Mechanism of Human Nucleocytoplasmic Hexosaminidase D

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ABSTRACT: Mammalian β-hexosaminidases have been shown to play essential roles in cellular physiology and health. These enzymes are responsible for the cleavage of the monosaccharides N-acetylgalosamine (GlcNAc) and N-acetylglactosamine (GlnNAc) from cellular substrates. One of these β-hexosaminidases, hexosaminidase D (HexD), encoded by the HEXDC gene, has received little attention. No mechanistic studies have focused on the role of this unusual nucleocytoplasmically localized β-hexosaminidase, and its cellular function remains unknown. Using a series of kinetic and mechanistic investigations into HexD, we define the precise catalytic mechanism of this enzyme and establish the identities of key enzymic residues. The preparation of synthetic aryl N-acetylglactosaminide substrates for HexD in combination with measurements of kinetic parameters for wild-type and mutant enzymes, linear free energy analyses of the enzyme-catalyzed hydrolysis of these substrates, evaluation of the reaction by nuclear magnetic resonance, and inhibition studies collectively reveal the detailed mechanism of action employed by HexD. HexD is a retaining glycosidase that operates using a substrate-assisted catalytic mechanism, has a preference for galactosaminide over glucosaminide substrates, and shows a pH optimum in its second-order rate constant at pH 6.5–7.0. The catalytically important residues are Asp148 and Glu149, with Glu149 serving as the general acid/base residue and Asp148 as the polarizing residue. HexD is inhibited by Gal-NAG-thiazoline (K = 420 nM). The fundamental insights gained from this study will aid in the development of potent and selective probes for HexD, which will serve as useful tools to improve our understanding of the physiological role played by this unusual enzyme.

Mammals possess four genes encoding β-hexosaminidases. These enzymes are responsible for the cleavage of the terminal monosaccharides N-acetylgalosamine (GlcNAc) and N-acetylglactosamine (GlnNAc) from cellular substrates. The HEXA and HEXB genes encode an α subunit and a β subunit, respectively, that dimerize to form three different isozymes: hexosaminidase A (HexA; comprising an α subunit and a β subunit), hexosaminidase B (HexB; comprising two β subunits), and hexosaminidase S (HexS; comprising two α subunits). All three isozymes localize to the lysosome, where they hydrolyze terminal GlcNAc and GlnNAc residues from glycoconjugates and oligosaccharides to aid in the disposal of unneeded glycoconjugates. Deficiencies in either of these genes lead to the genetic diseases known as Sandhoff and Tay-Sachs disease. O-GlcNcase (OGA), encoded by the HEXC gene, is a nucleocytoplasmic β-hexosaminidase. This enzyme removes GlcNAc residues from proteins bearing N-acetylgalosamine O-linked to serine and threonine residues of nucleocytoplasmic proteins (O-GlcNAc) and has been implicated in an array of physiological processes and also found to be essential for development of mammals. The fourth gene encoding a mammalian β-hexosaminidase (HEXDC) was identified recently. While the physiological substrate and function of the gene product, hexosaminidase D (HexD), remain unknown, studies have indicated a link between this enzyme and rheumatoid arthritis. The intracellular and extracellular roles played by this enzyme within cells are therefore topics of great interest.

Preliminary characterization of recombinantly expressed murine HexD, which is 80% similar to human HexD, shows it can hydrolyze substrates containing both GlnNAc and GlcNAc but with a preference for the former. This differs from OGA, which exclusively hydrolyses GlcNAc-containing substrates, whereas HexA, HexB, and HexS can hydrolyze substrates containing both GlnNAc and GlcNAc. HexD was shown, by reverse transcriptase polymerase chain reaction (RT-PCR) experiments, to be ubiquitously expressed in all murine tissues examined and localized to the nucleus and cytoplasm upon being overexpressed in mammalian cells. Interestingly, the biochemical properties of recombinantly expressed murine HexD are in accord with the characteristics of an enzyme observed in bovine tissue isolates decades earlier, however, the gene encoding it was unknown, and this observation received limited interest.

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Glycoside hydrolases (GHs) are classified by sequence homology into more than 100 families [Carbohydrate Active enZyme (CAZy) database]. In general, the enzyme mechanism and three-dimensional fold are conserved among members within any family. The CAZy database classifies HexD, along with the lysosomal β-hexosaminidases encoded by the HEXA and HEXB genes, into family GH20. Conversely, OGA bears no sequence similarity to the other β-hexosaminidases and falls into family GH84. Although there is a low degree of sequence conservation between these two families of mammalian β-hexosaminidases, studies have shown they share a similar, but unusual, substrate-assisted catalytic mechanism. The three-dimensional structures of human HexA, human HexB, and bacterial homologues of OGA have also provided insights into the active site architecture, revealing how these enzymes facilitate catalysis using key enzymic residues. Together, these mechanistic and structural studies have guided the development of potent and specific inhibitors against β-hexosaminidases, which have acted as powerful tools for probing the cellular function of HexD. While our knowledge of the physiological function of HexD lags behind that of the other mammalian β-hexosaminidases, given the increasing important roles played by β-hexosaminidases in health and disease, gaining a clear fundamental understanding of this unusual nucleocytoplasmically localized enzyme is important.

Figure 1. Substrate-assisted catalytic mechanism employed by GH20 enzymes. Binding of substrate to the enzyme leads to formation of a Michaelis complex in which the pyranose ring of the substrate adopts a conformation placing the leaving group in a pseudoaxial orientation. This allows nucleophilic attack of the anomeric center by the C2 acetamido group and departure of the leaving group. The resulting transient bicyclic oxazoline intermediate is then hydrolyzed by attack of a molecule of water at the anomeric center, releasing the sugar hemiacetal product. Two conserved enzymic carboxyl groups provide general acid and general base catalysis during each step of the reaction.

Materials and Methods

HexD Protein Expression and Purification. The plasmid containing the gene encoding human HexD in a pET30a vector, which has been reported previously, was transformed into Escherichia coli BL21(DE3) cells. Successful transformants were cultured in Luria-Bertani broth supplemented with 50 μg/mL kanamycin at 37 °C until an optical density of 0.6 absorbance units was reached. Protein expression was induced with 0.5 mM isopropyl β-D-thiogalactoside at 15 °C for 20 h.
Cells were harvested and resuspended in 20 mM HEPES (pH 7.4), 150 mM NaCl, and 5 mM imidazole and incubated in the presence of 1 mg/mL lysozyme, 0.02 mg/mL DNase, and an EDTA-free protease inhibitor tablet (Roche) for 20 min at 4 °C. Cells were lysed by high-pressure cell disruption (Constant Systems). Following clarification, the supernatant was applied to a 5 mL HisTrap nickel column (GE Healthcare), pre-equilibrated in the same buffer, and the protein was eluted from an imidazole gradient. The elution was concentrated and applied to a HiPrep 26/10 desalting column equilibrated in 20 mM HEPES (pH 7.4), 150 mM NaCl buffer to remove the imidazole. The elution was subsequently concentrated and applied to a S200 16/60 gel filtration column, pre-equilibrated in the same buffer, and fractions judged to be pure were pooled for subsequent kinetic studies.

**HexD Mutagenesis.** The primers listed in Table S1 were used to amplify the plasmid encoding the HexD gene with the desired mutation. Reaction mixtures were subjected to digestion with DpnI for 2 h at 37 °C and subsequently transformed into E. coli DH5α cells. Plasmid DNA was extracted from cells using standard procedures and subsequently sequenced (GATC sequencing) to ensure the mutation was successfully incorporated. Protein expression and purification were performed as described for wild-type HexD, and the enzymes were obtained in similar yields.

**General Procedures for Synthesis of Compounds.** All synthetic reagents used in this study were obtained from Sigma-Aldrich (Oakville, ON), Carbosynth (San Diego, CA), Alfa Aesar (Ward Hill, MA), or Acros Organics (Geel, Belgium). Anhydrous reactions were conducted in flame-dried glassware under a positive pressure of dry argon. Air- or moisture-sensitive reagents and anhydrous solvents were transferred with oven-dried syringes or cannulae. Flash chromatography was performed using E. Merck silica gel (230–400 mesh). Solution-phase reactions were monitored using analytical thin layer chromatography (TLC) with E. Merck precoated silica gel plates. Chemical shifts are reported in parts per million downfield as NMR solvents, unless otherwise stated. Chemical shifts are reported as δ (ppm) for 1H NMR and δ (ppm) for 13C NMR. When 19F NMR spectra are reported, chemical shifts are reported as δ (ppm) for 19F NMR. Infrared spectra were recorded as a KBr disk with a Nicolet Magna 560 FTIR (E. Biochemistry). 1H and 13C NMR spectra were recorded in CDCl3 or DMSO-d6 with TMS and/or DSS as internal standards.

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3-H), 2.06 (s, 3H), 2.02 (s, 3H); 13C NMR (125 MHz, CDCl3)

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General Procedure B: Synthesis of Aryl 2-Acetamido-2-deoxy-β-D-galactopyranoside (1e–j).

4-Nitrophenyl 2-Deoxy-2-fluoroacetamido-β-D-galactopyranoside (1b). 1H NMR (500 MHz, MeOD-δ6) δ 2.26 (s, 3H); 13C NMR (125 MHz, MeOD-δ6) δ 173.22, 144.08, 126.64, 117.79, 110.09, 102.40, 77.07, 73.02, 69.58, 62.46, 54.19, 22.98; HRMS (ESI+) [M + Na]+ calcd for C14H15N3O8Na 419.0678, found 419.0678.

4-Nitrophenyl 2-Deoxy-2-trifluoroacetamido-β-D-galactopyranoside (1d). 1H NMR (500 MHz, MeOD-δ6) δ 2.03 (s, 3H); 13C NMR (125 MHz, MeOD-δ6) δ 163.52, 144.13, 126.69, 117.72, 100.04, 77.58, 71.90, 69.58, 62.46, 54.19, 22.98; HRMS (ESI+) [M + Na]+ calcd for C14H15F3N3O8Na 454.1708, found 454.1708.

4-Nitrophenyl 2-Deoxy-2-fluoroacetamido-β-D-galactopyranoside (1e). 1H NMR (500 MHz, MeOD-δ6) δ 2.27 (s, 3H); 13C NMR (125 MHz, MeOD-δ6) δ 174.18, 159.27, 153.32, 119.18, 115.48, 102.40, 77.00, 73.02, 62.47, 54.19, 22.98; HRMS (ESI+) [M + H]+ calcd for C14H15NO9F 328.1389.
nitrophenyl GlcNAc (PNP-GlcNAc) and 3-fluoro-4-nitrophenyl GlcNAc (3F4NP-GlcNAc). Final concentrations of substrates ranged from 15.6 μM to 2 mM; final concentrations of enzyme were 100 nM for the wild type and 100 nM to 2 μM for the mutants depending on their activity. Reaction volumes of 200 μL were used in all cases. The wavelengths at which measurements were taken correspond to the absorbance maxima of the aryl leaving groups, which have been previously determined as shown in Table S2. Michaelis–Menten parameters were calculated from nonlinear regression of Michaelian saturation curves using GraphPad Prism. Relative activities are reported as a ratio of the second-order rate constants of wild-type HexD versus those of its mutants.

**Brønsted Analyses.** Brønsted analyses with N-acetylgalactosamine-derived substrates with a range of leaving groups were performed as described above with the exception that reactions were conducted in quartz cuvettes (Sarstedt, volume of 200 μL, path length of 1.0 cm). Substrate concentrations ranged from 15.6 μM to 2 mM; enzyme concentrations used for each substrate are listed in Table S2. Reactions were performed in duplicate and monitored using the wavelengths listed in Table S2. Michaelian saturation curves were observed for all substrates and were calculated as described above.

**pH Profiles.** Reactions were conducted in 96-well plates as described above, using PNP-GalNAc as the substrate (final concentration between 15.6 μM and 2 mM). 50 mM citrate, 100 mM NaCl buffer was used for reactions in the pH range of 5−5.5; 50 mM phosphate, 100 mM NaCl buffer was used for reactions in the pH range of 6.0−7.5, and 50 mM CHES, 100 mM NaCl buffer was used for reactions in the pH range of 8.0−9.0. The stability of HexD at each pH was verified. Absorbance values were corrected according to the proportion of protonated and deprotonated 4-nitrophenol at each pH value. A final enzyme concentration of 100 nM was used for each reaction, and saturation kinetics were observed in all cases; Michaelis–Menten parameters were calculated from saturation curves as described above. A bell-shaped ionization curve, described by eq 1, was fitted to the data.

\[
\frac{k_{cat}}{K_M} = \frac{k_{cat, max} \times 10^{pK_a - pK_{H^+}}}{10^{2pK_a - pK_{H^+} - pK_{H^+}^2} + 10^{pK_a - pK_{H^+}} + 1}
\]

Experimental kinetic pK_a values were obtained from the pH values corresponding to the half-maximal enzyme activity on each limb of the curve.

**NMR Experiments.** NMR-based monitoring of HexD-catalyzed hydrolysis was conducted using a Bruker AvanceII 500 MHz spectrometer equipped with a 5 mm TXI inverse probe. 0.6 mg of PNP-GalNAc was dissolved in 600 μL of deuterated phosphate-buffered saline [137 mM NaCl, 2.7 mM KCl, and 10 mM Na_2HPO_4 and KH_2PO_4 in D_2O (pH 7.4)], and a t_0 spectrum was recorded. HexD was exchanged into deuterated PBS by repeated concentration using a 10 kDa molecular mass cutoff centrifugation device and subsequent

![Figure 2](https://example.com/figure2.png)

**Figure 2.** Determination of substrate preference and stereochemical outcome for HexD hydrolysis. (A) Michaelis–Menten kinetics for HexD-catalyzed hydrolysis of PNP-GalNAc (1a, ●) and PNP-GlcNAc (2a, ▲). (B) Michaelis–Menten kinetics for HexD-catalyzed hydrolysis of 3F4NP-GalNAc (1f, ●) and 3F4NP-GlcNAc (2b, ■). Error bars represent the SEM. (C) 1H NMR spectrum showing the progress of HexD hydrolysis of PNP-GalNAc. The initial product formed was β-GalNAc, which was subsequently mutarotated to α-GalNAc (500 MHz, D_2O). (D) Structures of PNP-GalNAc (1a), PNP-GlcNAc (2a), 3F4NP-GalNAc (1f) and 3F4NP-GlcNAc (2b).
resuspension; 120 μL of HexD was added to the substrate at a
cfinal concentration of 1 μM. Reaction progress was monitored
at 5 min intervals for 1 h, with a final spectrum recorded 3 h
after enzyme addition. Spectra were processed using
MestReNova version 10.

**Inhibition Studies.** Inhibition studies of HexD with Gal-
NAG-thiazoline\(^{9,36}\) were performed in 96-well plates using
PNP-GalNAc as the substrate (62.5 μM to 1 mM) with 0.0063
mg/mL enzyme. Gal-NAG-thiazoline was tested at six
concentrations ranging from 5 times above to 5 times below
the apparent \(K_d\). Data were plotted using GraphPad Prism, and
\(K_d\) values were obtained using nonlinear regression analysis
with the equation for competitive binding:

\[
K_d(\text{app}) = K_d(1 + [I]/K_i);
\]

\[
Y = V_{max}X/[K_d(\text{app}) + X].
\]

**RESULTS**

**Substrate Selectivity.** Gutermigg et al. qualitatively
showed previously that HexD was active on synthetic substrates
containing both GalNAc and GlcNAc β-linked to the leaving
group, while it was not active on those substrates containing an
α-linkage or on gluco- or galacto-derived substrates lacking an
N-acetyl group.\(^7\) We first set out to establish the detailed kinetic
parameters for HexD acting on both gluco- and galactosami-
nides substrates. Kinetic parameters were determined with
substrates containing 4-nitrophenyl (PNP) or 3-fluoro-4-
nitrophenyl (3F4NP) leaving groups (Figure 2A,B,D). Measure-
ments showed that the catalytic efficiency as reflected in the
second-order rate constant (\(k_{cat}/K_M\)) was higher for
substrates containing GalNAc than for substrates containing
Glcnac: 21-fold higher for the 4-nitrophenyl leaving group and
substrates containing GalNAc than for substrates containing
hydrolysis catalyzed by HexD: \((\text{ Equation for competitive binding:})\)

\[K_M = \frac{1 + [I]}{K_i},\]

\[V_{max} = \frac{X}{K_d + X}.
\]

Table 1. Kinetic Parameters\(^{a}\) for HexD Hydrolysis of
Galactosaminides vs Glucosaminides

| substrate            | \(K_M\) (mM) | \(k_{cat}\) (min\(^{-1}\)) | \(k_{cat}/K_M\) (min\(^{-1}\) mM\(^{-1}\)) |
|----------------------|-------------|--------------------------|----------------------------------------|
| PNP-GalNAc (1a)      | 0.0721 ± 0.0048 | 79.3 ± 1.2 | 1090 ± 75                              |
| PNP-GlcNAc (2a)      | 0.6067 ± 0.053  | 30.4 ± 0.89 | 50.1 ± 4.6                             |
| 3F4NP-GalNAc (1f)    | 0.172 ± 0.019  | 122 ± 3.9   | 707 ± 81                               |
| 3F4NP-GlcNAc (2b)    | 0.852 ± 0.054  | 92.4 ± 2.7  | 109 ± 7.6                              |

\(^{a}\)Calculated kinetic parameters represent the mean ± SEM of triplicate values.

![Figure 3](image3.png)

Figure 3. HexD activity with N-fluoroacetyl PNP-GalNAc substrates. (A) Michaelis–Menten kinetics of N-fluoroacetyl PNP-GalNAc derivative
hydrolysis catalyzed by HexD: \((\bullet)\) PNP-GalNAc (1a), \((\square)\) PNP-GalNAc-F (1b), \((\Delta)\) PNP-GalNAc-F\(_2\) (1c), and \((\bigcirc)\) PNP-GalNAc-F\(_3\) (1d). Error
bars represent the SEM. (B) Taft-like linear free energy analysis plotting \(\log(k_{cat}/K_M)\) for HexD-catalyzed hydrolysis of substrates 1a–1d vs the Taft
electronic parameter (\(\sigma^e\)) of the N-fluoroacetyl groups. (C) Structures of N-fluoroacetyl PNP-GalNAc substrates (1b–1d).

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parameters governing their hydrolysis by HexD was determined. Panel A of Figure 3 show the Michaelis–Menten plots of data for HexD acting on PNP-GalNAc (1a) and the series of fluorinated derivatives (1b–1d) (Figure 3C). Incorporation of each successive fluorne atom significantly impairs the rate of hydrolysis, with a 34-fold reduction in $k_{cat}/K_M$ for trifluorinated substrate 1d relative to that of nonfluorinated PNP-GalNAc 1a (Table 2). A plot of the log $k_{cat}/K_M$ values against the Taft electronic parameter ($\sigma^*$) of the acyl groups reveals a Taft correlation (Figure 3B). The negative slope of this correlation is consistent with the carbonyl group of the acyl group acting as a nucleophile in the enzyme-catalyzed reaction. This provides strong evidence of HexD employing a substrate-assisted hydrolysis mechanism, similar to those of other mammalian $\beta$-hexosaminidases in GH families 20 and 84.

### Establishing the Importance of Residues Predicted To Be Involved in Catalysis

We next sought to identify key active site residues involved in the substrate-assisted mechanism of HexD. By examining sequence homology across enzymes of GH family 20 (Figure S1), we identified Asp148 and Glu149 as the predicted candidate residues acting as the catalytic polarizing and acid/base residues. Site-directed mutagenesis was used to produce HexD variants in which these residues were mutated to either alanine or the type (WT) HexD using PNP-GalNAc or 3F4NP-GalNAc as the substrate. Mutation of either residue to alanine or the E149A and E149Q mutations leading to a 1.6–2.2-fold decrease in $k_{cat}/K_M$ when PNP-GalNAc was used as the substrate.

We also tested the importance of His92, which is conserved on the basis of sequence alignments with the GH20 lysosomal $\beta$-hexosaminidases (Figure S1) and is predicted to be an active site residue on the basis of structural information from the $\beta$-hexosaminidases, for HexD activity. The $k_{cat}/K_M$ for the H92A mutant was 70-fold lower than that of the wild-type enzyme, suggesting it makes an important contribution to catalysis and may reside in the enzyme active site as seen for His262 in HexA and His294 in HexB.16,17 The overall decrease in $k_{cat}/K_M$ is contributed solely by a decrease in the rate of turnover, whereas the $K_M$ is consistent with the wild-type value.

### Brønsted Linear Free Energy Analyses

To further probe the importance of Asp148 and Glu149 as catalytic residues, Brønsted linear free energy analyses were conducted for WT HexD and the D148A and E149A mutants. A panel of aryl galactosaminide substrates (1e–1j) (Figure 4C) was synthesized, and Michaelis–Menten kinetic parameters were determined for each enzyme using these substrates as well as 4-methylumbelliferyl 2-acetamido-2-deoxy-$\beta$-D-glucopyranoside (3). The logarithm of the second-order rate constant measured for turnover of each substrate was plotted against the $pK_a$ of the corresponding phenol leaving group (Figure 4A). A $\beta_{lg}(V/K)$ value of $−0.44 \pm 0.05$ was obtained for WT HexD, indicating a moderate dependence on leaving group ability. In contrast, more negative $\beta_{lg}$ values of $−0.89$ and $−0.85$ were obtained for the D148A and E149A mutant enzymes, respectively. The steeper negative slopes of both mutants relative to the WT enzyme are indicative of a stronger dependency on leaving group ability. This perturbation may stem from the inability of the mutants to stabilize the accumulation of charge in the transition state as the anomeric substituent departs, leading to a transition state in which there is greater charge development on the departing glycosidic oxygen. Alternatively, the steeper slopes observed for these $\beta_{lg}$ values could be explained by the mutations causing a change in the position of the transition state along the reaction coordinate. In any event, these data provide strong direct evidence that D148 and E149 are key active site residues involved in transition state stabilization.

### Table 2. Kinetic Parameters for Hydrolysis of PNP-GalNAc and Fluorinated Derivatives

| Substrate          | $\sigma^*$ | $K_M$ (mM) | $k_{cat}$ (min$^{-1}$) |
|--------------------|------------|------------|------------------------|
| PNP-GalNAc         | 0          | 0.176 ± 0.014 | 145 ± 3.5              |
| PNP-GalNAc-F       | 0.8        | 0.533 ± 0.017 | 156 ± 1.9              |
| PNP-GalNAc-F$_{1c}$| 2.0        | 1.020 ± 0.016 | 162 ± 0.65             |
| PNP-GalNAc-F$_{1d}$| 2.8        | 1.030 ± 0.027 | 2.52 ± 0.17            |

$^a$Calculated kinetic parameters represent the mean ± SEM of triplicate values.

### Table 3. Kinetic Parameters for Hydrolysis by Wild-Type HexD and Various Mutants

| Substrate          | $K_M$ (mM) | $k_{cat}$ (min$^{-1}$) |
|--------------------|------------|------------------------|
| WT                 | 0.172 ± 0.019 | 122 ± 3.9              |
| D148A              | 0.566 ± 0.068 | 0.461 ± 0.023          |
| D148N              | 1.73 ± 0.27  | 0.293 ± 0.027          |
| E149A              | 0.0525 ± 0.0034 | 36.1 ± 0.52         |
| E149Q              | 0.175 ± 0.031 | 89.3 ± 4.6             |
| H92A               | 0.196 ± 0.028 | 3.09 ± 0.13            |

$^a$Calculated kinetic parameters represent the mean ± SEM of triplicate values.

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The pH dependence of HexD Activity. The pH dependence of HexD activity was determined by measuring $k_{cat}/K_M$ across a pH range of 5.0–9.0. To perform these studies, we first established that HexD was stable for at least 30 min over this pH range under the conditions used for the assay. A plot of the logarithm of the second-order rate constant versus pH (Figure 4B) displays a bell-shaped profile with a pH optimum of 6.5–7.0, which is not unexpected given the subcellular localization of HexD in the nucleus and cytoplasm. The pH profile for HexD is consistent with that observed for the GH84 β-hexosaminidase OGA,31,32 which is similarly localized to the nucleus and cytoplasm. When a bell-shaped ionization curve (eq 1) was fit to the data, the kinetic $pK_a$ values of the D148 and E149 residues can be estimated by identifying the pH values corresponding to half of the maximum $k_{cat}/K_M$ value on each limb of the curve. Using this technique, the $pK_a$ values of the acidic and basic limbs were estimated to be 5.6 and 7.8, respectively. The pH dependence of activity for the E149A mutant displays a similar bell-shaped curve but has an alkaline shift in comparison to wild-type HexD with a pH optimum of 7.5. This shift suggests E149 plays a key role in catalysis but indicates at least two other residues contribute to the pH profile. The kinetic $pK_a$ values of the E149A pH curve are 6.7 for the acidic limb and 7.8 for the basic limb. A small decrease in activity and a small upward shift in pH optimum were also observed for E149A and E149Q mutants of the murine HexD (data not shown). Unfortunately, we were unable to determine the pH dependence of activity for the D148A mutant as its activity was extremely low even at the pH optimum.

Inhibition of HexD. NAG-thiazoline was first designed as a mimic of the oxazoline intermediate that forms during substrate-assisted hydrolysis for hexosaminidases,29 and this thiazoline feature has been successfully incorporated into inhibitors of a range of hexosaminidases that employ a substrate-assisted catalytic mechanism.14,16,25,33 Given the preference of HexD for galacto-configured substrates, we decided to test the inhibition of HexD by galacto-configured NAG-thiazoline (Gal-NAG-thiazoline, 4; Figure 5B). Compound 4 is an effective inhibitor of HexD-catalyzed hydrolysis of PNP-GalNAC, and the Lineweaver–Burk plot confirms an expected competitive mode of inhibition (Figure 5A). Nonlinear regression reveals the $K_i$ value to be 420 nM. A very similar value (520 nM) has been determined in preliminary studies with the murine enzyme (data not shown). Curiously, we were unable to observe any inhibition with gluco-configured NAG-thiazoline at a concentration of 1 mM despite the significant activity we observe for this enzyme toward β-N-acetylglucosaminide substrates. Inhibition of HexD by 4 provides further evidence that it employs a catalytic mechanism involving anionic assistance with formation of a transient oxazoline intermediate.

**DISCUSSION**

Glycoside hydrolases from GH families 18, 20, 56, 84, and 85 have been shown to operate through a substrate-assisted catalytic mechanism (Figure 1) with some mechanistic variations but usually involving two carbonyl residues positioned within the active site.14,15,25,32–34 One of these residues acts as a polarizing residue, hydrogen bonding with the N–H bond of the acetamido group to both appropriately orient and promote attack of the carbonyl oxygen at the anomeric center. Concurrently, a general acid/base residue, acting as a general acid in the first step of the reaction, stabilizes the developing charge on the departing glycosidic oxygen to facilitate its departure, leading to formation of the bicyclic oxazoline intermediate. The oxazoline is hydrolyzed through the near microscopic reverse of the first step where the acid/base residue acts as a base; it is considered a “near” microscopic reverse because in the first step of the mechanism the leaving
group/nucleophile is an alcohol whereas in the second step it is a water molecule.

A distinguishing feature of the substrate-assisted catalytic mechanism employed by \( \beta \)-hexosaminidases is the participation of the 2-acetamido group of the substrate in which the carbonyl oxygen acts as a nucleophile. The presence of electron-withdrawing fluorine atoms on the acetamido group of a substrate is therefore expected to decrease the level of catalysis by reducing the electron density of the carbonyl oxygen. HexD activity with substrates possessing zero, one, two, or three fluorine atoms on the acetamido group was tested, and a trend (Figure 3A) is observed whereby the rate of hydrolysis is reduced as the degree of fluorination increases. The data were subjected to Taft-like linear free energy analysis, which accounts for the influence of polar effects on a reaction rate and can be used to determine whether the acetamido group is responsible for the variability in reaction rate observed between substrates. By plotting the log \( K_{cat}/K_M \) of each substrate against the Taft electronic parameter \( \rho \) of the corresponding \( N \)-acyl substituent, we obtained the Taft reaction constant \( \rho \) (Figure 3B), revealing a negative linear correlation with an increasing degree of fluorine substitution. The negative slope of the reaction constant \( \rho = -0.49 \pm 0.11 \) suggests this residue interacts with a positively charged center in the transition state, consistent with the hypothesis that the carbonyl oxygen acts in a nucleophilic capacity to intercept the electrophilic anomeric center. The magnitude of the reaction constant is similar to that calculated for OGA \( \rho = -0.42 \pm 0.08 \) (Table 4), in which the acetamido group has been unambiguously shown to participate in catalysis. In contrast, the value observed for HexD is smaller than the reaction constant obtained for HexB \( \rho = -1.0 \pm 0.1 \), which also operates using a substrate-assisted mechanism. An important consideration, however, is that the Taft reaction constant is governed by both a steric and electronic component according to

\[
\rho = \rho^* + \delta
\]

where \( \rho^* \) represents the sensitivity of the reaction to the electronic character of the substituents and \( \delta \) represents the sensitivity of the reaction to the Taft steric coefficients of the substrates. It is, therefore, conceivable that the decrease in reaction rate with an increasing degree of fluorine substitution is due in part to an increased number of steric interactions caused by the difference in size between fluorine (atomic radius of 147 pm) and hydrogen atoms (atomic radius of 120 pm). However, though \( K_M \) is a kinetic parameter and might not always reflect substrate dissociation, particularly in cases where an enzyme-bound intermediate accumulates, the clear dependence of the first-order rate constants \( k_{cat} \) on leaving group ability suggests that the rate-determining step for these substrates with WT HexD is the ring-forming step in which the leaving group is departing. Accordingly, the relatively constant \( K_M \) values observed among these fluorinated substrates suggest that binding ability is not substantially affected by substitution with fluorine and imply that the electronic effects of the fluorine atoms are the major contributor to the reduction in hydrolysis rate for HexD.

Sequence alignments of enzymes within a CAZy family have demonstrated that the catalytic residues, most commonly two carboxylic acid residues, are highly conserved. Structural and kinetic studies have demonstrated that GH families 18 and 56 possess a DXE motif, where the aspartate and glutamate residues are the critical residues used to effect catalysis. The lysosomal \( \beta \)-hexosaminidases HexA and HexB, members of GH family 20, use a conserved DE catalytic pair with two carboxylic acid residues adjacent to each other (Figure S1), whereas GH84 enzymes, such as OGA, use a DD catalytic pair. Unusually, GH85 enzymes possess an asparagine residue in place of the aspartate in an NXE motif; the asparagine has been proposed to exist as the imidic acid tautomer to accept a hydrogen bond from the substrate amide.

The data obtained in this study through kinetic analyses of HexD mutants strongly support the assignment of Asp148 and Glu149 as the key catalytic residues. Mutation of Asp148 to either alanine or asparagine significantly impairs the activity of the enzyme even in the presence of substrates with activated leaving groups. Notably, mutation of Glu149 to alanine or glutamine has a much less pronounced effect on the rate of hydrolysis of activated substrates such as PNP-GalNAc. These data suggest Glu149 functions as the catalytic general acid/base during catalysis, because the modest reductions in activity observed for the E149 mutants with activated substrates can be rationalized by the low pK values of the phenolic leaving groups of PNP-GalNAc and 3F4NP-GalNAc (7.18 and 6.42, respectively). These leaving groups cannot benefit greatly from general acid catalysis when the microscopic pK of the general acid is greater than the pK of the leaving group. These changes in activity in association with mutating the general acid/base catalytic residue are consistent with analysis of the corresponding mutants of OGA. Analysis of the pH dependence of wild-type HexD activity reveals a bell-shaped profile (Figure 4B), with an acidic limb pK of 5.6 and a basic limb pK of 7.8. Such pH profiles are notoriously hard to dissect fully, but the simplest interpretation is the bell shape stems from titration of the two catalytic residues. In such a case, the acidic limb would reflect titration of the polarizing residue and the basic limb titration of the acid/base residue. If this interpretation is correct, the pK values of the activated substrates are lower than the kinetic pK of the general acid/base catalytic residue. Accordingly, the general acid catalysis afforded by Glu149 is inefficient because it cannot donate a proton to the departing phenol. When tested against substrates bearing worse leaving

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Table 4. Comparison of the Kinetic and Inhibition Properties of Hexosaminidases HexA and HexB (GH20), OGA (GH84), and HexD (GH20)

| Substrate           | HexD (µM) | HexA (µM) | HexB (µM) | OGA (µM) |
|---------------------|-----------|-----------|-----------|----------|
| GlcNAc/GalNAc selectivity | 0.048<sup>a</sup> | 8.6<sup>b</sup> | 8.6<sup>b</sup> | >50<sup>d</sup> |
| Taft reaction constant (ρ) | −0.49 ± 0.1<sup>e</sup> | ND<sup>f</sup> | −1.0 ± 0.1<sup>f</sup> | −0.42 ± 0.08<sup>f</sup> |
| β<sub>0</sub> (V/K) | −0.44 ± 0.05<sup>g</sup> | ND<sup>f</sup> | −0.29<sup>i</sup> | −0.11 ± 0.01<sup>i</sup> |
| K<sub>i</sub> for NAG-thiazoline (µM) | >1000<sup>j</sup> | 0.27<sup>j</sup> | 0.19<sup>j</sup> | 0.070<sup>j</sup> |
| K<sub>i</sub> for Gal-NAG-thiazoline (µM) | 0.42<sup>a</sup> | 0.82<sup>e</sup> | 0.32<sup>d</sup> | poor<sup>i</sup> |

<sup>a</sup>From this work. <sup>b</sup>From ref 14. <sup>c</sup>From ref 28. <sup>d</sup>From ref 41. <sup>e</sup>From ref 42. <sup>f</sup>From ref 43. <sup>g</sup>From ref 44. <sup>h</sup>Not determined.

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groups, the catalytic ability of the E149A mutant is significantly impaired relative to that of wild-type HexD, as demonstrated by the steeper negative correlation observed in the Brønsted analyses (Figure 4A). It should be pointed out here, however, that the pH dependence for the E149A mutant also gave a bell-shaped profile, with an alkaline shift relative to the profile for wild-type HexD. This indicates there is titration of at least two residues other than Glu149 contributing to the pH dependence; whether this is true only in the case of the Glu149 mutant HexD, where the chemistry of the active site environment will have been perturbed by removal of the charged residue or is also true for the wild-type enzyme is difficult to know in the absence of structural data. The low activity observed for the D148A HexD mutant, even with activated substrates, as well as the steep negative correlation observed in the Brønsted analyses, is consistent with this residue playing a role in polarizing the N–H group on the acetamido group of the substrate to aid nucleophilic attack by the carbonyl oxygen at the anomic position. The impact of this mutation is also reminiscent of the effect of analogous mutations made in OGA and HexB, where impairments of 6700- and 2400-fold have been observed, respectively.27,31 The assignment of Asp148 as the polarizing residue and Glu149 as the general acid/base is consistent with previous conclusions for analogous residues in the DE motif of GH20 enzymes, such as the lysosomal β-hexosaminidases,27 and the DD motif of GH84 enzymes, such as OGA.31

The inhibitor NAG-thiazoline was originally designed to mimic the transient oxazoline intermediate that is formed during substrate-assisted catalysis (Figure 1).29 Interestingly, however, Whitworth and colleagues have since demonstrated using linear free energy relationships that NAG-thiazoline is a mimic of the transition state for human OGA.56 Incorporation of the thiazoline motif during design of inhibitors has proven to lead to effective inhibitors of a range of hexosaminidases that employ a substrate-assisted mechanism.14,16,17,25,33 On the basis of the mechanistic evidence obtained from kinetic analyses in this study, we expected the galacto-configured derivative of NAG-thiazoline, Gal-NAG-thiazoline (4), would inhibit HexD. We observe this to be the case, with Gal-NAG-thiazoline displaying potent competitive inhibition of HexD with a \( K_i \) value of 420 nM, which provided further experimental support for HexD using substrate-assisted catalysis.

Notably, functionalized derivatives of NAG-thiazoline have been developed as potent and selective inhibitors of the lysosomal β-hexosaminidases HexA/B and OGA, which have proven to be useful tools for interrogating the biological role of these enzymes.14,23,37,38 Unfortunately, Gal-NAG-thiazoline has limited use as a chemical tool for studying the physiological role played by HexD because it also inhibits the lysosomal β-hexosaminidases (Table 4). These studies, however, lay important groundwork for the rational design of selective inhibitors of HexD. The inhibition of HexD by Gal-NAG-thiazoline suggests this inhibitor scaffold may be suitable for adaptation and may allow creation of compounds that will selectively inhibit HexD without impacting the activity of lysosomal β-hexosaminidases. Given the limited tolerance of HexD toward glucose-based substrates, combined with its localization in the nucleus and cytoplasm, the enzyme likely serves a biological role that is distinct from that of the lysosomal β-hexosaminidases and OGA. Accordingly, the development of potent and specific probes will be useful for investigating the potential physiological function of HexD.

**CONCLUSIONS**

In summary, we have performed a detailed kinetic characterization of HexD using a combination of kinetic studies with wild-type and mutant enzymes, linear free energy analyses, NMR experiments, and inhibition studies. HexD is a retaining glycosidase, which operates through a substrate-assisted hydrolysis mechanism consistent with that of other mammalian β-hexosaminidases in GH families 20 and 84. The enzyme shows a moderate preference for galactosaminides over glucosaminides and operates optimally in the pH range of 6.5–7.0. We have dissected the catalytic roles played by Asp148 and Glu149, which we propose act as the polarizing residue and general acid/base, respectively. Furthermore, HexD is inhibited by Gal-NAG-thiazoline, which is consistent with the use of a substrate-assisted mechanism, and this observation will provide a starting point for further inhibitor development. Despite the mechanistic similarities it shares with the lysosomal β-hexosaminidases, which belong to family GH20, HexD is distinct as it localizes to the nucleus and cytoplasm and shows only weak activity against glucosaminides. We anticipate the insights gained from this study will assist in the development of potent and selective inhibitors of HexD as well as other research tools, which will serve to improve our understanding of the physiological role played by HexD in human health and disease.

**ASSOCIATED CONTENT**

1. Supporting Information
2. **Biochemistry** 2016, 55, 2735

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Notes
The authors declare no competing financial interest.

Abbreviations

GlCNac, N-acetylglucosamine; GalNAC, N-acetylgalactosamine; HexA, hexosaminidase A; HexB, hexosaminidase B; HexS, hexosaminidase S; OGA, O-GlcNAcase; HexD, hexosaminidase D; GH, glycoside hydrolase; CAzy, Carbohydrate Enzyme.
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