

A Fluorescence Polarization Activity-Based Protein Profiling Assay in the Discovery of Potent, Selective Inhibitors for Human Nonlysosomal Glucosylceramidase

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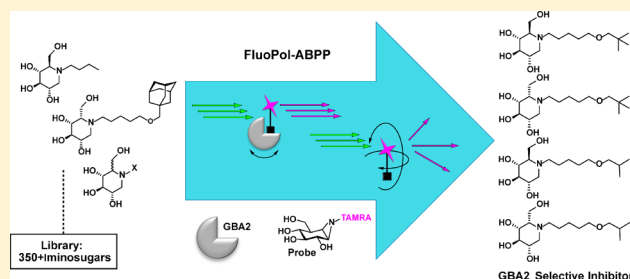
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Supporting Information

ABSTRACT: Human nonlysosomal glucosylceramidase (GBA2) is one of several enzymes that controls levels of glycolipids and whose activity is linked to several human disease states. There is a major need to design or discover selective GBA2 inhibitors both as chemical tools and as potential therapeutic agents. Here, we describe the development of a fluorescence polarization activity-based protein profiling (FluoPol-ABPP) assay for the rapid identification, from a 350+ library of iminosugars, of GBA2 inhibitors. A focused library is generated based on leads from the FluoPol-ABPP screen and assessed on GBA2 selectivity offset against the other glucosylceramide metabolizing enzymes, glucosylceramide synthase (GCS), lysosomal glucosylceramidase (GBA), and the cytosolic retaining β -glucosidase, GBA3. Our work, yielding potent and selective GBA2 inhibitors, also provides a roadmap for the development of high-throughput assays for identifying retaining glycosidase inhibitors by FluoPol-ABPP on cell extracts containing recombinant, overexpressed glycosidase as the easily accessible enzyme source.



INTRODUCTION

Human neutral glucosylceramidase (GBA2, nonlysosomal glucocerebrosidase), a member of the glycoside hydrolase (GH) family, GH116 (www.cazy.org), hydrolyzes glucosylceramide in the cytosol to produce glucose and ceramide.¹⁻³ GBA2 is located at the cytosolic leaflet of the endoplasmic reticulum, Golgi apparatus, and endosomes.^{1,4} GBA2 controls levels of glucosylceramide, which is synthesized at the cytosolic leaflet of the Golgi apparatus prior to translocation to the lumen of the organelle for elongation to complex glycosphingolipids. GBA2 activity is complementary to GH30 human acid glucosylceramidase (GBA, glucocerebrosidase), which processes glucosylceramide in lysosomes.⁵ Genetic mutations in GBA are at the basis of the lysosomal storage disorder Gaucher disease. Gaucher macrophages are loaded with massive amounts of glucosylceramide.⁵ Other tissues in Gaucher, however, have remarkably normal glucosylceramide levels, and it is likely that GBA2 can compensate for reduced GBA activity in these tissues.⁵ Glucosylceramide has been linked to inflammation as well,⁶ underscoring the biomedical

potential of compounds able to modulate glucosylceramide levels.

In addition to its interplay with GBA, GBA2 is itself implicated in many diseases. Mutations in the GBA2 gene cause hereditary spastic paraplegia and autosomal recessive cerebellar ataxia,⁷⁻⁹ indicating that compounds able to enhance GBA2 activity could have therapeutic potential. Conversely, compounds able to (partially) block GBA2 activity also hold therapeutic value; we recently found that lowering GBA2 activity in Niemann-Pick type C (NPC) mouse models led to elongated life-span and improved motor coordination.¹⁰ This finding we correlated to correction of GBA2 activity levels in Purkinje cells, which are among the most affected neuronal cells in NPC. Slowing-down of neuropathological processes in NPC mice was achieved by genetic ablation of the GBA2 gene and alternatively by pharmacological inhibition of the gene product, the latter with *N*-alkyldeoxynojirimycin derivatives such as *N*-butyldeoxynojirimycin or *N*-(5)-adamantane-1-yl-

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methoxypentyl deoxynojirimycin (respectively Zavesca (**1**) and AMP-DNM (**2**), Figure 1).¹⁰ Indeed, we and others have reported on the development of *N*-alkyldeoxynojirimycin derivatives as potent GBA2 inhibitors. The most potent GBA2 inhibitor at present is AMP-DNM (**2**), with a K_i of

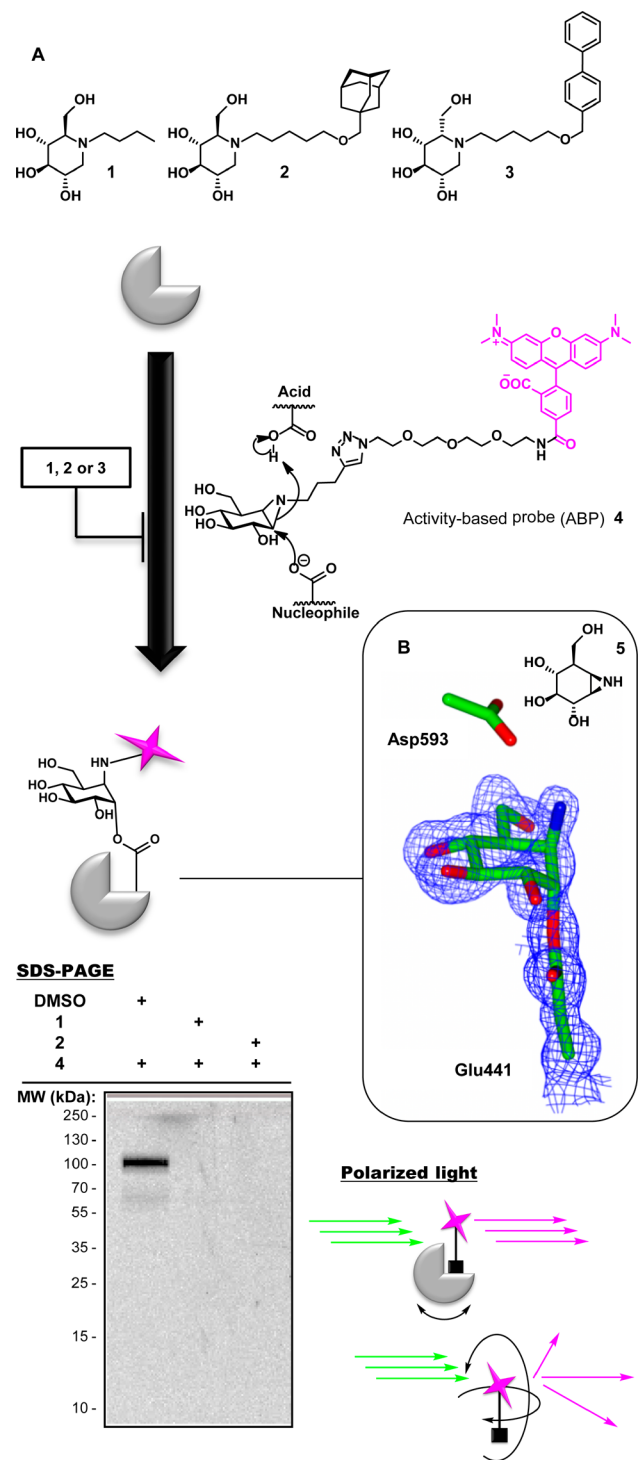


Figure 1. (A) Screening for GBA2 inhibitors (e.g., **1**, **2**, or **3**) using activity-based probe **4** as a read-out in a FluoPol-ABPP assay. (B) 3-D structure (PDB SNCX) of the TxGH116 β -glucosidase bound to cyclophellitol aziridine **5**; the enzymatic nucleophile Glu441 and putative acid/base Asp593 are shown for reference. The map shown is a maximum-likelihood weighted $2F_{\text{obs}} - F_{\text{calc}}$ at $0.4 \text{ e}^-/\text{\AA}^3$.

about 3 nM; this compound however also inhibits glucosylceramide synthase (GCS) and GBA at high nanomolar concentrations.^{11,12}

GBA2 inhibitors thus have considerable potential both as cellular probes/tools and as therapeutic agents. Often, though, potent GBA2 inhibitors have significant off-targets; not only promiscuity toward GBA but also the enzyme responsible for the biosynthesis of glucosylceramide: GCS. Selective *in vivo* GBA2 inhibition is therefore complicated. For this reason, and considering the pharmaceutical potential of GBA2 as a drug target, selective GBA2 inhibitors are desirable commodities. Similarly, in order to access specific inhibitors, there is an urgent need for assays that rapidly identify potential GBA2 inhibitors. We here report on the development of a fluorescence polarization activity-based protein-profiling (FluoPol-ABPP) assay for fast identification of GBA2 inhibitors in extracts from cells overexpressing GBA2. Screening of our iminosugar library, which contains over 350 entries, provided lead structures bearing a relatively small (compared to existing nanomolar GBA2 inhibitors, including **2** and **3**) apolar head groups. Building a focused library around this theme and assessment of the inhibitory potential of these against GBA2, GBA, and GCS yielded new leads for, and new insights into, future development of potent, selective GBA2 inhibitors. These leads, combined with our FluoPol-ABPP assay, are a useful addition to existing reagents and tools in unraveling GBA2 functioning in health and disease.

RESULTS AND DISCUSSION

Our FluoPol-ABPP assay is based on our activity-based glycosidase profiling methodology that capitalizes on the two-step double displacement mechanism employed by retaining glycosidases. *N*-Alkylcyclophellitol aziridines, with the aziridine nitrogen bearing reporter groups (fluorophores, biotin), are in class, broad-spectrum retaining β -glucosidase activity-based probes (ABPs) reporting on GBA, GBA2, and the cytosolic retaining β -glucosidase GBA3 in human tissue (Figure 1A).¹³ In order to support specific reaction and covalency, we solved the structure of the *Thermoanaerobacterium xyloxyticum* TxGH116 β -glucosidase, a close homologue of human GBA2 with a conserved active site, in complex with cyclophellitol aziridine **5** (the precursor of ABP **4**) to ascertain that cyclophellitol aziridines react with the active site nucleophile and thus that competitors of ABP **4** are true active site inhibitors, Figure 1B.¹⁴ The aziridine indeed binds to the invariant catalytic nucleophile Glu441, in an active site pocket on the enzyme surface. The active site pocket is freely accessible to solvent, with little steric hindrance for tagged, functionalized aziridines such as **4** (Supplementary Figure 4a). Binding of tagged ABP **4** to both GBA2 and TxGH116 can also be directly observed by fluorescent scanning of the respective proteins after ABP labeling (Figure 1A and Supplementary Figure 4B).

FluoPol-ABPP studies that have appeared in the literature make use of the tetramethylrhodamine (TMR) dye.¹⁵ In order to stay in tune with these studies, we therefore synthesized TMR-cyclophellitol aziridine **4** (for synthesis details see the Supporting Information, SI Schemes 1 and 2). GBA2 was stably overexpressed in HEK293 cells, constituting the majority of all β -glucosidase activity (the ratio of GBA2 levels over other targets labeled by **4** was established by prolonged imaging of the gel shown in Figure 1A; see the Supporting Information, SI Figure S1). Extracts of HEK293 overexpressing GBA2 were

treated with ABP 4 at 500 nM at pH 7, and the protein content was resolved by SDS PAGE. Fluorescence scanning of the wet gel slab (λ_{ex} 530 nm, λ_{em} 580 nm, Figure 1A) revealed a major band at about 102 kDa, which corresponds to the molecular weight of mature GBA2. Levels of active GBA2 in the overexpressing cells are such that endogenous GBA and GBA3, which are also targets of cyclophellitol aziridine ABPs, are hardly visible, even though labeling of GBA and GBA3 is saturating at the probe concentrations used. GBA2 labeling with 4 can be abolished by preincubation with the known GBA2 inhibitors Zavesca (1) (100 μM final concentration) and AMP-DNM (2) (10 μM final concentration) in a competitive ABPP experiment (Figure 1A, lanes 2 and 3, respectively). In fluorescence polarization assays, a fluorophore is excited at a specific wavelength with linear polarized light. The orientation of emitted light and thereby the degree of polarized light depend on the movement the excited fluorophore has undergone, which depends on its size: large molecules (such as GBA2-bound 4) move and rotate slower than small molecules (here: free 4). Samples are irradiated with linear polarized light (λ_{ex} 530 nm), and the degree of loss of polarized light emitted at λ_{em} 580 nm represents the degree of inhibition of GBA2 labeling with 4. Initial experiments using fluorescence polarization as a read-out were performed in 96-well plates. The pH optimum of GBA2 is reported¹ to be at pH 5.8, and contemporary assays for measuring GBA2 inhibitor activities are performed at or close to this pH. We (Figure 2A)

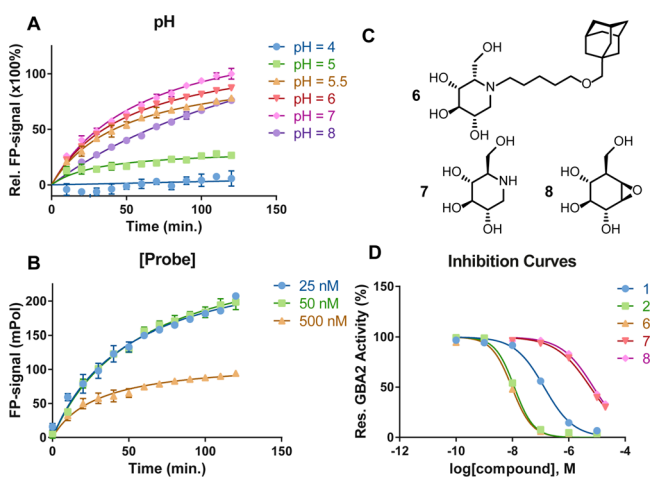


Figure 2. Optimization and validation of FluoPol-ABPP for GBA2. (A) Effect of pH. (B) Effect of probe concentration. (C) Structures of additional established inhibitors. (D) Inhibition by established inhibitors (1, 2, 6, 7, and 8). Error bars represent standard error of the mean (SEM).

established that, in a range from pH 4 to pH 8 (the range at which GBA2 possesses measurable activity), maximal return of polarized light occurs at pH 7. The optimal final probe concentration in the assay conditions used (protein concentration 0.5 mg/mL, V_{final} 75 μL), which intrinsically depends on the amount of active GBA2 present in the extracts (which we keep constant), was found to be 25–50 nM (Figure 2B); in this setting a suitable difference in polarized light emission (Δ about 180 mPol) was achieved between samples derived from GBA2-overexpressing cells and wild-type cells. IC_{50} values of a set of known GBA2 inhibitors, both competitive (Zavesca (1), AMP-DNM (2), *L*-ido-AMP-DNM (6), DNM (7)) and mechanism-based (cyclophellitol, 8), were next established using the

FluoPol-ABPP format using 50 nM ABP 4. Values obtained (Figure 2D) were in line with literature values on the same compound, for both the potent (2, 3, 6) and less potent (1, 7, 8), which were all established using fluorogenic substrate assays, thereby validating our FluoPol-ABPP assay.¹⁶

We next transferred our assay to a 384-well plate format (V_{final} 15 μL) and screened our complete iminosugar library, containing both previously published and until now unpublished compounds (the structures of all 358 compounds and the synthesis of new compounds are in the Supporting Information). All compounds were assessed at 100 nM final concentration, and the percentage of GBA2 inhibition calculated from the relative loss of fluorescence polarization was observed. Besides *N*-alkyldeoxyojirimycin, the corresponding *D*-galacto- and *L*-ido-configured iminosugars are known to be potent GBA2 inhibitors, and library entries adhering to this general description are picked up in our FluoPol-ABPP assay as well (Figure 3).¹⁶ Alternatively

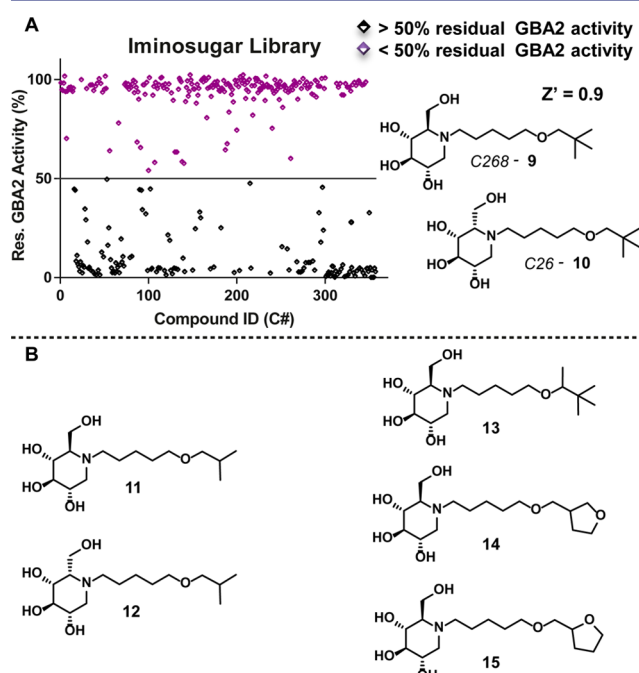


Figure 3. FluoPol-ABPP assay with an iminosugar library (358 entries). (A) Competition at 100 nM. (B) Examples of second-generation analogues of compounds 9 and 10.

configured piperidines, partially deoxygenated deoxyojirimycin derivatives, and hydroxylated pyrrolidines show up only rarely, if at all, as inhibitors in our assays (for a more detailed grouping of compound classes and GBA2 inhibition see the Supporting Information).

Among the most potent GBA2 inhibitors we identified neopentylloxypentyl-deoxyojirimycin 9 and its *L*-ido-congener 10 (Figure 3). Intriguingly, for these compounds the hydrophobic *N*-alkyl group is relatively small compared to that of 2 and 3. We previously found that large (as in 2) and preferably aromatic (as in 3) alkoxy substituents are preferred by GCS and reasoned that a reduced size of this part of the molecule may be favorable for achieving GBA2 selectivity over the ceramide glucosyltransferase. Accordingly, we synthesized a set of close analogues of compounds 9 and 10 and assessed their inhibitory potency against GBA2, GBA, GBA3, and GCS (examples of second-generation library 11–15 in Figure 3; for

Table 1. Inhibition Values and Selectivity Ratios of Deoxynojirimycin Derivatives on GCS, GBA, GBA2, and GBA3

compound	GCS ^a	GBA1 ^b	GBA2 ^b	GBA3 ^b	GCS/GBA2 ^c	GBA1/GBA2 ^c	GBA3/GBA2 ^c
1	50	675	0.326	>1000	153	2071	>3067
2	0.15	0.28	0.0035	27.6	43	80	7886
3	0.008	25.3	0.0027	27.8	3	9370	10296
6	0.15	12.4	0.0083	66.1	18	1494	7964
9	5.05	3.8	0.0051	27.6	990	745	5412
10	2.77	120	0.0077	119	360	15584	15455
11	3.87	10.8	0.0075	128	516	1440	17067
12	0.729	1223	0.0095	177	77	128737	18632
13	6	7.3	0.0059	116	1017	1237	19661
14	53	92	0.0536	180	989	1716	3358
15	>50	122	0.1771	145	>282	689	819

^aInhibition value for *in situ* assay is given as IC₅₀ (μM). ^bInhibition value for *in vitro* assay is given as K_i (μM). ^cRelative GBA2 offset (ratio)

synthesis and analytical details see the [Supporting Information](#)).

As can be seen in [Table 1](#), compounds **9** and **10** are highly potent and rather selective GBA2 inhibitors. Removal of one methyl group, as in isopropyl derivatives **11** and **12**, led to an increase of inhibitory potency against GBA2 with concomitant (slight) decrease of inhibitory potency against GCS. *L*-ido-configured compounds **10** and **12** are over 100 000-fold selective for GBA2 over GBA1, which is a significant improvement compared to previously reported inhibitors showing a maximal 10 000-fold selectivity (for selectivity ratios of the compounds in [Table 1](#) for GBA2 offset against GBA, GBA3, and GCS, respectively, see the [Supporting Information](#), SI 8). As mentioned earlier, the activity against GCS is much reduced when comparing to our previously reported adamantyl and biphenyl compounds, respectively **2** and **3**. Compounds **9**, **13**, and **14** are about 1000-fold more potent on GBA2 over GCS, while previously reported compounds are at best 150-fold more selective. While substitution of the neopentyl moiety for tetrahydrofuranylmethyl groups, as in **14** and **15**, had a detrimental effect on inhibitory potency toward GBA, GBA2, and GCS, it has no significant effect on the selectivity. Based on both potency and selectivity, neopentyl and isopropyl derivatives **9–12** are perhaps the most potent and selective GBA2 inhibitors reported to date.

In a final *in situ* target engagement experiment we performed competitive ABPP on live cells. Cells overexpressing GBA2 or GBA3 and containing endogenous GBA levels were treated with compounds **9–12** at various final concentrations, lysed, and then treated with ABP **4**. As can be seen from the images of the SDS PAGE gels ([Figure 4](#)), all compounds are cell permeable and selectively block GBA2 over GBA and GBA3 at the concentrations tested.

CONCLUSIONS

Collectively, this work describes the development of an effective FluoPol-ABPP assay for the rapid screening of GBA2 inhibitors. Fluorescence polarization (FluoPol) is often used for the analysis of the potency of small molecules to inhibit enzymes¹⁷ but has to the best of our knowledge not been used in the context of glycosidase inhibitor assays. This certainly holds true for the combination of FluoPol and ABPP,¹⁸ as we show here. FluoPol-ABPP can be conducted on target enzymes in complex mixtures, as long as the ABP used modifies with considerable selectivity the target enzyme. This can be achieved even with broad-spectrum ABPs such as the one used here, simply by bringing the target enzyme to

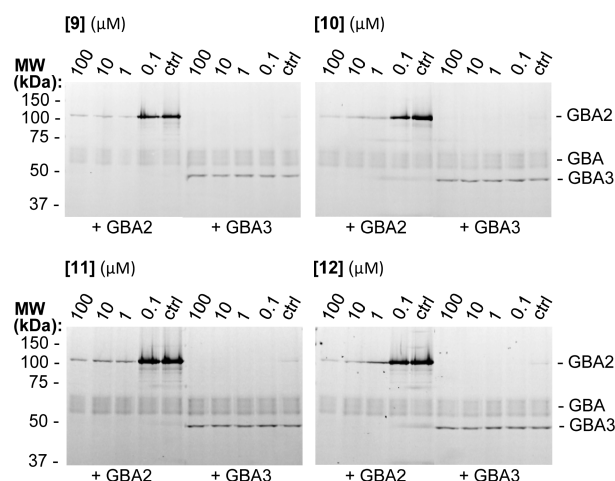


Figure 4. Competitive ABPP experiment of HEK293T cells overexpressing GBA2 or GBA3 (and expressing endogenous GBA) treated with compounds **9–12** at various final concentrations prior to cell lysis and ABPP profiling of remaining enzyme activity.

overexpression. Target selectivity is easily established by running an SDS PAGE on the ABP-labeled sample. One potential caveat of FluoPol-ABPP is the covalent and irreversible nature of the ABP used. Most studies on FluoPol-ABPP inhibitor screening entail the assessment of covalent inhibitors.¹⁸ Here, we show that competitive inhibitors can also be identified using FluoPol-ABPP as an initial screen, after which more in-depth kinetic studies can be executed on selected compounds using conventional assays using fluorogenic substrates. Here, we identified a few compounds from our 358-compound collection that possess considerable GBA2 selectivity, and on the basis of these lead structures we prepared a small focused library to add to our growing library of iminosugar glycosidase inhibitors. In-depth analysis revealed that, indeed, modulating the size of the hydrophobic nitrogen substituent yields inhibitors more potent, and crucially also more selective, for GBA2 compared to existing inhibitors. From a technical point of view, the FluoPol-ABPP assay can be adapted to allow high-throughput screening of large compound selections on GBA2 inhibition. Conceptually, with this study we show that FluoPol-ABPP, as a recently proposed strategy for high-throughput screening of various enzymes, is also eminently feasible for searching for glycosidase inhibitors, especially considering that the list of activity-based glycosidase probes targeting a variety of retaining glycosidases is growing.

EXPERIMENTAL SECTION

Cell Culture. Human embryonic kidney 293T (HEK293T) cells (Sigma) were cultured in DMEM high glucose (Gibco) supplemented with 10% NBS and 100 units/mL penicillin/streptomycin (Gibco) at 37 °C and 5% CO₂. RAW 264.7 cells (American Type Culture Collection) were cultured in RPMI (Gibco) supplemented with 10% fetal calf serum, 1 mM Glutamax, and 100 units/mL penicillin/streptomycin (Gibco) at 37 °C and 5% CO₂.

Transient and Stable Overexpression of Human GBA2 and GBA3. Primers used for transient GBA2-expressing HEK293T were designed based on NCBI reference sequence NM_020944.2. The full-length coding sequences were cloned into pcDNA3.1/Myc-His (Invitrogen, Life Technologies, Carlsbad, CA, USA). Subconfluent HEK293T cells were transfected with empty pcDNA3.1 or GBA2pcDNA3.1 (plasmid:PEI ratio 1:3). Media was refreshed 24 h later, and cells were collected 72 h after transfection in phosphate-buffered saline (PBS) buffer. Cells were centrifuged at 1000 rpm for 10 min, after which the supernatant was removed. Cell pellets were snap-frozen with liquid nitrogen and stored at -80 °C.

Stable GBA2-expressing HEK293T cells were generated as follows. The PCR-amplified human GBA2 (GBA2, acc. no. NM_020944.2) coding sequence (using the following primers: sense 5'-GGGGACAA-GTTTGTACA AAAAAGCAGGCTTAACCACCATGGGGACCC-AGGAT CCAG-3' and antisense 5'-GGGGACCACTTTGTAC AAGAAAGCTGGGTTTCACTCTGGGCTCAGTTTG-3') was cloned into pDNOR-221 and subcloned in pLenti6.3/TO/V5-DEST using the Gateway system (Invitrogen). Correctness of the construct was verified by sequencing. HEK293T cells were transfected with pLenti6.3-GBA2 in combination with the envelope and packaging plasmids pMD2G, pRRE, and pRSV. Subsequently, culture supernatant containing viral particles was collected and used for infection of HEK293T cells. Selection using blasticidin for several weeks rendered cells stably expressing human GBA2 as determined by activity assays. For stable expression of GBA3 in HEK293T cells, the PCR-amplified GBA3 (GBA3, acc. no. NM_020973.4) coding sequence (using the following oligonucleotides: sense 5'-GAATTCGCCGACCATGGCTTCCCTGCAGGATTTG-3' and antisense 5'-GCGGCCGC-AGATGTGCTTCAA GGCCATTG-3') was cloned into pcDNA3.1/Zeo and transfected into HEK293T cells using FuGene 6 Transfection Reagent (Promega Benelux, Leiden, The Netherlands). Selection using Zeocin for several weeks rendered cells stably expressing human GBA3 as determined by activity assays.

Preparation of Lysates. Cell pellets were resuspended in lysis buffer (20 mM hepes, 2 mM DTT, 0.25 M sucrose, 1 mM MgCl₂, 2.5 U/mL benzonase, pH 7.0). The homogenate was incubated on ice for 30 min after lysis and homogenization using SilentCrusher (Heidolph). Ultracentrifugation was performed at 32,000 rpm for 30 min at 4 °C. Supernatant fractions were aliquoted in appropriate volumes, after the total protein concentration was determined via a Bradford assay, using BSA (Sigma) for standards and BioRad Quickstart Bradford Reagents. Aliquots were snap-frozen with liquid nitrogen and stored at -80 °C.

Gel-Activity-Based Protein Profiling Experiments. Prepared lysate was diluted in assay buffer (20 mM Hepes, 2 mM DTT at pH 7.0) until appropriate final protein concentration was reached. Samples were incubated with 500 nM probe at 37 °C for 1 h ($V_{\text{final}} = 20 \mu\text{L}$). Protein content was denatured using Laemli buffer (4 \times) at 100 °C for 3 min. Reactions were resolved by 12.5% SDS-PAGE electrophoresis, and wet slabs were scanned for fluorescence (Molecular Imager Gel Doc XR, Biorad). For the *in situ* competitive ABPP experiments stable GBA2- and GBA3-overexpressing HEK293T cells were incubated for 1 h at 37 °C with various concentrations of the selected inhibitors. Cells were washed two times with PBS and then harvested by scraping in 25 mM potassium phosphate buffer (pH 6.5) containing 0.1% Triton X-100 (v/v). Cells were lysed via snap-freezing in liquid nitrogen. Protein concentration was determined via BCA assay (Thermo Fischer). Labeling of the glucosidases was conducted using 500 nM ABP 4 for 30 min at 37 °C on 20 μg of protein. Protein content was denatured using Laemli buffer (5 \times) at 100 °C for 5 min. Reactions were resolved

by 10% SDS-PAGE electrophoresis, and wet slabs were scanned as described above.

Optimization of the FluoPol-ABPP Assay. Optimal probe concentration on the FluoPol signal was determined by varying probe concentrations from 25 to 500 nM probe at a constant lysate concentration (0.5 mg/mL) and neutral pH. FluoPol-ABPP assays were also performed at different pH values. Hepes was replaced by 20 mM citric acid-citrate buffer for reactions at pH \leq 6. The pH experiments were performed at optimal probe concentration (50 nM). Competition experiments were conducted by 30 min of preincubation of compounds in the lysates at 37 °C (2.5% DMSO). All reactions ($V_{\text{final}} 75 \mu\text{L}$) containing bovine gamma-globulin (Sigma) and Chaps (Sigma), respectively 0.5 and 1 mg/mL, were carried out in 96-well plates (flat-black bottom, Greiner). FluoPol signals were monitored on an Infinite M1000Pro (Tecan) using λ_{ex} 530 nm and λ_{em} 580 nm. Mock-containing lysates were used as reference samples, samples without probe as blanks to correct for background polarization, and GBA2-containing lysates without inhibitor as controls. All samples were corrected for background polarization, and the residual enzyme activity was calculated based on the polarization signal from the controls. Polarization signals were plotted against time or inhibitor concentration and processed in GraphPad Prism 6.0. IC₅₀ values were calculated via nonlinear regression using the mentioned software ($N = 2$, $n = 3$).

FluoPol-ABPP Screen of the Iminosugar Library. The screen, using the optimized conditions as described above, was conducted in a 384-well black-bottom plate (Greiner) with reaction volumes set at 15 μL . Concentration of the iminosugars during the screen was 100 nM. The FluoPol signal was measured on a PHERAstar (BMG Labtech). Resulting polarization signals were processed as described above. Residual enzyme activities were plotted against the corresponding iminosugar.

Analysis of Selected Compounds as Inhibitors of Enzymatic Activity of GCS, GBA, GBA2, and GBA3. Enzyme assays of GCS, GBA1, GBA2, and GBA3 were carried out as previously described (refs 16 and 19), and inhibition by selected compounds was assessed. Briefly, for GBA1, pure recombinant human enzyme (Cerezyme from Genzyme) was used. Activity was measured with 3.7 mM 4-methylumbelliferone (4MU)- β -D-glucopyranoside (Sigma) in 150 mM McIlvaine buffer pH 5.2 supplemented with 0.2% sodium taurocholate (w/v), 0.1% Triton X-100 (v/v), and 0.1% bovine serum albumin (BSA) (w/v) for 30 min. The reaction was stopped with excess 1 M NaOH-glycine (pH 10.3), and liberated 4MU fluorescence was measured with an LS55 fluorimeter (PerkinElmer) using λ_{ex} 366 nm and λ_{em} 445 nm. For GBA2 measurements, cellular homogenates of a stable HEK293T-overexpressing GBA2 cell line preincubated for 30 min with an inhibitor of GBA1 (1 mM conduritol β epoxide CBE from Sigma) were used. Activity was measured with 3.7 mM 4MU- β -D-glucopyranoside in 150 mM McIlvaine pH 5.8 and 0.1% BSA (w/v) for 1 h. Reactions were stopped as described above for GBA1. For GBA3 measurements, cellular homogenates of a stable HEK293T-overexpressing GBA3 cell line also preincubated for 30 min with an inhibitor of GBA1 (1 mM conduritol β epoxide CBE from Sigma) were used. Reactions were conducted in 100 mM hepes pH 7.0 and 0.1% BSA (w/v) for 1 h. All IC₅₀ values were determined in triplicate, and the inhibitors tested were preincubated for 30 min at 37 °C. K_i values were determined in duplicate using a range from 0.05 to 5 mM 4MU- β -D-glucopyranoside in appropriate buffer containing inhibitor. Incubation time and temperature were the same as described above, but without the preincubation step. Observed fluorescence was curve-fitted against inhibitor or substrate concentrations using GraphPad Prism 6.0 in order to obtain IC₅₀ or K_i values.

IC₅₀ values for GCS were determined *in situ* with 6-[N-methyl-N-(7-nitrobenz-2-oxa-1,3-diazol-4-ylaminododecanoyl)]sphingosine (NBD-ceramide) as substrate. RAW 264.7 cells were grown to confluence in six-well plates and preincubated for 1 h with an inhibitor of GBA1 activity (300 μM CBE), followed by 1 h incubation at 37 °C with 1 μM C6-NBD-ceramide and in the presence of a range of inhibitor concentrations. The cells were washed 3 \times with PBS and harvested by scraping. After lipid extraction (described by Bligh and

Dyer²⁰), the C6-NBD lipids were separated and detected by high-performance liquid chromatography (λ_{Ex} 470 nm and λ_{Em} 530 nm). IC₅₀ values were determined in duplicate from the titration curves of observed formed C6-NBD-glucosylceramide, and data were curve-fitted via GraphPad Prism 6.0.

■ ASSOCIATED CONTENT

📄 Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.7b07352.

Synthetic procedures for ABP 4 and (unpublished) deoxynojirimycin analogues, ¹H and ¹³C NMR spectra, IR and LC-MS data; figures validating the FluoPol-ABPP method and additional figures for the screen showing a preference for *gluco*- and *ido*-configured iminosugars; compound structures and inhibitory potencies on GCS, GBA1, and GBA2 (PDF)

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Notes

The authors declare no competing financial interest.

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