



Amyloid β production is regulated by β 2-adrenergic signaling-mediated post-translational modifications of the ryanodine receptor

Received for publication, June 10, 2016, and in revised form, May 2, 2017. Published, Papers in Press, May 5, 2017, DOI 10.1074/jbc.M116.743070

Renaud Bussiere^{‡1}, Alain Lacampagne^{§1}, Steven Reiken^{¶1}, Xiaoping Liu[¶], Valerie Scheuerman[§], Ran Zalk[¶], Cécile Martin[‡], Frederic Checler^{‡2}, Andrew R. Marks^{¶2,3}, and Mounia Chami^{‡2,4}

From the [‡]Université Côte d'Azur, CNRS, IPMC, France, "Labex Distalz," 660 route des Lucioles, 06560 Sophia-Antipolis, Valbonne, France, [§]INSERM U1046, CNRS UMR9214, CNRS LIA1185, Université de Montpellier, CHRU Montpellier, 34295 Montpellier, France, and [¶]Department of Physiology and Cellular Biophysics, Clyde and Helen Wu Center for Molecular Cardiology, Columbia University College of Physicians and Surgeons, New York, New York 10032

Edited by Paul E. Fraser

Alteration of ryanodine receptor (RyR)-mediated calcium (Ca^{2+}) signaling has been reported in Alzheimer disease (AD) models. However, the molecular mechanisms underlying altered RyR-mediated intracellular Ca^{2+} release in AD remain to be fully elucidated. We report here that RyR2 undergoes post-translational modifications (phosphorylation, oxidation, and nitrosylation) in SH-SY5Y neuroblastoma cells expressing the β -amyloid precursor protein (β APP) harboring the familial double Swedish mutations (APP^{swe}). RyR2 macromolecular complex remodeling, characterized by depletion of the regulatory protein calstabin2, resulted in increased cytosolic Ca^{2+} levels and mitochondrial oxidative stress. We also report a functional interplay between amyloid β ($\text{A}\beta$), β -adrenergic signaling, and altered Ca^{2+} signaling via leaky RyR2 channels. Thus, post-translational modifications of RyR occur downstream of $\text{A}\beta$ through a β 2-adrenergic signaling cascade that activates PKA. RyR2 remodeling in turn enhances β APP processing. Importantly, pharmacological stabilization of the binding of calstabin2 to RyR2 channels, which prevents Ca^{2+} leakage, or blocking the β 2-adrenergic signaling cascade reduced β APP processing and the production of $\text{A}\beta$ in APP^{swe}-expressing SH-SY5Y cells. We conclude that targeting RyR-mediated Ca^{2+} leakage may be a therapeutic approach to treat AD.

Alzheimer disease (AD)⁵ is one of the leading neurodegenerative pathologies in the western countries. The two main neuropathological lesions of AD are amyloid plaques, composed mainly of amyloid β ($\text{A}\beta$) peptides, and neurofibrillary tangles, composed of hyperphosphorylated tau (1, 2). The $\text{A}\beta$ peptides, which form the core of the amyloid plaques, are produced by the sequential proteolytic cleavages of the β amyloid precursor protein (β APP). β APP can be processed by two alternative post-translational pathways: (i) an amyloidogenic route in which β APP is cleaved by β -secretase (BACE-1) to generate a soluble sAPP β fragment and a C-terminal fragment of 99 amino acids (C99), which is further cleaved by a presenilin (PS1 and PS2)-dependent γ -secretase complex to generate $\text{A}\beta$ and AICD (APP intracellular domain) and (ii) a non-amyloidogenic pathway in which β APP is sequentially hydrolyzed by α -secretase to produce a soluble sAPP α and a C-terminal fragment of 83 amino acids (C83). C83 is further cleaved by γ -secretase to generate a p3 peptide and AICD (3, 4). The amyloid cascade hypothesis is mainly supported by genetic studies indicating that autosomal dominant cases of AD (familial AD) are linked to mutations in β APP (5) and on PS1 and PS2 (6, 7), leading to modifications of $\text{A}\beta$ production.

Data are now converging to suggest an important contribution of endoplasmic reticulum (ER) Ca^{2+} homeostasis deregulation in AD pathological process (8, 9). This combines pathological ER Ca^{2+} release via the inositol 1,4,5-trisphosphate (IP_3R) (10–12), and the ryanodine (RyR) receptors (9, 13–16). Importantly, alteration of RyR-mediated Ca^{2+} release likely contributes to ER Ca^{2+} deregulation in AD (9).

This work was supported by National Institutes of Health Grants R01HL061503 (NHLBI; to A. R. M.), R01HL102040 (NHLBI; to A. R. M.), R01AR060037 (NIAMS; to A. R. M.), NIH T32 HL120826 (to A. R. M.), and R25NS076445 (NINDS; to A.R.M.). This work was also supported by Fondation Leducq and a generous donation from Carol Stix (A. R. M.) and grants from INSERM, the Philippe Foundation, and a Schaefer Award from Columbia University (to A. L.), LECMA (Ligue Européenne Contre la Maladie d'Alzheimer) (to M. C.), LABEX (Laboratory of excellence, program investment for the future), DISTALZ (Development of Innovative Strategies for a Transdisciplinary approach to Alzheimer's disease) (to F. C.), and Hospital-University Federation (FHU OncoAge) (to F. C.). A. R. M. is a board member and owns shares in ARMGO Pharma Inc., which is targeting RyR channels for therapeutic purposes. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health.

✂ Author's Choice—Final version free via Creative Commons CC-BY license.

This article contains supplemental Figs. 1.

¹ These authors should be considered as co-first authors.

² These authors should be considered as co-senior authors.

³ To whom correspondence may be addressed. Tel.: 212-851-5348; Fax: 212-851-5346; E-mail: arm42@columbia.edu.

⁴ To whom correspondence may be addressed. Tel.: 33-493953457; Fax: 33-493953408; E-mail: mchami@ipmc.cnrs.fr.

⁵ The abbreviations used are: AD, Alzheimer disease; $\text{A}\beta$, amyloid β ; β APP, β -amyloid precursor protein; AICD, APP intracellular domain; β 2-AR, β 2-adrenergic receptor; CTF, C-terminal fragments; PP1, serine/threonine protein phosphatase 1; RyR, ryanodine receptor; SERCA, sarco-endoplasmic reticulum Ca^{2+} -ATPase; C99, C-terminal fragment of 99 amino acids; C83, C-terminal fragment of 83 amino acids; PS, presenilin; ER, endoplasmic reticulum; IP_3R , inositol 1,4,5-trisphosphate; ICI, ICI 118–551; ROS, reactive oxygen species; EEA1, early endosome antigen 1; LC, locus coeruleus; CGP 20712A, (\pm)-2-hydroxy-5-[2-[[2-hydroxy-3-[4-[1-methyl-4-(trifluoromethyl)-1H-imidazol-2-yl]phenoxy]propyl] amino]ethoxy]-benzamide methanesulfonate salt; ICI 118,551, hydrochloride ((\pm)-1-[2,3-(dihydro-7-methyl-1H-inden-4-yl)oxy]-3-[(1-methylethyl)amino]-2-butanol hydrochloride; Tricine, N-[2-hydroxy-1,1-bis(hydroxymethyl)ethyl]glycine; ANOVA, analysis of variance; DNP, 2,4-dinitrophenylhydrazine.

RyR post-translational modifications enhance A β production

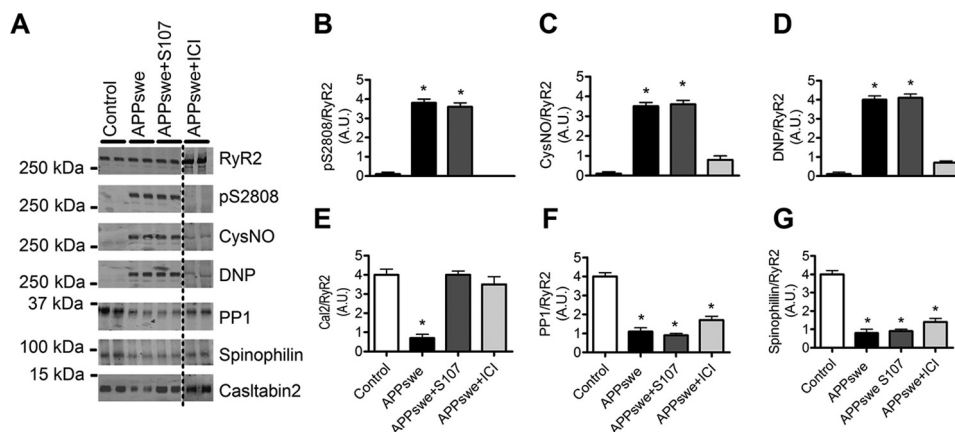


Figure 1. Remodeling of RyR2 macromolecular complex in SH-SY5Y neuroblastoma cell line stably expressing APPswe. A, representative SDS-PAGE analyses and quantification of RyR2 immunoprecipitated from the human SH-SY5Y neuroblastoma cell line stably expressing APPswe or mock vector used as control (*Control*). RyR2 was immunoprecipitated and immunoblotted for RyR2-pS2808 (RyR2 PKA phosphorylation site), S-nitrosylation (*CysNO*) and oxidation (*DNP*) as well as components of the RyR2 channel complex including: protein phosphatase 1 (*PP1*) and its anchoring protein spinophilin and calstabin2. Cells were treated with either S107 (10 μ M, for 12 h) or ICI118-551 (*ICI*) (1 μ M, for 12 h). APPswe+ICI treatment was conducted in the same set of experiments as the control, APPswe, and APPswe+S107 but was run on a separate membrane. B–G, graphs represent the mean \pm S.E. obtained from three independent experiments. *, $p < 0.05$ calculated versus control using one-way ANOVA and Bonferroni post-test. A.U., arbitrary units.

RyR dysfunction has been reported in AD models; however, the molecular mechanisms underlying RyR-mediated ER Ca^{2+} leak in AD are still not fully understood. RyR macromolecular complexes include four RyR protomers (565 kDa each) and several regulatory proteins including the RyR channel-stabilizing subunit calstabin. Recent studies have revealed that RyR-mediated ER Ca^{2+} depletion is linked to post-translational modifications (hypernitrosylation, hyperphosphorylation, oxidation) of the RyR macromolecular complexes resulting in calstabin depletion and “leaky RyR channels” (17–19).

β 2-Adrenergic receptors (β 2-ARs) have been implicated in the development of AD both in humans and in AD animal models (20–23). However, the association between β -adrenergic signaling and RyR-mediated Ca^{2+} deregulation in AD has not been reported.

Results

Remodeling of RyR2 macromolecular complexes in SH-SY5Y neuroblastoma cells stably expressing APPswe

We examined post-translational modifications of RyR2 channels consistent with the biochemical signature of leaky RyR2 channels (19, 24–28) in an *in vitro* AD study model. We used human SH-SY5Y neuroblastoma cells stably expressing pcDNA3.1 (control) or human β APP harboring the double Swedish mutations (APPswe: APPKM670/671NL) constructs. We previously reported that SH-SY5Y cells expressing APPswe yield increased levels of APP C-terminal fragments (CTFs) fragments (C99 and C83) and of A β peptides (13).

RyR2 was immunoprecipitated and immunoblotted for protein kinase A (PKA) phosphorylation (at residue Ser-2808), oxidation (2,4-dinitrophenylhydrazone (DNP)), nitrosylation (anti-Cys NO), and levels of the channel stabilizing subunit calstabin2 (FKBP12.6) in the RyR2 macromolecular complex. Neuronal RyR2 from control SH-SY5Y cells had no biochemical remodeling of the RyR2 macromolecular complex, whereas APPswe-expressing cells exhibited RyR2 PKA phosphorylation, oxidation, nitrosylation, and calstabin2 depletion (Fig. 1,

A–E). RyR2 macromolecular complexes from APPswe-expressing cells also exhibited reduced levels of serine/threonine protein phosphatase 1 (PP1) and spinophilin (Fig. 1, A, F, and G) (24, 25). We then tested whether the Rycal S107 can prevent depletion of calstabin2 from RyR2 (19, 26–28). S107 had no effect on PKA phosphorylation or oxidation/nitrosylation of RyR2 (Fig. 1, A–D) but significantly prevented dissociation of calstabin2 from RyR2 channels (Fig. 1, A and E). S107 did not restore PP1 and spinophilin levels in RyR2 complexes (Fig. 1, A, F, and G). PKA phosphorylation of RyR2 occurs downstream of β 2-AR activation (24, 25). In addition, we already reported that PKA, H_2O_2 , and Noc-12 (NO-donor) individually caused a \sim 2-fold and, in combination, caused an \sim 4-fold decrease in the binding affinity of calstabin2 to neuronal RyR2 (19). Because the β 2-AR has been reported to trigger AD-related biochemical and anatomical alterations (20–23), we examined the effects of β 2-AR blockade on RyR2 post-translational modifications and remodeling. The selective β 2-AR antagonist ICI118-551 (ICI) prevented RyR2 PKA phosphorylation, oxidation, nitrosylation, and depletion of calstabin2 (Fig. 1, A–E) without affecting PP1 and spinophilin levels on RyR2 complex (Fig. 1, A, F, and G).

We further performed several tests to ascertain that RyR2 post-translational remodeling occurs through β 2-AR in our study model. Indeed, SH-SY5Y cells express both β 1- and β 2-AR but not β 3-AR (29). APPswe cells were treated using increasing concentrations of the β 2-AR antagonist ICI (Fig. 2, A and B), the PKA inhibitor (H-89) (Fig. 2, C and D), or the β 1-AR antagonist CGP 20712A (Fig. 2, E and F). We show that complete blockade of RyR2 phosphorylation and calstabin2 dissociation from RyR2 was obtained with ICI at 10 nM, 0.1 μ M, and 1 μ M (Fig. 2, A and B). Accordingly, we also observed a dose-dependent blockade of RyR2 post-translational remodeling in APPswe cells with H-89 with a maximal effect at 1 μ M and 5 μ M (Fig. 2, C and D). Importantly, treatment of APPswe cells with the β 1-AR antagonist CGP 20712A up to 1 μ M did not modify either RyR2 phosphorylation or calstabin2 binding to RyR2 (Fig. 2, E and F). Although H-89 does not behave as a fully

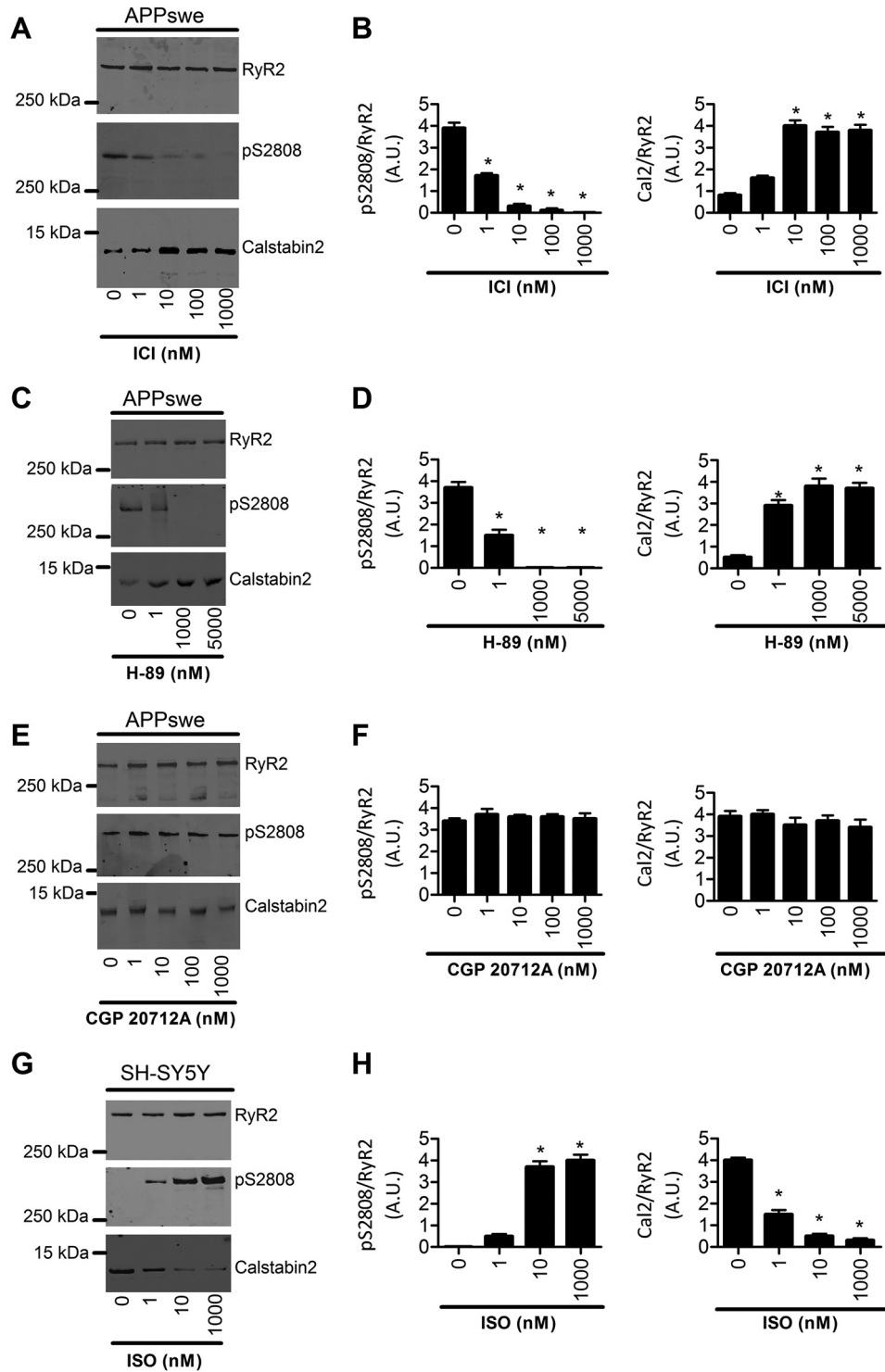


Figure 2. Remodeling of the RyR2 macromolecular complex in the SH-SY5Y neuroblastoma cell line stably expressing APP_{Swe} and in SH-SY5Y control cells. Representative SDS-PAGE analyses and quantification of RyR2 immunoprecipitated from the human SH-SY5Y neuroblastoma cell line stably expressing APP_{Swe} treated with different doses of ICI118–551 (ICI) (1 nM, 10 nM, 100 nM, and 1000 nM, for 12 h) (A), H-89 (1 nM, 1000 nM, and 5000 nM, for 12 h) (C), and CGP 20712A (1 nM, 10 nM, 100 nM, and 1000 nM, for 12 h) (E), and from SH-SY5Y control cells treated with different doses of Isoproterenol (1 nM, 10 nM, and 1000 nM, for 1 h) (G). RyR2 was immunoprecipitated and immunoblotted for RyR2-pS2808 (RyR2 PKA phosphorylation site) and for calstabin2. Graphs B, D, F, and H represent the mean \pm S.E. obtained from three independent experiments. *, $p < 0.05$ calculated versus control using one-way ANOVA and Bonferroni post-test. A.U., arbitrary units.

specific inhibitor of PKA (30), it supports our view that RyR2 post-translational remodeling in APP_{Swe}-expressing cells occurs specifically and in a dose-dependent manner through β 2-AR activation (Fig. 2, A–F).

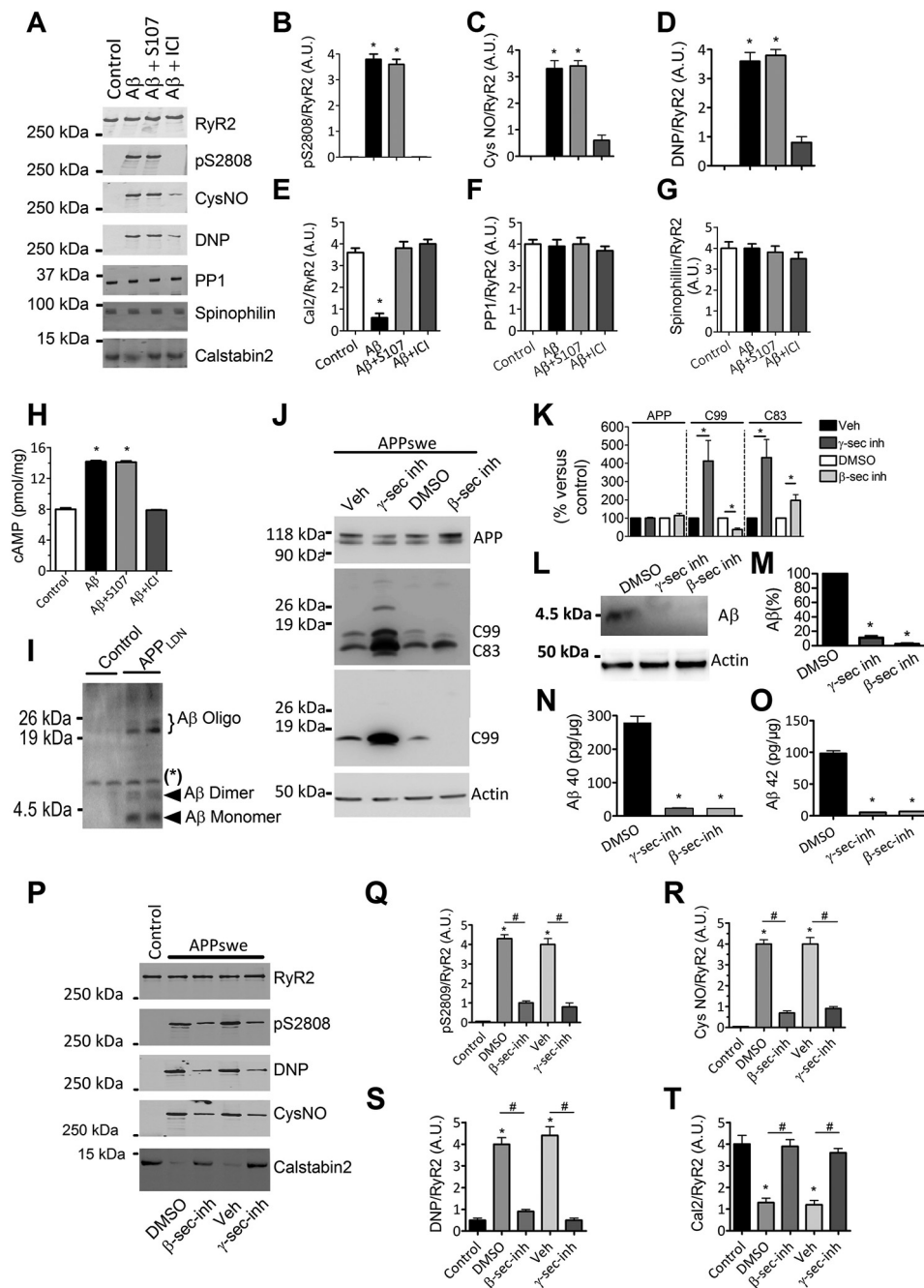
A β causes the biochemical signature of leaky RyR2 channels

We then sought to determine the potential influence of A β or other APP metabolites on the RyR2 macromolecular complex remodeling. We used an A β preparation containing soluble A β

RyR post-translational modifications enhance A β production

oligomers (considered as the most toxic forms of A β) (Fig. 3I and Ref. 31) and found that exposure of control human SH-SY5Y neuroblastoma cells to A β (1–5 nM, 30–60 min) resulted in RyR2 PKA phosphorylation, oxidation, nitrosylation, and depletion of calstabin2 from the RyR2 macromolecular complex (Fig. 3, A–E). However, in contrast to SH-SY5Y-APPsw cells, the amounts of PP1 and spinophilin in the RyR2 complex were unaffected by acute treatment with A β (Fig. 3, A, F, and G). Pretreatment with S107 reduced A β -induced calstabin2 depletion from the RyR2 macromolecular complex (Fig. 3, A and E). We also show that treatment of SH-SY5Y control cells with increasing concentrations of the β -AR agonist isoproterenol induces RyR2 phosphorylation associated with a dissociation of calstabin2 from RyR2 (Fig. 2, G and H). ICI inhibited

A β -induced RyR2 remodeling, thus preventing RyR2 PKA phosphorylation, oxidation, nitrosylation, and calstabin2 depletion (Fig. 3, A–E). Neither S107 nor ICI affected PP1 and spinophilin levels in the RyR2 complex. Acute A β exposure increased cAMP levels (14.2 ± 0.2 versus 8.0 ± 0.3 pmol/mg), which was inhibited by ICI (7.9 ± 0.1 pmol/mg) (Fig. 3H), suggesting that A β acts upstream of β 2-AR signaling. These data and those of Fig. 1 demonstrate that RyR2 remodeling is likely linked to A β (Figs. 1 and 3, A and H) and to β 2-AR activation (Fig. 2, A–H). Because our A β preparation contains monomeric and low molecular weight oligomeric forms of A β , we cannot attribute the observed effects to a specific A β species. We could detect only a moderate elevation of cAMP levels in APPsw model ($\sim 20\%$ increase over control cells considered as 100%, $n = 3$;



data not shown), but data were obtained at only one time point. Thus, other experiments will be necessary to unravel time-dependent modulation of cAMP production in the APPswe model.

We also used a pharmacological approach to modulate APP metabolism by using β -secretase (LY288672, 30 μ M) or γ -secretase (ELND006, 5 μ M) inhibitors and investigate the potential implication of other β APP metabolites in RyR2 remodeling.

Although inhibition of β -secretase reduces C99 fragment production and enhances the level of C83 fragment, γ -secretase inhibition enhances the levels of both C99 and C83 fragments (Fig. 3, *J* and *K*). As expected, β - and γ -secretase inhibitors fully block intracellular and extracellular A β production (Fig. 3, *L–O*). Interestingly, blocking β -secretase or γ -secretase reduces RyR2 PKA phosphorylation, oxidation, nitrosylation, and calstabin2 depletion in APPswe-expressing cells (Fig. 3, *P–T*).

A β -mediated remodeling of RyR2 channel increases cytosolic Ca²⁺ and mitochondrial ROS in SH-SY5Y neuroblastoma cells

We previously reported that SH-SY5Y cells expressing APPswe display increased cytosolic Ca²⁺ due to increased Ca²⁺ release from the ER via IP₃R and RyR and enhanced Ca²⁺ entry via voltage-dependent and voltage-independent plasma membrane Ca²⁺ channels (13). We now report that SH-SY5Y cells expressing APPswe exhibit increased basal cytosolic Ca²⁺ signals compared with control cells (Fig. 4, *A* and *B*). Accordingly, exogenous A β treatment of SH-SY5Y cells for 12 h (an experimental condition mimicking APPswe cells chronically overproducing A β) also results in elevated basal [Ca²⁺]_{cyt} (Fig. 4, *C* and *D*). As previously reported (35), acute application of A β to human neuroblastoma SH-SY5Y cells induced a slow rise of [Ca²⁺]_{cyt} (Fig. 4, *E* and *F*). Interestingly, A β -induced Ca²⁺ elevation was prevented by S107, ICI, or ryanodine pretreatment ($\Delta F/F_0 = 1.95 \pm 0.16$ ($n = 57$), 0.63 ± 0.08 ($n = 31$), 0.28 ± 0.08 ($n = 44$), 0.63 ± 0.09 ($n = 23$) in A β , A β +S107-, A β +ICI-, and A β +ryanodine-treated cells, respectively) (Fig. 4, *E* and *F*). Our

finding is in accordance with data showing that preincubation with inhibitory ryanodine virtually eliminated the generation of Ca²⁺ signals elicited by oligomeric A β in primary hippocampal neurons (36).

S107, ICI, and ryanodine did not completely reverse the rise in [Ca²⁺]_{cyt}. This may indicate that, in addition to leaky RyR2, there are other components that contribute to the A β -induced rise in [Ca²⁺]_{cyt}. Possibilities include: 1) intracellular Ca²⁺ release via the IP₃Rs and 2) Ca²⁺ entry through the plasma membrane as previously reported (13). Accordingly, A β -mediated Ca²⁺ elevation is lower in nominally Ca²⁺-free extracellular medium, and this elevation is completely reversed by ryanodine (data not shown).

We also analyzed basal [Ca²⁺]_{cyt} in APPswe cells untreated or treated with S107 or ICI (Fig. 4, *A* and *B*). We show that S107 or blocking β 2-adrenergic signaling by ICI significantly reduce basal [Ca²⁺]_{cyt} in APPswe-expressing cells (Fig. 4, *A* and *B*).

It is known that IP₃Rs are activated by Ins(1,4,5)P₃, a metabolic product of GPCR activity, and that the activation of these channels is amplified by Ca²⁺-induced Ca²⁺ release, a regenerative mechanism by which Ca²⁺ enhances its own release from IP₃Rs and RyRs (37). Accordingly, it was already reported that exacerbated IP₃R-evoked Ca²⁺ signals in the PS1_{M146V}- and the 3 \times Tg-AD-derived neurons occur through increased Ca²⁺-induced Ca²⁺ release through the RyR (15). We show herein that increased inositol 1,4,5-trisphosphate-mediated cytosolic Ca²⁺ signal (carbachol stimulation) in APPswe (Fig. 4, *G–J*) contributed to RyR-mediated Ca²⁺ release/leak. Thus, stabilization of calstabin2 on RyR2 channels by S107 or blockade of RyR2 phosphorylation by ICI (Fig. 4, *G0–J*), both, lowers carbachol-mediated cytosolic Ca²⁺ signals as revealed by reduced peak values and the area under the curve reflecting integrated Ca²⁺ response (Fig. 4, *G–J*). Similar results were obtained using cytosolic aequorin probe providing “calibrated” measurements of [Ca²⁺]_{cyt} (38) (supplemental Fig. S1). All over, these data demonstrate that cytosolic Ca²⁺ increase in APPswe cells and in

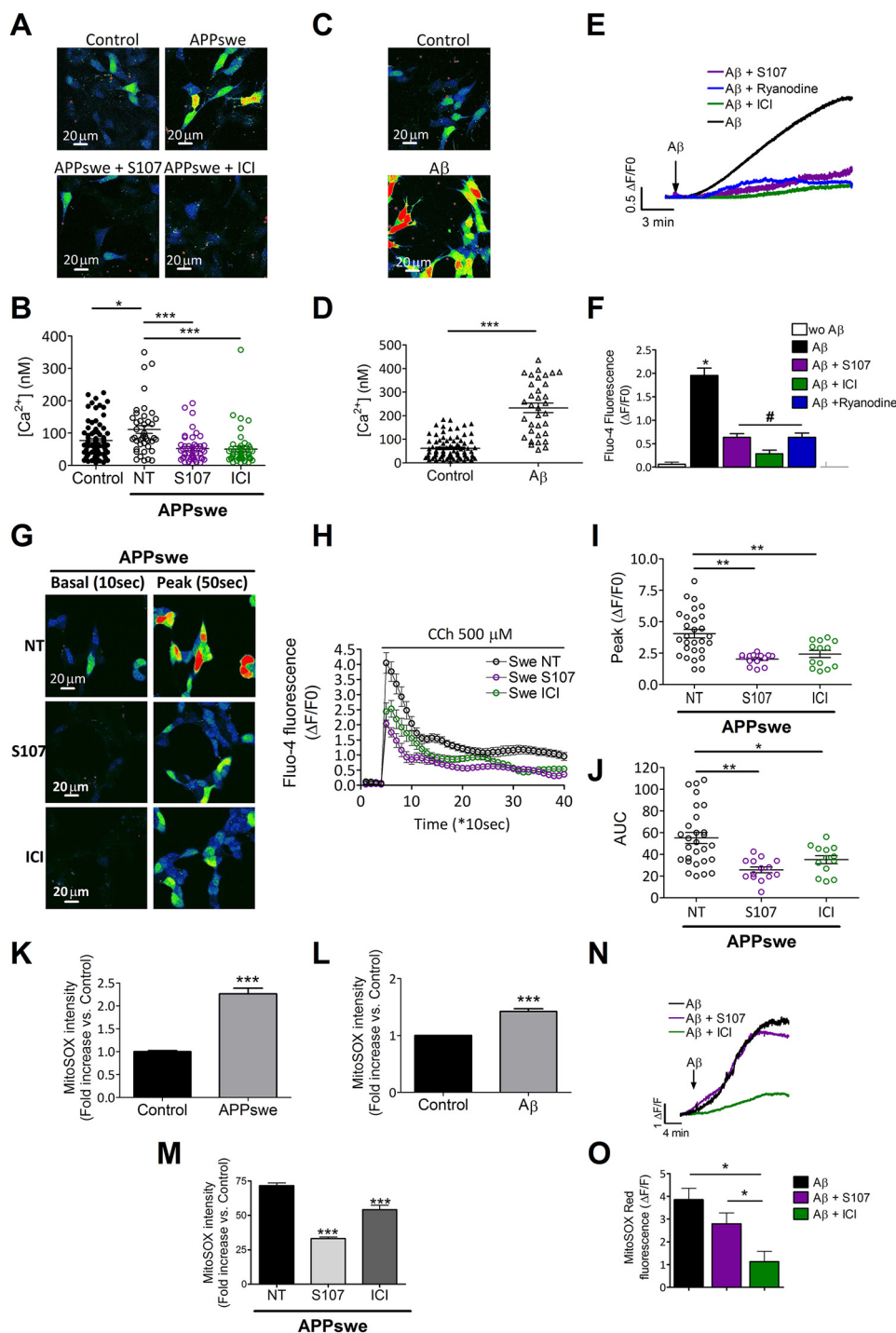
Figure 3. A β caused the biochemical signature of leaky RyR2 channels. *A*, representative SDS-PAGE analyses and quantification of RyR2 immunoprecipitated from the wild-type human SH-SY5Y neuroblastoma cell line treated with oligomeric A β (1–5 nM, 30–60 min) alone or in combination with either S107 (10 μ M, for 12 h) or ICI118–551 (ICI) (1 μ M, for 12 h). RyR2 was immunoprecipitated and immunoblotted for RyR2-pS2808 (RyR2 PKA phosphorylation site), S-nitrosylation (CysNO), and oxidation (DNP) as well as components of the RyR2 channel complex as in Fig. 1. *B–G*, graphs represent the mean \pm S.E. obtained from three independent experiments. *, $p < 0.05$ calculated versus SH-SY5Y untreated cells used as control (Control) using one-way ANOVA and Bonferroni post-test. *H*, A β treatment (1–5 nM) for 12 h caused a rise in intracellular [cAMP] (pmol/mg of total proteins) in the SH-SY5Y neuroblastoma cell line that was not reduced by S107 (10 μ M, for 12 h) and reduced by ICI (1 μ M, for 12 h) ($n = 5$ for each condition). Data are the mean \pm S.E. *, $p < 0.05$ calculated versus Control using one-way ANOVA and Bonferroni post-test. *I*, a representative Tris-Tricine gel showing oligomeric A β preparations obtained from conditioned media of CHO cells stably transfected with hAPP695 cDNA harboring London mutation (APP_{LDN}: APPV642I). Conditioned media of CHO cells stably transfected with pcDNA4 empty vector were used as control. *, nonspecific band. *J*, representative SDS-PAGE and Tris-Tricine gels showing the modulation of β APP metabolism in SH-SY5Y cells expressing APPswe treated for 12 h with β -secretase inhibitor (*inh*; LY288672, 30 μ M, for 12 h; Ref. 32) or γ -secretase inhibitor (ELND006, 5 μ M, for 12 h; Refs. 33 and 34) or with DMSO (control for β -secretase inhibitor) or vehicle (control for γ -secretase inhibitor) and revealed on total extracts. β APP was detected using the APP N-terminal antibody. β -CTF (C99) was detected using the 6E10 antibody. α - and β -CTFs (C83 and C99, respectively) were detected using the APP C-terminal antibody. Actin was used as loading control. *K*, graph represents the mean \pm S.E. of APP, C99, and C83 obtained from 5–8 independent experiments as shown in *J*. *, $p < 0.05$ calculated versus controls (DMSO or vehicle) using one-way ANOVA and Tukey's multiple comparisons test. *L*, representative Tris-Tricine gel showing total intracellular A β in the SH-SY5Y neuroblastoma cell line stably expressing APPswe, control (DMSO), or treated for 12 h with γ - or β -secretase inhibitors as in *J*. Actin was used as loading control. *M*, the graph represents the mean \pm S.E. obtained from 3 independent experiments. *, $p < 0.01$ calculated versus DMSO treated SH-SY5Y APPswe cells using one-way ANOVA and Tukey's multiple comparisons test. *N* and *O*, ELISA of A β 40 (*N*) and of A β 42 (*O*) done on cell culture media (50 μ l) of human SH-SY5Y neuroblastoma cell line stably expressing APPswe treated as described in *J*. DMSO-treated cells were used as control. A β was quantified using A β 40 or A β 42 standard curves. The graph represents the mean \pm S.E. from three experiments and is expressed in pg/mg protein. *, $p < 0.01$ calculated versus DMSO treated SH-SY5Y APPswe cells using one-way ANOVA and Tukey's multiple comparisons test. *P*, representative SDS-PAGE analyses and quantification of RyR2 immunoprecipitated from the human SH-SY5Y neuroblastoma cell line stably expressing mock vector used as control (Control) or APPswe treated as described in *J*. RyR2 was immunoprecipitated and immunoblotted for RyR2-pS2808 (RyR2 PKA phosphorylation site), S-nitrosylation (CysNO), and oxidation (DNP) as well as calstabin2. *Q–T*, the graphs represent the mean \pm S.E. obtained from three independent experiments. *, $p < 0.05$ calculated versus control; #, $p < 0.05$ calculated versus DMSO or vehicle using one-way ANOVA and Bonferroni post-test. *A.U.*, arbitrary units.

RyR post-translational modifications enhance A β production

A β -treated cells is largely contributed by RyR-mediated Ca²⁺ release.

Next we found that both endogenous production (*i.e.* APPsw cells) and exogenous application of A β (*i.e.* treatment of wild-type SH-SY5Y cells with A β for 12 h) resulted in increased mitochondrial reactive oxygen species (ROS) production (Fig. 4, *K* and *L*). Interestingly, mitochondrial ROS production stimulated by chronic A β production in SH-SY5Y-APPsw cells was partially inhibited by both ICI and S107 (Fig. 4*M*). We also show that acute application of A β enhanced MitoSOX fluorescence until reaching a steady state plateau

(Fig. 4*N*). *Bar graphs* show the mean plateau value of $\Delta F/F \pm$ S.E. obtained in each experimental condition and reveal a significant decrease of MitoSOX fluorescence intensity in A β +ICI cells *versus* A β alone and A β +S107 (Fig. 4*O*). S107 pretreatment reduced A β -mediated ROS production (plateau phase of representative data in Fig. 4*O*), but this reduction was not statistically significant. These data reveal that A β may contribute to mitochondrial ROS elevation that is significantly reversed by ICI and to a lesser extent with S107 (Fig. 4, *N* and *O*). These data may indicate that mitochondrial ROS elevation in APPsw likely results from both β -adrenergic and RyR-me-



diated Ca²⁺ leak. In A β -treated cells, mitochondrial ROS elevation likely occurs in a β -adrenergic-dependent manner and is only partially due to RyR-mediated ER Ca²⁺ leak.

Pharmacological inhibition of RyR2 leak reduces β APP metabolism and the production of A β in APPswe-expressing SH-SY5Y cells

Next, we determined whether S107-mediated inhibition of RyR2 leak could affect β APP processing and A β production in SH-SY5Y cells expressing APPswe. Interestingly, the levels of C83 and C99 were reduced by S107 treatment (1–5 μ M) (Fig. 5A, D, E, F, H, and I). As the control, we used dantrolene to block RyR-mediated Ca²⁺ release (Fig. 5, B, D, and E). C99 undergoes cleavage by γ -secretase to yield A β 40 and A β 42 peptides and AICD (39). AICD is also produced from C83 fragment upon γ -secretase cleavage (39). S107 treatment reduced the level of AICD (Fig. 5, F and K) and of total intracellular A β (Fig. 5, F and J). Interestingly, the total amount of secreted A β 42, but not of A β 40, was also significantly reduced by S107 treatment in SH-SY5Y cells expressing APPswe (Fig. 5L). Thus, S107 treatment reduces γ -secretase-mediated AICD production in an APPswe-expressing cells membrane preparation, similar to γ -secretase inhibition (Fig. 6, A and B).

We also evaluated whether S107 affects β APP trafficking (localization). This was performed using immunofluorescence and confocal imaging with antibodies recognizing specific cellular compartments involved in β APP trafficking. The cellular localization of β APP was assessed using Pearson's correlation and Mander's Overlap coefficients (Fig. 6, H–I). We show that S107 did not alter β APP localization in early endosome (EEA1) (Fig. 6, C, H, and I), ER (SERCA2b) (Fig. 6, D, H, and I), recycling endosomes (CD71) (Fig. 6, E, H, and I), late endosomal and lysosomal membranes (CD63) (Fig. 6, F, H, and I), and the trans-Golgi network (TGN46) (Fig. 6, G, H, and I).

Blockade of β 2- but not β 1-adrenergic signaling reduces β APP metabolism in APPswe-expressing SH-SY5Y cells

As with S107 treatment, blockade of β 2-AR with ICI reduced the production of the C99 fragment in APPswe-expressing SH-SY5Y cells (Fig. 7, A and B), whereas the blockade of β 1-AR

with CGP did not (Fig. 7, C and D). The effect of ICI and S107 on C99 production was not additive (Fig. 7, E and F). In accordance with ICI data, inhibition of PKA by H-89 reduced [Ca²⁺]_{cyt} (supplemental Fig. S1), and C99 production (Fig. 7, G and H). Thus, RyR2 channel remodeling occurring downstream of β 2-adrenergic signaling contributes to regulation of β APP processing.

Discussion

We show herein that RyR post-translational remodeling accounts for exacerbated RyR2-mediated Ca²⁺ release in an *in vitro* AD study model. Enhanced neuronal RyR2-mediated ER Ca²⁺ leak is linked to pathophysiological post-translational modifications in the macromolecular RyR complex (17). Interestingly, post-translational modifications of RyR2 were reported in cerebral ischemia (18), where endogenous RyRs undergo S-nitrosylation and S-gluthathionylation processes that resulted in high activity channels and ultimately lead to cortical neuronal death (18). Recently, Liu *et al.* (19) described the contribution of RyR2 post-translational remodeling to stress-related memory impairments.

We report that RyR2 undergoes PKA phosphorylation, oxidation, and nitrosylation in SH-SY5Y-overexpressing APPswe. Remodeled RyR2 macromolecular complex is depleted of calstabin2 and of both PP1 and its anchoring protein spinophilin. The equilibrium of phosphorylation and dephosphorylation of the channel is generally regarded as an important regulatory mechanism of RyR. Importantly, the amount of spinophilin (a postsynaptic marker) is decreased in the AD brain (40, 41). We report a depletion of PP1 and spinophilin in SH-SY5Y-overexpressing APPswe mutation producing chronically A β but not upon acute treatment of wild type SH-SY5Y with oligomeric A β . This may suggest that spinophilin depletion contributes to PKA phosphorylation of RyR2 by reducing the targeting of the phosphatase PP1 to the channel likely at advanced AD stages. We focused in this study on RyR2 and its associated regulatory protein calstabin2. We cannot exclude the possibility that RyR1 and RyR3 may also play a role in AD pathology.

Post-translational modifications of RyRs in AD patients have not been reported yet. However, AD brains manifest excessive

Figure 4. A β -mediated treatment elevates basal cytosolic [Ca²⁺]_{cyt} and mitochondrial ROS in human neuroblastoma cells. A, representative pseudo-colored images (projection of Z-stacks) of basal [Ca²⁺]_{cyt} in human SH-SY5Y neuroblastoma cell line stably expressing APPswe or mock vector used as control measured using Fluo-4, AM probe. The scale bar represents 20 μ m. B, the scatter plot represents calibrated [Ca²⁺]_{cyt} (nM) obtained in control cells (n = 84) and in APPswe-expressing cells untreated (n = 42) or treated with S107 (1 μ M, for 12 h) (n = 46) or with ICI (10 μ M, for 12 h) (n = 47). *, p < 0.05; ***, p < 0.001, calculated using one-way ANOVA and Tukey's multiple comparisons test. C, representative pseudo-colored images (projection of Z-stacks) of basal [Ca²⁺]_{cyt} measured using Fluo-4, AM in the human SH-SY5Y neuroblastoma cell line treated with A β (1–5 nM) for 12 h. The scale bar represents 20 μ m. D, the scatter plot represents calibrated [Ca²⁺]_{cyt} (nM) obtained in the control (n = 83) or A β -treated cells (n = 34). ***, p < 0.001 calculated versus control using the t test. E, acute A β application caused a rise in [Ca²⁺]_{cyt} in the SH-SY5Y neuroblastoma cell line (WT) (n = 42) that was partially inhibited by either S107 (10 μ M) (n = 31), ICI (1 μ M) (n = 44), or ryanodine (10 μ M) (n = 23) pretreatment. F, the graph shows the mean \pm S.E. of $\Delta F/F_0$ of Fluo-4 fluorescence in the plateau phase. *, p < 0.001 calculated versus control (wo A β); #, p < 0.001 calculated versus A β -treated cells using one-way ANOVA and Tukey's multiple comparisons test. G, representative pseudo-colored images (focal plane) of the Fluo-4 signal before (10 s, basal) and after carbachol stimulation (500 μ M) (50 s, peak) measured using Fluo-4 AM probe in SH-SY5Y neuroblastoma cell line stably expressing APPswe non-treated cells (NT) (n = 28) or treated for 12 h with either S107 (1 μ M) (n = 14) or ICI (10 μ M) (n = 13). The color scale is shown where the black/blue and the yellow/red represent low and high Fluo-4 fluorescence respectively. H, normalized curves showing carbachol (CCh)-mediated Ca²⁺ responses, presented as the mean $\Delta F/F_0 \pm$ S.E. I, scatter plots represent the mean of peak values presented as mean $\Delta F/F_0 \pm$ S.E. for non-treated (NT) SH-SY5Y APPswe (n = 28) or cells treated for 12 h with S107 (1 μ M, n = 14) or ICI (10 μ M, n = 13). J, scatter plots represent the mean area under the curve (AUC) \pm S.E. units of the normalized Ca²⁺ response for cells as in J. I and J: *, p < 0.05; **, p < 0.01 analyzed by one-way ANOVA and Dunnett's post-test versus APPswe NT. K–M, the graphs represent MitoSOX red dye-fold change in control (n = 228) and APPswe cells (n = 144) (K) and in control (n = 6) and A β -treated (n = 6) cells (N), and in SH-SY5Y APPswe non-treated (n = 293) and treated with S107 (n = 382) or ICI (n = 136) (M). ***, p < 0.001 analyzed by Student's t test (K and L) and one-way ANOVA and Dunnett's post-test versus APPswe NT (M). N and O, acute A β treatment caused a rise in mitochondrial ROS production, as detected by MitoSOX red dye, in the SH-SY5Y neuroblastoma cell line (n = 293 for A β treated cells) that was not inhibited by S107 (n = 382) and reduced by ICI (n = 136). The graph shows the mean \pm S.E. The graph shows $\Delta F/F_0$ of MitoSOX fluorescence in the plateau phase normalized to the basal level before A β stimulation. *, p < 0.05 using one-way ANOVA and Dunnett's post-test versus A β .

RyR post-translational modifications enhance A β production

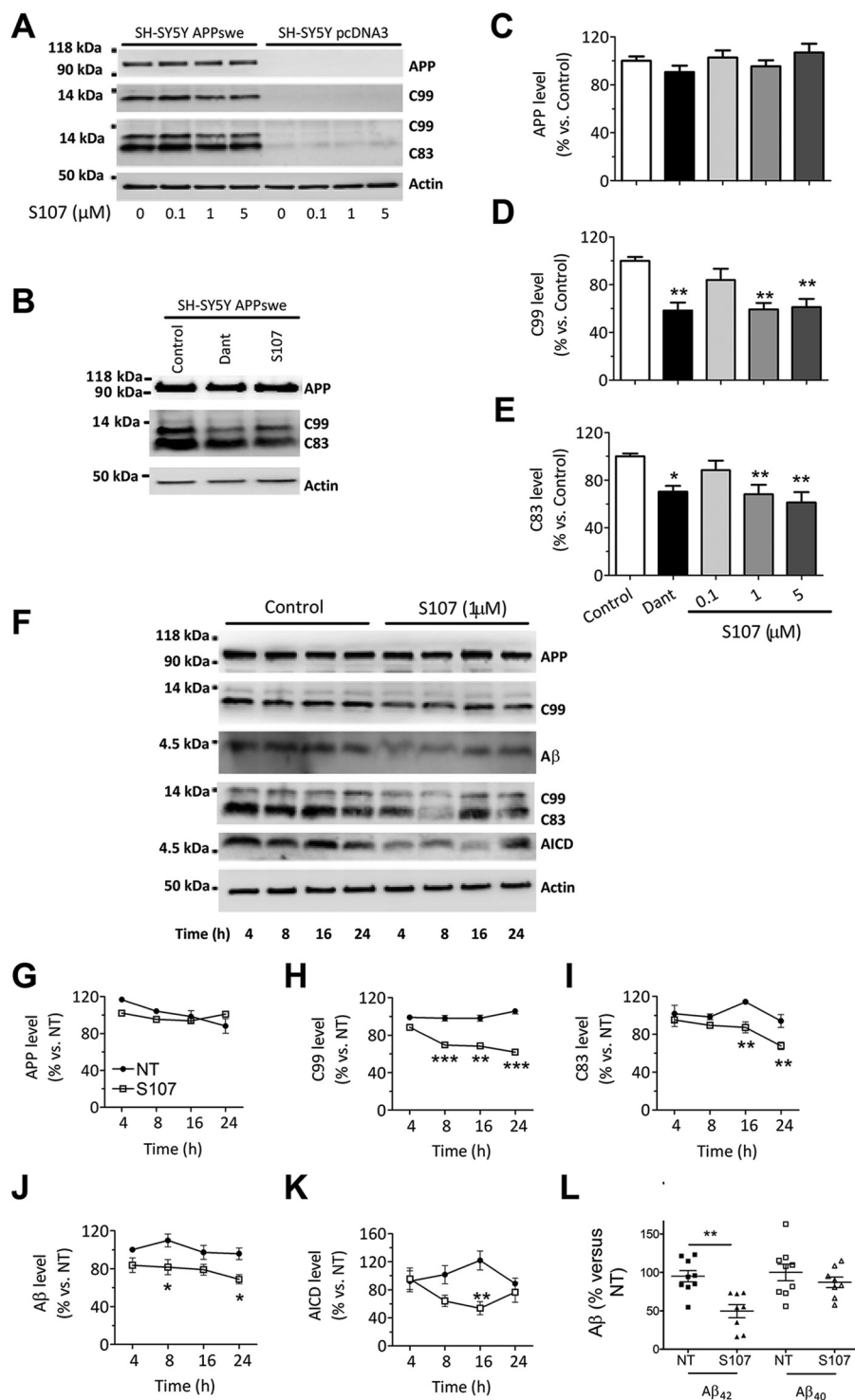


Figure 5. Pharmacologic inhibition of RyR2 reduced β APP metabolism and the production of CTFs and of A β in SH-SY5Y APPsw model. *A*, representative Tris-Tricine and SDS-PAGE analyses of full-length β APP, C83, and C99 levels (revealed as described in Fig. 2J) in SH-SY5Y cells expressing pcDNA3.1 or APPsw non-treated (0) or treated with S107 (0.1, 1, or 5 μ M for 12 h). *B*, representative Tris-Tricine and SDS-PAGE analyses of full-length β APP and its CTF (C99 and C83) in SH-SY5Y cells expressing APPsw non-treated (Control) or treated with dantrolene (50 μ M) or S107 (1 μ M) for 12 h. *C–E*, graphs show the mean \pm S.E. obtained from control ($n = 11$), S107 (0.1, 1, or 5 μ M, $n = 5, 3$, and 9, respectively), and dantrolene ($n = 5$). *, $p < 0.05$; **, $p < 0.01$, versus non-treated cells analyzed by one-way ANOVA and Dunnett's post-test versus APPsw control cells. *F*, representative Tris-Tricine and SDS-PAGE analyses of full-length β APP, C83 and C99, A β , and AICD levels in SH-SY5Y cells expressing APPsw non-treated (Control) or treated with S107 (1 μ M) for the indicated times (4, 8, 16, and 24 h). β APP was detected using APP N-terminal antibody. C99 and A β were detected using 6E10 antibody. C83 and C99 and AICD were detected using APP C-terminal antibody. *A* and *E*, actin was used as loading control. *G–K*, graphs show the mean \pm S.E. obtained from seven different experiments. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$ versus controls at the same time of analysis using two-way ANOVA and the Bonferroni post-test. *L*, ELISA analyses of extracellular A β ₄₀ and A β ₄₂ in APPsw cells treated with S107 (1 μ M, for 4 h) and presented as % versus APPsw control cells. **, $p < 0.01$ using *t* test, from 8 experiments. NT, non-treated cells.

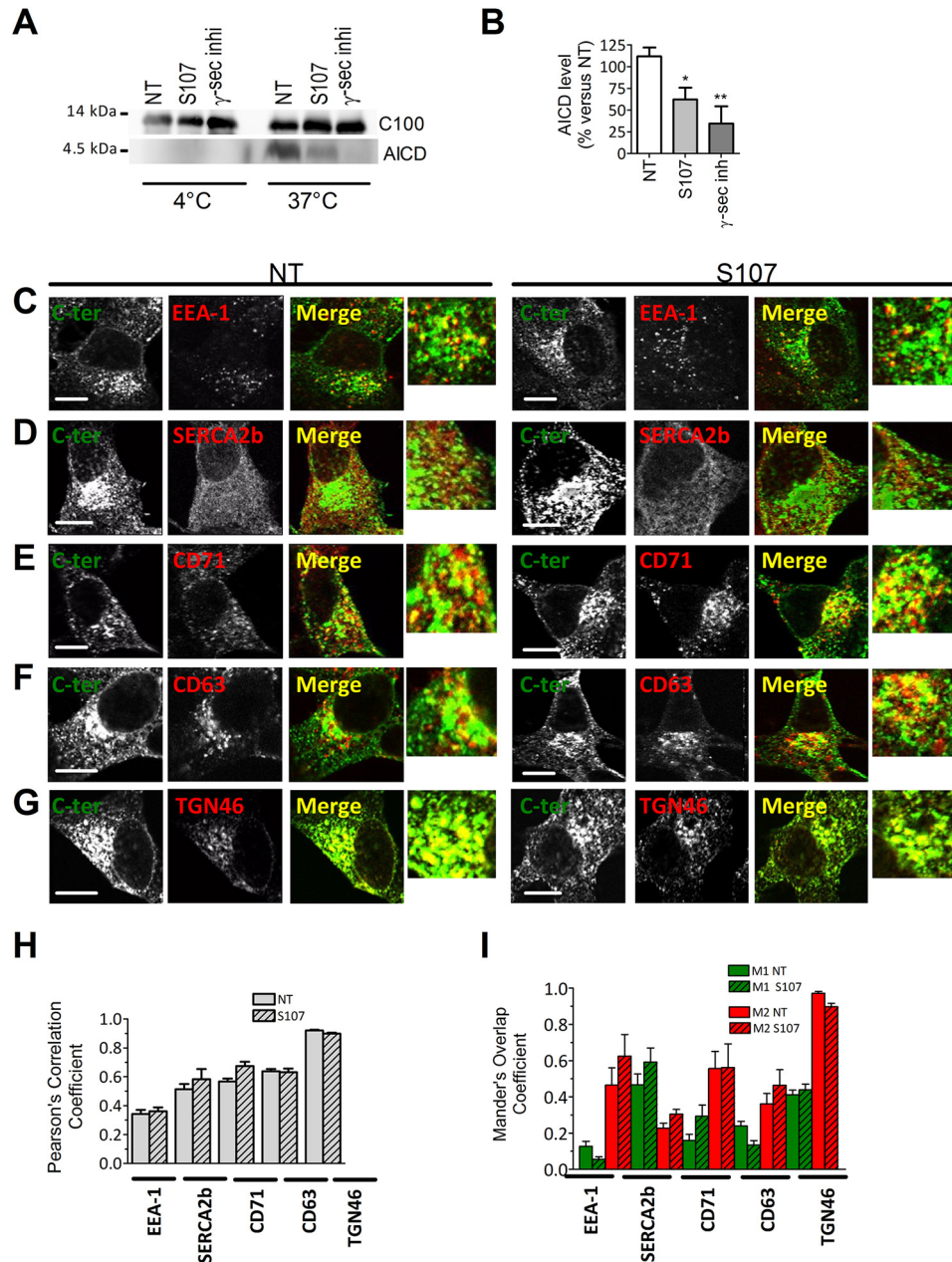


Figure 6. Pharmacologic inhibition of RyR2 leak reduced γ -secretase activity and does not alter β APP intracellular localization. *A*, cell-free AICD production from recombinant C100 peptide performed at 37°C or 4°C in the presence of membrane fractions isolated from APPsw cells treated with S107 (1 μ M, for 12 h) or with γ -secretase inhibitor (*inhi*; 5 μ M, for 12 h). C100 and AICD were detected using APP C-terminal antibody. *NT*, non-treated cells. *B*, graph shows the mean \pm S.E. obtained from seven different experiments. *, $p < 0.05$; **, $p < 0.01$ versus APPsw non-treated cells using one-way ANOVA and Dunnett's post-test versus APPsw NT. *C–G*, double-labeling immunofluorescence of SH-SY5Y cells expressing APPsw non-treated or treated with S107 (1 μ M) for 12 h using antibodies against C-terminal (C-ter) APP antibody (green signal) and EEA1 (*C*), Ca²⁺-ATPase of the endoplasmic reticulum (*SERCA2b*) (*D*), the recycling endosome marker transferrin receptor (*CD71*) (*E*), lysosome membrane and late endosome membrane (*CD63*) (*F*), and the trans-Golgi network marker (*TGN46*) (*G*). Green and red signals are depicted in white to have better contrast visualization. Merge and magnified overlay images show green and red signals co-localization (depicted as yellow signals). The scale bar represents 10 μ m. *H* and *I*, the graphs show Pearson's correlation coefficient (*H*) and Mander's Overlap Coefficient (*I*). *M1* corresponds to the green signal overlapping with the red signal. *M2* corresponds to the red signal overlapping with the green signal. Analyses were performed in at least four different fields in each condition obtained from three independent experiments. *H* and *I*, differences are statistically non-significant.

generation of reactive nitrogen and ROS, contributing to neuronal cell injury and death via a series of redox reactions (40, 42–44). cAMP levels are also significantly elevated in CSF from patients with Alzheimer dementia (45). In addition, PKA activation has been shown to contribute to the regulation of β APP processing, to A β -mediated cell death *in vitro* and *in vivo*, and to oxidative stress (46–50). Interestingly, A β stimulates PKA

activity and PKA-dependent signaling pathways by activating β -ARs (51–53). Blocking β 2-adrenergic signaling diminishes A β production and delays functional decline in AD (47, 48, 54). Accordingly, activation of β 2-ARs enhances γ -secretase activity and accelerates A β plaques formation (20, 22), a process that can be blocked by the specific β 2-AR antagonist (22). Moreover, it was reported that A β 42 peptide binds to the N terminus

RyR post-translational modifications enhance A β production

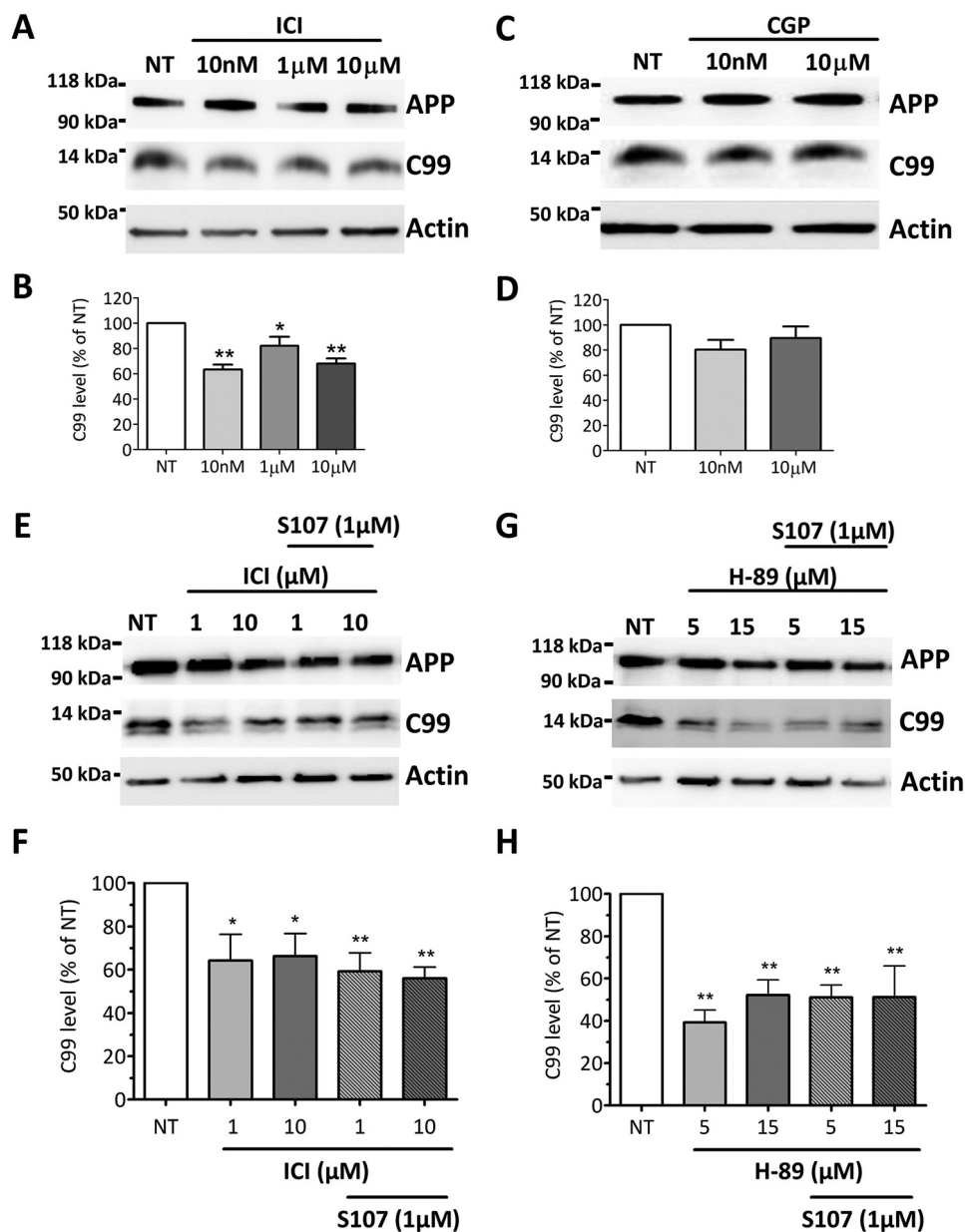


Figure 7. Blockade of β 2- but not of β 1-adrenergic, cascade reduced β APP metabolism in SH-SY5Y APPsw cells. A and C, representative Tris-Tricine and SDS-PAGE analyses of full-length β APP and C99 levels (determined as described in Fig. 5A) in SH-SY5Y cells expressing APPsw non-treated (NT) or treated with ICI (10 nM, 1 μ M, or 10 μ M) (A) or with CGP 20712A (10 nM or 10 μ M) (C) for 12 h. B and D, graphs show the mean \pm S.E. for ICI (B) and CGP (D) treatments, respectively, obtained from six different experiments. *, $p < 0.05$; **, $p < 0.01$ analyzed by one-way ANOVA and Dunnett's post-test versus APPsw NT. E and G, representative Tris-Tricine and SDS-PAGE analyses of full-length β APP and C99 levels in SH-SY5Y cells expressing APPsw non-treated (NT), treated for 12 h with ICI (1 μ M or 10 μ M) (E) or H-89 (5 μ M or 15 μ M) (G) alone or in combination with S107 1 μ M for 12 h. F and H, graphs show the mean \pm S.E. obtained from five different experiments. *, $p < 0.05$; **, $p < 0.01$, analyzed using one-way ANOVA and Dunnett's post-test versus APPsw NT.

of β 2-ARs, enhancing PKA-dependent AMPA receptor hyperactivity (55). Consistent with these observations, we show that acute application of A β to SH-SY5Y neuroblastoma cells increases cAMP levels and oxidative stress contributing to RyR2 remodeling that can be improved by treatment with β 2-AR antagonist ICI. On the other hand, we show that both β 2-AR antagonist ICI and PKA inhibitor H-89 reduce ER Ca²⁺ release and β APP processing. This supports a positive feedback loop where A β activates β 2-AR signaling cascade leading to RyR2 remodeling and ER Ca²⁺ leak that, in turn, amplifies β APP processing and A β production (Fig. 8). We show that the increase of A β levels is linked to β 2-AR signaling cascade. How-

ever, we cannot exclude the possibility that other G_s-coupled receptors in PKA/cAMP pathway may also play a role in AD. It has previously been demonstrated that isoproterenol, a β -adrenergic agonist, increases mitochondrial ROS production in cardiomyocytes in a concentration- and cAMP-PKA-dependent manner (56). Mitochondrial ROS production could contribute to RyR2 oxidation. RyR2 leak may in turn increase mitochondrial ROS production (57). This seems to be the scenario in SH-SY5Y-expressing APPsw cells, as both Ca²⁺ and ROS elevation was reversed by S107 and ICI. The situation in acute A β -treated cells seems different as mitochondrial ROS elevation was reversed by the β 2-AR antagonist and was only slightly

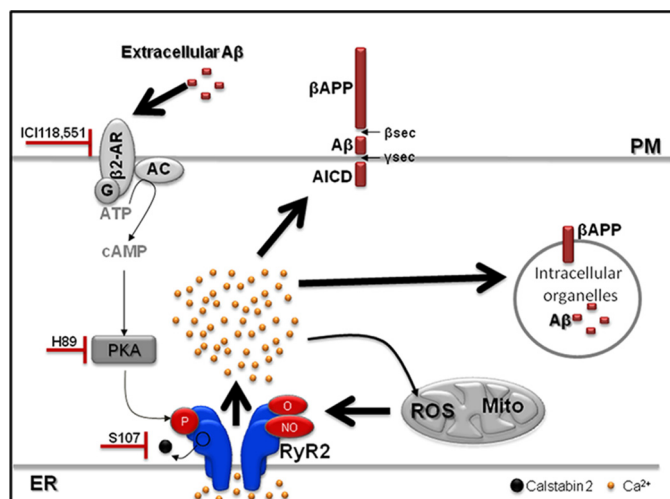


Figure 8. Scheme showing the cascade linking A β , β 2-adrenergic signaling, and leaky RyR2 channels in SH-SY5Y cells expressing APP_{sw}. β APP processing occurs within intracellular organelles and in the plasma membrane (PM), thus producing A β inside intracellular organelles and in the extracellular media. A β -mediated β 2-AR activation leads to RyR2 phosphorylation and calstabin2 dissociation, thereby enhancing RyR2-Ca²⁺ leak. RyR2-Ca²⁺ leak promotes Ca²⁺ entry into mitochondria (97) and mitochondrial ROS production. Mitochondrial ROS production leads to RyR2 oxidation and nitrosylation. Post-translational RyR2 remodeling (phosphorylation, oxidation, and nitrosylation) and calstabin2 dissociation enhances RyR2-Ca²⁺ leak. Blocking β 2-adrenergic signaling or RyR2 channel leak reduces β APP processing and A β production. AC, adenylate cyclase.

contributed by leaky RyR channels (positive feedback loop in Fig. 8).

A growing body of literature supports the clinical relevance of the locus coeruleus (LC), norepinephrine, and noradrenergic receptors in AD as the LC widespread projections terminate in areas important for learning and memory such as the hippocampus and the cortex. Neuronal cell death in the LC and other brain stem nuclei is a well defined characteristic of AD pathology (58). It has been proposed that the downstream consequences of LC degeneration are decreased levels of norepinephrine in terminal regions (59) and a compensatory up-regulation of adrenergic receptors (60). A synthesis of evidence suggests that LC destruction may contribute to reduced A β clearance (58, 61, 62). These studies support our data showing that activation of the β -AR cascade occurs in a cell model of AD and that this contributes to enhanced A β production at least in part through PKA-mediated RyR remodeling and Ca²⁺ leak (Fig. 8).

Targeting of β 2-adrenergic signaling to reverse/prevent AD pathology remains a controversial therapeutic track. Genetic knockdown or pharmacologic blockade of β 2-AR elicits beneficial effects on tauopathies (63) as well as in Tg2576 transgenic mice (64). Conversely, other studies have reported that administration of a β 2-AR antagonist exacerbates neuropathology and cognitive deficits in a mouse model of AD (65) and that pharmacological stimulation of β 2-AR improves cognitive function and restores synaptic density in a mouse model of Down syndrome (66). Recently it has also been reported that an enriched environment activates β 2-ARs and protects against A β -induced reduction in LTP and prevents hippocampal dysfunction by A β oligomers (67). PKA activation under adrenergic stimulation is important for learning and memory consoli-

ation. It appears, however, that A β -mediated activation of this cascade may be detrimental.

Increased [Ca²⁺]_{cyt} was observed by different laboratories in various AD models (68–72). Importantly, elevated basal Ca²⁺ has also been reported in peripheral blood mononuclear cells of AD patients (73). Our results contradict a previous study showing that transient expression of FAD APP mutations does not directly perturb intracellular Ca²⁺ homeostasis (74). The level and nature of β APP metabolites (*i.e.* in our model stably expressing β APP and the former study transiently expressing FAD APP mutants; Ref. 74) could explain such discrepancies. Altered RyR-mediated ER Ca²⁺ increase has also been observed in non-APP models such as PS1 and PS2 mutant mice and cell models (15, 75–78). However, post-translational remodeling of RyRs has not been reported in PS1/2 models. Nevertheless, co-immunoprecipitation has revealed a physical interaction between PS1–2 and RyR2 (79–81). Other recent studies demonstrated that the interaction of RyR with PS1 and PS2 N-terminal fragments strongly increased both mean currents and open probability of single RyR channels in a Ca²⁺-dependent manner (82, 83). Other groups demonstrated the molecular interaction of PS2 and sorcin (a modulator of RyR channel) (84).

These studies provide evidence that PS may also play an important role in the regulation of RyR channel activity. However, the question remains whether this regulation occurs only under physiological conditions or may also occur under pathological conditions, namely in cells expressing the mutated forms of PS1–2. Also unclear is whether β -AR-mediated RyR post-translational remodeling occurs in non-APP models, such as PS1 and PS2 mutant mice and cell models.

It has been reported that Ca²⁺ plays a role in the production of A β peptides (for review, see Ref. 9). Our data show that inhibiting RyR-mediated Ca²⁺ leak either with S107 or upstream by blocking β 2-adrenergic signaling pathway reduces β APP metabolism and A β production. In support of these findings the RyR active drug dantrolene has been shown to reduce the extent of β APP phosphorylation likely through Cdk5 and/or GSK3 β Ca²⁺-dependent activation (13, 85). RyR-mediated Ca²⁺ leaks may also enhance β - and γ -secretase activities through direct interaction with these enzymes (86, 87) or by a stabilized γ -secretase complex (86, 87). Accordingly, we show that pharmacological blockade of Ca²⁺ leak by S107 reduces γ -secretase activity. These data led us to conclude that RyR post-translational remodeling amplifies AD pathogenesis through increased γ -secretase-mediated β APP processing. The findings in the present study demonstrate that A β activates both β -adrenergic and oxidative stress. This leads to post-translational remodeling of RyR2 by PKA phosphorylation, oxidation, and nitrosylation and depletion of regulatory proteins calstabin2, PP1, and spinophilin. Data from this study and others (13, 36, 88, 89) are in agreement with a vicious cycle in which leaky RyR2 channels promote A β production and A β enhances RyR2 leak (Fig. 8). These data demonstrate that RyR channels could be envisioned as relevant candidates for a novel therapeutic approach for AD.

RyR post-translational modifications enhance A β production

Experimental procedures

SH-SY5Y APP_{swe} model and treatments

Human SH-SY5Y neuroblastoma cells (CRL-2266, ATCC) were cultured following the manufacturer's instructions. SH-SY5Y cells stably expressing empty pcDNA3.1 vector (Control) or pcDNA3.1 bearing APP_{swe} cDNA were generated as already described (13) and maintained in the presence of Geneticin (400 μ g, Gibco). In this study we used polyclonal stable cells.

Cells were treated overnight for 12 h with β - or γ -secretase inhibitors. γ -Secretase inhibitor ELND006 was used at a 5 μ M final concentration and vehicle (methylcellulose/polysorbate 80, Sigma) was used as control (33, 34). β -Secretase inhibitor (Eli Lilly inhibitor LY288672 (32), synthesized, and kindly provided by Elan Pharmaceuticals) was used at 30 μ M final concentration prepared in DMSO. In some experiments vehicle or DMSO were used as controls because no difference was observed between these two treatments. Cells were treated with various concentrations of: S107 (a benzothiazepine derivative) (19, 26, 28) (0.1, 1, 5, or 10 μ M) for the indicated times (4, 8, 12, 16, or 24 h); H-89 2HCl (Sigma) (1 nM, 1 μ M, 5 μ M, or 15 μ M), a PKA inhibitor; ICI 118,551, hydrochloride ((\pm)-1-[2,3-(dihydro-7-methyl-1H-inden-4-yl)oxy]-3-[(1-methylethyl)amino]-2-butanol hydrochloride) (Sigma) (1 nM, 10 nM, 100 nM, 1 μ M, or 10 μ M), a highly selective β 2-adrenoreceptor antagonist; CGP, 20712A, ((\pm)-2-hydroxy-5-[2-[[2-hydroxy-3-[4-[1-methyl-4-(trifluoromethyl)-1H-imidazol-2-yl]phenoxy]propyl]amino]ethoxy]-benzamide methanesulfonate salt), a highly selective and potent β 1-adrenoreceptor antagonist (Sigma) (1 nM, 10 nM, 100 nM, 1 μ M, or 10 μ M); isoproterenol hydrochloride ((-)-isoprenaline hydrochloride, (-)-*N*-isopropyl-L-noradrenaline hydrochloride, (*R*)-3,4-dihydroxy- α -(isopropylaminomethyl)benzyl alcohol hydrochloride), a β -adrenergic agonist (Sigma) (1 nM, 10 nM, 100 nM, or 1 μ M). ICI, CGP, and H-89 were applied for 12 h. Isoproterenol was applied for 1 h.

Mock-transfected or APP695_{LDN}-expressing CHO cells (individualized clones) were obtained by stable transfection of pcDNA4 empty vector (Control) and hAPP695 cDNA harboring London mutation (APP_{LDN}: APPV642I) and subcloned in pcDNA4 vector. Cells were maintained in DMEM containing 10% FBS, sodium hypoxanthine-thymidine supplement, and 300 μ M proline (31).

A β preparation

Mock-transfected or APP695_{LDN} CHO cells were grown in 150-mm-diameter dishes until reaching 80% confluence, then washed with PBS and allowed to secrete for 24 h into 15 ml of Neurobasal medium (Invitrogen). Secretions were centrifuged (1000 \times *g* for 10 min) and then concentrated into Amicon Ultra-15 3000 filters (4000 \times *g* for 30 min). One-milliliter aliquots of A β preparation concentrates were stored at -80 $^{\circ}$ C until use. A β preparations were controlled on Tris-Tricine gels as already described (see the representative blot in Fig. 2I, and Ref. 31). This A β preparation yields both monomeric and oligomeric A β species but has the advantage of being a "natural" source of A β produced through sequential cleavage of β APP.

A β 40 and A β 42 peptide measurements

Detection of A β 40 and A β 42 peptides secreted in cell media were measured by an ELISA kit (NOVEXTM, ThermoFisher Scientific, France) using human C-terminal A β antibodies.

Biochemical analyses of RyR2 channel remodeling

RyR2 was immunoprecipitated from cell lysates with an RyR2-specific antibody (2 μ g) in 0.5 ml of a modified radioimmune precipitation assay buffer (50 mM Tris-HCl, pH 7.2, 0.9% NaCl, 5.0 mM NaF, 1.0 mM Na₃VO₄, 1% Triton X-100, and protease inhibitors) overnight at 4 $^{\circ}$ C. The RyR2-specific antibody is an affinity-purified polyclonal rabbit antibody custom-made by Yenzym Antibodies (San Francisco, CA) using the peptide CKPEFNNHKDYAQEK corresponding to amino acids 1367–1380 of mouse RyR2 with a cysteine residue added to the N terminus. The immune complexes were incubated with protein A-Sepharose beads (Sigma) at 4 $^{\circ}$ C for 1 h, and the beads were washed three times with radioimmune precipitation assay buffer. The immunoprecipitates were size-fractionated on SDS-PAGE gels (6% for RyR, 15% for calstabin) and transferred onto nitrocellulose membranes for 2 h at 200 mA. Immunoblots were developed using the following primary antibodies: anti-RyR2 (Affinity Bioreagents, 1:2000), anti-phospho-RyR-Ser(P)-2808 (1:5000; Ref 90), anti-calstabin (FKBP12 C-19, 1:1000, Santa Cruz Biotechnology, Inc., Santa Cruz, CA), anti-Cys-NO (1:1,000, Sigma), anti-PP1 (Santa Cruz Biotechnology, 1:2500), and anti-Spinophilin (Abcam, 1:500). To determine channel oxidation, the carbonyl groups in the protein side chains were derivatized to DNP by reaction with 2,4-dinitrophenylhydrazine. The DNP signal associated with RyR was determined using a specific anti-DNP antibody according to the manufacturer's instructions (Millipore, Billerica, MA). We did not explore other RyR2 phosphorylation sites responsive to other kinases (*i.e.* Ca²⁺/calmodulin-dependent kinase (CaMKII)). All immunoblots were developed with the Odyssey system (LI-COR Biosciences, Lincoln, NE) using IR-labeled anti-mouse and anti-rabbit IgG (1:10,000 dilution) secondary antibodies.

Immunofluorescence analysis

Cells grown on 25-mm round coverslips were fixed in paraformaldehyde 4% solution for 10 min at room temperature. Cells were permeabilized with Triton 0.5%, and nonspecific binding sites were blocked for 1 h with BSA (3%). Cells were then incubated at 4 $^{\circ}$ C overnight with primary antibodies diluted in BSA (3%). After 3 washes, coverslips were incubated with secondary antibodies (fluorescent Alexa Fluor antibodies, Alexa 488- and Alexa 594-conjugated (Invitrogen; 1:1000)) at room temperature for 1 h. Immunofluorescence images were acquired on a Leica SP5 confocal microscope using excitation filters 488 and 594 nm. Images were analyzed using ImageJ software. Images were background-corrected, and the colocalization of red and green staining was determined using JACoP plugging. Pearson's correlation coefficient and Mander's Overlap Coefficient were used to evaluate the extent of colocalization as described (91).

We used β APP C-terminal antibody (Sigma) recognizing 676–695 residues of β APP. Other antibodies recognizing the

following proteins were: SERCA2b antibody (clone IID8, Thermo Scientific Pierce Products) targeting SR Ca²⁺-ATPase of the endoplasmic reticulum; EEA1 antibody (clone 14/EEA1, BD Biosciences) targeting early endosome antigen 1; CD63 antibody (clone CLB-180, Abcam) targeting antigens of lysosome membrane and late endosome membrane; CD71 antibody (clone H68.4, Thermo Fisher Scientific) targeting transferrin receptor; TGN46 antibody (AHP500G, AbD Serotec).

Measurements of mitochondrial superoxide concentration

We used MitoSOX (Invitrogen) Red fluorogenic dye to detect superoxide in the mitochondria of living cells (92). Cells grown on 24-mm coverslips were loaded with 5 μ M MitoSOX red prepared in a KRB (containing 125 mM NaCl, 5 mM KCl, 1 mM Na₃PO₄, 1 mM MgSO₄, 5.5 mM glucose, and 20 mM HEPES, pH 7.4) supplemented with 1 mM CaCl₂ (KRB/CaCl₂) at 37 °C for 30 min. After a brief wash, Z-stack images were acquired on a SP5 confocal microscope (Leica). Dye intensity was quantified on Z-stack projection of images after thresholding using Leica SP5 confocal microscope and ImageJ software. Atpenin A5, a specific potent inhibitor of mitochondrial respiratory chain complex II, was used as the positive control (data not shown) (93). We presented data as fluorescence intensity (-fold increase *versus* respective controls in each experiment). Kinetic measurements of MitoSOX are presented as $\Delta F/F$ to measure A β -mediated MitoSOX elevations correcting for differences in basal fluorescence or probe loading.

Cytosolic Ca²⁺ measurements

Cells grown on 24-mm coverslips were loaded with 5 μ M Fluo-4, AM (cytosolic Ca²⁺ probe), prepared in a KRB/CaCl₂ at 37 °C for 30 min. After a brief washout, Z-stack images were acquired on a SP5 confocal microscope (Leica). Cellular loading of dyes was quantified on maximal projection of Z-stack images after thresholding using ImageJ software.

We calibrated the fluorescence measurements, and the intensities of fluorescence for each condition were translated into nanomolar concentrations of Ca²⁺ according to the formula,

$$[Ca^{2+}] = K_d \left(\frac{F - F_{min}}{F_{max} - F} \right) \quad (\text{Eq. 1})$$

where K_d for Fluo4 is 345 nM, F is the background-corrected fluorescence intensity recorded from cells during the experiment, F_{min} is the Ca²⁺-free indicator fluorescence (determined in the ionophore (ionomycin 0.5 μ M)/40 mM MnCl₂ solution), and F_{max} is the Ca²⁺-saturated indicator fluorescence (determined in the ionophore (ionomycin 0.5 μ M)/1 mM Ca²⁺ rich solution) (94).

Kinetic measurement of Fluo4, AM fluorescence (Fig. 3, *E*, *F*, *H*, *I*, and *J*) changes were normalized and presented as $\Delta F/F_0$ to measure A β - or carbachol-mediated Ca²⁺ elevations correcting for differences in basal fluorescence or probe loading (95).

cAMP measurement

We used the cAMP direct immunoassay kit (Abcam) following the manufacturer's instruction.

β APP processing analysis

To detect CTFs (C99 and C83) and A β peptide, protein extracts (40 μ g) were incubated with 70% formic acid (Sigma) and SpeedVac-evaporated for 40 min. The pellets were dissolved in 1 M Tris, pH 10.8, 25 mM betaine and diluted in 2 \times Tris-Tricine loading buffer (125 mM Tris-HCl, pH 8.45, 2% SDS, 20% glycerol, 0.001% bromphenol blue, and 5% β -mercaptoethanol). Proteins were resolved by 16.5% Tris-Tricine SDS-PAGE and transferred onto PVDF membranes. Membranes were boiled in PBS and incubated overnight with specific antibodies.

Antibodies

A β , β -CTF (C99), and full-length β APP were detected using 6E10 antibody (Covance, Rueil-Malmaison, France), which recognizes 1–16 residues of A β . α - and β -APP CTFs (C83 and C99, respectively) were detected using the APP C-terminal antibody (Sigma) recognizing 676–695 residues of β APP. Full-length β APP was also detected using APP (N-terminal) antibody (22C11, Millipore, S.A.S. France). β -Actin (Sigma) was used as loading control.

In vitro γ -secretase assay

In vitro γ -secretase activity was assessed as described (13, 96). 20 μ g of each subcellular fraction were resuspended in solubilization buffer (150 mM sodium citrate, pH 6.4, containing 3-[(3-cholamidopropyl) and dimethylammonio]-2-hydroxy-1-propanesulfonate 1% (v/v) supplemented with protease inhibitor mixture. All steps were performed at 4 °C. Solubilized membranes were diluted once with sodium citrate buffer (150 mM pH 6.4) and with reaction buffer (150 mM sodium citrate, pH 6.4, 20 mM dithiothreitol, 0.2 mg/ml BSA, 1 mg/ml egg phosphatidylcholine and 50 μ g/ml recombinant C100-FLAG). The resulting reaction mixtures were then either incubated with constant agitation for 16 h at 37 °C or stored at 4 °C (negative controls). Samples were then supplemented with 2 \times Tris-Tricine loading buffer, boiled for 5 min, and subjected to 16.5% Tris-Tricine SDS-PAGE.

Statistical analyses

Data are expressed as the mean \pm S.E. Sample size for each experiment is stated in the figure captions. Statistical analyses were performed using one-way or two-way ANOVA and Dunnett's, Bonferroni's, or Tukey's post-tests or *t* test. Minimum statistically significant differences were established at $p < 0.05$. Non-statistically significant differences are not shown in the graphs.

Author contributions—A. R. M., A. L., and M. C. independently conceived the study. R. B., F. C., M. C., A. R. M., A. L., X. L., and S. R. designed the experiments. S. R. and V. S. conducted all the biochemistry assays of RyR2. R. B., M. C., and C. M. conducted the biochemistry assays of APP processing, immunofluorescence, and γ -secretase activity experiments. R. B., M. C., A. L., and X. L. conducted the Ca²⁺ experiments. M. C., A. L., X. L., and R. Z. analyzed ROS production measurements. A. R. M., M. C., F. C., and A. L. analyzed the data and wrote and edited the manuscript.

Acknowledgment—We thank Elan for providing us with ELND006 and LY288672.

References

- Barage, S. H., and Sonawane, K. D. (2015) Amyloid cascade hypothesis: pathogenesis and therapeutic strategies in Alzheimer's disease. *Neuropeptides* **52**, 1–18
- De Strooper, B., and Karran, E. (2016) The cellular phase of Alzheimer's disease. *Cell* **164**, 603–615
- Pardossi-Piquard, R., and Checler, F. (2012) The physiology of the β -amyloid precursor protein intracellular domain AICD. *J. Neurochem.* **120**, 109–124
- Selkoe, D. J., and Hardy, J. (2016) The amyloid hypothesis of Alzheimer's disease at 25 years. *EMBO Mol. Med.* **8**, 595–608
- Zhang, Y. W., Thompson, R., Zhang, H., and Xu, H. (2011) APP processing in Alzheimer's disease. *Mol. Brain* **4**, 3
- Golde, T. E., Cai, X. D., Shoji, M., and Younkin, S. G. (1993) Production of amyloid β protein from normal amyloid β -protein precursor (β APP) and the mutated β APPs linked to familial Alzheimer's disease. *Ann. N.Y. Acad. Sci.* **695**, 103–108
- Van Cauwenberghe, C., Van Broeckhoven, C., and Sleegers, K. (2016) The genetic landscape of Alzheimer disease: clinical implications and perspectives. *Genet. Med.* **18**, 421–430
- Mattson, M. P. (2010) ER calcium and Alzheimer's disease: in a state of flux. *Sci. Signal.* **3**, pe10
- Del Prete, D., Checler, F., and Chami, M. (2014) Ryanodine receptors: physiological function and deregulation in Alzheimer disease. *Mol. Neurodegener.* **9**, 21
- Stutzmann, G. E., Caccamo, A., LaFerla, F. M., and Parker, I. (2004) Dysregulated IP₃ signaling in cortical neurons of knock-in mice expressing an Alzheimer's-linked mutation in presenilin1 results in exaggerated Ca²⁺ signals and altered membrane excitability. *J. Neurosci.* **24**, 508–513
- Shilling, D., Müller, M., Takano, H., Mak, D. O., Abel, T., Coulter, D. A., and Foskett, J. K. (2014) Suppression of InsP3 receptor-mediated Ca²⁺ signaling alleviates mutant presenilin-linked familial Alzheimer's disease pathogenesis. *J. Neurosci.* **34**, 6910–6923
- Jensen, L. E., Bultynck, G., Luyten, T., Amije, H., Bootman, M. D., and Roderick, H. L. (2013) Alzheimer's disease-associated peptide A β 42 mobilizes ER Ca²⁺ via InsP3R-dependent and -independent mechanisms. *Front. Mol. Neurosci.* **6**, 36
- Oulès, B., Del Prete, D., Greco, B., Zhang, X., Lauritzen, I., Sevalle, J., Moreno, S., Paterlini-Bréchet, P., Trebak, M., Checler, F., Benfenati, F., and Chami, M. (2012) Ryanodine receptor blockade reduces amyloid- β load and memory impairments in Tg2576 mouse model of Alzheimer disease. *J. Neurosci.* **32**, 11820–11834
- Chakroborty, S., Goussakov, I., Miller, M. B., and Stutzmann, G. E. (2009) Deviant ryanodine receptor-mediated calcium release resets synaptic homeostasis in presymptomatic 3 \times Tg-AD mice. *J. Neurosci.* **29**, 9458–9470
- Stutzmann, G. E., Smith, I., Caccamo, A., Oddo, S., Laferla, F. M., and Parker, I. (2006) Enhanced ryanodine receptor recruitment contributes to Ca²⁺ disruptions in young, adult, and aged Alzheimer's disease mice. *J. Neurosci.* **26**, 5180–5189
- Stutzmann, G. E., Smith, I., Caccamo, A., Oddo, S., Parker, I., and Laferla, F. (2007) Enhanced ryanodine-mediated calcium release in mutant PS1-expressing Alzheimer's mouse models. *Ann. N.Y. Acad. Sci.* **1097**, 265–277
- Lanner, J. T., Georgiou, D. K., Joshi, A. D., and Hamilton, S. L. (2010) Ryanodine receptors: structure, expression, molecular details, and function in calcium release. *Cold Spring Harb. Perspect. Biol.* **2**, a003996
- Bull, R., Finkelstein, J. P., Gálvez, J., Sánchez, G., Donoso, P., Behrens, M. I., and Hidalgo, C. (2008) Ischemia enhances activation by Ca²⁺ and redox modification of ryanodine receptor channels from rat brain cortex. *J. Neurosci.* **28**, 9463–9472
- Liu, X., Betzenhauser, M. J., Reiken, S., Meli, A. C., Xie, W., Chen, B. X., Arancio, O., and Marks, A. R. (2012) Role of leaky neuronal ryanodine receptors in stress-induced cognitive dysfunction. *Cell* **150**, 1055–1067
- Ni, Y., Zhao, X., Bao, G., Zou, L., Teng, L., Wang, Z., Song, M., Xiong, J., Bai, Y., and Pei, G. (2006) Activation of β 2-adrenergic receptor stimulates γ -secretase activity and accelerates amyloid plaque formation. *Nat. Med.* **12**, 1390–1396
- Yu, J. T., Tan, L., Ou, J. R., Zhu, J. X., Liu, K., Song, J. H., and Sun, Y. P. (2008) Polymorphisms at the β 2-adrenergic receptor gene influence Alzheimer's disease susceptibility. *Brain Res.* **1210**, 216–222
- Yu, N. N., Wang, X. X., Yu, J. T., Wang, N. D., Lu, R. C., Miao, D., Tian, Y., and Tan, L. (2010) Blocking β 2-adrenergic receptor attenuates acute stress-induced amyloid β peptides production. *Brain Res.* **1317**, 305–310
- Wang, D., Fu, Q., Zhou, Y., Xu, B., Shi, Q., Igwe, B., Matt, L., Hell, J. W., Wisely, E. V., Oddo, S., and Xiang, Y. K. (2013) β 2 adrenergic receptor, protein kinase A (PKA) and c-Jun N-terminal kinase (JNK) signaling pathways mediate tau pathology in Alzheimer disease models. *J. Biol. Chem.* **288**, 10298–10307
- Marx, S. O., Reiken, S., Hisamatsu, Y., Gaburjakova, M., Gaburjakova, J., Yang, Y. M., Rosembly, N., and Marks, A. R. (2001) Phosphorylation-dependent regulation of ryanodine receptors: a novel role for leucine/isoleucine zippers. *J. Cell Biol.* **153**, 699–708
- Marx, S. O., Reiken, S., Hisamatsu, Y., Jayaraman, T., Burkhoff, D., Rosembly, N., and Marks, A. R. (2000) PKA phosphorylation dissociates FKBP12.6 from the calcium release channel (ryanodine receptor): defective regulation in failing hearts. *Cell* **101**, 365–376
- Lehnart, S. E., Terrenoire, C., Reiken, S., Wehrens, X. H., Song, L. S., Tillman, E. J., Mancarella, S., Coromilas, J., Lederer, W. J., Kass, R. S., and Marks, A. R. (2006) Stabilization of cardiac ryanodine receptor prevents intracellular calcium leak and arrhythmias. *Proc. Natl. Acad. Sci. U.S.A.* **103**, 7906–7910
- Bellinger, A. M., Reiken, S., Dura, M., Murphy, P. W., Deng, S. X., Landry, D. W., Nieman, D., Lehnart, S. E., Samaru, M., LaCampagne, A., and Marks, A. R. (2008) Remodeling of ryanodine receptor complex causes "leaky" channels: a molecular mechanism for decreased exercise capacity. *Proc. Natl. Acad. Sci. U.S.A.* **105**, 2198–2202
- Marks, A. R. (2006) Novel therapy for heart failure and exercise-induced ventricular tachycardia based on "fixing" the leak in ryanodine receptors. *Novartis Found. Symp.* **274**, 132–147
- Mikami, M., Goubaeva, F., Song, J. H., Lee, H. T., and Yang, J. (2008) β -Adrenoceptor blockers protect against staurosporine-induced apoptosis in SH-SY5Y neuroblastoma cells. *Eur. J. Pharmacol.* **589**, 14–21
- Lochner, A., and Moolman, J. A. (2006) The many faces of H89: a review. *Cardiovasc. Drug Rev.* **24**, 261–274
- Guillot-Sestier, M. V., Sunyach, C., Ferreira, S. T., Marzolo, M. P., Bauer, C., Thevenet, A., and Checler, F. (2012) α -Secretase-derived fragment of cellular prion, N1, protects against monomeric and oligomeric amyloid β (A β)-associated cell death. *J. Biol. Chem.* **287**, 5021–5032
- May, P. C., Dean, R. A., Lowe, S. L., Martenyi, F., Sheehan, S. M., Boggs, L. N., Monk, S. A., Mathes, B. M., Mergott, D. J., Watson, B. M., Stout, S. L., Timm, D. E., Smith Labell, E., Gonzales, C. R., Nakano, M., et al. (2011) Robust central reduction of amyloid- β in humans with an orally available, non-peptidic β -secretase inhibitor. *J. Neurosci.* **31**, 16507–16516
- Probst, G., Aubele, D. L., Bowers, S., Dressen, D., Garofalo, A. W., Hom, R. K., Konradi, A. W., Marugg, J. L., Mattson, M. N., Neitzel, M. L., Semko, C. M., Sham, H. L., Smith, J., Sun, M., Truong, A. P., et al. (2013) Discovery of (R)-4-cyclopropyl-7,8-difluoro-5-(4-(trifluoromethyl)phenylsulfonyl)-4,5-dihydro-1H-pyrazolo[4,3-c]quinoline (ELND006) and (R)-4-cyclopropyl-8-fluoro-5-(6-(trifluoromethyl)pyridin-3-ylsulfonyl)-4,5-dihydro-2H-pyrazolo[4,3-c]quinoline (ELND007): metabolically stable γ -secretase Inhibitors that selectively inhibit the production of amyloid- β over Notch. *J. Med. Chem.* **56**, 5261–5274
- Lauritzen, I., Pardossi-Piquard, R., Bauer, C., Brigham, E., Abraham, J. D., Ranaldi, S., Fraser, P., St-George-Hyslop, P., Le Thuc, O., Espin, V., Chami, L., Dunys, J., and Checler, F. (2012) The β -secretase-derived C-terminal fragment of β APP, C99, but not A β , is a key contributor to early intraneuronal lesions in triple-transgenic mouse hippocampus. *J. Neurosci.* **32**, 16243–16255
- Demuro, A., Mina, E., Kaye, R., Milton, S. C., Parker, I., and Glabe, C. G. (2005) Calcium dysregulation and membrane disruption as a ubiquitous neurotoxic mechanism of soluble amyloid oligomers. *J. Biol. Chem.* **280**, 17294–17300
- Paula-Lima, A. C., Adasme, T., SanMartín, C., Sebollela, A., Hetz, C., Carrasco, M. A., Ferreira, S. T., and Hidalgo, C. (2011) Amyloid β -peptide oligomers stimulate RyR-mediated Ca²⁺ release inducing mitochondrial

- fragmentation in hippocampal neurons and prevent RyR-mediated dendritic spine remodeling produced by BDNF. *Antioxid. Redox Signal.* **14**, 1209–1223
37. Finch, E. A., Turner, T. J., and Goldin, S. M. (1991) Calcium as a coagonist of inositol 1,4,5-trisphosphate-induced calcium release. *Science* **252**, 443–446
 38. Brini, M., Marsault, R., Bastianutto, C., Alvarez, J., Pozzan, T., and Rizzuto, R. (1995) Transfected aequorin in the measurement of cytosolic Ca²⁺ concentration ([Ca²⁺]_i). A critical evaluation. *J. Biol. Chem.* **270**, 9896–9903
 39. Flammang, B., Pardossi-Piquard, R., Sevalle, J., Debayle, D., Dabert-Gay, A. S., Thévenet, A., Lauritzen, I., and Checler, F. (2012) Evidence that the amyloid- β protein precursor intracellular domain, AICD, derives from β -secretase-generated C-terminal fragment. *J. Alzheimers Dis.* **30**, 145–153
 40. von Bernhardi, R., and Eugenin, J. (2012) Alzheimer's disease: redox dysregulation as a common denominator for diverse pathogenic mechanisms. *Antioxid. Redox Signal.* **16**, 974–1031
 41. Palavicini, J. P., Wang, H., Bianchi, E., Xu, S., Rao, J. S., Kang, D. E., and Lakshmana, M. K. (2013) RanBP9 aggravates synaptic damage in the mouse brain and is inversely correlated to spinophilin levels in Alzheimer's brain synaptosomes. *Cell Death Dis.* **4**, e667
 42. Lin, M. T., and Beal, M. F. (2006) Mitochondrial dysfunction and oxidative stress in neurodegenerative diseases. *Nature* **443**, 787–795
 43. Hidalgo, C., and Carrasco, M. A. (2011) Redox control of brain calcium in health and disease. *Antioxid. Redox Signal.* **14**, 1203–1207
 44. Ghosh, D., LeVault, K. R., Barnett, A. J., and Brewer, G. J. (2012) A reversible early oxidized redox state that precedes macromolecular ROS damage in aging nontransgenic and 3xTg-AD mouse neurons. *J. Neurosci.* **32**, 5821–5832
 45. Martínez, M., Fernández, E., Frank, A., Guaza, C., de la Fuente, M., and Hernandez, A. (1999) Increased cerebrospinal fluid cAMP levels in Alzheimer's disease. *Brain Res.* **846**, 265–267
 46. Su, Y., Ryder, J., and Ni, B. (2003) Inhibition of A β production and APP maturation by a specific PKA inhibitor. *FEBS Lett.* **546**, 407–410
 47. Marambaud, P., Chevillier, N., Ancolio, K., and Checler, F. (1998) Post-transcriptional contribution of a cAMP-dependent pathway to the formation of α - and β / γ -secretases-derived products of β APP maturation in human cells expressing wild-type and Swedish mutated β APP. *Mol. Med.* **4**, 715–723
 48. Marambaud, P., Ancolio, K., Alves da Costa, C., and Checler, F. (1999) Effect of protein kinase A inhibitors on the production of A β 40 and A β 42 by human cells expressing normal and Alzheimer's disease-linked mutated β APP and presenilin 1. *Br. J. Pharmacol.* **126**, 1186–1190
 49. Ueda, K., Yagami, T., Kageyama, H., and Kawasaki, K. (1996) Protein kinase inhibitor attenuates apoptotic cell death induced by amyloid β protein in culture of the rat cerebral cortex. *Neurosci. Lett.* **203**, 175–178
 50. Eftekharzadeh, B., Ramin, M., Khodagholi, F., Moradi, S., Tabrizian, K., Sharif, R., Azami, K., Beyer, C., and Sharifzadeh, M. (2012) Inhibition of PKA attenuates memory deficits induced by β -amyloid (1–42), and decreases oxidative stress and NF- κ B transcription factors. *Behav. Brain Res.* **226**, 301–308
 51. Echeverria, V., Ducatzenzeiler, A., Chen, C. H., and Cuello, A. C. (2005) Endogenous β -amyloid peptide synthesis modulates cAMP response element-regulated gene expression in PC12 cells. *Neuroscience* **135**, 1193–1202
 52. Igbavboa, U., Johnson-Anuna, L. N., Rossello, X., Butterick, T. A., Sun, G. Y., and Wood, W. G. (2006) Amyloid β -protein1–42 increases cAMP and apolipoprotein E levels which are inhibited by β 1 and β 2-adrenergic receptor antagonists in mouse primary astrocytes. *Neuroscience* **142**, 655–660
 53. Prapong, T., Uemura, E., and Hsu, W. H. (2001) G protein and cAMP-dependent protein kinase mediate amyloid β -peptide inhibition of neuronal glucose uptake. *Exp. Neurol.* **167**, 59–64
 54. Rosenberg, P. B., Mielke, M. M., Tschanz, J., Cook, L., Corcoran, C., Hayden, K. M., Norton, M., Rabins, P. V., Green, R. C., Welsh-Bohmer, K. A., Breitner, J. C., Munger, R., and Lyketsos, C. G. (2008) Effects of cardiovascular medications on rate of functional decline in Alzheimer disease. *Am. J. Geriatr. Psychiatry* **16**, 883–892
 55. Wang, D., Govindaiah, G., Liu, R., De Arcangelis, V., Cox, C. L., and Xiang, Y. K. (2010) Binding of amyloid β peptide to β 2 adrenergic receptor induces PKA-dependent AMPA receptor hyperactivity. *FASEB J.* **24**, 3511–3521
 56. Andersson, D. C., Betzenhauser, M. J., Reiken, S., Meli, A. C., Umanskaya, A., Xie, W., Shiomi, T., Zalk, R., Lacampagne, A., and Marks, A. R. (2011) Ryanodine receptor oxidation causes intracellular calcium leak and muscle weakness in aging. *Cell Metab.* **14**, 196–207
 57. Brookes, P. S., Yoon, Y., Robotham, J. L., Anders, M. W., and Sheu, S. S. (2004) Calcium, ATP, and ROS: a mitochondrial love-hate triangle. *Am. J. Physiol. Cell Physiol.* **287**, C817–C833
 58. Ross, J. A., McGonigle, P., and Van Bockstaele, E. J. (2015) Locus coeruleus, norepinephrine and A β peptides in Alzheimer's disease. *Neurobiol. Stress* **2**, 73–84
 59. Iversen, L. L., Rossor, M. N., Reynolds, G. P., Hills, R., Roth, M., Mountjoy, C. Q., Foote, S. L., Morrison, J. H., and Bloom, F. E. (1983) Loss of pigmented dopamine- β -hydroxylase positive cells from locus coeruleus in senile dementia of Alzheimer's type. *Neurosci. Lett.* **39**, 95–100
 60. Kalaria, R. N., Andorn, A. C., Tabaton, M., Whitehouse, P. J., Harik, S. I., and Unnerstall, J. R. (1989) Adrenergic receptors in aging and Alzheimer's disease: increased β 2-receptors in prefrontal cortex and hippocampus. *J. Neurochem.* **53**, 1772–1781
 61. Jardanhazi-Kurutz, D., Kummer, M. P., Terwel, D., Vogel, K., Dyrks, T., Thiele, A., and Heneka, M. T. (2010) Induced LC degeneration in APP/PS1 transgenic mice accelerates early cerebral amyloidosis and cognitive deficits. *Neurochem. Int.* **57**, 375–382
 62. Heneka, M. T., Nadrigny, F., Regen, T., Martinez-Hernandez, A., Dumitrescu-Ozimek, L., Terwel, D., Jardanhazi-Kurutz, D., Walter, J., Kirchhoff, F., Hanisch, U. K., and Kummer, M. P. (2010) Locus coeruleus controls Alzheimer's disease pathology by modulating microglial functions through norepinephrine. *Proc. Natl. Acad. Sci. U.S.A.* **107**, 6058–6063
 63. Wisely, E. V., Xiang, Y. K., and Oddo, S. (2014) Genetic suppression of β 2-adrenergic receptors ameliorates tau pathology in a mouse model of tauopathies. *Hum. Mol. Genet.* **23**, 4024–4034
 64. Dobarro, M., Gerenu, G., and Ramirez, M. J. (2013) Propranolol reduces cognitive deficits, amyloid and tau pathology in Alzheimer's transgenic mice. *Int. J. Neuropsychopharmacol.* **16**, 2245–2257
 65. Branca, C., Wisely, E. V., Hartman, L. K., Caccamo, A., and Oddo, S. (2014) Administration of a selective β 2 adrenergic receptor antagonist exacerbates neuropathology and cognitive deficits in a mouse model of Alzheimer's disease. *Neurobiol. Aging* **35**, 2726–2735
 66. Dang, V., Medina, B., Das, D., Moghadam, S., Martin, K. J., Lin, B., Naik, P., Patel, D., Nosheny, R., Wesson Ashford, J., and Salehi, A. (2014) Formoterol, a long-acting β 2 adrenergic agonist, improves cognitive function and promotes dendritic complexity in a mouse model of Down syndrome. *Biol. Psychiatry* **75**, 179–188
 67. Li, S., Jin, M., Zhang, D., Yang, T., Koeglsperger, T., Fu, H., and Selkoe, D. J. (2013) Environmental novelty activates β 2-adrenergic signaling to prevent the impairment of hippocampal LTP by A β oligomers. *Neuron* **77**, 929–941
 68. Mattson, M. P., Cheng, B., Davis, D., Bryant, K., Lieberburg, I., and Rydel, R. E. (1992) β -Amyloid peptides destabilize calcium homeostasis and render human cortical neurons vulnerable to excitotoxicity. *J. Neurosci.* **12**, 376–389
 69. Resende, R., Pereira, C., Agostinho, P., Vieira, A. P., Malva, J. O., and Oliveira, C. R. (2007) Susceptibility of hippocampal neurons to A β peptide toxicity is associated with perturbation of Ca²⁺ homeostasis. *Brain Res.* **1143**, 11–21
 70. Agostinho, P., and Oliveira, C. R. (2003) Involvement of calcineurin in the neurotoxic effects induced by amyloid- β and prion peptides. *Eur. J. Neurosci.* **17**, 1189–1196
 71. Ferreira, E., Oliveira, C. R., and Pereira, C. (2004) Involvement of endoplasmic reticulum Ca²⁺ release through ryanodine and inositol 1,4,5-trisphosphate receptors in the neurotoxic effects induced by the amyloid- β peptide. *J. Neurosci. Res.* **76**, 872–880

RyR post-translational modifications enhance A β production

72. Christensen, R. A., Shtifman, A., Allen, P. D., Lopez, J. R., and Querfurth, H. W. (2004) Calcium dyshomeostasis in β -amyloid and τ -bearing skeletal myotubes. *J. Biol. Chem.* **279**, 53524–53532
73. Adunsky, A., Diver-Haber, A., Becker, D., and Hershkowitz, M. (1995) Basal and activated intracellular calcium ion concentrations in mononuclear cells of Alzheimer's disease and unipolar depression. *J. Gerontol. A Biol. Sci. Med. Sci.* **50**, B201–B204
74. Stieren, E., Werchan, W. P., El Ayadi, A., Li, F., and Boehning, D. (2010) FAD mutations in amyloid precursor protein do not directly perturb intracellular calcium homeostasis. *PLoS ONE* **5**, e11992
75. Brunello, L., Zampese, E., Florean, C., Pozzan, T., Pizzo, P., and Fasolato, C. (2009) Presenilin-2 dampens intracellular Ca²⁺ stores by increasing Ca²⁺ leakage and reducing Ca²⁺ uptake. *J. Cell Mol. Med.* **13**, 3358–3369
76. Tu, H., Nelson, O., Bezprozvanny, A., Wang, Z., Lee, S. F., Hao, Y. H., Serneels, L., De Strooper, B., Yu, G., and Bezprozvanny, I. (2006) Presenilins form ER Ca²⁺ leak channels, a function disrupted by familial Alzheimer's disease-linked mutations. *Cell* **126**, 981–993
77. Shilling, D., Mak, D. O., Kang, D. E., and Foskett, J. K. (2012) Lack of evidence for presenilins as endoplasmic reticulum Ca²⁺ leak channels. *J. Biol. Chem.* **287**, 10933–10944
78. Bezprozvanny, I., Supnet, C., Sun, S., Zhang, H., and De Strooper, B. (2012) Response to Shilling et al. (10.1074/jbc.M111.300491). *J. Biol. Chem.* **287**, 20469; author reply 20470
79. Takeda, T., Asahi, M., Yamaguchi, O., Hikoso, S., Nakayama, H., Kusakari, Y., Kawai, M., Hongo, K., Higuchi, Y., Kashiwase, K., Watanabe, T., Taniike, M., Nakai, A., Nishida, K., Kurihara, S., Donoviel, D. B., et al. (2005) Presenilin 2 regulates the systolic function of heart by modulating Ca²⁺ signaling. *FASEB J.* **19**, 2069–2071
80. Chan, S. L., Mayne, M., Holden, C. P., Geiger, J. D., and Mattson, M. P. (2000) Presenilin-1 mutations increase levels of ryanodine receptors and calcium release in PC12 cells and cortical neurons. *J. Biol. Chem.* **275**, 18195–18200
81. Lee, S. Y., Hwang, D. Y., Kim, Y. K., Lee, J. W., Shin, I. C., Oh, K. W., Lee, M. K., Lim, J. S., Yoon, D. Y., Hwang, S. J., and Hong, J. T. (2006) PS2 mutation increases neuronal cell vulnerability to neurotoxicants through activation of caspase-3 by enhancing of ryanodine receptor-mediated calcium release. *FASEB J.* **20**, 151–153
82. Hayrapetyan, V., Rybalchenko, V., Rybalchenko, N., and Koulen, P. (2008) The N-terminus of presenilin-2 increases single channel activity of brain ryanodine receptors through direct protein-protein interaction. *Cell Calcium* **44**, 507–518
83. Rybalchenko, V., Hwang, S. Y., Rybalchenko, N., and Koulen, P. (2008) The cytosolic N terminus of presenilin-1 potentiates mouse ryanodine receptor single channel activity. *Int. J. Biochem. Cell Biol.* **40**, 84–97
84. Pack-Chung, E., Meyers, M. B., Pettingell, W. P., Moir, R. D., Brownawell, A. M., Cheng, L., Tanzi, R. E., and Kim, T. W. (2000) Presenilin 2 interacts with sorcin, a modulator of the ryanodine receptor. *J. Biol. Chem.* **275**, 14440–14445
85. Muresan, Z., and Muresan, V. (2007) The amyloid- β precursor protein is phosphorylated via distinct pathways during differentiation, mitosis, stress, and degeneration. *Mol. Biol. Cell* **18**, 3835–3844
86. Hayley, M., Perspicace, S., Schulthess, T., and Seelig, J. (2009) Calcium enhances the proteolytic activity of BACE1: An in vitro biophysical and biochemical characterization of the BACE1-calcium interaction. *Biochim. Biophys. Acta* **1788**, 1933–1938
87. Ho, M., Hoke, D. E., Chua, Y. J., Li, Q. X., Culvenor, J. G., Masters, C., White, A. R., and Evin, G. (2010) Effect of metal chelators on γ -secretase indicates that calcium and magnesium ions facilitate cleavage of Alzheimer amyloid precursor substrate. *Int. J. Alzheimers Dis.* **2011**, 950932
88. Chakroborty, S., Briggs, C., Miller, M. B., Goussakov, I., Schneider, C., Kim, J., Wicks, J., Richardson, J. C., Conklin, V., Cameransi, B. G., and Stutzmann, G. E. (2012) Stabilizing ER Ca²⁺ channel function as an early preventative strategy for Alzheimer's disease. *PLoS ONE* **7**, e52056
89. Peng, J., Liang, G., Inan, S., Wu, Z., Joseph, D. J., Meng, Q., Peng, Y., Eckenhoff, M. F., and Wei, H. (2012) Dantrolene ameliorates cognitive decline and neuropathology in Alzheimer triple transgenic mice. *Neurosci. Lett.* **516**, 274–279
90. Wehrens, X. H., Lehnart, S. E., Reiken, S. R., Deng, S. X., Vest, J. A., Cervantes, D., Coromilas, J., Landry, D. W., and Marks, A. R. (2004) Protection from cardiac arrhythmia through ryanodine receptor-stabilizing protein calstabin2. *Science* **304**, 292–296
91. Bolte, S., and Cordelières, F. P. (2006) A guided tour into subcellular colocalization analysis in light microscopy. *J. Microsc.* **224**, 213–232
92. Robinson, K. M., Janes, M. S., and Beckman, J. S. (2008) The selective detection of mitochondrial superoxide by live cell imaging. *Nat. Protoc.* **3**, 941–947
93. Mbaya, E., Oulès, B., Caspersen, C., Tacine, R., Massinet, H., Pennuto, M., Chrétien, D., Munnich, A., Rötig, A., Rizzuto, R., Rutter, G. A., Paterlini-Bréchet, P., and Chami, M. (2010) Calcium signalling-dependent mitochondrial dysfunction and bioenergetics regulation in respiratory chain Complex II deficiency. *Cell Death Differ.* **17**, 1855–1866
94. Bootman, M. D., Rietdorf, K., Collins, T., Walker, S., and Sanderson, M. (2013) Converting fluorescence data into Ca²⁺ concentration. *Cold Spring Harb. Protoc.* **2013**, 126–129
95. Bootman, M. D., Rietdorf, K., Collins, T., Walker, S., and Sanderson, M. (2013) Ca²⁺-sensitive fluorescent dyes and intracellular Ca²⁺ imaging. *Cold Spring Harb. Protoc.* **2013**, 83–99
96. Sevalle, J., Amoyel, A., Robert, P., Fournié-Zaluski, M. C., Roques, B., and Checler, F. (2009) Aminopeptidase A contributes to the N-terminal truncation of amyloid β -peptide. *J. Neurochem.* **109**, 248–256
97. Sanmartín, C. D., Paula-Lima, A. C., García, A., Barattini, P., Hartel, S., Núñez, M. T., and Hidalgo, C. (2014) Ryanodine receptor-mediated Ca²⁺ release underlies iron-induced mitochondrial fission and stimulates mitochondrial Ca²⁺ uptake in primary hippocampal neurons. *Front. Mol. Neurosci.* **7**, 13