Introduction

Niemann-Pick Type C disease (NPC) is a lethal, autosomal recessive disorder caused by mutations in the NPC1 and NPC2 cholesterol transport proteins. NPC’s hallmark symptoms include an accumulation of unesterified cholesterol and other lipids in the late endosomal and lysosomal cellular compartments, causing progressive neurodegeneration and death. Although the age of onset may vary in those affected, NPC most often manifests in juveniles, and is usually fatal before adolescence. In this study, we investigated the effects of various drugs, many of which modify the epigenetic control of NPC1/NPC2 gene expression, in lowering the otherwise harmful elevated intracellular cholesterol levels in NPC cells. Our studies utilized a previously described image analysis technique, which allowed us to make quantitative comparisons of the efficacy of these drugs in lowering cholesterol levels in a common NPC1 mutant model. Of the drugs analyzed, several that have been previously studied (vorinostat, panobinostat, and β-cyclodextrin) significantly lowered the relative amount of unesterified cellular cholesterol, consistent with earlier observations. In addition, a novel potential treatment, rapamycin, likewise alleviated the NPC phenotype. We also studied combinations of effective compounds with β-cyclodextrin; the addition of β-cyclodextrin significantly enhanced the cholesterol-lowering activity of vorinostat and panobinostat, but had mixed effects with rapamycin. Collectively, these results may provide a basis for the eventual development of improved NPC therapies.

Abstract

Niemann-Pick Type C disease (NPC) is a lethal, autosomal recessive disorder caused by mutations in the NPC1 and NPC2 cholesterol transport proteins. NPC's hallmark symptoms include an accumulation of unesterified cholesterol and other lipids in the late endosomal and lysosomal cellular compartments, causing progressive neurodegeneration and death. Although the age of onset may vary in those affected, NPC most often manifests in juveniles, and is usually fatal before adolescence. In this study, we investigated the effects of various drugs, many of which modify the epigenetic control of NPC1/NPC2 gene expression, in lowering the otherwise harmful elevated intracellular cholesterol levels in NPC cells. Our studies utilized a previously described image analysis technique, which allowed us to make quantitative comparisons of the efficacy of these drugs in lowering cholesterol levels in a common NPC1 mutant model. Of the drugs analyzed, several that have been previously studied (vorinostat, panobinostat, and β-cyclodextrin) significantly lowered the relative amount of unesterified cellular cholesterol, consistent with earlier observations. In addition, a novel potential treatment, rapamycin, likewise alleviated the NPC phenotype. We also studied combinations of effective compounds with β-cyclodextrin; the addition of β-cyclodextrin significantly enhanced the cholesterol-lowering activity of vorinostat and panobinostat, but had mixed effects with rapamycin. Collectively, these results may provide a basis for the eventual development of improved NPC therapies.

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Competing Interests: We have read the journal's policy and have the following potential conflicts to declare: co-authors P. Helquist and O. Wiest are included as co-inventors on US Patent Application No. 12/888,267 filed on September 22, 2010 for HISTONE DEACETYLASE INHIBITORS AS THERAPEUTIC AGENTS FOR LYSOSOMAL STORAGE DISORDERS. We have no other potential conflicts to declare. This patent application will not alter our adherence to all PLoS One data and material sharing policies.

* E-mail: hgoodson@nd.edu
United States—a variety of other drug treatments have been proposed to help fight the progression of NPC [7]. Currently, a major focus in the development of potential treatments is epigenetic modulators—compounds that regulate gene expression at the systemic and/or cellular levels. Theoretically, by increasing the expression of mutated genes that may retain some limited function, deleterious symptoms that otherwise result from decreased amounts of the fully functional protein could be diminished. With respect to NPC, this would mean increasing the amounts of mutated NPC1 or NPC2 protein with the hope that their suppressed functionality could be overcome by such an increase.

In support of this idea, the overexpression of mutated NPC1 has been shown to ameliorate the NPC phenotype in a cell line homoallelic for the mutation H1061T; this mutation represents roughly 15% of disease-causing alleles and is the most common mutation found in NPC patients from Western countries. [8,9]. Additional relief has been demonstrated through increased expression of other factors responsible for cholesterol transport and removal processes, including but not limited to the Rab proteins. For example, Choudhury, et al. have demonstrated that overexpression of Rab7 and Rab9 in NPC cells significantly lowers intracellular cholesterol levels, while others have shown that increased levels of Rab4 and Rab6 produce similar results [10,11].

Two classes of molecules that upregulate gene expression are histone deacetylase (HDAC) inhibitors and DNA methyl transferase (DNMT) inhibitors. The histone deacetylase family consists of enzymes involved in the modification of histones, proteins that regulate DNA condensation into chromosomes. In addition, HDAC’s act on a variety of other nuclear and cytoplasmic proteins [12]. HDAC inhibitors increase expression by preventing deacetylation of histone lysine residues, inhibiting DNA packing and thus enhancing transcription of many (but not all) genes [12]. DNMT inhibitors, in contrast, act to inhibit DNA methylation. When methylation is suppressed, expression of potentially silenced genes is increased [13]. Because both HDAC and DNMT inhibitors help increase gene expression, they are potential candidates for treating NPC; they may upregulate the expression of NPC1, NPC2, Rabs, or, more broadly, other factors that contribute to the management of intracellular cholesterol levels.

Previous work has indicated that the HDAC inhibitors vorinostat (suberoylanilide hydroxamic acid, SAHA; marketed as Zolinza for the treatment of cutaneous T cell lymphoma) and panobinostat (LBH-589; currently in Phase III trials for cutaneous T cell lymphoma) successfully alleviate the NPC phenotype in NPC mutant cells [14]. Vorinostat has also been shown to effectively cross the blood-brain barrier, a significant hurdle for NPC mutant cells [14]. Vorinostat has also been demonstrated to increase the expression of a variety of protein factors in myeloblastic leukemic cell lines with positive therapeutic results [16]. However, its effects on NPC cells have yet to be determined.

Potential treatments of NPC are not limited to these two specific classes of molecules, however. Several other drugs, many of which are already FDA-approved for other conditions, have been identified as agents that alter cholesterol homeostasis, and thus might have potential as NPC therapeutics. These FDA-approved compounds are especially attractive as potential therapeutic agents for NPC, as many drug development and approval phases have been bypassed, significantly expediting their introduction as NPC treatments.

β-cyclodextrin and its derivatives have been shown in previous studies to reduce intracellular cholesterol levels significantly in NPC1 and NPC2 mutants [17], and one derivative has been identified as a cholesterol-chelating agent that mimics the NPC2 protein [18,19]. It also has significant positive effects on a mouse model of NPC [20], and, in anecdotal instances, it has been shown to improve the neurological prognosis of NPC patients [21]. While the effects of β-cyclodextrin on NPC cells are well established, its cooperative effects with other therapies are not fully understood.

In addition, evidence has been presented that chloroquine and chlorpromazine may be potentially useful in treating NPC. Chloroquine has been implicated in inhibiting lysosomal proteolysis, which may help maintain NPC1 and NPC2 protein levels [8]. Chlorpromazine, meanwhile, has been identified as an agent that facilitates cellular cholesterol efflux to serum-based acceptors, and has been suggested as having value in the treatment of NPC [22]. However, both compounds have been shown to inhibit cholesterol esterification [23,24], raising concerns over their potential to treat NPC.

Finally, rapamycin (sirolimus), an immunosuppressant, has been previously shown by Sharpe and Brown to decrease LDL-receptor gene expression, which may inhibit cholesterol uptake in treated cells [25]. It has also been demonstrated to stimulate cholesterol efflux in human mesangial cells previously induced by the cytokine IL-1β to retain cholesterol, suggesting a role in cellular lipid homeostasis [26]. Rapamycin has not yet been studied as a potential NPC therapy, but these observations suggest that it may have utility.

In light of these previous findings, we investigated the effect of each of these compounds—vorinostat, panobinostat, β-cyclodextrin, rapamycin, decitabine, chloroquine, and chlorpromazine—in intracellular cholesterol levels in NPC1 mutant fibroblasts. We determined their efficacy in lowering NPC cell cholesterol levels using a previously described procedure that quantifies cholesterol levels in lysosomes and late endosomes (referred to as lysosome storage organelles, or LSO’s) through fluorescence image analysis [27]. After identifying successful compounds, further studies were performed on the cholesterol-lowering effects of the successful drugs in combination with an effective concentration of β-cyclodextrin.

Materials and Methods

The methods used in this study closely model those previously described by Pipalia, et al. to be useful in determining the cholesterol-lowering efficacy of various compounds in NPC mutant cells [27]. The process quantitatively determines intracellular cholesterol levels by means of fluorescent microscopy and imaging, and produces an “LSO Ratio” value indicating the amount of unesterified cholesterol relative to cholesterol levels in untreated solvent controls.

Materials

Unless otherwise stated, all materials, including Eagle’s Minimal Essential Media (MEM), L-glutamine, fetal bovine serum (FBS), phosphate buffered saline (PBS), streptomycin, penicillin, DMSO, EtOH, filipin, paraformaldehyde (FA), chloroquine, chlorpromazine, rapamycin, decitabine (5-aza-2′-deoxycytidine) and β-cyclodextrin were purchased from Sigma-Aldrich, Inc. (St. Louis, MO). Metamorph imaging software was from Molecular Devices (Downington, PA), ImageJ image processing and analysis software
was from the National Institutes of Health (Bethesda, MD), and OriginPro statistical software was from Origin Lab Corp. (Northampton, MA).

Cell Culture

Healthy GM00498 human fibroblasts (male, non-fetal skin cells aged 3 years) were acquired from the Coriell Institute for Medical Research (Camden, NJ). The mutant GM03123 human fibroblast cell line (aged 9 years, female) was also acquired from Coriell. GM03123 cells are compound heterozygotes that possess two point missense mutations—one CT at nucleotide 709, and another TC at nucleotide 3182 of the NPC1 gene. These correspond to the substitutions P237S and I1061T (as mentioned, a very prevalent NPC1 mutation) in the luminal domains of the folded NPC1 protein, respectively [9]. Cells were cultured in 10% FBS/Eagle MEM solution, supplemented with L-glutamine, penicillin, and streptomycin, at 37°C and 5% CO₂.

Drug Solutions

The drugs studied—vorinostat (SAHA), panobinostat (LBH-589), decitabine, chloroquine, chlorpromazine, rapamycin, and β-cyclodextrin—were prepared in stock solutions of 20 mM, 300 μM, 20 mM, 200 mM, 80 mM, 240 μM, and 200 mM, respectively, and stored at −20°C. Vorinostat, decitabine, chlorpromazine, rapamycin, and β-cyclodextrin were dissolved in DMSO, while panobinostat was solubilized in EtOH, and chloroquine was dissolved in PBS. Stock solutions were diluted in media to the appropriate working solution concentrations.

Cell Plating/Treatment Doses

Cells were treated in black-walled Costar 96-well polystyrene plates (Corning, Inc., Corning, NY). Prior to use, the plate wells were treated with 0.1% gelatin in PBS for 1 h at room temperature in order to increase cell adherence and thus prevent cells from detaching or washing off in later steps. Cells of interest were removed from culture conditions and diluted so that approximately 1750 cells would be present in each well, in order to achieve roughly 50–60% confluence upon imaging. 100 μL of drug solution in media to reach 100 μL 0.1 mg/mL filipin was added to 100 μL of drug solution in media to reach a total well volume of 200 μL, with a range of concentrations for each tested compound. Proper solvent controls were run in parallel. The drugs that were identified as successful in the initial screen were tested in combination with β-cyclodextrin, at all but the highest individually-tested concentration for each.

Fixation and Staining

After incubation in drug treatments for the 48 h, cells were washed 2×200 μL with PBS and incubated for 45 minutes at 4°C in 3% PFA. Afterwards, the fixed cells were washed 1×200 μL with PBS, and treated with 50mM NH₄Cl in PBS at room temperature to terminate fixation. They were then washed 2×200 μL with PBS, and treated with 100 μL 0.1 mg/mL filipin staining solution in PBS for 2 hours at room temperature. Filipin binds to unesterified cholesterol and fluorescences under UV excitation, allowing for identification of intracellular cholesterol inclusions. After staining, the cells were washed with 3×200 μL PBS prior to image acquisition.

Fluorescent Microscopy

Images of each condition were acquired in 200 μL PBS using a Nikon TE2000 microscope (IX optivar, 10X/0.30NA bright field objective) and CCD Cascade 512B camera under UV excitation (340–380 nm, 40 nm dichronic, 430 nm long pass filter) using Metamorph (100 ms acquisition time, 1x binning) software. Twelve representative images of each condition (avoiding hyperconfluent clusters of cells) were acquired in each experiment.

Image Processing and Analysis

A quantitative measure of intracellular cholesterol levels in drug-treated cells relative to untreated solvent controls was determined, and averaged across the 12 images of each condition, in the manner described by Pipalia, et al. [27]. ImageJ software was used to process and normalize images. After 5th percentile background subtraction, contrasts on all images were adjusted to between 0 and 5000 units. Following these adjustments, low intensity thresholds—above which all cellular membranes are displayed—were set at 220 units/pixel, while the high thresholds—above which only cholesterol inclusions are visible—were set at 1200 units/pixel for all images. The total integrated pixel intensity above the high threshold was then divided by the total image area above the low threshold to yield the numerical LSO values. LSO ratio values were obtained by dividing the LSO value for a given treatment by the LSO value for that treatment’s solvent control. The resultant LSO ratio values represent the proportion of cholesterol levels in treated cells relative to levels in the controls. An LSO ratio of “1” represents cholesterol levels equal to those in the solvent controls, while an LSO ratio of “0” signifies minimal unesterified cellular cholesterol—approximates levels in healthy fibroblasts.

Statistical Analysis

The statistical significance of the cholesterol-lowering effects of the tested compound doses were determined using the one-way analysis of variance and Tukey’s Honest Significant Difference (HSD) post-hoc means comparison features of OriginPro software. Efficacies of the drugs tested in combination with β-cyclodextrin were analyzed using the two-way analysis of variance and Tukey’s HSD post-hoc means comparisons features of OriginPro.

Results

The methods used allowed us to determine the extent to which the tested drugs lowered intracellular unesterified cholesterol levels. During our initial screen of the drugs, in which their individual cholesterol-lowering efficacies were determined, several were found or reaffirmed to significantly lower intracellular cholesterol levels as demonstrated by their calculated LSO ratios. Representative images of NPC1 mutants exposed to treatments of each compound are presented in Figure 1.

Effects of Vorinostat, Panobinostat, and Decitabine

The compounds implicated in the modulation of gene expression displayed varying efficacies, with the two HDAC inhibitors—vorinostat and panobinostat—demonstrating the strongest effects (Figure 2 and Table 1). Vorinostat, acting alone, was found to lower intracellular cholesterol levels to approximately 46% of those found in the parallel solvent controls at a concentration of 10 μM. Even at its least effective tested dose, 1.25 μM, vorinostat lowered levels by almost 35%. Panobinostat, meanwhile, lowered cholesterol levels by nearly 75% at a concentration of 75 nM, and, even at its least effective dose, 300 nM, lowered levels approximately 48%. It appeared that at concentrations greater than 75 nM, the cholesterol-lowering efficacy of panobinostat was diminished. The lone DNMT inhibitor tested, decitabine, displayed no significant effects at any concentration. Complete numerical and statistical
Effects of 200 μM β-Cyclodextrin on Treatment with Vorinostat, Panobinostat, Rapamycin, and Decitabine

As stated earlier, due to their success in lowering NPC1 mutant cholesterol levels acting alone, vorinostat, panobinostat, and rapamycin were tested in combination with 200 μM β-cyclodextrin. For comparison purposes, the ineffective decitabine was also tested with β-cyclodextrin. Combinations of β-cyclodextrin and these compounds were significantly more effective than these compounds alone at lowering cholesterol levels (Table 2, Reduction from “Drug-Only”). More specifically, β-cyclodextrin enhanced the decreases in cholesterol observed through treatment with vorinostat alone by 26%–40%, and did so for panobinostat by 9%–22%. In addition, it increased the cholesterol-lowering efficacies of rapamycin and decitabine by 36%–40% and 35%–51%, respectively.

Effects of Vorinostat and Panobinostat on Treatment with 200 μM β-Cyclodextrin

Combinations of vorinostat and panobinostat with β-cyclodextrin appeared to be slightly more effective at reducing cholesterol levels than β-cyclodextrin alone (which, as mentioned earlier, lowered cholesterol levels at 200 μM by 68%). A vorinostat/β-cyclodextrin treatment lowered intracellular cholesterol levels from control levels by 69%–81%, while a panobinostat/β-cyclodextrin combination did so by 74%–84% (Table 2, Vorinostat, thus, appeared to increase the efficacy of β-cyclodextrin by 1%–14%, and panobinostat did so by 7%–16% (Table 2, Reduction from 200 μM β-Cyclodextrin). However, while vorinostat and panobinostat appear to enhance cholesterol reductions by these small amounts when added to β-cyclodextrin, the differences that were observed between the cholesterol levels in these treatments and the β-cyclodextrin-only treatment are not statistically significant.

Effects of Rapamycin and Decitabine on Treatment with 200 μM β-Cyclodextrin

Curiously, treatments with β-cyclodextrin and either rapamycin or decitabine, respectively, were not as effective as treatment with solely β-cyclodextrin. Rapamycin/β-cyclodextrin combinations only lowered cholesterol levels by 54%–60%, while decitabine/β-cyclodextrin combinations did so by only 40%–45% (Table 2). Though rapamycin and decitabine each demonstrated a slight reduction or no effect on cholesterol levels acting alone, when used in combination with β-cyclodextrin, they served to elevate cholesterol levels by 8%–14% (rapamycin) and 25%–28% (decitabine) relative to those found with treatment with only 200 μM β-cyclodextrin (Table 2, Reduction from 200 μM β-
NPC1 mutant fibroblasts were treated with various drugs for 48 h, after which they were fixed, filipin stained, imaged, and processed to give an LSO ratio value as described in Materials and Methods. Vorinostat, panobinostat, and β-cyclodextrin were very effective at lowering cholesterol levels, obtaining LSO ratios below 0.5 at several doses. Rapamycin was somewhat effective, consistently obtaining LSO ratios of ~0.8. Decitabine had no significant effect on intracellular cholesterol levels. Chloroquine and chlorpromazine were not effective; they elevated cholesterol levels at all tested concentrations. The red lines
indicate solvent controls (LSO ratio = 1). Each data point represents the average of at least three independent experiments totaling 36 images (three experiments x twelve images/experiment) and error bars are +/- SE. Data points without error bars reflect the results of one experiment; cells were not viable in two of three trials at chlorpromazine concentrations of 40 and 80 μM.

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Table 1. Dose Dependence Summary Statistical Data.

| Compound       | Treatment Concentration | Mean LSO Ratio | Percent Cholesterol Reduction |
|----------------|-------------------------|----------------|-------------------------------|
| Vorinostat     | 20 μM                   | 0.563          | 43.7 **                       |
|                | 10 μM                   | 0.456          | 54.5 ***                      |
|                | 5 μM                    | 0.483          | 51.7 ***                      |
|                | 2.5 μM                  | 0.521          | 47.9 ***                      |
|                | 1.25 μM                 | 0.652          | 34.8                          |
|                | 0.625 μM                | 0.709          | 29.1                          |
| Panobinostat   | 300 nM                  | 0.522          | 47.8 **                       |
|                | 150 nM                  | 0.346          | 65.5 ***                      |
|                | 75 nM                   | 0.252          | 74.8 ***                      |
|                | 37.5 nM                 | 0.282          | 71.8 **                      |
|                | 18.75 nM                | 0.369          | 63.1 ***                      |
|                | 9.375 nM                | 0.468          | 53.2                          |
| β-Cyclodextrin | 400 μM                  | 0.345          | 65.5 **                       |
|                | 200 μM                  | 0.321          | 67.9 **                       |
|                | 100 μM                  | 0.337          | 66.3 **                       |
|                | 50 μM                   | 0.388          | 61.2 **                       |
|                | 25 μM                   | 0.432          | 56.8 **                       |
|                | 12.5 μM                 | 0.545          | 45.5 **                       |
| Rapamycin      | 240 nM                  | 0.925          | 7.5                           |
|                | 120 nM                  | 0.862          | 13.8                          |
|                | 60 nM                   | 0.813          | 18.7 **                       |
|                | 30 nM                   | 0.823          | 17.7 **                       |
|                | 15 nM                   | 0.825          | 17.5 **                       |
|                | 7.5 nM                  | 0.787          | 21.3 **                       |
| Decitabine     | 20 μM                   | 1.178          | −17.8                         |
|                | 10 μM                   | 1.101          | −10.1                         |
|                | 5 μM                    | 1.046          | −4.6                          |
|                | 2.5 μM                  | 0.948          | 5.2                           |
|                | 1.25 μM                 | 0.950          | 5.0                           |
|                | 0.625 μM                | 0.923          | 7.7                           |
| Chloroquine    | 200 μM                  | 1.577          | −57.7                         |
|                | 100 μM                  | 1.481          | −48.1                         |
|                | 50 μM                   | 2.158          | −115.8                        |
|                | 25 μM                   | 1.830          | −83.0                         |
|                | 12.5 μM                 | 2.047          | −104.7                        |
|                | 6.25 μM                 | 1.791          | −79.1                         |
| Chlorpromazine | 80 μM                   | 1.074          | −7.4                          |
|                | 40 μM                   | 1.105          | −10.5                         |
|                | 20 μM                   | 1.561          | −56.1                         |
|                | 10 μM                   | 1.557          | −55.7                         |
|                | 5 μM                    | 1.296          | −29.6                         |
|                | 2.5 μM                  | 1.560          | −56.0                         |

* = significantly reduced from solvent control at p<0.05.
** = significant at p<0.01.
*** = significant at p<0.001.
Cyclodextrin). Rapamycin and decitabine thus appear to unexpectedly counteract β-cyclodextrin. Representative images from the combination treatments are also presented in Figure 1, and numerical data on all tested β-cyclodextrin combinations are displayed in Figure 3 and Table 2.

**Discussion**

In this report, we demonstrate the efficacy of several compounds and novel combinations thereof in treating NPC. Vorinostat and panobinostat, as expected, were confirmed to be agents that significantly lower intracellular cholesterol levels in NPC cells. Similarly, we confirmed β-cyclodextrin as a compound able to alleviate the NPC phenotype. More significantly, we observed that rapamycin, a drug that has not previously been tested in the context of NPC, lowered cholesterol levels in our tests. We also identified two drugs—chloroquine and chlorpromazine—that had previously shown some promise as cholesterol-lowering agents in this assay. It remains to be seen whether decitabine has any effect on NPC cells at lower cholesterol levels opens up a new perspective from which to investigate NPC pathology and treatment. Rapamycin acts by inhibiting a well-conserved and extensively investigated serine-threonine kinase known as mTOR (molecular target of rapamycin). Inhibition of mTOR is achieved after rapamycin binds to another protein factor, FKBP12 [28]. mTOR plays multiple roles in cell physiology, regulating cell growth and cell proliferation, and responding to cellular nutrient and energy levels [28,29].

There are multiple possible mechanisms by which rapamycin could affect cholesterol homeostasis in NPC cells. For instance, many mTOR downstream targets are involved with transcription and translation processes, such as the recruitment of ribosomes to mRNA [29]. As such, mTOR may play a role in modulating expression of proteins involved in cholesterol homeostasis. As noted above, rapamycin decreases LDL-receptor gene expression [25], which would be expected to reduce cholesterol uptake in treated cells. And, as mentioned, it stimulates cholesterol efflux in mesangial cells that have been induced to retain cholesterol [26]. Moreover, recent studies have shown that mTOR complex 1 (mTORC1) activates SREBP-1c (sterol regulatory element-binding protein 1c), which stimulates lipogenesis by upregulating the expression of genes associated with lipid synthesis [30,31]. This observation suggests that rapamycin may inhibit intracellular cholesterol accumulation by reducing SREBP activity. The ability of rapamycin to reduce the intracellular cholesterol load, as seen in our assays, could result from any or all of these mechanisms, or alternatively, from the variety of other metabolic alterations induced by rapamycin. In any case, its observed cholesterol-lowering activity suggests that rapamycin might have therapeutic value.

| Compound | Treatment Concentration (+200 μM β-Cyclodextrin) | Mean LSO Ratio | Percent Cholesterol Reduction from Solvent Control | Percent Cholesterol Reduction from 200 μM β-Cyclodextrin | Percent Cholesterol Reduction from "Drug-Only" Result |
|----------|-----------------------------------------------|--------------|---------------------------------|---------------------------------|-------------------------------|
| Vorinostat | 10 μM | 0.186 | 81.4 *** | 15.9 | 70.0 |
| | 5 μM | 0.227 | 77.7 *** | 10.0 | 60.0 |
| | 25 μM | 0.227 | 79.7 *** | 10.0 | 30.0 |
| | 1.25 μM | 0.264 | 74.6 *** | 10.0 | 38.8 |
| | 0.625 μM | 0.312 | 68.8 *** | 9.9 | 39.7 |
| Panobinostat | 150 nM | 0.256 | 74.4 *** | 10.0 | 0.0 |
| | 75 nM | 0.163 | 83.7 *** | 15.8 | 9.0 |
| | 37.5 nM | 0.179 | 82.1 *** | 14.2 | 10.0 |
| | 18.75 nM | 0.208 | 79.2 *** | 11.3 | 15.1 |
| | 9.375 nM | 0.246 | 75.4 *** | 7.5 | 22.2 |
| Rapamycin | 120 nM | 0.463 | 53.7 *** | -14.2 | 39.9 |
| | 60 nM | 0.453 | 54.7 *** | -13.2 | 36.0 |
| | 30 nM | 0.445 | 55.5 *** | -12.4 | 37.8 |
| | 15 nM | 0.447 | 55.3 *** | -12.6 | 37.8 |
| | 7.5 nM | 0.410 | 59.9 *** | -8.0 | 38.6 |
| Decitabine | 10 μM | 0.587 | 41.3 *** | -26.6 | 51.4 |
| | 5 μM | 0.599 | 40.1 *** | -27.8 | 44.7 |
| | 2.5 μM | 0.603 | 39.7 *** | -28.2 | 34.5 |
| | 1.25 μM | 0.574 | 42.6 *** | -25.3 | 37.6 |
| | 0.625 μM | 0.546 | 45.4 *** | -22.5 | 37.7 |

*** = significantly reduced from solvent control at p < 0.001.
1–LSO Ratio = 0.321; 67.9% cholesterol reduction.
2–See Table 1 for “Drug-Only” Results.
3–200 μM β-cyclodextrin significantly enhanced the cholesterol-lowering activity of all four compounds at p < 0.001.

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Nonetheless, the dose dependence relationship observed for rapamycin treatments is somewhat unexpected. As is shown in Figure 2 and Figure 3, the rapamycin dose dependence curve is nearly flat across the range of tested concentrations, and the cholesterol-lowering effects of rapamycin are actually more pronounced at lower dose levels (decrease of 21.3% at 7.5 nM), than they are at higher concentrations (insignificant decrease at 240 nM). If, however, rapamycin’s cholesterol-lowering effects are induced via the mTORC1/SREBP1c pathway, these observations are consistent with recent proposals that suggest mTORC1 is most strongly inhibited at low nanomolar (0.5–100 nM) concentrations [32]. It remains to be determined, though, whether or not the observed trend continues beyond our range of concentrations. These results indicate that further investigation of the impact of rapamycin on intracellular NPC cholesterol levels is warranted.

The second major finding of this study is that the addition of β-cyclodextrin to vorinostat, panobinostat, and rapamycin, significantly enhanced cholesterol reductions relative to the use of these three compounds individually. Though the specific manner of β-cyclodextrin’s cholesterol-lowering action remains to be fully elucidated—aside from its potential to mimic the NPC2 protein through a cholesterol-chelating mechanism [18,19]—its observed effect may also be partly attributable to its ability to act as a drug carrier, facilitating the entry of vorinostat, panobinostat, and rapamycin into cells [33].

However, the β-cyclodextrin/rapamycin and β-cyclodextrin/decitabine combination therapies also had surprising aspects: these treatments resulted in higher cholesterol levels than those found from treatment with β-cyclodextrin alone. Given that rapamycin lowered cholesterol levels by approximately 20% at most concentrations when used alone, and that decitabine, individually, had no significant effect on cholesterol levels at any concentration, it would seem logical that their addition to a β-cyclodextrin treatment would be slightly beneficial or, at worst, have no impact. The observed results suggest that rapamycin and decitabine may in some way counteract the action of β-cyclodextrin, or somehow adversely affect lipid homeostasis in the presence of β-cyclodextrin. Given the intricacy of the mTOR pathway discussed above, it is possible that β-cyclodextrin interacts in some way with a downstream target of mTOR to affect cholesterol homeostasis. And, though decitabine’s mechanism of action— inhibition of DNA methylation—is not nearly as complicated as that of rapamycin and mTOR, it is also possible that there is some underlying opposing interaction between β-cyclodextrin and DNA methylation machinery. Our results indicate that future studies on these apparent counteractive effects are justified.

In summary, we have reaffirmed the efficacy of vorinostat, panobinostat, and β-cyclodextrin in reducing intracellular cholesterol levels in NPC fibroblasts, and have shown that a new compound, rapamycin, alleviates the NPC phenotype. While the effects of rapamycin were relatively small, this observation could have clinical relevance because of its established use as an immunosuppressant and its many well-studied cellular interactions. We have also identified chloroquine, chlorpromazine, and decitabine as unsuccessful cholesterol-lowering agents in this assay. Finally, we have demonstrated that β-cyclodextrin significantly increases the cholesterol-lowering effectiveness of various compounds in vitro. Given that many of these compounds are already

Figure 3. Dose Response Curves for Cholesterol-Lowering Efficacy of Tested Drugs in Combination with β-Cyclodextrin. NPC1 mutant fibroblasts were treated with vorinostat, panobinostat, rapamycin, and decitabine, both with and without 200 μM β-cyclodextrin, for 48 h, after which they were fixed, filipin stained, imaged, and processed to give an LSO ratio value as described in Materials and Methods. β-Cyclodextrin increased the impact vorinostat, panobinostat, rapamycin, and decitabine had on late endosomal and lysosomal cholesterol levels at all concentrations. The red lines indicate vehicle controls (LSO ratio = 1), while green lines indicate the average result of treatment with only 200 μM β-cyclodextrin (LSO ratio = 0.321). Each point represents the average of at least three independent experiments totaling 36 images (three experiments x twelve images/experiment) and error bars are +/- SE.

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FDA-approved for other conditions, we hope that these findings can help to justify and guide further research and trials, and contribute to the eventual discovery of an effective treatment for this devastating disease.

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Author Contributions

Conceived and designed the experiments: ZTW TWH KLH KTV OW. Performed the experiments: ZTW TWH. Analyzed the data: ZTW. Contributed reagents/materials/analysis tools: HG PH OW. Wrote the paper: ZTW HG PH.