Pharmacological Analysis of *Drosophila melanogaster* γ -Secretase with Respect to Differential Proteolysis of Notch and APP

Casper Groth, W. Gregory Alvord, Octavio A. Quiñones, and Mark E. Fortini

Department of Biochemistry and Molecular Biology, Thomas Jefferson University, Philadelphia, Pennsylvania (C.G., M.E.F.); Data Management Services, Inc., National Cancer Institute, Frederick, Maryland (W.G.A., O.A.Q.)

Received November 16, 2009; accepted January 4, 2010

ABSTRACT

The γ -secretase aspartyl protease is responsible for the cleavage of numerous type I integral membrane proteins, including amyloid precursor protein (APP) and Notch. APP cleavage contributes to the generation of toxic amyloid β peptides in Alzheimer's disease, whereas cleavage of the Notch receptor is required for normal physiological signaling between differentiating cells. Mutagenesis studies as well as in vivo analyses of Notch and APP activity in the presence of pharmacological inhibitors indicate that these substrates can be differentially modulated by inhibition of mammalian γ -secretase, although some biochemical studies instead show nearly identical dose-response inhibitor effects on Notch and APP cleavages. Here, we examine the dose-response effects of several inhibitors on Notch and APP in *Drosophila melanogaster* cells,

which possess a homogeneous form of γ -secretase. Four different inhibitors that target different domains of γ -secretase exhibit similar dose-response effects for both substrates, including rank order of inhibitor potencies and effective concentration ranges. For two inhibitors, modest differences in inhibitor dose responses toward Notch and APP were detected, suggesting that inhibitors might be identified that possess some discrimination in their ability to target alternative γ -secretase substrates. These findings also indicate that despite an overall conservation in inhibitor potencies toward different γ -secretase substrates, quantitative differences might exist that could be relevant for the development of therapeutically valuable substrate-specific inhibitors.

The γ -secretase complex is a multisubunit aspartyl protease that executes the intramembrane proteolysis of certain type I integral membrane proteins, including amyloid precursor protein (APP), the Notch receptor, cluster of differentiation 44 (CD44), erythroblastosis oncogene B-4 (ErbB4), and Neuregulin (Selkoe and Wolfe, 2007; reviewed in McCarthy et al., 2009). Cleavage of APP by γ -secretase also contributes to the generation and secretion of amyloid- β pep-

tide, a major constituent of the neurotoxic amyloid plaques found in Alzheimer's disease. In the case of Notch signaling, which regulates a diverse array of developmental processes in many organisms (reviewed in Kopan and Ilag, 2009; Tien et al., 2009), γ -secretase-mediated cleavage of the Notch receptor is a key step in its activation and signal transduction. Consequently, the development of pharmacological compounds to treat Alzheimer's disease by reducing γ -secretase cleavage of APP is complicated by the need to avoid adverse effects on Notch and other physiologically essential γ -secretase substrates.

Despite the overall mechanistic similarity between APP and Notch cleavage by γ -secretase, subtle differences might exist that could be exploited therapeutically. Several studies involving mutationally altered forms of γ -secretase, including Alzheimer's disease-associated mutant forms, argue that

This work was supported by National Institutes of Health National Institute of General Medical Sciences [Grant GM087650]; Data Management Services contract HHSN2612008000016C; the Department of Biochemistry and Molecular Biology, Thomas Jefferson University; and the Intramural Research Program of the National Institutes of Health, National Cancer Institute, Center for Cancer Research.

Article, publication date, and citation information can be found at <http://molpharm.aspetjournals.org>.
doi:10.1124/mol.109.062471.

ABBREVIATIONS: APP, amyloid precursor protein; A β 42, amyloid β 42 peptide; PS1, Presenilin-1; PS2, Presenilin-2; Aph-1, anterior pharynx defective-1; APPL, amyloid precursor protein-like; APLP, amyloid precursor-like protein; ADAM, A Disintegrin And Metalloproteinase enzyme; NTF, N-terminal fragment; CTF, C-terminal fragment; NICD, Notch intracellular domain; AICD, amyloid precursor protein intracellular domain; DFK167, difluoro-ketone peptidomimetic inhibitor-167; DAPT, *N*-[*N*-(3,5-difluorophenylacetyl)-*L*-alanyl]-*S*-phenylglycine *t*-butyl ester; CpnE, compound E [(*S,S*)-2-[2-(3,5-difluorophenyl)acetylaminol]-*N*-(1-methyl-2-oxo-5-phenyl-2,3-dihydro-1*H*-benzo[*e*][1,4]diazepin-3-yl)-propionamide]; DBZ, dibenzazepine [(*S,S*)-2-[2-(3,5-difluorophenyl)acetylaminol]-*N*-(5-methyl-6-oxo-6,7-dihydro-5*H*-dibenzo[*b,d*]azepin-7-yl)propionamide]; BB94, batimastat; GM6001, *N*-[(2*R*)-2-(hydroxamidocarbonylmethyl)-4-methylpentanoyl]-*L*-tryptophan methylamide; Tricine, *N*-[2-hydroxy-1,1-bis(hydroxymethyl)ethyl]glycine.

these variants possess differential proteolytic activities toward Notch and APP (Capell et al., 2000; Kulic et al., 2000; Nakajima et al., 2000; Zhang et al., 2000; Moehlmann et al., 2002). Lewis et al. (2003) and Schroeter et al. (2003), using pharmacological approaches in which Notch and APP intramembrane proteolysis was monitored in parallel to determine their sensitivities to different γ -secretase inhibitors, reported nearly identical dose-dependent inhibitor effects on both substrates, whereas Yang et al. (2008) found that some inhibitors can partially discriminate between the APP and Notch cleavages. Preclinical animal studies on these compounds have generally revealed significant toxic side effects attributable to impaired Notch signaling in the mouse gut and immune system (Milano et al., 2004; van Es et al., 2005). Proprietary sulfonamide-based γ -secretase inhibitors have been reported to show significantly higher selectivity toward APP cleavage relative to Notch in cell-based assays and animal models (Barten et al., 2005; Best et al., 2007; Cole et al., 2009; Pu et al., 2009). Overall, the results of previous mutational and pharmacological studies suggest that different assay conditions, drug dosage regimens, or other experimental variations might account for the contradictory outcomes of these analyses.

An additional complication is that mammalian γ -secretase is heterogeneous, with functionally redundant genes encoding two different Presenilins (PS1 and PS2) and two Aph-1 proteins (Aph-1a and Aph-1b), generating several different biologically active complexes (Shirotani et al., 2004). The alternative subunits of mammalian γ -secretase show differential and partially overlapping tissue expression patterns (Hebert et al., 2004), suggesting that different complexes might exhibit distinct biological activities in specific tissues. Indeed, biochemical studies have shown that PS1-containing γ -secretase shows higher activity regarding APP cleavage compared with PS2-containing γ -secretase (Lai et al., 2003), and although Aph-1b-containing complexes contribute significantly to A β production in the brain (Serneels et al., 2009), genetic elimination of Aph-1b in mice leads to behavioral deficits associated with schizophrenia (Dejaegere et al., 2008). Thus, results obtained from previous studies of γ -secretase inhibitor effects on APP versus Notch could, in part, reflect the activities of functionally heterogeneous γ -secretase complexes in the various mammalian cell lines and assay systems employed.

To address this issue, we investigated the substrate specificity of several well characterized γ -secretase inhibitor compounds using *Drosophila melanogaster* cells, in which γ -secretase has a homogeneous composition. The fly genome possesses a single gene for each of the four γ -secretase components Presenilin, Nicastrin, Aph-1, and presenilin enhancer-2 (Hu and Fortini, 2003). Selecting a panel of γ -secretase inhibitors that target different regions of the complex, we performed dose-response studies on their ability to inhibit the intramembrane cleavage of *D. melanogaster* Notch and APP in *D. melanogaster* S2 cells. In general, the effective dose range and rank potencies of the different inhibitors were found to be nearly identical toward both substrates, in agreement with two mammalian studies that performed a similar parallel substrate analysis (Schroeter et al., 2003). However, we also observed that two of the γ -secretase inhibitors (DAPT and DFK167) show a modest but statistically significant discrimination be-

tween Notch and APP cleavages in *D. melanogaster*, similar to one mammalian study that reported increased substrate specificity toward APP versus Notch for DAPT (Yang et al., 2008). Our findings also demonstrate that γ -secretase enzyme sensitivity to several inhibitors that were developed to target mammalian γ -secretase is conserved in the invertebrate *D. melanogaster*, emphasizing the high degree of evolutionary conservation in the regulatory features of this proteolytic mechanism.

Materials and Methods

S2 Cell Culture Studies. The *APPL* gene was obtained by polymerase chain reaction from an embryonic cDNA library. An epitope-tagged form was constructed by inserting a V5-tag at its C terminus in the vector pMT/V5-His (Invitrogen, Carlsbad, CA), cotransfected into S2 cells with pCoHygro using Effectene, and subjected to Hygromycin B antibiotic selection (Invitrogen) to establish the metal-inducible stable cell line S2-MT-APPL. The Notch cell line S2-MT-Notch, S2 cell culture conditions, and preparation of cell extracts for immunoblot analysis were as described previously (Fehon et al., 1990). Notch lysates were resolved on 3 to 8% Tris-acetate gels, whereas APPL lysates were resolved on 12% Bis-Tris or 16.5% Tris-Tricine gels (Bio-Rad Laboratories). Immunoblots were probed with 1:5000 mouse anti-V5 mAb (Invitrogen), 1:1000 mouse anti-Notch mAb C17.9C6 (Fehon et al., 1990), and 1:2000 mouse anti- β -tubulin mAb (Developmental Hybridoma Bank, University of Iowa, Iowa City, IA). Protein bands were visualized using Kodak BioMax XAR film (Carestream Health, Rochester, NY) and quantified using Photoshop CS3 (Adobe Systems, Mountain View, CA).

Pharmacological Inhibition of γ -secretase Activity. Commercially available γ -secretase inhibitors DFK167 (MP Biomedicals), DAPT, DBZ, and Compound E (Calbiochem, San Diego, CA) were used at the concentrations indicated in the text and figure legends. For each inhibitor, pilot experiments were performed with different drug concentrations ranging from 0.1 nM to 150 μ M to determine the effective linear range and maximal inhibition dose for each compound (see Figure 4 legend). Inhibitors were added at the required concentrations to the S2 cell medium upon induction of Notch or APPL expression, 6 h before protein harvesting. For each sample, the same inhibitor was also included at the corresponding concentration in the lysis buffer for protein extraction and immunoblot analysis.

Statistical Analysis. Data were analyzed using a four-parameter nonlinear logistic regression model to determine IC₅₀ values, standard errors, and 95% confidence intervals (DeLean et al., 1978; Pinheiro and Bates, 2002) with S, S-Plus, and R software (TIBCO Software, Palo Alto, CA). Inhibitor potencies were calculated using standard nonlinear mixed-effects modeling techniques (Pinheiro and Bates, 2002). The regression model used was $Y \sim ((A - D)/(1 + (X/C)^B)) + D$, where Y is the response in relative intensity; X is the concentration in nanomoles; A and D are the lower and upper asymptotes, respectively; C is the concentration giving a response halfway between A and D , referred to as IC₅₀; and B is the "slope" factor corresponding to slope of the logit-log plot, when X is given by natural logarithms; specifically, $dY/d \ln(X) = [(D - A) \cdot B]/4$ when $X = C$, or $d \text{ logit } \{(Y - D)/(A - D)\}/d \ln(X) = B$.

Results

***D. melanogaster* APPL Is Cleaved by γ -Secretase to Generate a Released Intracellular Domain.** APP is a member of a multiprotein family with a complex evolutionary history. True APP orthologs are encoded by a single gene in mammals (Coulson et al., 2000) and duplicate genes in the zebrafish (Musa et al., 2001). More distantly related family

members include human APLP1 and -2, *D. melanogaster* APPL, and *C. elegans* APL-1 (reviewed in Coulson et al., 2000). In mammals, all three APP/APLP family members undergo similar processing by α -, β -, and γ -secretase (Eggert et al., 2004) (Fig. 1A). Although *D. melanogaster* APPL is subject to ectodomain shedding (Luo et al., 1990), it has not been directly demonstrated to be processed via proteolytic pathways similar to those of mammalian APP, including intramembrane proteolysis by endogenous *D. melanogaster* γ -secretase.

To examine APPL biochemical processing, we generated a stable inducible *D. melanogaster* S2 cell line expressing APPL with a V5-tag fused to its C terminus and analyzed APPL cleavage products on immunoblots after treatments with inhibitors against α - and γ -secretase. APPL holoprotein was readily detected under all conditions as two bands of differing intensities (Fig. 2A, lane 2) that might correspond to immature and glycosylated APPL species (Vingtdeux et al., 2005). Cells treated with the γ -secretase inhibitor Compound E (CpnE) revealed an overaccumulation of two low-molecu-

lar-mass APPL C-terminal fragment (CTF) bands of 14 and 16 kDa (Fig. 2B, topmost blot, lanes 3 and 5). These APPL CTFs were also found in lysates from untreated cells at longer exposures (Fig. 2B, middle blot, lanes 2 and 4), implying that the APPL CTFs are rapidly degraded in S2 cells. Inhibiting γ -secretase activity interferes with the turnover of these fragments, leading to their accumulation in a manner similar to that for processing of mammalian APP (Xia et al., 1998). To further analyze the APPL CTFs, we treated cells with combinations of α - and γ -secretase inhibitors and resolved the products using 16.5% Tris-Tricine SDS-polyacrylamide gel electrophoresis, which enhances the separation of low-molecular-mass species (Fig. 2C, lanes 2–5). With this method, three distinct APPL CTF bands were detected in cells treated with γ -secretase inhibitor for 6 h, termed APPL CTFa, -b, and -c (Fig. 2C, topmost blot, lane 4). Cells treated with both α - and γ -secretase inhibitors displayed a slight but reproducible accumulation of APPL CTFb and concomitant decrease in CTFc compared with γ -secretase inhibitor-treated cells alone (Fig. 2C, topmost blot, lanes 4 and 5).

Fig. 2. Processing of *D. melanogaster* APPL. Immunoblot analysis of epitope-tagged APPL expressed in *D. melanogaster* S2 cells treated with different inhibitor compounds that target either γ -secretase, endosome acidification, or metalloprotease activity (see *Materials and Methods*). Lane numbers, expression constructs, and drug combinations are indicated above each immunoblot. A, full-length APPL is processed into CTFs that can be detected more readily by preventing their processing by γ -secretase using the inhibitor CpnE. B, APPL also releases a highly unstable γ -secretase-dependent fragment (AICD) that can be detected in longer exposures. C, extended gel electrophoresis identifies three APPL CTFs (a, b, and c) with CTFc generation requiring metalloprotease activity. ConA, endosome acidification inhibitor Concanamycin A; Epo, proteasome inhibitor epoximycin; GM6001, α -secretase metalloprotease inhibitor. β -Tubulin serves as the loading control in all immunoblots.

Longer treatments for 24 h with CpnE resulted in a sharp reduction in APPL CTFa and CTFb, accompanied by highly elevated levels of CTFc (Fig. 2C, third blot, lane 4), whereas CTFb showed a preferential accumulation compared with CTFc when cells were treated simultaneously with α - and γ -secretase inhibitors (Fig. 2C, third blot, lane 5). The ability of the α -secretase metalloprotease inhibitor GM6001 to reduce levels of the APPL CTFc fragment at both time points indicates that generation of this product involves α -secretase activity in addition to γ -secretase.

Mammalian APP metabolism is also regulated by endocytic trafficking. Interfering with the normal process of endosomal acidification has been shown to cause accumulation of APP CTFs in multivesicular bodies of the endosomal compartment (Vingthong et al., 2007). Other studies have implicated the proteasome in regulating the turnover of γ -secretase-generated intracellular domain fragments (Tomita et al., 2006). In an effort to detect directly the *D. melanogaster* APPL AICD fragment, analogous to the released intracellular AICD fragment from mammalian APP, we treated S2 cells with concanamycin A and epoxomicin, inhibitors of endosome acidification and proteasomal degradation, respectively. In cells treated with both compounds, we detected an ~8-kDa APPL CTF fragment with the predicted size of the putative AICD (Fig. 2B, middle blot, lane 4). Production of this fragment was blocked by addition of CpnE to the concanamycin A/epoxomicin treatment regimen, confirming that this fragment is generated by γ -secretase-mediated proteolysis and thus corresponds to the APPL AICD (Fig. 2B, middle blot, lane 5).

Dose-Response Analysis of *D. melanogaster* APPL and Notch Intramembrane Cleavage with γ -Secretase Inhibitors. To investigate whether γ -secretase inhibitors

can differentially influence APPL and Notch intramembrane proteolysis in *D. melanogaster*, we performed dose-response studies of these two substrates in S2 cells. Metal-inducible transgenic S2 lines expressing full-length Notch (Fehon et al., 1990) or APPL (this study; see *Materials and Methods*) were incubated for 6 h in the presence of different concentrations of γ -secretase inhibitors. Cells were subsequently lysed in EDTA-containing hypotonic buffer for 30 min, which promotes ectodomain shedding of γ -secretase substrates and efficiently converts the substrate pool to the membrane-bound CTF forms that are optimal γ -secretase cleavage precursors (Rand et al., 2000). Cell lysates were resolved on SDS-polyacrylamide gel electrophoresis protein gradient gels and immunoprobed with antibodies recognizing either the Notch intracellular domain or the tagged intracellular domain of expressed APPL.

We have previously used this assay to study the role of γ -secretase core components in *D. melanogaster* with Notch receptor cleavage as the measure of γ -secretase activity (Hu et al., 2002; Hu and Fortini, 2003). In this assay, the Notch intracellular fragment generated by γ -secretase proteolysis (NICD; Fig. 3A) is sufficiently stable to be detected by immunoblot analysis as a distinct fragment of which production is blocked when cells are treated with high doses of γ -secretase inhibitors (Fig. 3B, lanes 2 and 4). Previous characterization of *D. melanogaster* Notch biochemical processing (Hu et al., 2002) has established that the three comigrating fragments in the ~100-kDa mobility range correspond to the membrane-bound CTFs generated by the extracellular furin-like and ADAM cleavages of Notch (Fig. 3B, upper bands in enlarged panel) and the released NICD fragment generated by γ -secretase cleavage (Fig. 3B, lower band in enlarged panel). In addition, a faster migrating species that is not

Fig. 3. Relationship of metalloprotease and γ -secretase Notch cleavage in *D. melanogaster* cells. **A**, the Notch receptor is initially synthesized as an ~300-kDa precursor that is processed by furin-like enzyme(s) in the *trans*-Golgi compartment. This cleavage leads to the production of a heterodimeric Notch receptor, which is further processed at the cell surface in a ligand-dependent manner by ADAM and γ -secretase. The ADAM cleavage occurs at an extracellular site and removes the Notch ectodomain, whereas γ -secretase cleavage (GS) occurs at multiple positions of the Notch transmembrane domain, leading to the release of NICD that functions as a nuclear transcriptional regulator. **B**, immunoblot analysis of Notch biochemical processing after treatment of Notch-expressing S2 cells with pharmacological inhibitors metalloproteases involved in Notch ectodomain shedding (BB94), γ -secretase Notch cleavage (CpnE), and both processes together (BB94 and CpnE). Expression constructs and drug combinations are shown above the immunoblot. The panel at bottom is an enlargement of the fragments detected in the ~100-kDa range, indicating the membrane-bound fragments produced by the extracellular furin-like and ADAM cleavages, including the putative heterodimeric Notch CTF generated by furin-like cleavage in the *trans*-Golgi (put het CTF; upper right arrow), the Notch extracellular truncation (NEXT) fragment generated by ADAM-mediated proteolysis (lower right arrow), and the NICD fragment produced by γ -secretase cleavage of NEXT (left arrow). The asterisk indicates a non-surface-derived Notch breakdown product that is often detected in Notch-expressing S2 cells (Hu et al., 2002).

derived from the cell-surface pool of Notch and represents a breakdown product is also observed in S2 cells (Hu et al., 2002) (Fig. 3B, asterisk in enlarged panel).

Because the corresponding γ -secretase-generated APPL AICD fragment was found to be highly unstable (Fig. 2B, middle blot, lanes 2 and 4), it could not be used reliably to monitor γ -secretase activity in determining inhibitor dose-response curves. We instead monitored APPL CTF levels, because the APPL CTFs are the direct stoichiometric precursors for the production of AICD. Indeed, mammalian APP CTF levels and AICD production exhibit a direct inverse relationship (Lewis et al., 2003), validating this approach. We therefore quantified the amount of APPL CTF α , the most discrete APPL CTF product detected on immunoblots (Fig. 2C, topmost blot, lanes 2–5), as an indicator of γ -secretase activity toward *D. melanogaster* APPL.

For the dose-response studies, we selected a panel of four γ -secretase inhibitors that interact directly with the γ -secretase complex and target different functional domains of the complex (Fig. 1B). DFK167 binds to the catalytic site of the Presenilin aspartyl protease subunit (Wolfe et al., 1998), whereas CpnE and DBZ target the N-terminal Presenilin fragment (Fuwa et al., 2007) and DAPT targets the C-terminal Presenilin fragment (Morohashi et al., 2006). Representative immunoblots for the Notch and APPL dose-response studies with the γ -secretase inhibitors DBZ, CpnE, DAPT, and DFK167 are shown in Fig. 4A. Increasing concentrations of these compounds administered to APPL- or Notch-expressing cells leads to the progressive accumulation of APPL CTF fragments and a decrease in NICD production in a strictly dose-dependent manner (Fig. 4A). It is worth noting that the accumulation of APPL CTFs reached a plateau at high inhibitor concentrations, indicating that the maximal inhibitory concentration for each drug was reached. The data obtained were used to plot dose-response curves and derive IC₅₀ values for the proteolysis of APPL and Notch (Fig. 4B). Both substrates exhibit the same rank order of potencies for all four inhibitors, DBZ being most potent, followed by CpnE, DAPT, and finally DFK167. It is noteworthy that DBZ and CpnE, the two most potent drugs, showed no apparent discrimination in blocking APPL and Notch cleavages, displaying statistically equivalent IC₅₀ values for both substrates ($p > 0.05$). In contrast, both DAPT and DFK167 display an approximately 4-fold selectivity in their cleavage preferences ($p < 0.05$), DAPT showing an increased effectiveness in blocking APPL cleavage relative to Notch proteolysis, whereas DFK167 inhibited Notch cleavage more effectively than APPL proteolysis in this *D. melanogaster* S2 assay (Fig. 4C).

Discussion

In this study, we demonstrate that APPL, the *D. melanogaster* ortholog of the Alzheimer's disease-associated protein APP, is rapidly metabolized and processed by α - and γ -secretase activities in *D. melanogaster* S2 cells. The transmembrane domain and adjacent regions of APPL have not been conserved in evolution, and no sequence homologous to the A β region of mammals is present in *D. melanogaster* APPL (Coulson et al., 2000). Our finding that APPL is nevertheless processed by γ -secretase supports the notion that its three-dimensional structure rather than specific residues near the cleavage site(s) renders it accessible to γ -secretase proteoly-

sis (Tanii et al., 2006). Consistent with this idea, overexpression in *D. melanogaster* of the APPL holoprotein, or the portion of APPL analogous to the A β peptide region, produced pathological lesions resembling those in Alzheimer's disease, including age-dependent formation of amyloid deposits, neurotoxic effects, and behavioral deficits (Carmine-Simmen et al., 2009).

The biology of mammalian γ -secretase is complicated by the fact that this multisubunit enzyme is heterogeneous in composition; duplicate genes encode both the Presenilin and Aph-1 subunits (Shirotani et al., 2004). The different complexes have poorly understood and potentially complex expression patterns, with overlapping yet distinct cell- and tissue-specific components (Hebert et al., 2004). Furthermore, the different complexes exhibit distinctive enzymatic activities regarding APP cleavage. Presenilin-2-containing complexes show a reduced ability to process APP compared with Presenilin-1-containing complexes (Lai et al., 2003), and both Presenilin-2- and Aph-1B-containing complexes exhibit a bias toward producing the more amyloidogenic A β 42 peptide (Placanica et al., 2009; Serneels et al., 2009). To complicate matters further, individual cells might express different γ -secretase complexes that are in equilibrium with one another (Placanica et al., 2009) and subject to strict regulatory control (Thinakaran et al., 1997). The functional heterogeneity in mammalian γ -secretase is a potential confounding factor in studies on the selectivity of various γ -secretase inhibitors with respect to the intramembrane proteolysis of alternative substrates, a critical issue in the search for inhibitors that can therapeutically reduce A β peptide generation and amyloid plaque formation but avoid toxic side effects on Notch and other substrates.

In the current study, we circumvented the problem of γ -secretase heterogeneity by performing dose-response inhibitor studies in parallel on APP and Notch in *D. melanogaster* cells, which possess a homogeneous γ -secretase complex. The fruit fly genome encodes a single variant for each of the four core γ -secretase components Presenilin, Nicastrin, Aph-1, and presenilin enhancer-2, and functional reconstitution studies have demonstrated that all of these components are required for mature γ -secretase assembly and proteolytic activity in *D. melanogaster* (Hu and Fortini, 2003; Takasugi et al., 2003). Expressing full-length APPL and Notch proteins in this *D. melanogaster* assay, we performed dose-response analyses to compare the potencies of four different γ -secretase inhibitors in blocking cleavage of APPL versus Notch. Our results indicate that all of the inhibitors generally have similar effects on APPL and Notch cleavage, including identical rank orders of potency toward each substrate and identical IC₅₀ values for two of the inhibitors, namely CpnE and DBZ, which interact with the N-terminal fragment of Presenilin (Fuwa et al., 2007). Despite the overall conservation in inhibitor activity toward APPL and Notch γ -secretase-mediated cleavage in *D. melanogaster*, two other inhibitors did show modest differences, suggesting that substrate-specific differences in cleavage exist and might be subject to pharmacological targeting. Both DAPT and DFK167, which target the C-terminal fragment of Presenilin and the catalytic site of γ -secretase, respectively (Wolfe et al., 1998; Morohashi et al., 2006), display quantitative differences in their ability to block APPL versus Notch cleavage in *D. melanogaster*. For DAPT, when APPL cleavage is ~50% inhibited, Notch cleav-

age is only slightly inhibited; conversely, for DFK167, ~50% inhibition of Notch is accompanied by relatively minor effects on APPL proteolysis. The increased potency of DAPT in blocking APPL cleavage as opposed to Notch proteolysis in *D.*

melanogaster S2 cells is similar to effects of DAPT in human HEK293 cells, as reported in a recent study (Fuwa et al., 2007). Although the IC₅₀ differences we observed with DAPT and DFK167 regarding APPL versus Notch proteolysis are

Fig. 4. Dose-response analysis of Notch and APPL cleavage sensitivity to γ -secretase inhibitors. A, representative examples of immunoblots used for dose-response studies of Notch and APPL with four different γ -secretase inhibitors. The substrate under analysis is shown in the black box in the upper left corner of each immunoblot; the inhibitor drug used is shown at upper left above each immunoblot set, with drug concentrations indicated above each lane. Hypotonic lysates from S2 cells were resolved on 3 to 8% Tris-acetate or 16.5% Tris-Tricine gels and immunoprobed with antibodies to the C termini of the expressed constructs (see *Materials and Methods*). Increasing drug concentrations led to a progressive accumulation of APPL CTFs and a progressive decrease in Notch NICD levels. Because of difficulties in detecting the highly unstable APPL AICD, immunoblot quantifications to establish the pharmacological potencies of inhibitors are based on CTFa abundance for the APPL studies, whereas NICD levels were monitored for the Notch dose-response studies. Replicate numbers of dose-response immunoblots performed for each substrate/inhibitor pair were as follows: Notch/DBZ (5), Notch/CpnE (5), Notch/DAPT (7), Notch/DFK167 (7), APPL/DBZ (4), APPL/CpnE (4), APPL/DAPT (4), and APPL/DFK167 (5). B, dose-response curves for Notch and APPL are shown in black and orange, respectively, for the γ -secretase inhibitors DBZ, CpnE, DAPT, and DFK167, as indicated. The tested concentration range for each compound analyzed was as follows: DBZ, 0.1–250 nM; CpnE, 1–1000 nM; DAPT, 25–2500 nM; DFK167, 1–150 μ M. C, the estimated IC₅₀ values show that DAPT and DFK167 exhibit modest selectivity in blocking Notch and APPL intramembrane proteolysis. Notch and APPL IC₅₀ estimates for DAPT and DFK167 differ statistically ($p < 0.05$). The two most potent inhibitors, DBZ and CpnE, do not discriminate between the Notch and APPL cleavages, their IC₅₀ estimates being statistically equivalent ($p > 0.05$). The IC₅₀ parameter estimates are reported along with their standard errors. Replicate numbers of dose-response quantitative datasets used for IC₅₀ calculations for each substrate/inhibitor pair were as follows: Notch/DBZ (3), Notch/CpnE (3), Notch/DAPT (2), Notch/DFK167 (3), APPL/DBZ (3), APPL/CpnE (2), APPL/DAPT (2), and APPL/DFK167 (3).

modest, our results suggest that some aspects of γ -secretase activity might be modulated by the action of small compounds to yield substrate-specific outcomes. Our findings, together with similar results from some mammalian γ -secretase inhibitor studies, lend credence to the idea that further development of compounds that can partially inhibit human APP cleavage, reducing A β load without deleterious toxic effects on Notch signaling, could offer compelling therapeutic benefits in Alzheimer's disease (Selkoe, 2001).

The tertiary structure of γ -secretase is still poorly understood, but recent findings suggest that the C-terminal fragment of the Presenilin aspartyl protease subunit is closely associated with the Aph-1 subunit within the active complex (Steiner et al., 2008). Indeed, these two subunits might be sufficiently tightly linked that mammalian Aph-1 isoforms can differentially regulate γ -secretase through allosteric effects on Presenilin (Serneels et al., 2009). The possibility that Aph-1 might regulate γ -secretase activity makes it an especially attractive potential drug target with the aim of developing drugs with higher discriminatory specificities toward APP rather than Notch proteolysis. In this context, incorporating our *D. melanogaster* cell-based γ -secretase activity assay in new inhibitor screening and evaluation protocols might prove useful because of the reduced regulatory complexity and uniform subunit composition of the *D. melanogaster* γ -secretase complex.

Acknowledgments

We thank Spyros Artavanis-Tsakonas (Harvard University) for providing the stable Notch-expressing cell line S2-MT-Notch.

References

- Barten DM, Guss VL, Corsa JA, Loo A, Hansel SB, Zheng M, Munoz B, Srinivasan K, Wang B, Robertson BJ, et al. (2005) Dynamics of β -amyloid reductions in brain, cerebrospinal fluid, and plasma of β -amyloid precursor protein transgenic mice treated with a γ -secretase inhibitor. *J Pharmacol Exp Ther* **312**:635–643.
- Best JD, Smith DW, Reilly MA, O'Donnell R, Lewis HD, Ellis S, Wilkie N, Rosahl TW, Laroque PA, Boussiquet-Leroux C, et al. (2007) The novel γ -secretase inhibitor *N*-[cis-4-[(4-Chlorophenyl)sulfonyl]-4-(2,5-difluorophenyl)cyclohexyl]-1,1,1-trifluoromethanesulfonamide (MRK-560) reduces amyloid plaque deposition without evidence of Notch-related pathology in the Tg2576 mouse. *J Pharmacol Exp Ther* **320**:552–558.
- Capell A, Steiner H, Romig H, Keck S, Baader M, Grim MG, Baumeister R, and Haass C (2000) Presenilin-1 differentially facilitates endoproteolysis of the β -amyloid precursor protein and Notch. *Nat Cell Biol* **2**:205–211.
- Carmin-Simmen K, Proctor T, Tschäpe J, Poeck B, Triphan T, Strauss R, and Kretschmar D (2009) Neurotoxic effects induced by the *Drosophila* amyloid- β peptide suggest a conserved toxic function. *Neurobiol Dis* **33**:274–281.
- Cole DC, Stock JR, Kreft AF, Antane M, Aschmies SH, Atchison KP, Casebier DS, Comery TA, Diamantidis G, Ellingboe JW, et al. (2009) (S)-*N*-(5-Chlorothiophene-2-sulfonyl)- β , β -diethylalaninol a Notch-1-sparing γ -secretase inhibitor. *Bioorg Med Chem Lett* **19**:926–929.
- Coulson EJ, Paliga K, Beyreuther K, and Masters CL (2000) What the evolution of the amyloid protein precursor supergene family tells us about its function. *Neurochem Int* **36**:175–184.
- Dejaegere T, Serneels L, Schäfer MK, Van Biervliet J, Horré K, Depoylu C, Alvarez-Fischer D, Herreman A, Willem M, Haass C, et al. (2008) Deficiency of Aph-1B/C- γ -secretase disturbs Nrg1 cleavage and sensorimotor gating that can be reversed with antipsychotic treatment. *Proc Natl Acad Sci USA* **105**:9775–9780.
- DeLean A, Munson PJ, and Rodbard D (1978) Simultaneous analysis of families of sigmoidal curves: application to bioassay, radioligand assay, and physiological dose-response curves. *Am J Physiol* **235**:E97–E102.
- Eggert S, Paliga K, Soba P, Evvin G, Masters CL, Weidemann A, and Beyreuther K (2004) The proteolytic processing of the amyloid precursor protein gene family members APLP-1 and APLP-2 involves α -, β -, γ -, and ϵ -like cleavages. Modulation of APLP-1 processing by N-glycosylation. *J Biol Chem* **279**:18146–18156.
- Fehon RG, Kooh PJ, Rebay I, Regan CL, Xu T, Muskavitch MA, and Artavanis-Tsakonas S (1990) Molecular interactions between the protein products of the neurogenic loci Notch and Delta, two EGF-homologous genes in *Drosophila*. *Cell* **61**:523–534.
- Fuwa H, Takahashi Y, Konno Y, Watanabe N, Miyashita H, Sasaki M, Natsugari H, Kan T, Fukuyama T, Tomita T, et al. (2007) Divergent synthesis of multifunctional molecular probes to elucidate the enzyme specificity of dipeptidic γ -secretase inhibitors. *ACS Chem Biol* **2**:408–418.
- Hébert SS, Serneels L, Dejaegere T, Horré K, Dabrowski M, Baert V, Annaert W, Hartmann D, and De Strooper B (2004) Coordinated and widespread expression of γ -secretase in vivo: evidence for size and molecular heterogeneity. *Neurobiol Dis* **17**:260–272.
- Hu Y and Fortini ME (2003) Different cofactor activities in γ -secretase assembly: evidence for a nicastrin-Aph-1 subcomplex. *J Cell Biol* **161**:685–690.
- Hu Y, Ye Y, and Fortini ME (2002) Nicastrin is required for γ -secretase cleavage of the *Drosophila* Notch receptor. *Dev Cell* **2**:69–78.
- Kopan R and Ilagan MX (2009) The canonical Notch signaling pathway: unfolding the activation mechanism. *Cell* **137**:216–233.
- Kulic L, Walter J, Multhaup G, Teplow DB, Baumeister R, Romig H, Capell A, Steiner H, and Haass C (2000) Separation of presenilin function in amyloid β -peptide generation and endoproteolysis of Notch. *Proc Natl Acad Sci USA* **97**:5913–5918.
- Lai MT, Chen E, Crouthamel MC, DiMuzio-Mower J, Xu M, Huang Q, Price E, Register RB, Shi XP, Donoviel DB, et al. (2003) Presenilin-1 and Presenilin-2 exhibit distinct yet overlapping γ -secretase activities. *J Biol Chem* **278**:22475–22481.
- Lewis HD, Pérez Revuelta BI, Nadin A, Neduvilil JG, Harrison T, Pollack SJ, and Shearman MS (2003) Catalytic site-directed γ -secretase inhibitors do not discriminate pharmacologically between Notch S3 and β -APP cleavages. *Biochemistry* **42**:7580–7586.
- Luo LQ, Martin-Morris LE, and White K (1990) Identification, secretion, and neural expression of APPL, a *Drosophila* protein similar to human amyloid protein precursor. *J Neurosci* **10**:3849–3861.
- McCarthy JV, Twomey C, and Wujek P (2009) Presenilin-dependent regulated intramembrane proteolysis and γ -secretase activity. *Cell Mol Life Sci* **66**:1534–1555.
- Milano J, McKay J, Dagenais C, Foster-Brown L, Pognan F, Gadiant R, Jacobs RT, Zacco A, Greenberg B, and Ciaccio PJ (2004) Modulation of Notch processing by γ -secretase inhibitors causes intestinal goblet cell metaplasia and induction of genes known to specify gut secretory lineage differentiation. *Toxicol Sci* **82**:341–358.
- Moehlmann T, Winkler E, Xia X, Edbauer D, Murrell J, Capell A, Kaether C, Zheng H, Ghetti B, Haass C, et al. (2002) Presenilin-1 mutations of leucine 166 equally affect the generation of the Notch and APP intracellular domains independent of their effect on A β 42 production. *Proc Natl Acad Sci USA* **99**:8025–8030.
- Morohashi Y, Kan T, Tominari Y, Fuwa H, Okamura Y, Watanabe N, Sato C, Natsugari H, Fukuyama T, Iwatsubo T, et al. (2006) C-terminal fragment of Presenilin is the molecular target of a dipeptidic γ -secretase-specific inhibitor DAPT (*N*-[N-(3,5-difluorophenyl)-L-alanyl]-S-phenylglycine *t*-butyl ester). *J Biol Chem* **281**:14670–14676.
- Musa A, Lehrs H, and Russo VA (2001) Distinct expression patterns of two zebrafish homologues of the human APP gene during embryonic development. *Dev Genes Evol* **211**:563–567.
- Nakajima M, Shimizu T, and Shirasawa T (2000) Notch-1 activation by familial Alzheimer's disease (FAD)-linked mutant forms of presenilin-1. *J Neurosci Res* **62**:311–317.
- Pinheiro JC and Bates DM (2002) *Mixed-Effects Models in S- and S-Plus*. Springer, New York.
- Placanica L, Tarassishin L, Yang G, Peethumongsin E, Kim SH, Zheng H, Sisodia SS, and Li YM (2009) Pen2 and presenilin-1 modulate the dynamic equilibrium of presenilin-2 and presenilin-2 γ -secretase complexes. *J Biol Chem* **284**:2967–2977.
- Pu J, Kreft AF, Aschmies SH, Atchison KP, Berkowitz J, Caggiano TJ, Chlenov M, Diamantidis G, Harrison BL, Hu Y, et al. (2009) Synthesis and structure-activity relationship of a novel series of heterocyclic sulfonamide γ -secretase inhibitors. *Bioorg Med Chem* **17**:4708–4717.
- Rand MD, Grimm LM, Artavanis-Tsakonas S, Patriub V, Blacklow SC, Sklar J, and Aster JC (2000) Calcium depletion dissociates and activates heterodimeric Notch receptors. *Mol Cell Biol* **20**:1825–1835.
- Schroeter EH, Ilagan MX, Brunkan AL, Hecimovic S, Li YM, Xu M, Lewis HD, Saxena MT, De Strooper B, Conrod A, et al. (2003) A presenilin dimer at the core of the γ -secretase enzyme: insights from parallel analysis of Notch 1 and APP proteolysis. *Proc Natl Acad Sci USA* **100**:13075–13080.
- Selkoe DJ (2001) Presenilin, Notch, and the genesis and treatment of Alzheimer's disease. *Proc Natl Acad Sci USA* **98**:11039–11041.
- Selkoe DJ and Wolfe MS (2007) Presenilin: running with scissors in the membrane. *Cell* **131**:215–221.
- Serneels L, Van Biervliet J, Craessaerts K, Dejaegere T, Horré K, Van Houtvin T, Esselmann H, Paul S, Schäfer MK, Berezoyska O, et al. (2009) γ -secretase heterogeneity in the Aph-1 subunit: relevance for Alzheimer's disease. *Science* **324**:639–642.
- Shirokani K, Edbauer D, Prokop S, Haass C, and Steiner H (2004) Identification of distinct γ -secretase complexes with different APH-1 variants. *J Biol Chem* **279**:41340–41345.
- Steiner H, Winkler E, and Haass C (2008) Chemical cross-linking provides a model of the γ -secretase complex subunit architecture and evidence for close proximity of the C-terminal fragment of presenilin with APH-1. *J Biol Chem* **283**:34677–34686.
- Takasugi N, Tomita T, Hayashi I, Tsuruoka M, Niimura M, Takahashi Y, Thinakaran G, and Iwatsubo T (2003) The role of presenilin cofactors in the γ -secretase complex. *Nature* **422**:438–441.
- Tanai H, Jiang J, Fukumori A, Tagami S, Okazaki Y, Okochi M, and Takeda M (2006) Effect of valine on the efficiency and precision at S4 cleavage of the Notch-1 transmembrane domain. *J Neurosci Res* **84**:918–925.
- Thinakaran G, Harris CL, Ratovitski T, Davenport F, Slunt HH, Price DL, Borchelt DR, and Sisodia SS (1997) Evidence that levels of presenilins (PS1 and PS2) are coordinately regulated by competition for limiting cellular factors. *J Biol Chem* **272**:28415–28422.
- Tien AC, Rajan A, and Bellen HJ (2009) A Notch updated. *J Cell Biol* **184**:621–629.
- Tomita T, Tanaka S, Morohashi Y, and Iwatsubo T (2006) Presenilin-dependent intramembrane cleavage of ephrin-B1. *Mol Neurodegener* **1**:2.
- van Es JH, van Gijn ME, Riccio O, van den Born M, Vooijs M, Begthel H, Cozijnsen M, Robine S, Winton DJ, Radtke F, et al. (2005) Notch/ γ -secretase inhibition turns

- proliferative cells in intestinal crypts and adenomas into goblet cells. *Nature* **435**:959–963.
- Vingtdeux V, Hamdane M, Gompel M, Bégard S, Drobecq H, Ghestem A, Grosjean ME, Kostanjevecki V, Grognet P, Vanmechelen E, et al. (2005) Phosphorylation of amyloid precursor carboxy-terminal fragments enhances their processing by a γ -secretase-dependent mechanism. *Neurobiol Dis* **20**:625–637.
- Vingtdeux V, Hamdane M, Loyens A, Gelé P, Drobeck H, Bégard S, Galas MC, Delacourte A, Beauvillain JC, Buée L, et al. (2007) Alkalizing drugs induce accumulation of amyloid precursor protein by-products in luminal vesicles of multivesicular bodies. *J Biol Chem* **282**:18197–18205.
- Wolfe MS, Citron M, Diehl TS, Xia W, Donkor IO, and Selkoe DJ (1998) A substrate-based difluoro ketone selectively inhibits Alzheimer's γ -secretase activity. *J Med Chem* **41**:6–9.
- Xia W, Zhang J, Ostaszewski BL, Kimberly WT, Seubert P, Koo EH, Shen J, and Selkoe DJ (1998) Presenilin 1 regulates the processing of β -amyloid precursor protein C-terminal fragments and the generation of amyloid β -protein in endoplasmic reticulum and Golgi. *Biochemistry* **37**:16465–16471.
- Yang T, Arslanova D, Gu Y, Augelli-Szafran C, and Xia W (2008) Quantification of γ -secretase modulation differentiates inhibitor compound selectivity between two substrates Notch and amyloid precursor protein. *Mol Brain* **1**:15.
- Zhang DM, Levitan D, Yu G, Nishimura M, Chen F, Tandon A, Kawarai T, Arawaka S, Supala A, Song YQ, et al. (2000) Mutation of the conserved N-terminal cysteine (Cys92) of human presenilin 1 causes increased A β 42 secretion in mammalian cells but impaired *Notch/lin-12* signalling in *C. elegans*. *Neuroreport* **11**:3227–3230.

Address correspondence to: Mark E. Fortini, Department of Biochemistry and Molecular Biology, Thomas Jefferson University, Bluemle Life Sciences Building 830A, 233 South 10th Street, Philadelphia, PA 19107, E-mail: mark.fortini@jefferson.edu