significant	#N/A	#N/A
Rolipram	0.68385337	0.927557076	0.096625159	1.642222006	non-significant	12.5418827	non-significant	#N/A	#N/A
TTFA	0.68435442	0.928236686	0.095718688	1.626815816	non-significant	13.8616193	non-significant	#N/A	#N/A
Trehalose	0.68465151	0.928639656	0.095181202	1.617680815	non-significant	13.4904015	non-significant	#N/A	#N/A
Thiourea	0.6856344	0.929972821	0.093403012	1.587459049	non-significant	13.3179702	non-significant	#N/A	#N/A
AlCAR	0.6856387	0.929978654	0.093395232	1.587326824	non-significant	13.5079836	non-significant	#N/A	#N/A
Sodium Butyrate	0.68682089	0.931582134	0.091256493	1.550977228	non-significant	12.9650352	non-significant	#N/A	#N/A
Anacardic Acid	0.68702815	0.931863258	0.090881526	1.544604376	non-significant	13.0864165	non-significant	#N/A	#N/A
AS-605240	0.68745431	0.93244129	0.090110542	1.531500871	non-significant	12.8567112	non-significant	#N/A	#N/A
Z36	0.6880273	0.933218471	0.089073929	1.513882806	non-significant	12.0544662	non-significant	#N/A	#N/A
DTT	0.68908666	0.934655626	0.087157036	1.481303669	non-significant	14.2507243	non-significant	#N/A	#N/A
GLUTATHIONE	0.68911867	0.93469877	0.08709949	1.480325637	non-significant	13.8548174	non-significant	#N/A	#N/A
Phenidone	0.68975754	0.935565309	0.08594369	1.460681895	non-significant	13.4284451	non-significant	#N/A	#N/A
Tanshinone IIA	0.69043782	0.936488021	0.084712968	1.439764774	non-significant	17.4146317	inhibitor	#N/A	#N/A
Ambroxol	0.69100905	0.937262825	0.083679525	1.422200592	non-significant	13.9069992	non-significant	#N/A	#N/A
SU11652	0.69151048	0.937942945	0.082772374	1.406782826	non-significant	48.1363782	inhibitor	#N/A	#N/A
Glucosamine HCl	0.69337356	0.940469964	0.079401809	1.349497382	non-significant	13.2917278	non-significant	#N/A	#N/A
CAFFEIC ACID	0.69455608	0.942073893	0.07726247	1.313137603	non-significant	13.501902	non-significant	#N/A	#N/A
2',5'-Dideoxyadenosine	0.69455661	0.942074617	0.077261505	1.313121203	non-significant	13.1340798	non-significant	#N/A	#N/A
PMSF	0.69489604	0.942535005	0.076647435	1.30268458	non-significant	13.4715604	non-significant	#N/A	#N/A
BHT	0.69497004	0.94263538	0.076513553	1.300409159	non-significant	13.2234212	non-significant	#N/A	#N/A

Quinine HCl?2H2O	0.69512952	0.9428517	0.076225024	1.295505372	non-significant	13.0277028	non-significant	#N/A	#N/A
EX-527	0.69546196	0.943302609	0.075623596	1.285283626	non-significant	14.076725	non-significant	#N/A	#N/A
MS-275	0.69650748	0.944720718	0.073732107	1.253136243	non-significant	13.7133919	non-significant	#N/A	#N/A
Deoxycholate?Na	0.69776753	0.946429811	0.0714525	1.214392494	non-significant	13.3777872	non-significant	#N/A	#N/A
UNCO321 (trifluoroacetate salt)	0.69812082	0.946908997	0.070813357	1.203529745	non-significant	12.7081402	non-significant	#N/A	#N/A
1-Naphthoic Acid	0.69833878	0.947204633	0.070419035	1.196827929	non-significant	13.5562773	non-significant	#N/A	#N/A
CPD000466364_IDEBENONE	0.69848838	0.947407546	0.070148386	1.19222804	non-significant	15.7112306	non-significant	#N/A	#N/A
6-Gingerol	0.69853882	0.947475958	0.070057137	1.190677192	non-significant	13.3581181	non-significant	#N/A	#N/A
Sinefungin	0.69931674	0.948531109	0.068649766	1.166757788	non-significant	13.5767857	non-significant	#N/A	#N/A
HNHA	0.70005412	0.949531265	0.067315747	1.144085075	non-significant	13.0524942	non-significant	#N/A	#N/A
Ascorbic acid	0.70038397	0.949978672	0.066718991	1.133942721	non-significant	13.8484586	non-significant	#N/A	#N/A
JGB1741	0.70044766	0.950065049	0.06660378	1.13198461	non-significant	12.8790914	non-significant	#N/A	#N/A
Protocatechuic acid	0.70109301	0.950940384	0.065436249	1.112141493	non-significant	13.7039536	non-significant	#N/A	#N/A
NSC 18804	0.70179442	0.95189175	0.064167306	1.09057479	non-significant	12.1779246	non-significant	#N/A	#N/A
Diludin	0.70224595	0.952504193	0.063350424	1.07669122	non-significant	14.0624813	non-significant	#N/A	#N/A
CPD000449326_4-Thiazolidinecarboxylic acid	0.70361788	0.954365045	0.060868398	1.034507209	non-significant	13.5338642	non-significant	#N/A	#N/A
CPD000466276_1H-Imidazol-2-amine, N-oxide	0.70437383	0.955390382	0.059500793	1.011263661	non-significant	13.2004545	non-significant	#N/A	#N/A
D-1?-Tocopherylquinone	0.70461042	0.955711287	0.059072766	1.003989002	non-significant	14.383259	non-significant	#N/A	#N/A
MINOXIDIL	0.7047903	0.955955268	0.058747343	0.998458167	non-significant	13.8902216	non-significant	#N/A	#N/A
NSC 170984	0.70560369	0.957058535	0.057275793	0.973447997	non-significant	13.7215421	non-significant	#N/A	#N/A
EHNA?HCl	0.70584303	0.957383161	0.056842803	0.966088991	non-significant	13.7230966	non-significant	#N/A	#N/A
CPD00058212_NICOTINAMIDE	0.70624825	0.957932787	0.056109706	0.953629421	non-significant	13.3101562	non-significant	#N/A	#N/A
Canthaxanthin	0.70671491	0.958565759	0.05526544	0.939280451	non-significant	12.6254161	non-significant	#N/A	#N/A
L-Ergothioneine	0.7070655	0.959041276	0.054631191	0.928500879	non-significant	13.5619043	non-significant	#N/A	#N/A
2-Deoxyglucose	0.70711867	0.959113397	0.054534996	0.926865957	non-significant	13.8085296	non-significant	#N/A	#N/A
NSC 4972	0.70775561	0.95997732	0.053382685	0.907281517	non-significant	14.6920467	non-significant	#N/A	#N/A
Licocalcone A	0.70838406	0.960829732	0.05245729	0.887958036	non-significant	15.4279024	non-significant	#N/A	#N/A
Bakuchiol	0.70908538	0.961780983	0.05097694	0.866393945	non-significant	14.106679	non-significant	#N/A	#N/A
I-BET	0.70962659	0.962515062	0.049997817	0.849752976	non-significant	14.5300526	non-significant	#N/A	#N/A
PHENETHYL CAFFEATE (CAPE)	0.71007093	0.963117745	0.049193952	0.836090639	non-significant	13.3447358	non-significant	#N/A	#N/A
Diubutylr?l cAMP?Na	0.71048586	0.963680545	0.048443283	0.823332424	non-significant	15.1235361	non-significant	#N/A	#N/A
Cl-Amidine	0.71214363	0.965929094	0.045444145	0.77235966	non-significant	13.3279646	non-significant	#N/A	#N/A
EBSELEN	0.71257927	0.966519982	0.044656012	0.758964719	non-significant	14.1914293	non-significant	#N/A	#N/A
RETINYL PALMITATE	0.71300411	0.967096228	0.043887408	0.745901679	non-significant	14.4117909	non-significant	#N/A	#N/A
Tetramethylpyrazine	0.71410563	0.968590295	0.041894406	0.7120324	non-significant	13.6924222	non-significant	#N/A	#N/A
DMOG	0.71455333	0.969197541	0.041084655	0.698266638	non-significant	13.8074478	non-significant	#N/A	#N/A
EUGENOL	0.71457078	0.969221201	0.041053097	0.697730281	non-significant	14.4600912	non-significant	#N/A	#N/A
Suramin (sodium salt)	0.71458816	0.969244774	0.041021655	0.6971959	non-significant	14.4920601	non-significant	#N/A	#N/A
Pimelic Diphenylamide 106	0.71477383	0.969496613	0.040685749	0.691486904	non-significant	12.9250407	non-significant	#N/A	#N/A
THIOTIC ACID	0.71490771	0.969678204	0.040443542	0.6873704	non-significant	12.9765694	non-significant	#N/A	#N/A
CAPSAICIN	0.71511771	0.969963046	0.040063616	0.680913263	non-significant	15.8027153	non-significant	#N/A	#N/A
F-Amidine (trifluoroacetate salt)	0.71701416	0.972535325	0.036632682	0.622601781	non-significant	14.3585486	non-significant	#N/A	#N/A
MELATONIN	0.71899951	0.975228202	0.033040893	0.561556463	non-significant	13.296126	non-significant	#N/A	#N/A
Tenovin-1	0.72013119	0.976763166	0.030993542	0.526760085	non-significant	14.1870485	non-significant	#N/A	#N/A
S-Adenosylhomocysteine	0.72086133	0.97775351	0.02967261	0.504309782	non-significant	13.4924455	non-significant	#N/A	#N/A
D-1?-Tocopherol	0.72095143	0.977875719	0.029509606	0.501539407	non-significant	14.2380186	non-significant	#N/A	#N/A
Anethole trithione	0.72131544	0.978369444	0.02885107	0.490347061	non-significant	13.2788198	non-significant	#N/A	#N/A
Apigenin	0.72192713	0.979199125	0.027744444	0.471538886	non-significant	15.9947006	non-significant	#N/A	#N/A
CPD000466338_TEMOZOLOMIDE	0.72256178	0.980059946	0.026596261	0.452024769	non-significant	13.7405808	non-significant	#N/A	#N/A
Tocopherol succinate	0.72350181	0.981334968	0.024895623	0.423121064	non-significant	15.1103487	non-significant	#N/A	#N/A
(?)-Neplanocin A	0.72369287	0.981594114	0.024549971	0.417246425	non-significant	15.0920961	non-significant	#N/A	#N/A
8-CPT-cAMP?Na	0.72591605	0.984609572	0.02052792	0.348888457	non-significant	13.717633	non-significant	#N/A	#N/A
CARBAMAZEPINE	0.72662666	0.985573416	0.019242334	0.327038886	non-significant	15.4322247	non-significant	#N/A	#N/A
CAY10603	0.72810166	0.987574069	0.016573841	0.281685718	non-significant	20.7188449	inhibitor	#N/A	#N/A
NITRENDIPINE	0.72835712	0.987920558	0.016111691	0.273831108	non-significant	13.7014868	non-significant	#N/A	#N/A
Niguldipine	0.7285959	0.988244441	0.015679693	0.266488945	non-significant	14.7454201	non-significant	#N/A	#N/A
2',3',5'-triacetyl-5-Azacytidine	0.7294329	0.989379719	0.014165446	0.240753114	non-significant	13.2754286	non-significant	#N/A	#N/A
TOLAZAMIDE	0.73001594	0.990170538	0.013110643	0.222825878	non-significant	13.586633	non-significant	#N/A	#N/A
Splitomicin	0.73179126	0.992578516	0.009898855	0.16823898	non-significant	13.9822229	non-significant	#N/A	#N/A
Rilmenidine	0.73293856	0.994134687	0.007823218	0.132961865	non-significant	14.1453568	non-significant	#N/A	#N/A
ISOLQUIRITIGENIN	0.73322013	0.994516591	0.00731383	0.124304402	non-significant	13.6893361	non-significant	#N/A	#N/A
Ferulic acid ethylester	0.73377835	0.995273748	0.006303926	0.107140274	non-significant	14.1609667	non-significant	#N/A	#N/A
NDGA	0.735594	0.997736439	0.003019162	0.051313081	non-significant	13.2951775	non-significant	#N/A	#N/A
3-METHYL-1-PHENYL-2-PYRAZOLIN-5-ONE	0.73622825	0.998596717	0.001871714	0.031811284	non-significant	13.1101855	non-significant	#N/A	#N/A
SB-202190	0.73657807	0.999071204	0.001238839	0.021055057	non-significant	15.0528509	non-significant	#N/A	#N/A
DMSO	0.73726284	1	0 NA	NA	NA	14.1185499	non-significant	0.763576208	NO
L-690,330	0.73753204	1.000365131	-0.000487015	-0.008277218	non-significant	16.1993273	non-significant	#N/A	#N/A
Dacinosat	0.73842541	1.001576868	-0.002103244	-0.035746311	non-significant	17.1974639	inhibitor	#N/A	#N/A
Zebularine	0.7389284	1.002259108	-0.003013223	-0.051212145	non-significant	14.2841895	non-significant	#N/A	#N/A
CPD000469175_IMATINIB MESYLATE	0.73894809	1.002285817	-0.003048847	-0.051817603	non-significant	17.3947537	inhibitor	#N/A	#N/A
NSC 62794	0.74018775	1.003967255	-0.005291567	-0.089934429	non-significant	14.6048426	non-significant	#N/A	#N/A
Trolox	0.74019388	1.003975567	-0.005302655	-0.09012287	non-significant	14.0792243	non-significant	#N/A	#N/A
EPIGALLOCATHECHIN-3-MONOGALLATE	0.74191426	1.006309032	-0.008415056	-0.143020615	non-significant	15.1981258	non-significant	#N/A	#N/A
VERAPAMIL HYDROCHLORIDE	0.74216863	1.006654052	-0.008875247	-0.150841936	non-significant	14.7491465	non-significant	#N/A	#N/A
TEMPOL	0.74227701	1.006801056	-0.009071323	-0.154174404	non-significant	13.7859052	non-significant	#N/A	#N/A
Suramin?6Na	0.74418851	1.009393755	-0.012529492	-0.212948762	non-significant	15.017659	non-significant	#N/A	#N/A
HBEDA?HCl?H2O	0.74451277	1.009833578	-0.013116133	-0.22291919	non-significant	13.811366	non-significant	#N/A	#N/A
PX105684	0.74573859	1.011496244	-0.01533815	-0.260610476	non-significant	17.1309648	inhibitor	#N/A	#N/A
2,4-DPD	0.74745393	1.013822872	-0.018437097	-0.313353242	non-significant	13.6702338	non-significant	#N/A	#N/A
VALPROATE SODIUM	0.7477982	1.014289829	-0.019059929	-0.323938772	non-significant	13.1617751	non-significant	#N/A	#N/A
5147472	0.74790419	1.014433594	-0.019251685	-0.327197814	non-significant	14.7554661	non-significant	#N/A	#N/A
Metformin?HCl	0.74813514	1.014746845	-0.019669502	-0.334298943	non-significant	12.6720438	non-significant	#N/A	#N/A
THIOCTIC ACID	0.74905018	1.015987973	-0.021324932	-0.362434301	non-significant	15.0661687	non-significant	#N/A	#N/A
Chidamide	0.74934781	1.016391668	-0.021863385	-0.371585745	non-significant	16.529833	non-significant	#N/A	#N/A
CDC	0.75359838	1.022157015	-0.029553267	-0.502281453	non-significant	14.9593311	non-significant	#N/A	#N/A

3-Deazaneplanocin A	0.75381872	1.022455865	-0.029951876	-0.509056142	non-significant	15.300576	non-significant	#N/A	#N/A
Scriptaid	0.75599961	1.025413962	-0.033897419	-0.576113804	non-significant	20.230442	inhibitor	#N/A	#N/A
GENISTEIN	0.7570347	1.026817921	-0.035770034	-0.607940391	non-significant	13.7567165	non-significant	#N/A	#N/A
Seratrodist	0.75850285	1.028809278	-0.038426129	-0.65308285	non-significant	12.7799624	non-significant	#N/A	#N/A
4-iodo-SAHA	0.76008636	1.030957099	-0.041290916	-0.7017722	non-significant	16.2404314	non-significant	#N/A	#N/A
NSC 401077	0.76032395	1.031279361	-0.041720752	-0.70907761	non-significant	13.008035	non-significant	#N/A	#N/A
NVP-LBH589	0.76161097	1.033025026	-0.044049139	-0.74865042	non-significant	18.7133963	inhibitor	#N/A	#N/A
CAFFEINE	0.76509427	1.037749664	-0.050350912	-0.855754107	non-significant	14.1991437	non-significant	#N/A	#N/A
Trichostatin A	0.76618214	1.039225222	-0.052319027	-0.889203797	non-significant	18.5364429	inhibitor	#N/A	#N/A
HYDROQUINONE	0.76692274	1.040229743	-0.053658868	-0.911975461	non-significant	13.0605182	non-significant	#N/A	#N/A
NIMODIPINE	0.76823131	1.042004648	-0.056026256	-0.952211115	non-significant	14.0669637	non-significant	#N/A	#N/A
LY 294002	0.77558726	1.051982032	-0.069334199	-1.178390271	non-significant	13.7020925	non-significant	#N/A	#N/A
Hydroxychloroquine	0.77717596	1.054136897	-0.072208381	-1.227239308	non-significant	14.0450336	non-significant	#N/A	#N/A
ETYA	0.77964957	1.057492019	-0.076683479	-1.303297181	non-significant	15.0260388	non-significant	#N/A	#N/A
CBHA	0.79275085	1.075262178	-0.100385511	-1.706132206	non-significant	20.8654275	inhibitor	#N/A	#N/A
ETHOXYQUIN	0.79663177	1.080526141	-0.107406642	-1.825461958	non-significant	14.5066695	non-significant	#N/A	#N/A
NSC 26326	0.80297134	1.089124925	-0.118875794	-2.020389384	non-significant	15.1105917	non-significant	#N/A	#N/A
SAHA	0.8036859	1.090094133	-0.120168534	-2.042360538	non-significant	20.3880498	inhibitor	#N/A	#N/A
n-Octyl caffeate	0.80578661	1.092943473	-0.123969014	-2.106952745	non-significant	16.9668871	inhibitor	#N/A	#N/A
AGK2	0.81345014	1.103338048	-0.137833412	-2.342589295	non-significant	16.3257095	non-significant	#N/A	#N/A
CYCLOHEXIMIDE	0.84299465	1.143411287	-0.191283533	-3.251016934	inhibitor	18.8695629	inhibitor	0.826837469	NO
U74389G maleate	0.86352956	1.171264187	-0.228434033	-3.88241946	inhibitor	16.4015569	non-significant	0.926349295	YES
Tubastatin A (trifluoroacetate salt)	0.87325607	1.184456911	-0.246030633	-4.181487775	inhibitor	20.3912902	inhibitor	0.798336888	NO
ALLN	0.88442532	1.199606531	-0.26623736	-4.524917304	inhibitor	28.3709302	inhibitor	0.857586629	NO
MG132	0.91965387	1.247389418	-0.329970694	-5.608116384	inhibitor	49.5576448	inhibitor	0.789265	NO
Curcumin	0.97726982	1.32553788	-0.434205961	-7.379678291	inhibitor	16.6637492	non-significant	2.697478274	YES
NICLOSAMIDE	1.10130446	1.493774533	-0.658601837	-11.19346604	inhibitor	-49.302226	inhibitor	0.675069842	NO
Bafilomycin A1	1.21508581	1.648103969	-0.864448116	-14.69198851	inhibitor	106.214287	inhibitor	0.865698976	NO
CPD000469152_TERBINAFINE HCl	1.42060712	1.926866562	-1.236264691	-21.01130918	inhibitor	-20.11502	inhibitor	0.759322914	NO
GF 109203X	3.41064939	4.62609697	-4.836527537	-82.20066154	inhibitor	-40.013818	inhibitor	10.40788771	YES

Figure 5-source data

Compound	Primary GFP	Primary Group	Primary Z-score	RFP_9H	Secondary Z_score	Standard Error	Secondary Group	plate	rank	Label	RFP 9H unlabelled	intrinsic fluorescence
Torin1	#N/A	enhancer puncta	#N/A	0.16264	12.77375092	0.004073812	enhancer	plate 1	1	Torin1	#N/A	#N/A
Digoxigenin	0.25636422	enhancer	3.495227252	0.47716	10.46955572	0.010182837	enhancer	plate 2	2	Digoxigenin	#N/A	NO
CICLOPROX OLAMINE	0.41788536	non-significant	2.96994629	0.53314	4.38673894	0.008868595	enhancer	plate 1	3	Cicloprox Olamine	0.826837469	NO
Lanatoside C	0.41026219	enhancer	5.593446654	0.53402	7.475024566	0.006147924	enhancer	plate 2	4		0.62992903	NO
MITOXANTHRONE HYDROCHLORIDE	0.84579253	enhancer	6.011118446	0.54139	7.08712015	0.014294841	enhancer	plate 2	5		0.709758632	YES
Clofilium tosylate	0.62546852	enhancer	6.50961628	0.57243	3.497538865	0.028649703	enhancer	plate 1	6	Clofilium tosylate	#N/A	NO
Proscillaridin A	0.44034228	enhancer	7.245694819	0.57332	5.405245334	0.002805423	enhancer	plate 2	7		0.724563422	NO
ASTEMIZOLE	0.36700274	enhancer	3.81960912	0.57944	3.338753735	0.017360156	enhancer	plate 1	8	Astemizole	#N/A	NO
CLOQUINOL	0.32256505	enhancer	4.397798441	0.5915	3.065859359	0.028559644	enhancer	plate 1	9		#N/A	NO
Irinotecan hydrochloride trihydrate	0.4780429	enhancer puncta	6.51755767	0.59903	2.89538224	0 n=1 due to art	non-significant	plate 1	11		#N/A	#N/A
CAMPTOTHECIN	0.6053243	enhancer	6.299959084	0.61347	2.568519237	0.013899469	non-significant	plate 1	12		#N/A	#N/A
Benidipine hydrochloride	0.48895442	enhancer	8.045592404	0.61504	3.208069089	0.036972787	enhancer	plate 2	13		5.691125987	NO
PERHEXILINE MALEATE	0.47856857	enhancer	4.980739055	0.61811	2.463469972	0.03632359	non-significant	plate 1	14		#N/A	#N/A
Thiethylperazine dimalate	0.48550856	enhancer puncta	7.988891758	0.61837	3.033011481	0.003838942	enhancer	plate 2	15		0.703863488	NO
Hemicholinium bromide	-0.4531873	inhibitor	-3.220840366	0.62241	2.366101252	0.034648778	non-significant	plate 1	16		#N/A	#N/A
Pinaverium bromide	0.42260348	enhancer	5.761705784	0.62308	2.784758671	0.007543822	non-significant	plate 2	17		0.739777393	#N/A
DEQUALINIUM CHLORIDE	0.41039736	non-significant	2.916728428	0.62308	2.350888112	0.020482559	non-significant	plate 1	18		#N/A	#N/A
METHYLBENZETHONIUM CHLORIDE	0.91268008	enhancer	12.44332903	0.62409	2.731418006	0.009049988	non-significant	plate 2	19		0.783967327	#N/A
PAROXETINE HYDROCHLORIDE	0.34198959	enhancer	4.662629355	0.631	2.171653741	0.008318707	non-significant	plate 1	20		#N/A	#N/A
FLUPHENAZINE HYDROCHLORIDE	-0.2896035	inhibitor	-3.014070173	0.63136	2.163477921	0.007016697	non-significant	plate 1	21		#N/A	#N/A
MONOBENZONE	-0.2440303	inhibitor	-3.327068637	0.63386	2.106808577	0.005542692	non-significant	plate 1	22		#N/A	#N/A
Chlorhexidine	0.75528286	enhancer	7.860664256	0.63522	2.076073695	0.017631397	non-significant	plate 1	24		#N/A	#N/A
Mometasone furoate	0.49105052	enhancer puncta	3.489937207	0.63897	1.991259899	0.012581924	non-significant	plate 1	26		#N/A	#N/A
MFLOQUINE	0.3319713	enhancer	3.455016753	0.6419	1.924889799	0.012698719	non-significant	plate 1	27		#N/A	#N/A
Melengestrol acetate	0.20525549	enhancer	3.377414955	0.64303	1.899280564	0.018564834	non-significant	plate 1	28		#N/A	#N/A
CPD00059165 BRESTATIN	0.23670364	enhancer	3.894884553	0.64317	1.896118184	0.014925094	non-significant	plate 1	29		#N/A	#N/A
CLEMASTINE	0.75579044	enhancer	7.865946968	0.6433	1.893174968	0.006520221	non-significant	plate 1	30		#N/A	#N/A
ScJeticlopride hydrochloride	0.24723435	enhancer	3.37075221	0.64352	1.888199675	0 n=1 due to art	non-significant	plate 1	31		#N/A	#N/A
DIRITHROMYCIN	-0.5164544	inhibitor	-3.670484376	0.64372	1.883691041	0.005478418	non-significant	plate 1	32		#N/A	#N/A
THIORIDAZINE HYDROCHLORIDE	0.92542359	enhancer	9.631416999	0.64833	1.779263611	0.00935112	non-significant	plate 1	33		#N/A	#N/A
Gefitinib	0.31400324	enhancer	4.281068174	0.64863	1.772553124	0.017491893	non-significant	plate 1	34		#N/A	#N/A
CPD00469186 NELFINAVIR MESYLA	0.60961928	enhancer	10.03109499	0.64872	1.434189959	0.005796663	non-significant	plate 2	35		0.845214443	#N/A
PromethazineA7HCl	0.33508446	enhancer	4.568486038	0.65235	1.688334175	0.008751662	non-significant	plate 1	36		#N/A	#N/A
Penbutolol sulfate	0.28373084	enhancer	4.668702478	0.6529	1.675885365	0.023065261	non-significant	plate 1	37		#N/A	#N/A
Ethynodiol diacetate	0.27073023	enhancer	3.691091086	0.65298	1.674142663	0.012057412	non-significant	plate 1	38		#N/A	#N/A
Cytosporin A	0.53868835	enhancer	3.82850326	0.65708	0.994190768	0.002990588	non-significant	plate 2	39		#N/A	#N/A
GBR 12909 dihydrochloride	0.46558538	enhancer	3.308954362	0.65874	0.906674981	0.00830192	non-significant	plate 2	40		#N/A	#N/A
IDAZOXAN HYDROCHLORIDE	0.43701528	enhancer	5.958193892	0.65879	0.904110495	0.006644269	non-significant	plate 2	41		0.703511717	#N/A
AMINACRINE	0.58954185	enhancer puncta	9.700727096	0.65934	0.874869912	0.00767841	non-significant	plate 2	42		#N/A	#N/A
Halofantrine hydrochloride	0.23789425	enhancer puncta	3.914475583	0.65976	1.520618611	0.010880006	non-significant	plate 1	43		#N/A	#N/A
TENOXCAM	-0.4628417	inhibitor	-3.289454428	0.66007	0.83671378	0.007413766	non-significant	plate 2	44		#N/A	#N/A
FLUNARIZINE HYDROCHLORIDE	0.64299025	enhancer	6.691970378	0.6622	1.465464387	0.024622534	non-significant	plate 1	45		#N/A	#N/A
TOLBRAMYCIN	-0.5303392	inhibitor	-3.769165056	0.66424	1.419107467	0 n=1 due to art	non-significant	plate 1	46		#N/A	#N/A
Tetramisole hydrochloride	-0.5162222	inhibitor	-3.668834442	0.66447	1.414032045	0.008947285	non-significant	plate 1	47		#N/A	#N/A
BIFONAZOLE	0.21800455	enhancer	3.587196739	0.66468	1.409117127	0.025988512	non-significant	plate 1	48		#N/A	#N/A
FLUVOXAMINE MALEATE	0.32104921	enhancer	5.282764564	0.66519	0.566827818	0.005992063	non-significant	plate 2	49		0.793409498	#N/A
(R)-Duloxetine hydrochloride	0.57712202	enhancer puncta	9.496362611	0.66606	0.521208207	0.008358478	non-significant	plate 2	50		#N/A	#N/A
CHLOROQUINE DIPHOSPHATE	-0.5833479	inhibitor	-4.145902448	0.6677	1.340827563	0.00579674	non-significant	plate 1	51		#N/A	#N/A
CPD00469290 SAQUINAVIR MESYLA	0.21267557	enhancer	3.499510193	0.668	1.334098643	0.026679059	non-significant	plate 1	52		#N/A	#N/A
DONEPEZIL HYDROCHLORIDE	0.22495944	enhancer	3.701637312	0.66801	1.333920386	0.017939594	non-significant	plate 1	53		#N/A	#N/A
NSC 257473	0.44433951	enhancer	4.614089982	0.6688	1.316006399	0.008627397	non-significant	plate 1	54		#N/A	#N/A
Azacytidine-5	0.79363653	enhancer	10.82030898	0.67014	0.306090694	0.004012635	non-significant	plate 2	55		#N/A	#N/A
PROADIFEN HYDROCHLORIDE	0.26017043	enhancer	3.547120518	0.67015	1.285456158	0.025991715	non-significant	plate 1	56		#N/A	#N/A
Dipivefrin hydrochloride	-0.4963286	inhibitor	-3.527448987	0.67104	1.265283638	0.020205056	non-significant	plate 1	57		#N/A	#N/A
PERPHENAZINE	0.61889574	enhancer	6.441204919	0.67111	1.263577994	0.014966306	non-significant	plate 1	58		0.763576208	#N/A
Bisoprolol fumarate	-0.4232085	inhibitor	-3.007778244	0.67297	1.221470404	0.016814371	non-significant	plate 1	59		#N/A	#N/A
Quipazine dimaleate salt	-0.4779633	inhibitor	-3.396925118	0.673	1.22091623	0.018289153	non-significant	plate 1	60		#N/A	#N/A
CPD004466343 LETROZOL	0.5367188	enhancer puncta	8.831540259	0.67301	0.155220623	0.006300384	non-significant	plate 2	61		#N/A	#N/A
ASPIRIN	0.21685811	enhancer	3.56833246	0.67357	1.207939524	0.007620018	non-significant	plate 1	62		#N/A	#N/A
CPD004466368 GLIMEPIRIDE	0.43146808	enhancer	5.63715525	0.67404	1.19743859	0.025925192	non-significant	plate 1	63		#N/A	#N/A
CPD004466395 RITONAVIR	0.24017315	enhancer	3.951974192	0.6742	1.193731544	0.012312513	non-significant	plate 1	64		#N/A	#N/A
Carvedilol	0.26235371	enhancer	4.316948325	0.67424	1.192778327	0.02024943	non-significant	plate 1	65		#N/A	#N/A
CHLOROPYRAMINE HYDROCHLORIDE	0.42076681	enhancer puncta	5.736664948	0.67506	1.174276758	0.028189188	non-significant	plate 1	66		#N/A	#N/A
Tegaserod maleate	0.87268382	enhancer	14.35973987	0.67541	0.028516983	0.00809559	non-significant	plate 2	67		#N/A	#N/A
DMSO	#N/A	non-significant	#N/A	0.67549	0.024568329	0.002384526	non-significant	plate 2	68		1.179184516	NO
Hesperidine	-0.5090356	inhibitor	-3.617758548	0.67568	1.160131989	0.01214672	non-significant	plate 1	69		#N/A	#N/A
MITOTANE	0.26411353	enhancer	4.34590566	0.6759	1.155127465	0.012310531	non-significant	plate 1	70		#N/A	#N/A
Mirtazapine	-0.463698	inhibitor	-3.295540233	0.67657	-0.032160421	0.007560238	non-significant	plate 2	71		#N/A	#N/A
RACECADOTRIL	-0.4590654	inhibitor	-3.262616133	0.67675	-0.041885034	0.010522678	non-significant	plate 2	72		#N/A	#N/A
Amnikan hydrate	-0.4277658	inhibitor	-3.04016759	0.6768	-0.044592514	0.006383934	non-significant	plate 2	73		#N/A	#N/A
Kanamycin A sulfate	-0.4872137	inhibitor	-3.462668889	0.6775	-0.081286417	0.011569435	non-significant	plate 2	74		#N/A	#N/A
CPD004466360 FLUBENDAZOLE	0.63493224	enhancer	8.6565609	0.6785	-0.134107445	0.007243105	non-significant	plate 2	75		#N/A	#N/A
NSC 170984	0.44019962	enhancer puncta	4.916136748	0.6795	1.073749974	0.013548053	non-significant	plate 1	76		#N/A	#N/A
Topotcan	0.31810767	enhancer	5.234362465	0.67983	-0.204124039	0.005738607	non-significant	plate 2	77		#N/A	#N/A
Dexfenfluramine hydrochloride	-0.5762799	inhibitor puncta	-4.095669882	0.68013	1.059391628	0.01818349	non-significant	plate 1	78		0.774482364	#N/A
Terazosin hydrochloride	-0.295138	inhibitor	-4.023862949	0.68109	-0.270595863	0.003097366	non-significant	plate 2	79		#N/A	#N/A
GLICLAZIDE	-0.4935736	inhibitor	-3.507868763	0.68122	-0.277264189	0.004194259	non-significant	plate 2	80		#N/A	#N/A
Anethole trithione	0.24132769	enhancer	3.970971859	0.68136	-0.284348293	0.00563113	non-significant	plate 2	81		#N/A	#N/A
ACAMPROSATE CALCIUM	-0.4614512	inhibitor	-3.279572315	0.68141	1.030555459	0.015448197	non-significant	plate 1	82		#N/A	#N/A
ETHOPROPAZINE HYDROCHLORIDE	0.3289878	enhancer	4.485365192	0.68222	-0.329856087	0.00872658	non-significant	plate 2	83		#N/A	#N/A
CPD004466303 LEVETIRACETAM	-0.476691	inhibitor	-3.387883008	0.68294	0.995921752	0.01276144	non-significant	plate 1	84		#N/A	#N/A
PIRACETAM	-0.4740587	inhibitor	-3.3691748	0.68303	-0.3725452	0.006429157	non-significant	plate 2	85		#N/A	#N/A
Ribostamycin sulfate salt	0.27566664	enhancer puncta	3.758393291	0.68415	0.968470696	0.003258565	non-significant	plate 1	86		#N/A	#N/A
BEZAFIBRATE	-0.9902702	inhibitor	-7.03793361	0.68429	-0.438727938	0.006144875	non-significant	plate 2	87		#N/A	#N/A
Sertaconazole nitrate	0.4769561	enhancer	7.848163822	0.68667	-0.564111372	0.008085587	non-significant	plate 2	89		#N/A	#N/A
Algestone acetophenide	0.											

LOVASTATIN	-0.5262054	inhibitor	-3.739785549	0.69939	-1.234096598	0.009119661	non-significant	plate 2	103		#N/A	#N/A
Dinoprost trometamol	-0.4602376	inhibitor	-3.270947516	0.6996	0.618663797	0.008638988	non-significant	plate 1	104		#N/A	#N/A
Atorvastatin	-0.2042638	inhibitor	-3.361097864	0.70119	-1.329020137	0.007061163	non-significant	plate 2	105		#N/A	#N/A
SULCONAZOLE NITRATE	0.5005188	enhancer	6.823990351	0.7015	-1.345210792	0.005644672	non-significant	plate 2	106		#N/A	#N/A
SULFAMETER	0.40672644	enhancer	5.54524086	0.70257	0.551422953	0.008216233	non-significant	plate 1	107		#N/A	#N/A
DO 897/99	-0.4865664	inhibitor	-3.458068531	0.7039	-1.471513704	0.006413292	non-significant	plate 2	108		#N/A	#N/A
OXICONAZOLE NITRATE	0.51386189	enhancer	7.00590784	0.70466	-1.511580608	0.008578999	non-significant	plate 2	109		#N/A	#N/A
Etoricoxib	-0.5956954	inhibitor	-4.233657404	0.70469	0.503599328	0.020796188	non-significant	plate 1	110	0.726130427	#N/A	#N/A
PROPARACAINE HYDROCHLORIDE	0.46583228	enhancer	6.351080169	0.70655	0.461385483	0.010145724	non-significant	plate 1	111		#N/A	#N/A
Estramustine	0.45171892	enhancer	7.432893826	0.70711	-1.6407886	0.009831745	non-significant	plate 2	112		#N/A	#N/A
NAFTIFINE HYDROCHLORIDE	0.21016172	enhancer	3.458145536	0.70755	0.438886633	0.043722179	non-significant	plate 1	113		#N/A	#N/A
Gestrinone	-0.2598281	inhibitor	-4.275390281	0.70759	-1.665919394	0.01395934	non-significant	plate 2	114		#N/A	#N/A
CPD000336944 mevastatin	-0.4510806	inhibitor	-3.205867895	0.71014	-1.800045968	0.004180073	non-significant	plate 2	115		#N/A	#N/A
Tyloxapol	0.42839157	enhancer	5.840619681	0.71024	-1.805697914	0.007311768	non-significant	plate 2	116		#N/A	#N/A
HEXACHLOROPHENE	0.67843993	enhancer	11.16351732	0.71162	-1.878132333	0.009325885	non-significant	plate 2	118		#N/A	#N/A
Amphotericin B	-0.5038243	inhibitor	-3.580721237	0.71165	0.345873361	0.012610234	non-significant	plate 1	119	0.706443472	#N/A	#N/A
MEBENDAZOLE	0.37593186	enhancer	3.912539619	0.71488	-2.049803369	0.004454286	non-significant	plate 2	120		#N/A	#N/A
DIOXANIDE FUROATE	0.23836408	enhancer	3.249816349	0.71754	0.212573974	0.022902366	non-significant	plate 1	121		#N/A	#N/A
CHLORTETRACYCLINE HYDROCHLORIDE	0.46579662	enhancer	4.847814096	0.71768	0.209452285	0.019309703	non-significant	plate 1	122		#N/A	#N/A
PRAZOSIN HYDROCHLORIDE	-0.2837255	inhibitor	-3.868266743	0.71855	-2.243398148	0.008318949	non-significant	plate 2	123		#N/A	#N/A
TRICLOSAN	0.57623554	enhancer	9.481775926	0.7209	-2.366996431	0.01784983	non-significant	plate 2	124		#N/A	#N/A
AMOROLFINE HYDROCHLORIDE	0.45026197	enhancer	7.408920167	0.72105	-2.374978195	0.026561846	non-significant	plate 2	125		#N/A	#N/A
Bromocryptine mesylate	0.41298894	enhancer	4.298214103	0.72336	0.080958069	0.024958997	non-significant	plate 1	126		#N/A	#N/A
Avermectin B1a	0.36715902	enhancer	5.00578527	0.72552	-2.610317469	0.01880213	non-significant	plate 2	127		#N/A	#N/A
DMSO	#N/A	non-significant	#N/A	0.72569	0.028082996	0.004960111	non-significant	plate 1	128	0.675069842	NO	
DILAZEP DIHYDROCHLORIDE	-0.5391178	inhibitor	-5.610909504	0.73351	-0.14892721	0.01791535	non-significant	plate 1	129		#N/A	#N/A
Ivermectin	0.31826256	enhancer	3.31234195	0.7349	-3.104485911	0.008257464	inhibitor	plate 2	130		#N/A	NO
Haloproglin	0.55120193	enhancer	3.917438312	0.73786	-3.260059023	0.01481143	inhibitor	plate 2	131		#N/A	NO
Estropipate	0.29819842	enhancer	4.06558783	0.73851	-0.262091035	0.020159141	non-significant	plate 1	132		#N/A	#N/A
Alfacalcidol	0.44851495	enhancer	3.187633392	0.75281	-4.047774907	0.009385267	inhibitor	plate 2	133		#N/A	NO
Clinidipine	0.28226936	enhancer	4.644654177	0.7576	-4.299936095	0.011337818	inhibitor	plate 2	134		#N/A	NO
CYCLOHEXIMIDE	0.63325095	enhancer	8.633638512	0.84598	-2.694714613	0.028425129	non-significant	plate 1	135	Cyclohexamide	#N/A	NO
Quinacrine dihydrochloride hydrate	-2.1091858	inhibitor puncta	-21.95151267	0.85718	-2.948365052	0 n=1 due to art	non-significant	plate 1	136	Quinacrine	0.723005469	#N/A
DIACERIN	0.29801344	enhancer	4.063065895	0.92739	-4.537680299	0.074528679	inhibitor	plate 1		Diacerin	#N/A	YES
Nicosamide	0.27756754	enhancer	3.099866314	1.10624	-22.66078712	0.026593144	inhibitor	plate 2	137	Nicosamide	#N/A	NO