Alteration in synaptic nanoscale organization dictates amyloidogenic processing in Alzheimer’s disease

Endocytic Membrane of an Excitatory Synapse

Endocytic Vesicle
Healthy

APP
β Secretase
CTFβ
Aβ

Endocytic Vesicle
Alzheimer's Disease

50 nm
Article

Alteration in synaptic nanoscale organization dictates amyloidogenic processing in Alzheimer’s disease

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Summary

Despite intuitive insights into differential proteolysis of amyloid precursor protein (APP), the stochasticity behind local product formation through amyloidogenic pathway at individual synapses remain unclear. Here, we show that the major components of amyloidogenic machinery namely, APP and secretases are discretely organized into nanodomains of high local concentration compared to their immediate environment in functional zones of the synapse. Additionally, with the aid of multiple models of Alzheimer’s disease (AD), we confirm that this discrete nanoscale chemical map of amyloidogenic machinery is altered at excitatory synapses. Furthermore, we provide realistic models of amyloidogenic processing in unitary vesicles originating from the endocytic zone of excitatory synapses. Thus, we show how an alteration in the stochasticity of synaptic nanoscale organization contributes to the dynamic range of C-terminal fragments β (CTFβ) production, defining the heterogeneity of amyloidogenic processing at individual synapses, leading to long-term synaptic deficits as seen in AD.

Introduction

Enzymatic hydrolysis of peptide bonds (proteolysis) induces an irreversible alteration of the molecular structure and biological function of a protein. Proteolysis is a post-translational modification, generating functionally relevant and stable cleaved proteins referred to as proteoforms, which are pivotal regulators of many physiological and pathological processes (Klein et al., 2018; Rawlings et al., 2012). Sequential proteolysis is a targeted event where multiple enzymes act one after another on a single substrate resulting in several proteoforms (Klein et al., 2018). These alterations of the substrate molecule are controlled both spatially and temporally such that a change in the combination of proteases can result in proteoforms with antagonistic properties. Secretases are a class of proteases involved in precise proteolytic processing of amyloid precursor protein (APP), a single-pass transmembrane protein that is ubiquitously expressed throughout the body (Chow et al., 2010; Muller et al., 2017). APP can be processed by both canonical and non-canonical secretases, resulting in numerous proteoforms, mediating distinct and even opposing functions (Muller et al., 2017).

Decades of research indicate that alteration in proteolytic processing of APP is a crucial element toward progression of Alzheimer’s disease (AD) (Brunholz et al., 2012; DeBoer et al., 2014; Haass et al., 2012; Sun and Roy, 2018). The major focus on APP proteolysis processing is due to its importance in the generation of a peptide proteoform referred to as amyloid beta (Aβ), an essential component of Amyloid plaques found in the brain of patients diagnosed with AD. Extensive biochemical and molecular biology studies have identified that Aβ is generated by amyloidogenic processing through sequential cleavage by β- and γ-secretases. Despite several years of focus on AD, a lack of understanding still exists on how the equilibrium is shifted toward the amyloidogenic pathway or how this shift alters the molecular machinery involved in synaptic transmission and plasticity (Montagna et al., 2017). Over the last decade, groundwork has been laid to understand AD as a disease beginning with alteration of molecular properties of individual synapses (Lesne et al., 2006; Neuman et al., 2015; Opazo et al., 2018; Wei et al., 2010; Wilhelm et al., 2014). Recent studies have indicated the subcellular segregation of APP into regulatory nanodomains on the...
plasma membrane in both neuronal and non-neuronal cells (de Coninck et al., 2018; Kedia et al., 2020). Further, the nanoscopic fingerprints of these domains have been shown to be altered in neuronal processes and within functional subcompartments of an excitatory synapse and between different variants of APP implicated in AD (Kedia et al., 2020).

Although AD is considered to begin as a synaptopathy, it is not yet understood how the segregation of APP and secretases at nanoscale contributes toward the progression of AD (De Strooper and Karran, 2016; Haass et al., 2012; Selkoe et al., 2012; Selkoe, 2002). This is largely due to the lack of information on (1) the heterogeneity of localization of the amyloidogenic machinery within/outside functional zones of individual synapses (Harris and Stevens, 1989; Harris and Weinberg, 2012) and (2) a lack of intuitive understanding of the mechanisms that control the heterogeneity of diffusional collisions between APP and secretases resulting in different proteoforms (Ben Halima et al., 2016; Escamilla-Ayala et al., 2020; Kedia et al., 2020). Here, we have employed super-resolution imaging and analysis to reveal the subsynaptic organization of these molecules. We confirm that in addition to APP, both β- and γ-secretases are also organized into segregated domains of few tens of nanometers. This discrete association of nanodomains resemble high molecular weight multi-protein complexes with varying compositionality of secretases and APP (Chen et al., 2015; Liu et al., 2019a). We used this nanoscale heterogeneity in the molecular distribution in empirical *insilico* experiments of reconstructed vesicles to simulate the interactions between APP and secretases as diffusional collisions resulting in the product formation. We focused our efforts to understand the association of APP with β-secretases in specialized subsynaptic regions and how this stoichiometry of association directly influences the processing of APP through the amyloidogenic pathway. We present a unique data-driven realistic model for synaptic amyloidogenic processing from several thousands of synapses, wherein we identify a set of molecular determinants that decide the fate of APP proteolysis. We further demonstrate that even minor alterations in the molecular fingerprints of this synaptic nanorganization can yield significant changes in the local product formation through amyloidogenic processing.

Furthermore, with the aid of transgenic mouse models for AD and postmortem human brain tissues from AD patients, we validate the competency of this molecular model. Thus, we entail a nanoscale synaptic reaction-diffusion model of amyloidogenic processing with realistic numbers and geometrical constraints to understand molecular mechanisms that alter synaptic amyloidogenic processing.

Results

Differential nanoorganization of amyloidogenic proteolytic machinery in the functional domains of an excitatory synapse

The amyloidogenic processing of APP is the result of sequential proteolysis by β- and γ-secretases (Cole and Vassar, 2007; Yang et al., 2017). The spatial association of β- and γ-secretases is vital for amyloidogenic processing in different neuronal subcompartments. To comprehend this spatial variability of β- and γ-secretases within the neuronal processes and synaptic compartments, we evaluated the relative nanoscale distribution of the secretases in neuronal processes and within different functional zones of an excitatory synapse. We relied on the nanoscopic association of secretases with a marker for postsynaptic density (PSD) and a perisynaptic marker for endocytic zone (EZ), PSD95 and dynamin, respectively. Similar experimental paradigms have been previously used to segregate the fractional contribution of synaptic molecules localized to both synaptic compartments and within functional compartments of individual synapses (Kedia et al., 2020).

The distribution of β-secretase, BACE1 was evaluated for the quantitative estimation of the association of β-secretase in PSD and EZ (Figures 1 and S1). Confocal and stimulated emission depletion (STED) microscopy were performed sequentially to evaluate the diffraction limited and nanoscale distribution of these molecules. A gallery of representative images of individual synapses obtained by confocal and STED microscopy using PSD/EZ markers and the associated molecular domains of β-secretase (nanodomainβ) are presented (Figures 1, S1A, S1B, S1C, S1D, S1E, and S1F). The morphological and biophysical properties of nanodomainβ were characterized (Table 1). Interestingly, resolution scaled Pearson’s (RSP) coefficient and resolution scaled error (RSE) of β-secretase in PSD and EZ were significantly different. The colocalization with β-secretase was significantly higher in PSD, while the variability was more in EZ (Figures 1i and 1iv).

The nanodomainβ associated with PSD and EZ is referred to as nanodomainβPSD and nanodomainβEZ respectively. The distribution of length, area, intensity, and normalized intensity with respect to the median of the nanodomainβ intensity is indicated in Figures 1ii, 1iii, 1v, 1vi, S2i, and S2ii and Table 1.
Figure 1. Nanoscale distribution of β-secretase in the functional zones of excitatory postsynapse using STED microscopy

(A and B) Overlay of STED images of postsynaptic density marker PSD95 and a marker for endocytic zone Dynamin (green) with β-secretase (Magenta). The black contour indicates the automated detection of neuronal processes. Inset 1–6 indicate a gallery of synapses where the black contour within inset represents automatically detected regions for confocal marker for postsynapse and endocytic zone (black). Black in the overlay images represents the overlap between the corresponding green and magenta images. The scale bars in B represent 7 μm and inset corresponds to 1.4 μm.

(i and iv) Comparison of RSP and RSE for quantifying colocalization of β-secretase for functional zones of an excitatory postsynapse. Data are represented as mean ± SEM. Significance was determined by unpaired two-tailed Student’s t test. *p < 0.05, **p < 0.01, and ***p < 0.001, ns p > 0.05.

(ii and iii) Indicate the distribution of length of all β-secretase nanodomains obtained by STED microscopy in post and perisynaptic compartments, respectively.

(v and vi) Indicate the distribution of intensity of all β-secretase nanodomains obtained by STED microscopy in post and perisynaptic compartments, respectively. n = 5669 puncta (post) and 3798 puncta (peri).
For analyzing the association of γ-secretase in PSD and EZ, the distribution of the catalytic subunit of γ-secretase namely, presenilin1 (PS1) was evaluated against PSD95 and Dynamin, respectively (Figures 2 and S3). We analyzed the PSD95 and Dynamin positive regions (Figures 2, S3A, S3B, S3C, S3D, S3E, and S3F), as well as characterized the morphological and biophysical properties of molecular domains of γ-secretase (nanodomainγ) in these regions (Table 1). The colocalization of γ-secretase in PSD and EZ was similar, while variability was significantly higher in EZ (Figures 2i and 2iv). The nanodomainγ associated with PSD and EZ is referred to as nanodomainγ/PSD and nanodomainγ/EZ, respectively. The distribution of length, area, intensity, and normalized intensity with respect to the median of the nanodomainγ intensity is indicated in Figures 2ii, 2iii, 2v, 2vi, S4i, and S4ii and Table 1.

Our results show significant differences in the morphological and biophysical characteristics of the β- and γ-secretase nanodomains associated with PSD and EZ (Figures S2 and S4). The length, area, and intensity were significantly higher for nanodomainγ/PSD in comparison to nanodomainγ/EZ. However, these parameters were significantly lower for nanodomainγ/EZ when compared to nanodomainγ/PSD. The distribution of normalized intensity of nanodomainγ/PSD, nanodomainγ/EZ and nanodomainγ/EZ showed a higher variability when compared to nanodomainγ/PSD (Figures S2i and S4i). To determine the proximity of spatial association of β- and γ-secretases to PSD and EZ, we quantified nearest neighborhood distance (NND) for the secretases with respect to these functional zones. The β-secretases were proximal to PSD in comparison to EZ, while the γ-secretases associated closely with EZ (Figures S2ii and S4ii). This indicated a potential difference in the distribution of secretases associated with these domains, signifying the relevance of nanoscale association of APP, β-secretase and γ-secretase in synaptic subcompartments for the amyloidogenic processing of APP.

Further, we assessed the association of β- and γ-secretases with APP. We examined regions where APP colocalized with either β- or γ-secretase. A gallery of confocal and STED images representing this association are shown. Where APP nanodomains colocalizing with β- or γ-secretases are referred to as nanodomainβ/APP and nanodomainγ/APP, respectively (Figures 3A, 3B, 3C, 3D, 3E, and 3F).
Figure 2. Nanoscale distribution of γ-secretase in the functional zones of excitatory postsynapse using STED microscopy

(A and B) Overlay of STED images of postsynaptic density marker PSD95 and a marker for endocytic zone Dynamin (green) with γ-secretase (Magenta). The black contour indicates the automated detection of neuronal processes. Inset 1–6 indicate a gallery of synapses where the black contour within inset represents automatically detected regions for confocal marker for postsynapse and endocytic zone (black). Black in the overlay images represents the overlap between the corresponding green and magenta images. The scale bars in B represent 7 µm and inset corresponds to 1.4 µm.

(i and iv) Comparison of RSP and RSE for quantifying colocalization of γ-secretase for functional zones of an excitatory postsynapse. Data are represented as mean ± SEM. Significance was determined by unpaired two-tailed Student’s t test. *p ≤ 0.05, **p ≤ 0.01, and ***p ≤ 0.001, ns p > 0.05.

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(v and vi) Indicate the distribution of intensity of all γ-secretase nanodomains obtained by STED microscopy in post and perisynaptic compartments, respectively. n = 4936 puncta (post) and 5921 puncta (peri).
Figure 3. Nanoscale association of β/γ-secretase and with APP in the neuronal processes using STED microscopy

(A and D) Gallery of confocal images of neuronal processes identified by automatic detection of confocal marker for APP (magenta) puncta with pseudocolour overlay of β/γ-secretase (green). (B and E) STED image of the respective regions identified from (A, D). (C and F) Represent automatically detected regions for confocal marker for β/γ-secretase (black) along with nanoscale representation of β/γ-secretase (green) and APP (magenta). Black in the overlay images represents the overlap between the corresponding green and magenta images. (i and iii) Indicate the distribution of length of all β/γ-secretase nanodomains on APP obtained by STED microscopy. (ii and iv) Indicate the distribution of intensity of all β/γ-secretase nanodomains on APP obtained by STED microscopy. (G) Confocal image of the individual γ-secretase identified by automatic detection of confocal marker for γ-secretase (red) with pseudocolour overlay of β-secretase (blue).
To validate this, we evaluated the relative distribution of β- and γ-secretases in synapses. We analyzed regions marked positive for β- and γ-secretases by confocal and STED microscopy (Figures 3G, 3H, and 3I). The sub-diffraction limited clusters, where β- and γ-secretases were associated, were referred to as nanodomain$_{\beta\gamma}$. The distribution of length, area, intensity, and normalized intensity with respect to the median of the nanodomain$_{\beta\gamma}$ is presented (Figures 3iv, 3v, and Table 1). Similar to nanodomain$_{\beta\gamma}$, nanodomain$_{\gamma\beta}$ displayed a large variability in the intensity, consistent with our hypothesis that the number of β-secretase molecules associated with synaptic nanodomains might be a limiting factor for the amyloidogenic processing of APP. This was also validated by the observation that although the colocalization of the two secretases with APP remained similar, variability was higher for γ-secretases/APP compared to β-secretases/APP (Figure S5iii).

Similar to the association of secretases with functional zones of the synapse, we performed NND analysis between APP and secretases. The distance of β-secretase to APP was proximal and less variable in comparison to γ-secretases to APP and β-to γ-secretase (Figure S5iv). This further confirmed that multiple molecular parameters could define the spatial association of secretases with APP, thus directly influencing the local proteolysis of APP. APP is known to be processed sequentially by β-secretase followed by γ-secretase. The catalytic activity of β-secretase is deemed to occur in the acidic pH range (~4.5) which corresponds to membrane-bound organelles involved in the secretory pathway, such as vesicles budding from the endocytic zone of the synapse (Cole and Vassar, 2007; Lu et al., 2007; Watanabe et al., 2014). Here, quantitative parameters obtained from super-resolution microscopy on the distribution and association of APP and β/γ-secretase were used to simulate the dynamics of APP processing in the internalized membrane of endocytic zones of dendritic spines reconstructed from EM slices of CA3-CA1 region of the hippocampus.

**Nanoscale alteration of molecular fingerprints of amyloidogenic machinery in multiple models of AD**

To test the potential association of the nanoscale fingerprints of amyloidogenic machinery and AD, we evaluated the morphological and biophysical traits of β-secretase and APP in multiple models of AD. We recently reported that the lateral diffusion and nanoscale aggregation properties of APP-wildtype differed considerably from its detrimental variant namely, APP-Swedish (Kedia et al., 2020). APP/PS1 mice contain humanized APP within the Aβ region bearing the Swedish mutation, as well as PSEN1 encoding deltaE9 mutation, under the control of the mouse prion promoter (Jankowsky et al., 2001). These mice do not harbor any mutations in BACE. APP/PS1 transgenic mouse model has been well characterized for cognitive impairment (Lalonde et al., 2005; Volianskis et al., 2010), with the reduction of transient long-term potentiation by 3 months of age (Volianskis et al., 2010). Also, age-dependent loss of classical synaptic proteins such as Synaptophysin, Synaptotagmin, Homer, and PSD95 is observed in these mice as early as 4 months (Hong et al., 2016). Thus, APP/PS1 mice harboring APP-Swe mutation has been very well characterized for early synaptic deficits. In our paradigm, we selected APP/PS1 mice which were 3-4 months old. This timeline coincides with the earliest appearance of synaptic impairment which precedes the detectable levels of Aβ deposits. Thus, APP/PS1 model was chosen to evaluate the role of nanoscale synaptic deficits in the early onset of AD. Global levels of APP and C-terminal fragment (CTF) was increased in APP/PS1 (Tg) mice in comparison to wildtype (WT) due to overexpression, while the β-secretase levels remained
unaltered when evaluated by immunoblotting (Figures 4A, 4B, 4i, and 4ii). Since the foresaid experiment only identifies human variant of APP, we verified the expression of the total APP pool using an antibody that recognized both human and murine APP variants (Figure S6 and Table S3). Next, we evaluated the nanoscale organization of β-secretase in different functional zones of the synapse. The quantitative association of β-secretase with PSD and EZ was assessed by imaging with STED microscopy using Shank2 and Clathrin as markers, respectively (Figures 4C and 4D). We found that the length and intensity of nanodomain APP/PSD and nanodomain APP/EZ were significantly altered between WT and Tg mice (Figures 4iii and 4iv and Table S1). Furthermore, on comparison of the integrated intensity of β-secretase in PSD/EZ (Figure S7i), we found that in Tg mice, the cumulative β-secretase levels were decreased in PSD, while it increased in EZ. Additionally, we found that the proximity of β-secretase to PSD and EZ were altered antagonistically. The NND of β-secretase to PSD increased, while a significant decrease was observed for EZ in Tg mice (Figure S7ii). Taken together, the β-secretase levels in Tg mice were augmented significantly in PSD/EZ both inside and outside of nanodomains, along with a decrease in the length of nanodomains. This nanoscale alteration of β-secretase was pronounced at EZ together with a reduction in NND, reflecting an increase in the β-secretase load per endocytic event originating from EZ in Tg mice.

In order to evaluate if such alterations were also consistent in the human brain, we determined the length and intensity of nanodomain APP/PSD, nanodomain APP/EZ, nanodomain APP/PSD, and nanodomain APP/EZ with similar markers for PSD and EZ in three sets of post-mortem human brain tissues from patients with AD and their corresponding controls (control) using Airyscan microscopy, providing sub-diffraction limited resolution (Figures S8A, S8B, S8C, and S8D, and Table S4). Similar to rodent models, we randomly chose 5–10 non-overlapping regions from the radiatum layer of the hippocampus (Figures S8E and S8F). The quantitative association of β-secretase and APP with PSD and EZ was assessed by Airyscan microscopy using Shank2 and Clathrin as markers, respectively (Figures 5 and 6). Airyscan images of β-secretase association with EZ and PSD in control and AD human brains were evaluated for the sub-diffraction limited zones of enrichment (Figures 5A and 5B). Although, there was no significant alteration in the size of nanodomain APP/PSD and nanodomain APP/EZ, the content of β-secretase enriched in these domains were significantly higher in the brain of patients with AD compared to the control samples (Figures 5i and 5ii). A similar strategy was employed to characterize nanodomain APP/PSD and nanodomain APP/EZ in control and AD samples from the brain sections of the same patients investigated for the alteration of subsynaptic organization of β-secretase (Figures 6A and 6B). In this case, the APP content in the nanodomains increased significantly in both EZ and PSD (Figures 6i and 6ii). However, comparing AD samples to the control, the size of the nanodomain APP/PSD increased in contrast to a small yet significant decrease in the size of the nanodomain APP/EZ (Figures 6i and 6ii).

Next, to understand heterogeneity in organization of amyloidogenic machinery at finer spatial scales, we performed STED microscopy on one set of age- and gender-matched AD and control samples to investigate if we could enhance the spatial separation between EZ and PSD, and performed an extensive analysis of the nanodomains of β-secretase and APP associated with these functional zones (Figures 9A, 9B, 9C, and 9D and Table S4). We found that the intensity of nanodomains of APP and β-secretase in both PSD and EZ increased in AD (Figures 9i and 9ii and Table S1). While the length of nanodomains of APP increased in PSD, it was reduced in EZ in AD (Figures 9i and 9ii and Table S1). In contrast to APP, the length of nanodomains of β-secretase increased in EZ, while remaining similar at PSD in AD (Figures 9i and 9ii and Table S1). The investigations related to the biophysical characteristics of the nanodomain APP/PSD, nanodomain APP/EZ, nanodomain APP/PSD, and nanodomain APP/EZ between AD and control.
Figure 5. Alteration of distribution of β-secretase in human brain slices using Airyscan microscopy
(A and B) Compartmentalization of β-secretase clusters in post/perisynapse in human brain slices from control (A) and AD (B) using Airyscan microscopy. White in the overlay images represents the overlap between the corresponding blue, green and red images. Scale bar at (B) indicates 30 μm.

(i and ii) Diversity (median/IQR 25%–75% interval) in nanodomain length (μm) and intensity for β-secretase clusters in perisynapse (i) or postsynapse (ii) in human brain slices from AD and control represented as (median/IQR 25%–75% interval).
In silico experiments in unitary vesicles predict the biophysical determinants of CTFβ production

Recent studies have indicated unhindered diffusion of APP on the membrane in contrast to a confined motion in functional nanodomains of APP (nanodomain\textsuperscript{APP}), as well as differential localization of APP molecules in PSD and EZ (Kedia et al., 2020). Concurrent to the observation of secretases that we report here, an EZ can have different permutations and combinations of components of β-amyloidogenic machinery. To estimate physiologically the realistic dynamics of CTFβ production, we systematically simulated the diffusion of APP molecules on a membrane that includes, (1) single molecule mobility and (2) reduced mobility represented by nanodomain\textsuperscript{APP} of varying content (Table S2 (Kedia et al., 2020)). These simulations were carried out in a confined volume representative of a unit endocytic process. This reconstructed endosomal compartment where the activity of β-secretase was optimal (diameter 0.120 μm) is referred to as a “unitary vesicle”. The characteristics of unitary vesicles are shown in Table S2.

An immediately relevant question in this context was how the spatial distribution and biophysical characteristics of a simulated unitary vesicle affect CTFβ production. The resulting number of CTFβ in the reaction is a result of competing dynamics of reaction rates, diffusion and numbers of reactants and geometrical constraints of the unitary vesicle. We investigated two contrasting hypotheses as advantageous for the production of CTFβ. We asked if (1) slow-moving-large and densely populated APP clusters enabled more productive encounters with β-secretase (referred to as “the sitting duck hypothesis”) or (2) rapidly diffusing APP monomers (referred to as “movers and shakers hypothesis”). We simulated APP and β-secretase binding on a unitary vesicle for the physiologically realistic range of mobility parameters and typical numbers of each species (Table S2). The binding rates for the kinetic scheme (Figures 7A and 7B) were modified from those measured independently in experiments (Ben Halima et al., 2016), refer to methods for calculation of binding rates.

The simulations were carried out for unitary vesicles with APP clusters of sizes 5, 9, and 13 as illustrative instantiations of clusters seen in vesicles. For simplicity, BACE1 was always simulated as a monomer in these control “in silico” experiments. The complete distribution of cluster sizes and an entire range of diffusion coefficients from APP-WT to APP-Swe within and outside the nanodomains have been characterized and described previously (Kedia et al., 2020). These values are summarized in Table S2. The simulated single molecule diffusion rates were an order of magnitude faster than nanodomain\textsuperscript{APP} and was consistent with experimental observations (Figures 7 and S11 and Table S2). The simulation also included the distribution of single molecules and nanodomains as seen in Figures 7C and 7D. For the unitary vesicles, the mobility of the β-secretase was considered similar to APP monomers. The freely diffusing β-secretase monomers interacted with APP molecules to produce CTFβ (Figure 7i). Each simulation was carried out for 5 s to mimic the time scales over which endocytosis takes place. The simulation results suggested that the production of intermediates was not sensitive to the possible range of rates of diffusion (including both APP-WT and APP-Swe) inside unitary vesicles (Figures 7j and S11i).

Next, we quantified the differences in the production of CTFβ corresponding to changes in k\textsubscript{1} and k\textsubscript{2} in APP-WT and APP-Swe conditions (Figures 7ii, 7iii, S11ii, and S11iii). A significant biophysical property that distinguishes APP-Swe from APP-WT is the reaction rate of intermediate product formation and CTFβ production. Both the forward rates k\textsubscript{1} and k\textsubscript{2} were seen to be higher for APP-Swe (Ben Halima et al., 2016). The resulting number of CTFβ production is a result of competing dynamics of reaction rates, diffusion and numbers of reactants and geometrical constraints of the unitary vesicle.

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Figure 6. Alteration of distribution of APP in human brain slices using Airyscan microscopy
(A and B) Compartmentalization of APP clusters in post/perisynapse in human brain slices from control (A) and AD (B) using Airyscan microscopy. White in the overlay images represents the overlap between the corresponding blue, green and red images. Scale bar at (B) indicates 30 μm.
Simulations showed up to a 20–25% increase in CTFβ comparable to a unitary vesicle with nanodomain and integrated content of Aβ42 neurons of APP/PS1 Tg mice compared to that of the wildtype (Figures S14A and S14B). Here, both the average were selected as per the APP-WT affinity values. Each simulation was carried out for 5 s and CTFβ production was maintained (data not shown). The propensity to form CTFβ in this case was comparable to a unitary vesicle with nanodomain. In summary, the increase in the rate of CTFβ production was correlated with an increase in the content of APP or β-secretase in nanodomainAPP or nanodomainAPP. This was also consistent with our observations of augmented levels of Aβ42 in primary hippocampal neurons of APP/PS1 Tg mice compared to that of the wildtype (Figures S14A and S14B). Here, both the average and integrated content of Aβ42 was significantly enhanced in Dynamin enriched compartments (Figure S14i). In these simulations, clusters of various sizes were instantiated, and their binding affinities were selected as per the APP-WT affinity values. Each simulation was carried out for 5 s and CTFβ present at the end of each simulation was considered (see Transparent Methods and Table S2 for details). Our simulations showed up to a 20–25% increase in CTFβ formation for vesicles with an increased size of nanodomainAPP or nanodomainAPP. When mimicking the unitary vesicle with the forward reaction rates corresponding to APP-Swe, there was a multi-fold increase in the probability of production of CTFβ (Figures S11iv and S13i). These simulations predict that inside the confined volume of a unitary vesicle where the molecular density of APP and β-secretase are close to realistic estimation, the forward reaction rates to produce CTFβ take a dominant role in amyloidogenic processing.

**Discussion**

Over the last decade, several studies have shown that individual synapses are heterogeneous structures where nanoscale segregation of synaptic molecules on the membrane plays a crucial role in synaptic transmission and plasticity (Chen et al., 2018; Dani et al., 2010; Nair et al., 2013; Venkatesan et al., 2020). Despite the functional overlap of amyloidogenic machinery, such evaluations on its subsynaptic organization have not yet been addressed in detail (Almeida et al., 2005; Muller et al., 2017; Snyder et al., 2005). Here, we show the nanoscale compartmentalization and differential association of components of β-amyloidogenic machinery in functional zones of the excitatory synapse. We have identified that within the functional zones of the synapse (Harris and Weinberg, 2012), the constituents of this machinery are clustered into nanoscale structures called nanodomains. The nanodomains of APP, β- and γ-secretases overlap discretely at
Figure 7. **Insilico** evaluation of CTFβ production for APP-WT/Swe show differential processing kinetics within unitary vesicles

(A) Schematic of nanoscale lateral organization of components of amyloidogenic machinery indicating the variability of free and segregated molecules of APP, β- and γ-secretases in the functional Zones of an excitatory synapse. Scale bar at A and D indicates 250 and 100 nm respectively.

(B) Reaction -Kinetics -Scheme

\[ \text{APP + BACE1} \xrightarrow{\text{Intermediate (k_i, k_2)}} \text{CTFβ + APP,}\beta + \text{BACE1 (k_2)} \]

(C) Lateral Projection

(D) Azimuthal Projection

Endocytic Vesicles

- Diffusional Collision by Lateral movement
- Reversible immobilization in Nanodomains

**Figure 7.** *iScience* evaluation of CTFβ production for APP-WT/Swe show differential processing kinetics within unitary vesicles

(A) Schematic of nanoscale lateral organization of components of amyloidogenic machinery indicating the variability of free and segregated molecules of APP, β- and γ-secretases in the functional Zones of an excitatory synapse. Scale bar at A and D indicates 250 and 100 nm respectively.

(B) Two-step reaction model for amyloidogenic processing of APP by β-secretase through lateral diffusion and single molecule collisions. The collision can either result in a metastable intermediate or process APP into CTF fragment which will stay on the membrane. Lateral diffusion can aid in the formation or elimination of APP nanodomains on the membrane.
neuronal processes with varying compositionality. These nanodomains and their association is similar to mega-Dalton sized secretase complexes which contain necessary molecules involved in the proteolysis of APP (Chen et al., 2015; Liu et al., 2019a). Consistent with this observation, these secretase rich complexes were also found to be very heterogeneous in their composition in neuronal synapses. The heterogeneity in composition is a cumulative outcome of (i) morphological and biophysical properties of the nanodomains, (ii) the distribution of the freely diffusible pool and nanodomains in the functional zones of synapse, and (iii) the proximity of nanodomain of APP to secretases. The molecular signatures that we identified varied between functional zones of the synapse, as well as between individual synapses. This implied that the variability in compositionality directly impacts the association/dissociation of these domains, thus controlling the locus of APP processing.

Recent evidences indicate that APP and γ-secretase molecules are transiently immobilized on the plasma membrane with the existence of an equilibrium between the nanodomain and extra-nanodomain APP and secretase molecules (Escamilla-Ayala et al., 2020; Kedia et al., 2020). The existence of APP nanodomains in neuronal and non-neuronal cells have been confirmed (de Coninck et al., 2018; Kedia et al., 2020). However, the segregation of γ-secretase into nanodomains in non-neuronal cells is still a matter of debate (Escamilla-Ayala et al., 2020; Liu et al., 2019a). The results presented here are consistent with previous reports of γ-secretase segregated into nanodomains in the synaptic compartments of neurons (Schedin-Weiss et al., 2016). Additionally, immobilization kinetics, morphology and the amount of APP molecules immobilized in nanodomains differed between well characterized wild-type (APP-WT), detrimental (APP-Swedish) and protective (APP-Icelandic) variants of APP (Kedia et al., 2020). All these parameters influence the availability of APP and secretases molecules per unit area on the neuronal membrane. The spatial proximity of β-amyloidogenic machinery is a pre-requisite for the proteolysis of APP. Since the spatial availability of components of amyloidogenic machinery is a limiting factor in the sequential cleavage of APP, we evaluated the heterogeneity in the localization by investigating the compositionality within subsynaptic compartments. We found a high variability in the intensity of β-secretase nanodomain with PSD, EZ, APP and with γ-secretase. The γ-secretase nanodomains were tightly associated with APP and PSD and were variable at EZ. This implied that in addition to the availability of β-secretase, how these secretase molecules associate with different pools of APP in the functional compartments of the synapse is a limiting factor for amyloidogenic processing of APP. Our results on the nanoscale variability of β-secretase are consistent with various evidences which associate a minor but enzymatically active portion of β-secretase with γ-secretase in functional complexes of high molecular weight resulting in the generation of Aβ (Liu et al., 2019a). However the mechanism of this association is unclear since only around a maximum of 10% of the total cellular β-secretase seems to be incorporated into such high molecular weight complexes resulting in a rather high variability of colocalization between β- and γ-secretases (Liu et al., 2019a, 2019b).

APP, β- and γ-secretases are transmembrane molecules which are distributed heterogeneously both in the functional compartments of synapses and on the neuronal membrane. Although the segregation of these molecules happens in both PSD and EZ, their environment and local distribution determine the association between them. This heterogeneity can be regulated by (1) molecular organization of APP and secretases within EZ at the time of internalization and (2) competing timescales for APP processing and diffusional collisions of these molecules on the membrane. As described previously, the synaptic endosomes of the
postsynaptic compartment are derived from the membrane of the EZ (Watanabe et al., 2014), where APP and secretases can coexist either in nanodomains or outside as a diffusive pool. This implies that a vesicle recycled from EZ can have either a diffusive population or a mix of diffusive population recruited along with regulatory nanodomains of APP and secretases. Thus, the amount of APP processed through an endocytic compartment is dictated by the compositionality of the individual compartments. These pools of APP/secretases or nanodomains of APP or secretases in varying combinations result in a dynamic range of Aβ production at the level of individual synapses. To include these dynamics, we instantiated the compositionality of amyloidogenic machinery in endocytic compartments, where only the compositionality of APP was altered. We systematically investigated if the “Sitting duck hypothesis” wherein APP molecules are confined in nanodomainAPP would be favorable for APP processing over the “Movers and shakers hypothesis” wherein single APP molecules are diffusing randomly. Interestingly, within the confined volume of the unitary vesicle, physiologically relevant changes in diffusion of molecules did not have a significant effect on the formation of CTFβ. The current estimation of secretases in the vesicle puts them at a very high concentration on the vesicular membrane. At such a high concentration in a confined volume, the diffusive effects of APP can be a minor factor in the production of CTFβ. Previous observations on the proteolysis of APP in multisecretase complexes have confirmed the formation of Aβ. Although the mechanism remains unclear, it poses an alternative to overcome product formation through stochastic collision of molecules on the plasma membrane. Thus, the spatial association of these complexes might regulate the local processing of APP resulting in an instantaneous increase in the rate of production of Aβ, as observed by increased molecular presence of components of amyloidogenic machinery in nanodomains during the progression of AD.

The processing of APP by secretases in the unitary vesicle is considered as a two-step reaction model, where a reversible intermediate complex is formed that can further result in the formation of CTFβ with an irreversible forward rate of reaction. Previous observations have indicated differences in rate of reaction of both APP-WT and APP-Swe (Ben Halima et al., 2016). It was, therefore, important to address the implication of these differences on APP processing by secretases. Within a unitary vesicle, the irreversible forward rate of reaction showed a significant effect on CTFβ production in comparison to the rate of formation of reversible intermediate product upon collision. Our data showed that within a unitary vesicle, the total number of APP molecules internalized per process was positively correlated with the probability of occurrence of nanodomainAPP. Subsequently, as the amount of APP and/or β-secretase immobilized in nanodomains increased, there was a significant increase in the amount of CTFβ being produced. The high affinity of collisions and cluster formation of APP and β-secretase molecules within a unitary vesicle ensured this arrangement. Through super-resolution imaging and analysis, it was observed that the secretase molecules were also found in nanodomains inside EZ. It remains elusive how acidic intraluminal pH within unitary vesicles can lead to altered kinetics of diffusional collisions between APP and β-secretase molecules. The propensity to form clusters of APP and β-secretase is found to be increased in multiple models of AD; our simulation demonstrates how it can contribute to the molecular progression of AD. A potential alternate mechanism could be that APP, β- and γ-secretase clusters could dissociate into monomers when exposed to acidic pH in the lumen, increasing the probability to collide with its substrate molecules. However, this remains to be experimentally verified and thus remains an open question.

Here, we have characterized molecular determinants that control the rate of formation of products in ascending order: diffusion of molecules, size of the nanodomain, the onward reaction rate of formation of reversible APP-secretase complex and the forward reaction rate for formation of CTFβ from the APP-secretase complex. This was consistent with the diffusive behavior of APP-Swe (Kedia et al., 2020) and with the biochemical rate of reaction of the formation of CTFβ (Ben Halima et al., 2016). Though diffusion itself does not have a notable influence within the unitary vesicle, it may have an important role on the synaptic membrane where the lateral diffusion of transmembrane molecules can control their local density through nanoscale association and segregation in short time scales. Several scaffolding molecules are known to interact at various levels with components of the β-amyloidogenic machinery (Perreau et al., 2010; Sisodia and St George-Hyslop, 2003). It would be interesting to see if the confinement kinetics of APP or secretases are affected by these molecules. Since the number of APP and secretases molecules in the synaptic compartment is very high, the retrieval of these molecules by endocytosis would happen in a timescale of seconds to minutes (Kumari et al., 2010; Lu et al., 2007; Watanabe et al., 2014). These variables confirm that individual synapses have their own dynamic range for Aβ production arising from the molecular composition of
the EZ membrane. This is in line with our observation of higher Aβ42 content in Dynamin enriched compartments, and is also consistent with the previous reports of increased Aβ content in both synaptic and endosomal compartments during the molecular progression of AD (Abramov et al., 2009; Gouras et al., 2010; Pickett et al., 2016; Sannerud et al., 2011, 2016).

Most of the modulators have been shown to affect APP processing and alter the profile of the Aβ peptides (De Strooper et al., 2010; Liu et al., 2019b). In a recent study, a close evaluation of multiple BACE inhibitors indicated that they extended the protein’s half-life (Liu et al., 2019b). This would generate additional Aβ or process neuronal substrates different from APP, affecting both synaptic function and non-amyloidogenic processing in long-term. The molecules that could dissociate β/γ secretase complexes (e.g. rocurbic acid) without altering the secretase levels would be better suited to diminish the amyloidogenic processing of APP (Liu et al., 2019b). Thus, we envision controlling the molecular properties of APP and secretases such as the association and dissociation of the nanoclusters or ability to control their lateral exchange instead of modulating the enzymatic properties of the secretases which could be the focus for next generation therapeutic targets (Kedia and Nair, 2020).

APP and secretases are present in multiple compartments of neurons and are known to interact among themselves as well as with several other molecules crucial for synaptic maintenance and neuronal function (Chen et al., 2015; De Strooper et al., 2010; Haapasalo and Kovacs, 2011; Muller et al., 2017). The selective interactions with these molecules could alter the localization and availability of APP and secretases involved in both amyloidogenic and non-amyloidogenic pathways. It is also known that association of APP among themselves and its family of molecules can influence the generation of Aβ (Eggert et al., 2018; Gorman et al., 2008; Khalifa et al., 2010). In addition to the foresaid variables, post-translational modifications of APP and secretase as well as its association with lipid rafts is known to interfere with the rate of formation of different proteoforms (Grimm et al., 2017; Hicks et al., 2012; Rajendran and Annaert, 2012). However, the intracellular conditions that decide the probability of occurrence of specific pathways could be determined by the availability of molecules at a specific functional zone of the synapse. It remains to be seen how each of these interactions could contribute to the local trafficking and confinement rates that would decide the nature of rate of formation of both canonical and non-canonical proteoforms of Aβ.

Our study sheds light on how the changes in the nanoorganization of amyloidogenic machinery at PSD and EZ could contribute to the alterations in dynamic load of amyloidogenic processing. The local increase in concentration of APP and secretases directly influences the load of these molecules in each vesicle, contributing to the overall increase in Aβ production. Moreover, a gradual increase in such an association would also be able to influence endocytic processes occurring distant from synapses, that could affect the production of Aβ not only in synapses but also in extra-synaptic compartments. Finally, the existence of mega-Dalton rich complexes containing machinery for canonical and non-canonical processing of Aβ has already shown to generate Aβ in vitro (Liu et al., 2019a). Our study confirms that at least few of these nanodomains could be a potential locus for these mega-Dalton-rich complexes with the ability to influence Aβ production. Though, the mechanism of Aβ generation in these complexes remains vague, it would still support the existence of different Aβ pools.

Further, recent observations demonstrate the nanoscale architecture of human Aβ plaques revealing a dense core with a peripheral halo (Querol-Vilaseca et al., 2019). The spatial elevation of APP within nanodomains would complement either the amyloidogenic or non-amyloidogenic processing of APP. This local increase has the potential to create a nucleus for forming a dense core with higher order Aβ species aggregating in specific patterns with higher packing density. In such a scenario, the molecular density would be higher at the center and the unorganized binding and/or unspecific aggregation of the smaller Aβ structures outside would be loosely bound, creating the halo for the observed dense core of plaques. The recent technical advances to evaluate the nanoscale aggregation properties of soluble and insoluble proteins in vivo and in vitro would allow us to examine this in real-time at molecular scales (Balzarotti et al., 2017; Sezgin et al., 2019). Combination of high-resolution studies with labeling strategies enabling better detection of subcellular structures in 3D and super-resolution imaging of endogenous molecules tagged by single cell gene editing strategies would provide insights on the nanoscale variability of chemical reactions at single excitatory synapses (Kedia et al., 2020; Nishiyama et al., 2017; Tang et al., 2016; Willems et al., 2020). These observations with minimal perturbations on the signaling pathways would enable us to better
understand the mechanistic intricacies of the molecular pathology underlying the neurodegenerative diseases such as AD.

The different models used to evaluate the nanoscale organization of β-amyloidogenic machinery converge to provide evidence that the compositional nature of this machinery is altered at synapses, and is a critical determinant in deciding the shift in equilibrium toward β-amyloidogenic pathway. This is in resonance with our investigations in multiple models of AD, wherein we show an increase in the content of β-secretase and APP in nanodomains. This supports our hypothesis that the availability of β-secretase and APP in nanodomains of subsynaptic compartments can be a limiting factor for β-amyloidogenic processing of APP. Each synapse integrates to form a system that is regulated by both local and global homeostasis, where the local signature of the machinery becomes the decisive factor for β-amyloidogenic processing. Altogether, we describe a holistic approach for the systematic investigation of AD as a synaptopathy. This approach uncovers a fundamental nanomachinery, where alteration in real-time molecular interactions in the scale of milliseconds to minutes can contribute toward long-term deficits such as those seen in AD.

Limitations of the study
We focused our observations based on 2D super-resolution imaging paradigms. Though the resolution is improved laterally, we believe a finer paradigm that improves axial resolution can be used such as 3D super-resolution imaging. In addition to this, employing volumetric labeling in combination with 3D super-resolution imaging would define the geometrical representation of sub neuronal structures with better accuracy. Secondly, APP/PS1 mutant mice expresses humanized APP within Aß region (that bears the Swedish mutation, APP-Swe) along with PS1ΔE9. Both are expressed under the control of the mouse prion promoter (Jankowsky et al., 2001). We believe that the use of better models where transgenes are not over-expressed or selective enrichment of neurons with single gene editing methods would allow us to quantify the molecular changes happening precisely at individual synapses. The final limitation of the study is associated with modeling, where we used the mobility and immobilization kinetics of APP molecules to mimic the reaction-diffusion kinetics in the vesicles. The mobility of β-secretase molecules was assumed to be similar to APP. These parameters could vary in membrane bound compartments where the intraluminal pH is acidic in nature. It is an open question if the endocytosed clusters would remain as such or would break apart inside the vesicles. Despite these predictions, both the cases (presence of a cluster or dissociation of cluster into monomers) would result in elevated CTFß levels in membrane bound compartments.

Resource availability

Lead contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Deepak Nair (deepak@iisc.ac.in).

Materials availability
The study did not yield any new reagents.

Data and code availability
The study did not generate any additional data sets other than provided in the manuscript.

Methods
All methods can be found in the accompanying Transparent Methods supplemental file.

Supplemental information
Supplemental Information can be found online at https://doi.org/10.1016/j.isci.2020.101924.

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Author contributions
SK and DN designed research, SK performed all the experiments unless otherwise indicated, SK and DN performed analysis, PRN and MJ prepared neuronal cell cultures, SK² and SSS shared reagents and SK² performed immunoblotting. PR performed computational model simulations, SK, PR, DN, and SN designed the model and wrote the relevant results and methods for the simulations. AM provided the human brain samples and performed neuropathological characterizations, SK and PRN performed immunohistochemistry for human samples. NR provided mouse brain samples, SK and NR performed immunohistochemistry for mouse brain samples. NS, SK, SK², and DN performed antibody validation. SK and DN wrote the manuscript. All the authors read, provided critical inputs, and approved the final version of the manuscript. (SK (Shekhar Kedia) and SK² (Satish Kumar)).

Declaration of interests
The authors declare no competing interests.

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Supplemental Information

Alteration in synaptic nanoscale organization dictates amyloidogenic processing in Alzheimer's disease

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Figure S1. Nanoscale distribution of β-secretase in functional zones of an excitatory post-synapse. Related to Figure 1: (A) A gallery of confocal images of the individual synapses identified by automatic detection of postsynaptic marker PSD95 (green) with pseudocolour overlay of β-secretase (magenta). (B) STED image of the same synapses identified in (A). (C) Represent automatically detected regions for confocal marker identified for postsynaptic marker (black), PSD (green) and β-secretase (magenta). (D) A gallery of confocal images of the regions identified by automatic detection of perisynaptic marker Dynamin (green) puncta with pseudocolour overlay of β-secretase (magenta). (E) STED image of the regions identified in (D). (F) Represent automatically detected regions for confocal marker identified for perisynaptic compartment (black), EZ (green) and β-secretase (magenta). Black in the overlay images represents the overlap between the corresponding green and magenta images. Scale bar at (F) indicates 0.6 µm.
Figure S2. Nanoscale architecture of β-secretase clusters in different functional zones of a synapse using STED microscopy. Related to Figure 1 and Table 1: (i) (left to right) A heatmap of the nanodomain intensity of β-secretase in post and perisynapse normalized with respect to the median of the global β-secretase nanodomain intensity. (ii) (left to right) Diversity (median/IQR 25% to 75% interval) in β-secretase clusters for nanodomain length, area and intensity in post and perisynapse. (iii) (left to right) Indicate the distribution (left) and diversity (median/IQR 25% to 75% interval) (right) in observed nearest neighbor distances from β-secretase to post/perisynapse. Significance was determined by unpaired two-tailed Mann–Whitney test. n= 5669 puncta (post) and 3798 puncta (peri). *P ≤ 0.05, **P ≤ 0.01, and ***P ≤ 0.001, ns P> 0.05.
Figure S3. Nanoscale distribution of γ-secretase in functional zones of an excitatory post-synapse. Related to Figure 2: (A) A gallery of confocal images of the individual synapses identified by automatic detection of postsynaptic marker PSD95 (green) with pseudocolour overlay of γ-secretase (magenta). (B) STED image of the same synapses identified in (A). (C) Represent automatically detected regions for confocal marker identified for postsynaptic marker (black), PSD (green) and γ-secretase (magenta). (D) A gallery of confocal images of the regions identified by automatic detection of perisynaptic marker Dynamin (green) puncta with pseudocolour overlay of γ-secretase (magenta). (E) STED image of the regions identified in (D). (F) Represent automatically detected regions for confocal marker identified for perisynaptic compartment (black), EZ (green) and γ-secretase (magenta). Black in the overlay images represents the overlap between the corresponding green and magenta images. Scale bar at (F) indicates 0.6 µm.
Figure S4. Nanoscale architecture of γ-secretase clusters in different functional zones of a synapse using STED microscopy. Related to Figure 2 and Table 1: (i) (left to right) A heatmap of the nanodomain intensity of γ-secretase in post and perisynapse normalized with respect to the median of the global γ-secretase nanodomain intensity. (ii) (left to right) Diversity (median/IQR 25% to 75% interval) in γ-secretase clusters for nanodomain length, area and intensity in post and perisynapse. (iii) (left to right) Indicate the distribution (left) and diversity (median/IQR 25% to 75% interval) (right) in observed nearest neighbor distances from γ-secretase to post/perisynapse. Significance was determined by unpaired two-tailed Mann–Whitney test. n = 4936 puncta (post) and 5921 puncta (peri). *P ≤ 0.05, **P ≤ 0.01, and ***P ≤ 0.001, ns P > 0.05.
Figure S5. Nanoscale architecture of β/γ-secretase clusters with APP in the neuronal processes using STED microscopy. Related to Figure 3 and Table 1: (i) (left to right) A heatmap of the nanodomain intensity of β/γ-secretase clusters with APP normalized with respect to the median of the global β-secretase nanodomain intensity (BACE on APP and BACE on PS) or with the median of the global γ-secretase nanodomain intensity (PS on APP). (ii) (left to right) Diversity (median/IQR 25% to 75% interval) in β/γ-secretase clusters with APP in neuronal processes for nanodomain length, area and intensity. Significance was determined by unpaired two-tailed Mann–Whitney test. (iii) Comparison of RSP and RSE for quantifying colocalization of β/γ-secretase and with APP in neuronal processes. Data are represented as mean +/- SEM. Significance was determined by one-way ANOVA followed by Tukey’s multiple comparison test. (iv) (left to right) Indicate the distribution (left) and diversity (median/IQR 25% to 75% interval) (right) in observed nearest neighbor distances from β-secretase to APP, γ-secretase to APP and β-secretase to γ-secretase. Significance was determined by Kruskal-Wallis test followed by Dunn’s multiple comparison test. n= 13484 puncta (β-secretase on APP), 6033 puncta (γ-secretase on APP) and 4762 puncta ((β-secretase on γ-secretase). *P ≤ 0.05, **P ≤ 0.01, and ***P ≤ 0.001, ns P> 0.05.
Figure S6. Global levels of APP and C-terminal fragments (CTFs) is increased in APP/PS1 (Tg) mice in comparison to wildtype (WT) mice. Related to Figure 4: SDS-PAGE and western blotting of the transgenic (APP/PS1) and non-transgenic (WT) mouse brain homogenates immunoblotted with anti-APP antibody (CT15) which recognizes both murine and human APP variants and with 6E10 antibody which recognizes only human APP variants shows the overexpression of APP in the transgenic (APP/PS1) as compared to non-transgenic (WT) mice. In addition to APP, higher levels of APP-CTFs are also observed in the transgenic mice (APP/PS1) due to APP overexpression.
Figure S7. Compartmentalization of β-secretase in different functional zones of the synapse in mice brain slices using STED microscopy. Related to Figure 4: (i) (left to right) Diversity in β-secretase (median/IQR 25% to 75% interval) integrated intensity in post and perisynapse for WT and APP/PS1 Tg mice. (ii) (left to right) Diversity (median/IQR 25% to 75% interval) in observed nearest neighbor distances from β-secretase to post/perisynapse for WT and APP/PS1 Tg mice. n= 1585 (WT BACE on post), 1948 (Tg BACE on post), 1276 (WT BACE on peri), 1381 (Tg BACE on peri) puncta from 3 animals. Significance was determined by unpaired two-tailed Mann–Whitney test. *P ≤ 0.05, **P ≤ 0.01, and ***P ≤ 0.001, ns P > 0.05.
Figure S8. Heterogeneous distribution of components of β-amyloidogenic machinery in human brain slices using Airyscan microscopy. Related to Figure 5 and 6: (A, B, C, D) Visualization of a section of human brain immunostained for a marker for post/perisynapse with β-secretase (A, B) or APP (C, D) from control (A, C) and AD (B, D) using Airyscan microscopy. (F) Magnified view of the boxed regions in (E, same region marked as A*) showing discrete localization of β-secretase in different functional zones of the synapse. White in the overlay images represents the overlap between the corresponding blue, green and red images. Scale bar at (D) indicates 1.2 mm, at (E) 1 mm and at (F) 25 μm. The regions depicted in B* and B** is the same as the regions in Fig. 5A.
Figure S9. Alteration of nanoscale molecular determinants involved in β-amyloidogenic machinery in human brain slices using STED microscopy. Related to Figure 5 and 6: (A, B, C, D) Nanoscale colocalization of APP (A, B) and β-secretase (C, D) clusters in post/perisynapse in human brain slices from AD and control using STED microscopy. The intensity of APP and BACE is pseudocolour coded from white (minimum) to black (maximum) with black contours representing PSD/EZ. Scale bar in B and D indicates 0.75 µm. (i, ii) Diversity in nanodomain length and intensity for APP and β-secretase clusters in post/perisynapse in human brain slices from AD and control represented as (median/IQR 25% to 75% interval). n= 1637 (control APP on post), 1197 (AD APP on post), 1574 (control APP on peri), 1733 (AD APP on peri), 2461 (control BACE on post), 1539 (AD BACE on post), 1627 (control BACE on peri), 1595 (AD BACE on peri) puncta from 1 set of human brain of patient with AD and their corresponding control. Significance was determined by unpaired two-tailed Mann–Whitney test. *P ≤ 0.05, **P ≤ 0.01, and ***P ≤ 0.001, ns P > 0.05.
Figure S10. Compartmentalization of APP/β-secretase in different functional zones of a synapse in human brain slices using STED microscopy. Related to Figure 5 and 6: (i, ii) Diversity in APP (i) and β-secretase (ii) integrated intensity in post/perisynapse (left to right) in human brain slices from AD and control represented as (median/IQR 25% to 75% interval). (iii, iv) Diversity (median/IQR 25% to 75% interval) in observed nearest neighbor distances from APP/β-secretase to post/perisynapse (left to right) in human brain slices from AD and control. n= 1637 (control APP on post), 1197 (AD APP on post), 1574 (control APP on peri), 1733 (AD APP on peri), 2461 (control BACE on post), 1539 (AD BACE on post), 1627 (control BACE on peri), 1595 (AD BACE on peri) puncta from 1 set of human brain of patient with AD and their corresponding control. Significance was determined by unpaired two-tailed Mann–Whitney test. *P ≤ 0.05, **P ≤ 0.01, and ***P ≤ 0.001, ns P > 0.05.
Figure S11. Insilico evaluation of CTFβ production for APP-Swe show differential processing kinetics within unitary vesicles. Related to Figure 7: (i) The number of CTFβ intermediates formed in each endocytic vesicle as a function of experimentally observed diffusion coefficients for APP-Swe. The diffusion coefficients (median) corresponding to APP-WT and APP-Swe are demarcated by a vertical dotted line. The intermediate product formation is not affected by the rate of diffusion of APP within a vesicle. All other parameters were kept constant for this plot. (ii) The number of CTFβ formed in an endocytic vesicle as a function of forward reaction rate for forming CTFβ intermediates ($k_1$). (iii) Amount of CTFβ formed in an endocytic vesicle as a function of irreversible forward reaction rate for intermediates to final products ($k_2$). The other simulation parameters such as diffusion coefficients (median of APP-Swe) and reaction rates are kept constant and set to APP-Swe values, while $k_1$ and $k_2$ are varied independently in (ii) and (iii) respectively. CTFβ production substantially increases with increase in both $k_1$ and $k_2$. CTFβ formation is more sensitive to variations in $k_2$ than $k_1$ (for APP-Swe conditions). (iv) Probability to produce CTFβ is correlated with the APP nanodomain size internalized per unitary vesicle.
Figure S12. Instantaneous distribution of the density of APP molecules in an endocytic vesicle as a function of APP nanodomain size. Related to Figure 7: Increase in APP cluster size at the endocytic zone results in cumulative increase in APP per unitary endocytic vesicle.

Figure S13. β-secretase clustering increases the probability of production of CTFβ. Related to Figure 7: (i) Probability to produce CTFβ is correlated with the β-secretase nanodomain size internalized per unitary vesicle for APP-WT (left). The results obtained for APP-Swe is indicated in right.
Figure S14. Visualization of Aβ42 in Dynamin enriched regions using confocal microscopy. Related to Figure 7: (A, B) Distribution of Aβ42 in Dynamin enriched regions using confocal microscopy in primary hippocampal neurons obtained from wild type (WT) (A) and APP/PS1 transgenic (Tg) (B) mice. The intensity of Aβ42 is pseudocolour coded from white (minimum) to black (maximum) with black contours representing Dynamin enriched regions. Scale bar at (B) indicate 2.5 µm. (i) Diversity in Aβ42 (median/IQR 25% to 75% interval) average and integrated intensity in Dynamin enriched regions for neurons obtained from WT and APP/PS1 Tg mice. n= 613 (WT Aβ42/Dynamin) and 1068 (Tg Aβ42/Dynamin). Significance was determined by unpaired two-tailed Mann–Whitney test. *P ≤ 0.05, **P ≤ 0.01, and ***P ≤ 0.001, ns P > 0.05.
Table S1: Summary of quantitative estimation of morphological and biophysical properties of different nanodomains for mouse and human brain slices imaged by STED microscopy (Related to Figure 4, 5 and 6)

| Category/Parameter | Length (µm) | Intensity (a.u.) |
|-------------------|-------------|-----------------|
| **WT nanodomain** | **0.152±/0.003** | 167.10+/7.57 (107.90, 53.24-216.70) |
| **β/PSD** | **(0.133, 0.079-0.192)** | |
| **Tg nanodomain** | **0.137±/0.003** | 168.90+/5.44 (118.60, 76.70-206.00) |
| **β/PSD** | **(0.116, 0.093-0.162)** | |
| **WT nanodomain** | **0.113±/0.004** | 60.63+/5.08 (34.57, 16.27-71.54) |
| **β/EZ** | **(0.101, 0.066-0.147)** | |
| **Tg nanodomain** | **0.092±/0.003** | 127.90+/8.74 (82.55, 37.33-146.60) |
| **β/EZ** | **(0.072, 0.033-0.120)** | |
| **Control nanodomain** | **APP/PSD** | **0.055±/0.003** | 18.27+/6.16 (5.97, 4.96-8.43) |
| **APP/EZ** | **0.094±/0.001** | 13.32+/1.03 (7.65, 5.77-11.09) |
| **AD nanodomain** | **APP/PSD** | **0.074±/0.004** | 17.52+/2.73 (10.22, 8.62-14.47) |
| **APP/EZ** | **0.094±/0.001** | 13.32+/1.03 (7.65, 5.77-11.09) |
| **Control nanodomain** | **APP/PSD** | **0.068±/0.003** | 96.13+/13.06 (27.43, 11.95-95.32) |
| **APP/EZ** | **0.068±/0.003** | 96.13+/13.06 (27.43, 11.95-95.32) |
| **AD nanodomain** | **β/PSD** | **0.069±/0.002** | 20.42+/3.68 (7.79, 6.19-10.83) |
| **β/EZ** | **0.070±/0.003** | 37.82+/4.50 (8.38, 6.27-13.72) |
| **Control nanodomain** | **β/PSD** | **0.086±/0.002** | 24.11+/3.57 (11.03, 8.77-14.98) |
| **β/EZ** | **0.086±/0.002** | 24.11+/3.57 (11.03, 8.77-14.98) |

*Mice slices and **human brain slices. Values indicated are Mean+/SEM while values in brackets represent median, IQR from 25% percentile to 75% percentile*
Table S2: Summary of parameters employed to instantiate the unitary vesicle (Related to Figure 7)

| Parameter                                      | Value                  |
|-----------------------------------------------|------------------------|
| Diameter of Unitary Vesicle*                  | 0.120 µm               |
| Area of Unitary Vesicle*                      | 0.045 µm²              |
| Density of APP molecules ‡                   | 1000 molecules/µm²     |
| Density of β-secretase molecules              | 5600 molecules/µm²     |
| Number of APP molecules‡                      | 45                     |
| Number of β-secretase molecules               | 252                    |
| Number of APP clusters‡                       | 1                      |
| Number of β-secretase clusters                | 0                      |
| Area of APP nanodomain†                       | 0.0056 µm²             |
| Area of β-secretase nanodomain                | No nanodomain used     |
| Diffusion coefficient of APP-WT monomers‡     | 0.083 µm²/S            |
| Diffusion coefficient of immobilized APP-WT molecules‡ | 0.013 µm²/S         |
| Diffusion coefficient of APP-Swe monomers‡    | 0.020 µm²/S            |
| Diffusion coefficient of immobilized APP-Swe molecules‡ | 0.002 µm²/S         |
| $K_1$ APP-WT                                  | $4.0 \times 10^{-6}$ µm²·Number·s⁻¹ |
| $K_1$ APP-WT**                                | 0.1 s⁻¹                |
| $K_2$ APP-WT**                                | 0.002 s⁻¹              |
| $K_1$ APP-Swe                                 | $1.25 \times 10^{-5}$ µm²·Number·s⁻¹ |
| $K_1$ APP-Swe**                               | 0.1 s⁻¹                |
| $K_2$ APP-Swe**                               | 0.025 s⁻¹              |
| Real-time duration of simulation              | 5 seconds              |

**(Ben Halima et al., 2016), *(Kumari et al., 2010; Watanabe et al., 2014), ‡(Kedia et al., 2020)**
Table S3: Summary of primary and secondary antibodies used for different experiments (Related to Figure 1, 2, 3, 4, 5 and 6)

| Antibody/Experiment          | Dilution | Catalogue Number | Company                      |
|------------------------------|----------|------------------|------------------------------|
| **Immunocytochemistry**      |          |                  |                              |
| (Figures 1, 2, 3 & S1, S3, S14) |          |                  |                              |
| Anti-PSD95                   | 1:500    | MA1-046          | Thermo Scientific            |
| Anti-Dynamin                 | 1:1000   | 05-319           | Upstate/Millipore            |
| Anti-APP-CT                  | 1:500    | 171610           | Calbiochem/Millipore         |
| Anti-APP-CT                  | 1:500    | MAB343-C         | Merck Millipore              |
| Anti-Presenilin1             | 1:500    | MAB232           | Merck Millipore              |
| Anti-Presenilin1             | 1:500    | PRS4203          | Sigma-Aldrich                |
| Anti-BACE1*                  | 1:500    | 840101           | Biolegend/Covance            |
| Anti-β-Amyloid (1-42)        | 1:50     | 805509           | Biolegend/Covance            |
| Alexa Fluor 594              | 1:400    | A11037           | Life Technologies            |
| Alexa Fluor 594              | 1:400    | A11032           | Life Technologies            |
| Abberior Star Red            | 1:400    | 2-0002-011-2     | Abberior                     |
| **Immunohistochemistry**     |          |                  |                              |
| (Figures 4, 5, 6 & S8, S9)   |          |                  |                              |
| Anti-Shank2                  | 1:500    | 162204           | Synaptic Systems             |
| Anti-Claathrin               | 1:100    | ab2731           | Abcam                        |
| Anti-APP-NT*                 | 1:200    | 3207             | Inhouse Developed            |
| Anti-BACE1*                  | 1:200    | 840101           | Biolegend/Covance            |
| Alexa Fluor 594              | 1:200    | A11037           | Life Technologies            |
| Alexa Fluor 488              | 1:200    | A11029           | Life Technologies            |
| Alexa Fluor 555              | 1:200    | A21435           | Life Technologies            |
| Alexa Fluor 647              | 1:200    | A21245           | Life Technologies            |
| Abberior Star Red            | 1:200    | 2-0002-011-2     | Abberior                     |
| Abberior Star Red            | 1:200    | 2-0112-011-8     | Abberior                     |
| **Immunoblotting**           |          |                  |                              |
| (Figures 4 & S6)             |          |                  |                              |
| 6E10 antibody (1-16)         | 1:1000   | 803004           | Biolegend/Covance            |
| Anti-BACE                    | 1:500    | 5606S            | Cell Signaling Technology    |
| Anti-APP-CT (CT15)**         | 1:250    | CT15             | Inhouse Developed            |
| Anti-β-Actin                 | 1:5000   | AC-15 (A1978)    | Sigma                        |

*Specificity of the BACE1 antibody (840101) and APP-NT antibody (3207) was in-house validated
**(Wahle et al., 2006)
**Table S4:** Summary of neuropathological and clinical data for control and AD human case (Related to Figure 5 and 6)

| ID     | Age (yrs.) | Gender | PMI  | Braak Stage | Aβ Phase | Clinical Diagnosis                              | Cause of Death           |
|--------|------------|--------|------|-------------|----------|------------------------------------------------|--------------------------|
| R493/16 (Control)* | 92 | Female | 15 h | B2-Braak stage III/IV | A2-Thal phase III | No history of cognitive decline/dementia | Cardiorespiratory arrest |
| 15/B271 (AD)* | 92 | Female | 5 h  | B2-Braak stage III/IV | A2-Thal phase III | Alzheimer's disease | Pneumonia |
| 12/B243Z (AD)** | 82 | Female | Not available | B1-Braak stage I | A1-Thal phase I | Alzheimer's disease | Pneumonia |
| 19/B352Q (AD)** | 76 | Female | 8 hrs | B2-Braak stage III/IV | A2-Thal phase III | Alzheimer's disease | Myocardial infarction |
| 14/B258T (AD)** | 83 | Male | Not available | B2-Braak stage IV | A2-Thal phase III | Alzheimer's disease | Cardiorespiratory arrest |
| 8L2L (Control)** | 88 | Male | Not available | B0 | A0 | No history of cognitive decline/dementia | Pneumonia |
| 4L2K (Control)** | 81 | Female | Not available | B2-Braak stage III | A2-Thal phase II | No history of cognitive decline/dementia | Cardiorespiratory arrest |
| 9L2R (Control)** | 75 | Male | Not available | B1-Braak stage I | A1-Thal phase I | No history of cognitive decline/dementia | Pneumonia |

Post-mortem brain interval time (PMI), *for STED microscopy, **for Airy scan microscopy
Transparent Methods

Experimental Animals
Wild type (WT) C57BL/6 mice or APPswe/PS1ΔE9 double transgenic (Tg) mice (JAX Stock# 004462) were bred at the Institutional Central Animal Facility and were maintained in a temperature-controlled room on 12 h light/12 h dark cycle under pathogen free environment with ad libitum access to water and food. All experiments involving animals were carried out in accordance with institutional guidelines for the use and care of animals after approval from the Institutional Animal Ethics Committee (IAEC), Indian Institute of Science, Bangalore, India.

Primary Hippocampal Culture
Mixed sex primary hippocampal neurons cultured from postnatal day 0 or 1 (P0-P1) wild type C57BL/6 mice or APPswe/PS1ΔE9 double transgenic (Tg) mice (JAX Stock# 004462) were prepared and maintained (Kedia et al., 2020). The cells were seeded at a density of 0.1×10⁶ cells/mL in 18 mm #1.5 (corrected for 0.17+/- 0.01) glass coverslips (coated with poly-D-lysine at a concentration of 100 µg/mL) in a 12-well cell culture plate. Primary hippocampal neurons were used for immunocytochemical evaluation at DIV 20-21.

Antibodies
The details of the primary and secondary antibodies used in this study are summarized (Table S3).

Immunocytochemistry
Cells were fixed with 4% paraformaldehyde plus 4% sucrose in PBS at 4°C for 10 minutes, followed by quenching with 0.1M glycine in PBS at room temperature and permeabilization with 0.25% Triton X-100 for 5 minutes and then blocked with 10% BSA in PBS for 30 minutes at room temperature. This was followed by incubation with the appropriate primary antibody for 1-2 hr. Following washing, cells were then incubated with a suitable secondary antibody for 45 minutes (Kedia et al., 2020). Following washing, cells were mounted with prolong (Molecular Probes, cat. no. MAN0010261) for confocal or STED imaging.

Mouse Brain Lysates Preparation and Immunoblotting
APPswe/PS1ΔE9 double transgenic (Tg) mice were obtained from Jackson Laboratories, USA (JAX Stock# 004462). The institutional and national guidelines for the care and use of laboratory animals were followed. Brains from 3 months old Tg and or wild type (WT) mice were perfused transcardially with ice-cold saline and removed from the skull. Brain hemispheres were snap-frozen in liquid nitrogen and stored at -80°C until further use. Frozen brains were homogenized with a douncer followed by sonication in 0.32 M sucrose in 50 mM Tris buffer (pH 7.3) containing protease and phosphatase inhibitors (Roche Diagnostics, Germany). Mouse brain homogenates were cleared by centrifugation at 16,100 g and 4°C for 30 minutes. After centrifugation, the resulting supernatant sucrose extract containing the soluble proteins was collected and stored at -80°C (Kumar et al., 2016). The total protein concentration was determined by the BCA protein assay kit (Thermo Scientific, USA).

Mouse brain lysates were separated by 4–12% NuPAGE (ThermoFisher Scientific) using MES buffer and transferred on to nitrocellulose membranes. The blots were blocked in 3% BSA (diluted in 1x TBS/T) for 2 hours. APP, APP-CTFs, BACE and β-actin Proteins were detected with 6E10 (BioLegend), D10E5 (Cell Signaling Tech) and AC15 (Sigma/Merck) the indicated primary antibodies and respective horseradish peroxidase-conjugated secondary antibodies or conjugated fluorescent antibodies (IRDye 680RD and IRDye 800CW). The same immunoblots were reprobed with anti-β-actin antibody and used as loading control. Immunoreactivity was detected by enhanced chemiluminescence reaction (Biorad) or near infrared detection (LI-COR Odyssey). Band intensities were analyzed using Chemidoc XRS documentation system (Bio-Rad) and Image Studio 5.x CLx software (LI-COR). The bands were compared with an overlay of See Blue prestained ladder (#LC5295, Invitrogen) to compare the bands obtained in the immunoblot.

Immunohistochemistry (Mice)
Mice (3-4 months old) were transcardially perfused with 15 ml of ice-cold PBS followed by 50 ml of ice-cold 4% paraformaldehyde (pH 7.4). The brains were post-fixed in 4% PFA overnight, cryoprotected in 30% sucrose, frozen and stored at -80°C until use. For labelling, 25 µm coronal cryosections from Tg (APPswe/PS1ΔE9, JAX Stock# 004462) mice and control littermates were cut in a cryostat and incubated in blocking/permeabilization solution containing 1% normal goat serum, 3% BSA and 0.3% Triton-X-100 in PBS. Free floating sections were incubated overnight with primary antibodies at 4°C. The sections were washed and incubated with the secondary antibodies at room temperature for 1 h. Following washing, the sections were mounted with prolong with DAPI (Molecular Probes, cat. no. P36962) for confocal/STED imaging. The primary and secondary antibodies used are indicated (Table S3) (Ramanan et al., 2005). The imaging was performed from CA1 to CA2 stratum radiatum layer of the hippocampus.

Human Tissue Collection and Preparation
The brain was removed at autopsy following the written informed consent of next of kin. It was fixed in 10% neutral buffered formalin for 3-4 weeks following which it was sliced serially in the coronal plane. The findings in the gross examination were recorded (cortical atrophy, hippocampal atrophy, infarcts, atherosclerosis). Neuroanatomical areas were sampled in
Immunohistochemistry (Human)
The tissues were processed for routine paraffin embedding. 4 μm thick serial sections were cut and stained with haematoxylin & eosin stains for morphological assessment. Serial sections were collected on positively charged slides for performing immunohistochemistry by indirect immunoperoxidase method, following antigen retrieval by heat and DAB/H$_2$O$_2$ as the chromogen to visualize immunolabelling (DAKO Envision Detection System) to detect neurodegenerative changes using specific antibodies to Paired Helical Filaments (PHF), phosphorylated Tau to detect Neurofibrillary Tangles (NFT), neuritic plaques and neuropil threads (Clone Tau 5, mouse monoclonal antibody, 1:50 dilution, BioGenex, USA) and β-amyloid antibody to detect senile plaques and vascular amyloid deposition (Clone 6F/3D, mouse, monoclonal antibody, 1:100 dilution, Leica Biosystems, USA. The number of Tau positive Neurofibrillary tangles (NFTs), Aβ positive Senile Plaques (SPs) and Diffuse Plaques (DPs) were counted in accordance with published studies (Purohit et al., 2011).

The AD neuropathological changes were ranked according to three to three parameters: the Aβ plaque score (Thal et al., 2002), Braak and Braak NFT stage (Braak et al., 2006) and the CERAD NP score (Mirra et al., 1991) to obtain an “ABC score” reported as 4 levels: not, low, intermediate low, intermediate and high (Montine et al., 2012). The score of the blessed dementia rating scale of the control case was 1.5/17. The summary of staging and pathological details is available (Table S4).

Immunofluorescence (Human)
Paraffin embedded tissue sections were obtained from control and AD cases. Sections were first deparaffinized in xylene and then dehydrated through a series of grades of ethanol. Following quenching and washing, heat induced antigen retrieval was performed using sodium citrate buffer (pH 6.0). Sections were then blocked using universal blocking reagent (cat. no. HK085-5K, BioGenex, USA) for 1 h. Sections were further incubated overnight with primary antibodies at 4°C. Following washing and incubation with the secondary antibodies at room temperature for 1 h and washing, sections were mounted with prolong with DAPI (Molecular Probes, cat. no. P36962) for confocal/STED imaging. Slides were stored at 4°C in dark. The primary and secondary antibodies used are indicated (Table S3). The imaging was performed from CA1 to CA2 radiatum layer of the hippocampus.

Stimulated Emission Depletion microscopy (STED)
A commercial STED inverted microscope (Abberior Expert Line 775 nm, Abberior Instruments GmbH, Göttingen, Germany) was used to obtain confocal and super-resolved images of the same region with a sampling of 15 nm. The microscope was equipped with two pulsed excitation lasers at 561 nm and 640 nm and a pulsed depletion laser at 775 nm. The laser powers were adjusted to 70%, 50% and 40% of their respective total power for 561 nm, 640 nm and 775 nm, respectively (Kedia et al., 2020).

Confocal Microscopy
Confocal microscopy imaging was performed on Zeiss LSM 780 or Zeiss LSM Zeiss 880 at a sampling of 35-100nm/pixel. For each experimental conditions the acquisition and illumination criteria were kept constant (Kedia et al., 2020; Venkatesan et al., 2020).

Airyscan super-resolution Microscopy
Immunohistochemical samples from AD or control human brain co-labelled with nuclear, post synaptic, and endocytic markers together with either APP or BACE were used for super-resolution imaging using Airyscan microscopy. Airyscan was performed on Zeiss LSM 880 equipped with 32 array detectors for acquisition of super-resolution images. We first performed confocal imaging of hippocampal regions using a 40X objective of numerical aperture of 1.3 at a sampling of 11 μm/pixel. Each tile was 350X350 μm$^2$. We then created a mosaic from the individual tiles to reconstruct the hippocampal region. This was performed by tiling images to generate a mosaic image of 3.5X3.5 mm$^2$ to 4.5X4.5 mm$^2$. We then performed Airyscan imaging on 5-10 non-overlapping regions from the radiatum layer of the hippocampus with an effective field size of 75 X75 μm$^2$ sampled at 35 nm/pixel using a 63X objective with a numerical aperture of 1.4. For image acquisition, 405, 488, 543 and 633 nm lasers were used. The illumination intensities, sampling of the images, digital and analogue gain of the detectors, emission window for each fluorescent channel and their corresponding pinhole sizes were maintained constant across acquisition. The raw images acquired using Airyscan mode were processed using Zeiss acquisition and analysis software of the microscope to generate final super-resolution images. The reconstruction parameters were kept constant throughout the samples.

Semiautomated detection of dendritic compartments and functional zones of an excitatory synapse
The active dendritic area of protein of interest and synapses were distinguished from the rest of the dendrite using a custom defined segmentation protocol. The intensity of the epifluorescence/confocal images of markers for different functional zones of the synapse (post/peri) was thresholded to generate the mask of the puncta. A spine morphometry analysis was then performed, and masks were filtered using various morphological filters like length, breadth and area through IMA plugin.
running inside MetaMorph software (Molecular Devices). Similar analysis was performed on super-resolution images to detect functional zones of an excitatory synapse (post/peri) that correspond to PSD/EZ functional zones (Kedia et al., 2020).

**Resolution Scaled Pearson’s Coefficient (RSP) and Resolution Scaled Error (RSE) Analysis**

RSP and RSE were calculated using NanoJ-SQUIRREL, a plugin supported by Fiji (Culley et al., 2018; Venkatachalapathy et al., 2019). The reference image was either a marker for functional zones of the synapse for β/γ-secretase on post/peri analysis or APP or γ-secretase for β/γ-secretase on APP and β-secretase on γ-secretase analysis (Kedia et al., 2020). The subject image was of the protein of interest (secretases). Both reference and subject images were super-resolution images.

**Super-resolution Cluster Analysis**

Clusters of molecular aggregation (nanodomains) were identified from super-resolution images by a custom algorithm written as a plug-in supported by MetaMorph (Molecular Devices) (Izaddin et al., 2012; Keckhar et al., 2013; Nair et al., 2013). Nanodomains were detected from super-resolution images using the Palm-Tracer plugin. Nanodomains were analyzed using bi-dimensional Gaussian fitting, from which the principal (2.3σ_long) and the auxiliary axes (2.3σ_short, data not shown) were determined for STED while for the data from Airyscan, length0 (σ_long) was plotted. The fitting was performed on each cluster that was identified as a domain. Several parameters like area and intensity of nanodomain were computed for each experimental category. The intensity of the nanodomain was normalized with the global median/mean value to allow relative comparison of the content of nanodomains either between different pools of nanodomains when associated with functional zones of the synapse or with other components of amyloidogenic machinery.

**Nearest Neighbor Distance Measurement**

The nearest neighbor distances (NND) were calculated using interaction analysis, a plugin supported by Fiji (Helmuth et al., 2010; Kedia et al., 2020; Shivanandan et al., 2013). The reference image (Y) was post/perisynaptic marker and image (X) was the protein of interest in all the categories where analysis is performed between the synaptic marker and the protein of interest. While for BACE-APP, PS-APP and BACE-PS, the NNDs were calculated with Y as APP for BACE-APP, PS-APP and as PS for BACE-PS. The ROUT method of identifying outliers with Q= 1% was used for NND measurements.

**Model Components and Geometry**

Simulations were performed in a spherical vesicle of diameter 120 nm to reflect a typical endocytic vesicle. The details of variables used to instantiate the unitary vesicle is summarized in Table S2. MCell, version 3, a Monte Carlo Cell simulator, was used to carry out the simulations. Monte Carlo algorithms were used by MCell to simulate the diffusion of individual molecules present either on a surface or in a confined volume (Kerr et al., 2008). MCell carries out user-specified molecular reactions stochastically. These simulations track each molecule and the relevant reactions to calculate spatiotemporal trajectories. Simulations were performed on a cluster with 1464 processing units. Several thousands of trajectories (1000-5000, the higher number of trajectories simulated for reactions with low reaction rates for greater confidence) were simulated to compute the average reaction diffusion trajectory for APP with β-secretase.

**Model Configurations**

Amyloidogenic processing of APP occurs due to sequential activities of β- and γ-secretases. Processing by β-secretase is a critical and necessary condition for the generation of Aβ. To understand this processing step, β-secretase and APP were modelled with varying compositionality inside endocytic vesicles. The origin of these endocytic vesicles was randomly chosen from perisynaptic compartments, where the distribution of these molecules was quantified through nanoscopy. The following three assumptions were then used to define the characteristics of endocytic vesicles. 1) Endocytic vesicles originate from EZ and are instantiated as confined spherical membrane bound organelles with a luminal pH of 5.5 and a diameter of 20 nm. 2) β-secretase is distributed uniformly in the vesicular membrane while the compositionality of APP changes from a vesicle is populated with 36 single molecules of APP with a combination of no nanodomains or nanodomain_APP of 5, 9 or 13 confined APP molecules residing in an area of 0.0056 µm². 3) The vesicle has no other substrates for β-secretase apart from APP. Approximately 250 β-secretase molecules were distributed uniformly on the vesicles. The diffusion coefficient of the confined APP molecules within nanodomain_APP (Table S2) was taken to be 1×10⁻² µm²/s, while that of APP monomer was taken to be 7×10⁻² µm²/s (Kedia et al., 2020). Since β-secretase is a single pass transmembrane protein like APP and its diffusion coefficient has been estimated to be of the same order of magnitude as the APP monomer, we assumed the diffusion coefficient of β-secretase monomer to be 7×10⁻² µm²/s as well. To simulate APP- β-secretase reaction, we assumed the APP processing to follow Michaelis- Menten kinetics (Ben Halima et al., 2016). Both the rates k_cat (catalytic rate) and the k_M (the Michaelis constant) for APP-WT and APP-Swe have been measured by Ben Halima et al. at. al and were used here for the simulations (Ben Halima et al., 2016). As these constants were obtained experimentally under conditions that facilitated 3D diffusion of molecules, only the k_cat value was taken from their data while the k_1 and k⁻¹ were set to values such that a reasonable number of intermediate molecules were formed. k_M for APP-Swe was approximately 7 times that of APP-WT and was simulated appropriately. The k_cat value for APP-Swe was used as reported in Ben Halima et al., 2016 (Ben Halima et al., 2016). The details of variables used for the simulation is summarized (Table S2).
Statistics
Statistical analysis and significances were performed using GraphPad Prism version 7.04 for Windows, GraphPad Software, La Jolla California USA, (www.graphpad.com). D’Agostino-Pearson Omnibus normality test and Shapiro-Wilk normality test were used to test normal distribution. All statistical values were shown as mean +/- SEM for normally distributed data or median (IQR 25% to 75% interval) for non-normally distributed data, unless otherwise indicated. Tabulated summary data are presented as mean +/- SEM (median, IQR 25% to 75% interval). Normally distributed datasets were compared using two-tailed unpaired Student’s t-test (for two-group), one-way analysis of variance (ANOVA) test followed by Tukey’s multiple comparison test (for multi-group). Non-normally distributed datasets were tested by non-parametric two-tailed Mann-Whitney test (for two-group) or Kruskal-Wallis test followed by Dunn’s multiple comparison test (for multi-group). Indications of significance correspond to P values *P ≤ 0.05, **P ≤ 0.01, and ***P ≤ 0.001, ns P > 0.05. The calculation of the required sample sizes was obtained from the power and sample size calculator from statistical solutions (Nair et al., 2013). To account for variability, data were obtained from 2-4 independent cultures/animals.

Schemes
All the schemes and graphical abstract were prepared using Biorender.com

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