A Novel Action of Alzheimer's Amyloid β -Protein ($A\beta$): Oligomeric $A\beta$ Promotes Lipid Release

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Interactions between amyloid β -protein ($A\beta$) and lipids have been suggested to play important roles in the pathogenesis of Alzheimer's disease. However, the molecular mechanism underlying these interactions has not been fully understood. We examined the effect of $A\beta$ on lipid metabolism in cultured neurons and astrocytes and found that oligomeric $A\beta$, but not monomeric or fibrillar $A\beta$, promoted lipid release from both types of cells in a dose- and time-dependent manner. The main components of lipids released after the addition of $A\beta$ were cholesterol, phospholipids, and monosialoganglioside (GM1). Density-gradient and electron microscopic analyses of the conditioned media demonstrated that these $A\beta$ and lipids formed particles and were recovered from the fractions at densities of

~1.08–1.18 g/ml, which were similar to those of high-density lipoprotein (HDL) generated by apolipoproteins. The lipid release mediated by $A\beta$ was abolished by concomitant treatment with Congo red and the PKC inhibitor, H7, whereas it was not inhibited with *N*-acetyl-L-cysteine. These $A\beta$ -lipid particles were not internalized into neurons, whereas HDL-like particles produced by apolipoprotein E were internalized. Our findings indicate that oligomeric $A\beta$ promotes lipid release from neuronal membrane, which may lead to the disruption of neuronal lipid homeostasis and the loss of neuronal function.

Key words: amyloid β -protein; cholesterol release; phospholipid; high-density lipoprotein; cultured neurons; Alzheimer's disease

The mechanism underlying the initiation of the clinicopathological process in Alzheimer's disease (AD) is assumed to be the age-related aggregation of amyloid β -protein ($A\beta$) (Selkoe, 1994; Esiri et al., 1997). This assumption has been supported in part by the findings that highly aggregated $A\beta$ fibrils, but not $A\beta$ monomers, induce neurodegeneration (Mattson et al., 1993; Pike et al., 1993; Lorenzo and Yankner, 1994). This assumption has also been challenged by recent evidence indicating that $A\beta$ oligomers also play an important role in AD pathogenesis (Walsh et al., 1997; Hartley et al., 1999) and that neurodegeneration is induced in mouse brain without amyloid plaque formation (Chui et al., 1999; Hsia et al., 1999).

The role of lipid metabolism in the pathogenesis of AD has been highlighted by the finding that apolipoprotein E (apoE) epsilon 4 is a strong risk factor for the development of AD (Corder et al., 1993; Saunders et al., 1993; Strittmatter et al., 1993). The findings that monosialoganglioside (GM1)-bound $A\beta$ is the initially deposited $A\beta$ species in AD brain (Yanagisawa et al., 1995), and that amyloid fibril formation is induced by $A\beta$ binding to membrane vesicles containing ganglioside (Choo-Smith et al., 1997) and phospholipids (Terzi et al., 1995), suggest that interactions of $A\beta$ with lipids play a crucial role in the pathogenesis of AD. In addition, it has been reported that $A\beta$ modulates cholesterol metabolism in the plasma membrane (Liu et al., 1998) and membrane functions by altering the physico-

chemical properties of membrane constituents including lipids (Muller et al., 1995; Mason et al., 1996; McLaurin and Chakrabarty, 1996; Avdulov et al., 1997). Previous studies have shown that the properties of this interaction of $A\beta$ with lipids are dependent on the aggregation state of $A\beta$ (Avdulov et al., 1997; Mason et al., 1999). These lines of evidence indicate that $A\beta$ interacts with neuronal membranes, disrupting its lipid environment. The interactions of $A\beta$ with lipids have also been demonstrated in physiological conditions; $A\beta$ has been shown to be associated with high-density lipoproteins (HDLs) in the cerebrospinal fluid (Koudinov et al., 1996; Fagan et al., 2000) and human plasma (Koudinov et al., 1994; Biere et al., 1996; Matsubara et al., 1999). However, the mechanism underlying the formation of these $A\beta$ -lipid complexes is poorly understood, and their significance in cholesterol metabolism in the CNS or in the pathogenesis of AD remains to be elucidated.

We have recently shown that apoE modulates cellular cholesterol metabolism in an isoform-specific manner (Michikawa et al., 2000), and tau phosphorylation is enhanced in cholesterol-deficient neurons (Fan et al., 2001) and in the brains of Niemann-Pick disease type C mice (Sawamura et al., 2001), suggesting that cholesterol is the key molecule in the pathogenesis of tauopathy. These lines of evidence have led us to the question of whether $A\beta$ affects cholesterol homeostasis in neurons, leading to the abnormal phosphorylation of tau. In this study, we conducted experiments to examine the effect of $A\beta$ on cholesterol metabolism in cultured neurons.

MATERIALS AND METHODS

Cell culture. All experiments were performed in compliance with existing laws and institutional guidelines. Neuron-rich cultures were prepared from cerebral cortices as described previously (Michikawa and Yanagisawa, 1998), with some modifications. In brief, uteri of gravid rats at embryonic days (E) 17–18 were removed under anesthesia. Cerebral cortices from fetal rat brains were dissected, freed of meninges, and diced

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into small pieces; the cortical fragments were incubated in 0.25% trypsin and 20 mg/ml DNase I in PBS (8.1 mM Na_2HPO_4 , 1.5 mM KH_2PO_4 , 137 mM NaCl, and 2.7 mM KCl, pH 7.4) at 37°C for 20 min. The fragments were then dissociated into single cells by pipetting. The cells were suspended in the feeding medium and plated onto poly-D-lysine-coated 24-well plates at a cell density of $2 \times 10^4/\text{cm}^2$. The feeding medium consisted of DMEM nutrient mixture (DMEM/F12, 50:50) and N_2 supplements. More than 99% of the cultured cells were identified as neurons by immunocytochemical analysis using monoclonal antibody against microtubule-associated protein 2, a neuron-specific marker, at 3 d in culture. For astrocyte-rich cultures, mixed glial cells were prepared according to a previously described method (Isobe et al., 1999). In brief, dissociated cells were prepared from E17–18 rat cerebral cortices as described above and seeded in 75 cm^2 flasks at a cell density of 1×10^7 in DMEM containing 10% FBS. After 2 weeks of incubation *in vitro*, the cells in the astrocytic monolayer were removed by vigorously shaking the flasks. The medium with floating cells was removed, and the remaining monolayer cells were trypsinized (0.1%) and reseeded onto 12-well plates.

Preparation of oligomeric $\text{A}\beta$. Synthetic $\text{A}\beta$ 1–40 (TFA salt) was purchased from Peptide Institute (Osaka, Japan; lot numbers 490703, 491131, 500324, 500520, 500701, and 501001), Bachem (Bubendorf, Switzerland; lot numbers 518765 and 519600), and Sigma (St. Louis, MO; lot number 38H49581). $\text{A}\beta$ was dissolved in dimethyl sulfoxide (DMSO) at 13.3 M and diluted with PBS to obtain a 350 μM stock solution. $\text{A}\beta$ solution was then incubated for 24 hr at 37°C (i $\text{A}\beta$). After incubation, $\text{A}\beta$ solution was filtered with a 0.45 μM Millipore filter (Millipore, Bedford, MA). Peptide concentrations of both nonfiltered and filtered $\text{A}\beta$ were determined using a bicinchoninic acid protein assay kit (Pierce, Rockford, IL). The aggregation state of $\text{A}\beta$ in both solutions was monitored by measuring the intensity of thioflavin-T fluorescence. Fresh $\text{A}\beta$ 1–40 was dissolved in the same manner to obtain 350 μM solution and used for experiments immediately after determination of its peptide concentration. For electron microscopic analysis, each sample was centrifuged in PBS in a SW 41-Ti swing rotor (at 4°C) for 48 hr at 34,200 rpm using a Beckman TL-70. After centrifugation, electron microscopic analysis of the lower part of each solution containing resuspended pellet, if any, was performed.

Quantification of released and intracellular cholesterol and phosphatidylcholine. Neurons in 6- or 12-well plates were labeled in DMEM supplemented with N_2 supplements containing 37 Bq/ml of [^{14}C]acetate (DuPont NEN) for 48 hr, the time period necessary to achieve isotopic steady state in these cells, as has been described previously (Michikawa et al., 2000). Astrocytes in 12-well plates were labeled in DMEM with 5% FBS containing 37 Bq/ml of [^{14}C]acetate (DuPont NEN) for 48 hr. The labeled neurons and astrocytes were rinsed three times with fresh DMEM and treated with the reagents that were examined. Aliquots of 1.0 ml of each conditioned medium were filtered with a 0.45 μM Millipore filter and then transferred into clean glass tubes containing 4.0 ml of chloroform/methanol (2:1 v/v). The organic phase was separated from the aqueous phase, washed twice by vigorous shaking with 3 ml of chloroform/water (1:1 v/v), separated from the aqueous phase by centrifugation, and dried under N_2 gas. For extraction of intracellular lipids, dried cells were incubated in hexane/isopropanol (3:2 v/v) for 1 hr at room temperature. The solvent from each plate was removed and dried under N_2 gas. The organic phases were redissolved in 50 μl of chloroform, and 10 μl of each sample was spotted on activated silica gel high-performance thin-layer chromatography (HPTLC) plates (Merck, Darmstadt, Germany). The lipids were separated by sequential one-dimensional chromatography using the chloroform/methanol/acetic acid/water (25:15:4:2, v/v/v/v) solvent system, followed by another run in hexane/diethyl ether/acetic acid (80:30:1). [^{14}C]cholesterol, [^{14}C]sphingomyelin, and [^{14}C]phosphatidylcholine were used as standards. The chromatography plates were exposed to radiosensitive films, and each lipid was visualized and quantified with BAS2500 (Fuji Film, Tokyo, Japan). The amount of lipid release was calculated as the percentage of released lipid relative to the total lipid content (released plus intracellular lipid).

Density gradient ultracentrifugation. After incubation with i $\text{A}\beta$ at 8 $\mu\text{g}/\text{ml}$ for 24 or 48 hr, the neuronal or astrocyte culture medium was filtered with a 0.45 μM Millipore filter. A discontinuous KBr gradient was prepared in a 14 $\times$ 89 mm ultracentrifuge tube (Ultraclear, Beckman) from the bottom to the top with 2 ml of sample at a density of 1.30 g/ml , 3 ml at 1.21 g/ml , 2 ml at 1.063 g/ml , 2 ml at 1.19 g/ml , and 4 ml at 1.006 g/ml KBr solution (all salt solutions contained 0.1% disodium-EDTA and 0.002% sodium azide, pH 7.4). The sample in the

KBr gradient was centrifuged using a Beckman TL-70 ultracentrifuge in a SW 41-Ti swing rotor (at 4°C) for 48 hr at 34,200 rpm. After density gradient centrifugation, 12 fractions (1.0 ml each) were collected from the top of the gradient using a micropipette. The densities of the fractions were determined by measuring the weight of each 100 μl fraction collected. The last fraction was stirred to dissolve the pellet. The cholesterol and phospholipid contents were determined in each fraction. Five milliliters of chloroform/methanol (2:1 v/v) solvent were added to 1 ml of each fraction, and the mixture was stirred vigorously. After centrifugation, the organic phase was removed from each fraction and dried under N_2 gas. The organic residue was dissolved in a small volume of chloroform, and the total cholesterol and phospholipid contents were determined using cholesterol and phospholipid determination kits (Kyowa Medix, Tokyo, Japan and Wako, Osaka, Japan, respectively).

Viability assay. The release of the cytoplasmic enzyme, lactate dehydrogenase (LDH), into culture medium was determined for the quantification of cell death. Fifty microliters of culture medium were transferred to a fresh 96-well flat-bottomed plate and a colorimetric LDH-release assay was performed according to the instructions of the manufacturer (Promega, Madison, WI); absorbances were read at 490 nm immediately thereafter. For determination of total LDH, the neuronal cultures were incubated with 100 mM H_2O_2 for 10 min at room temperature and released LDH was determined, and the percentage of released LDH per total LDH in each culture was calculated.

Measurement of thioflavin-T binding to aggregated $\text{A}\beta$. Determination of the aggregated $\text{A}\beta$ state in solution was performed on the basis of a previously established method (LeVine, 1995; LeVine, 1999). A 350 μM stock solution of $\text{A}\beta$ 1–40 was prepared as described above and incubated for 24 hr at 37°C. The solution was then diluted to threefold with PBS. One-half the amount of the solution was filtered with a 0.45 μM pore-sized Millipore filter. The protein concentration of nonfiltered and filtered solution was determined using a bicinchoninic acid protein assay kit. Freshly dissolved $\text{A}\beta$ was prepared as described above and diluted to threefold with PBS, and its protein concentration was determined. The concentration of each solution was then adjusted to 50 μM with PBS. Steady-state fluorescence measurements for $\text{A}\beta$ were performed with a multiplate reader (Fluoroskan Ascent, Labsystems Inc., Franklin, MA) (excitation 446 nm, emission 490 nm) in 48-well plates. Each well contained 10 μl of 50 μM $\text{A}\beta$ in 1000 $\mu\text{l}/\text{well}$ of 5 μM thioflavin-T in 50 mM glycine-NaOH, pH 8.5.

Immunoblot analysis of $\text{A}\beta$, apoE, apoJ, and GM1 ganglioside. For immunoblot analysis of $\text{A}\beta$, samples of each fraction isolated by density gradient ultracentrifugation were dissolved in sample solubilizing buffer consisting of 50 mM Tris-HCl, pH 6.8, 10% glycerol, 4% SDS, 10% mercaptoethanol, 8 M urea, and 0.01% bromophenol blue. For analysis of apoE and GM1 ganglioside, samples of each fraction were dissolved in equal volumes of Laemmli buffer. They were then subjected to 4–20% gradient Tris/tricine SDS-PAGE (Dai-ichi Pure Chemical Co., Tokyo, Japan). The separated proteins were transferred onto an immobilized or polyvinylidene difluoride membrane (Millipore) with a semidry electrophoretic transfer apparatus (Nihon Eido, Tokyo, Japan) using a transfer buffer (0.1 M Tris, 0.192 M glycine, and 20% methanol). The blots were blocked with 100% Block Ace (Dainippon Pharmaceutical Co., Osaka, Japan) for 1 hr and incubated with primary antibodies overnight at 4°C. The primary antibodies used were monoclonal antibodies specific for the $\text{A}\beta$ 1–40 ending site, BA27 (Suzuki et al., 1994) (at a final concentration of 3.1 $\mu\text{g}/\text{ml}$), specific for human $\text{A}\beta$ 1–17, 6E10 (Kim et al., 1990) (Senetek, St. Louis, MO) (at a final concentration of 1 $\mu\text{g}/\text{ml}$), polyclonal anti-apoE antibody, AB947 (Chemicon, Temecula, CA) (1:1,000), and anti-apoJ antibody (Rockland, Gilbertsville, PA) (at a final concentration of 0.2 $\mu\text{g}/\text{ml}$). The blots were washed four times with PBS-T (PBS containing 0.05% Tween 20) within a period of 60 min and then incubated with secondary antibodies (horseradish peroxidase-conjugated anti-goat or anti-mouse antibodies, used at a final concentration of 0.4 $\mu\text{g}/\text{ml}$) for 1 hr. For the detection of GM1 ganglioside, the membrane was probed with horseradish peroxidase-conjugated cholera toxin B (Sigma) (final concentration at 42 ng/ml) overnight at 4°C. In between steps, the blots were washed four times with PBS-T for 15 min. Bound antibodies or cholera toxin was detected using ECL (Amersham Pharmacia Biotechnology).

Immunoprecipitation of lipids in association with $\text{A}\beta$ and apoE. The neurons and astrocytes were labeled with 37 Bq/ml of [^{14}C]acetate (DuPont NEN) for 48 hr, followed by three washes in DMEM, and treated with 8 μM i $\text{A}\beta$. The conditioned media, in which neurons and astrocytes were cultured in the presence of i $\text{A}\beta$ for 24–48 hr, were

filtered. The filtered conditioned media were then incubated with 1 μ l of mouse monoclonal antibody, 6E10, or goat polyclonal antibodies, AB947, and anti-apoJ antibody and mouse normal mouse IgG, together with 100 μ l of 20% protein G-Sepharose (Amersham Pharmacia Biotechnology) slurry under rotation at 4°C overnight. The immunoprecipitated lipids associated with protein G-Sepharose were washed in PBS-T three times and solubilized in a solution of chloroform/methanol (2:1 v/v), and the solution was evaporated by N_2 gas. The organic phases were redissolved in 20 μ l of chloroform, and all samples were spotted on activated silica gel HPTLC plates; the lipids were separated by sequential one-dimensional chromatography using the chloroform/methanol/acetic acid/water (25:15:4:2) solvent system, followed by another run in hexane/diethyl ether/acetic acid (80:30:1). The chromatography plates were exposed to radiosensitive films, and each lipid was visualized and quantified with BAS2500. The amount of immunoprecipitated lipid was calculated as the percentage of lipid relative to the total lipid in the media.

Immunoelectron microscopy. The solutions containing A β -lipid particles were obtained by density gradient ultracentrifugation, diluted with H_2O to a density of 1.006, and centrifuged again at 46,000 rpm in a SW 50.1-Ti rotor for 24 hr at 4°C. The 400 μ l portion at the bottom was used for electron microscopic study. The solutions were placed on a carbon-coated electron microscopy grid. Nonspecific binding was blocked by incubation in PBS with 1% bovine serum albumin (BSA) for 10 min. The grids were then placed on a droplet of either the A β -specific antibody, 6E10 (at final concentration of 5 μ g/ml), or normal mouse IgG for 60 min (both diluted in PBS, 0.1% BSA), and passed over seven droplets of washing solution (PBS) for 1 min each. The grids were placed on a droplet of anti-mouse IgG conjugated to 5 nm colloidal gold particles for 60 min (Sigma; diluted 1:20 in PBS, 0.1% BSA), passed over seven droplets of washing solution (PBS), and passed over another seven droplets of distilled water. The specimens were then negatively stained with 2% sodium phosphotungstate.

Statistical analysis. Statistical analysis was performed using StatView computer software (Macintosh), and multiple pairwise comparisons among the sets of data were performed using ANOVA and the Bonferroni *t* test.

RESULTS

Characterization of A β used in this study

We studied the effect of A β 1–40 on lipid metabolism from the point of view of lipid release from neurons and astrocytes in culture. Because A β 1–42 is known to be highly amyloidogenic and assumed to play a critical role in the pathogenesis of AD, the effect of A β 1–42 on cellular lipid metabolism is also an important issue that needs to be addressed. However, the fact that synthetic A β 1–42 is very difficult to handle and that oligomerized A β 1–40 as well as A β 1–42 can be associated with lipids led us to use A β 1–40 in the present study. To characterize A β used in this study, A β 1–40 incubated for 24 hr at 37°C at 350 μ M (iA β -nonfiltered), A β 1–40 incubated in the same way followed by filtration through a 0.45 μ M Millipore filter (iA β -filtered), and freshly dissolved A β (fresh A β) were subjected to thioflavin-T assay, Western blot analysis, and electron microscopy. Determination of A β peptide concentration in each sample was performed using a bicinchoninic acid protein assay kit (Pierce, Rockford, IL). The concentration of A β in each solution was then adjusted to 100 μ M using PBS, and the solutions were used for the experiments. As we reported previously (Isobe et al., 2000), the intensity curve of thioflavin-T reaction with A β , which was incubated at 350 μ M at 37°C, was saturated at 24 hr of incubation. The fluorescence intensity of iA β -filtered was similar to that of A β -nonfiltered, whereas that of fresh A β was as low as background levels of PBS (Fig. 1*a*). Immunoblot analysis showed that tetrameric, trimeric, dimeric, and monomeric A β were found in the samples of iA β -nonfiltered and iA β -filtered, whereas only dimeric and monomeric A β were found in fresh A β (Fig. 1*b*). Electron microscopy showed that fibrils were formed in the samples from iA β -nonfiltered, 7.9 $\pm$ 0.5 nm in diameter and 118 $\pm$ 14 nm in

Figure 1. Characterization of A β 1–40. A β 1–40 was prepared as described in Materials and Methods. *a*, The aliquots of iA β -nonfiltered, iA β -filtered, fresh A β , and PBS were subjected to thioflavin-T assays as described in Materials and Methods. Three independent experiments were performed, and similar results were obtained. *b*, The equal volume of 2 $\times$ sample solubilizing buffer was added to each A β solution, of which the concentration was normalized with PBS. The samples were then subjected to 4–20% Tris/tricine SDS-PAGE, followed by Western blot analysis. *c*, Electron micrograph of each sample is shown. The samples were centrifuged at 34,500 rpm for 48 hr using a SW 41-Ti rotor. Electron microscopic analysis of the lower part of each solution containing the resuspended pellet was performed. Results of negative staining show that fibrillar structures are found in the sample of iA β -nonfiltered (*c*); however, no fibril is detected in the samples of iA β -filtered (*d*) or fresh A β (*e*). Scale bar, 50 nm.

length for measurable fibrils (Fig. 1*c*), although the length was difficult to determine because of the twisted configuration. The morphological characteristics of these fibrils are identical to those of protofibrils with curvilinear structures of 4–11 nm in diameter and <200 nm in length as has been reported previously (Walsh et al., 1997). In contrast, electron microscopy of iA β -filtered and fresh A β did not reveal any structures such as protofibrils or A β -derived diffusible ligands (ADDLs) (Fig. 1*d,e*).

Oligomeric A β promotes lipids release from neurons

When we analyzed lipids released from the cells, all the conditioned media of the cultures treated with iA β were examined after filtration with a 0.45 μ M Millipore filter. Electron microscopic study did not reveal any structures such as fibrils, protofibrils, or ADDLs in the conditioned media of the neuronal cultures incubated with iA β -nonfiltered or iA β -filtered for 24 hr (data not shown). The dose-response curves for the release of cholesterol and phosphatidylcholine from neurons at 4 hr of incubation with iA β -filtered are shown in Figure 2*a*. Incubation

Figure 2. Effect of iA β on cholesterol and phosphatidylcholine release from neurons in culture. Neuron-rich cultures were labeled with [^{14} C]acetate for 48 hr as described in Materials and Methods. Cells were then washed three times with DMEM and incubated with iA β at various concentrations for 4 hr. Synthetic A β 1–40 dissolved at high concentration and incubated at 37°C for 24 hr, followed by filtration, was used. The released lipids in the media and the cellular lipids were extracted and analyzed as described in Materials and Methods. The iA β -mediated release of cholesterol and phosphatidylcholine (PC) (a) was significantly increased in a dose-dependent manner. Each data point represents mean $\pm$ SE for three samples. For the time course study of iA β -mediated lipid release from neurons, cultured neurons were labeled with [^{14} C]acetate for 48 hr and then washed three times with DMEM and incubated with iA β at a final concentration of 8 μ M. The iA β -mediated release of cholesterol and PC (b) increased with incubation time. Each data point represents mean $\pm$ SE for three samples. Effect of Congo red on iA β -mediated lipid release and effect of fresh A β on lipid release from neurons were investigated using labeled neurons with [^{14} C]acetate for 48 hr. c, Cells were washed three times with DMEM and then incubated with iA β (10 μ M), iA β (10 μ M) with Congo red (CR) (10 μ M), CR alone (10 μ M), freshly dissolved A β (frA β) (10 μ M), and frA β plus CR (10 μ M) in serum-free N₂ medium for 24 hr. The release of cholesterol and PC in iA β -treated culture medium was abolished by concurrent treatment with Congo red. Freshly dissolved A β 1–40 did not promote lipid release from these cells. Each data point represents mean $\pm$ SE for four samples. * p < 0.005 versus CONT, iA β + CR, frA β , and frA β + CR. CONT, Control cultures; iA β , incubated A β 1–40; CR, Congo red; frA β , fresh A β 1–40. d, Cells were washed three times with DMEM and then incubated with none (CONT), iA β (5 μ M), iA β (5 μ M) + NAC (1 mM), NAC (1 mM), H₂O₂ (2 mM), and H₂O₂ (2 mM) + NAC (1 mM). * p < 0.001 versus CONT and NAC; ** p < 0.0001 versus H₂O₂ + NAC; # p < 0.06 versus CONT and NAC. NAC, N-acetyl-L-cysteine. e, The cultures were washed three times with DMEM and then incubated with none (CONT), iA β (5 μ M), and iA β (5 μ M) + H7 (30 nM) for 16 hr at 37°C, and the lipids in the medium and the cells were quantified as described in Materials and Methods. * p < 0.004 versus CONT and iA β + H7.

with iA β promoted the release of cholesterol (essentially all unesterified) and phosphatidylcholine from neurons in a dose-dependent manner. Saturation of A β -mediated lipid release was observed at A β concentrations higher than 1 μ M, which can be expected to be present in local extracellular spaces *in vivo*. The time course of cholesterol and phosphatidylcholine release from cultured neurons into the medium in the presence of 8 μ M A β is shown in Figure 2b. The cholesterol and phosphatidylcholine release mediated by A β increased in a time-dependent manner. We used six batches of A β 1–40 obtained from Peptide Institute, one batch of A β 1–40 from Sigma, and two batches of A β 1–40 from Bachem. The iA β prepared from these batches promoted lipid release from neurons. Because among these A β peptides tested, A β peptide from Peptide Institute has the strongest ability

to promote lipid release (data not shown), we used A β peptide obtained from Peptide Institute.

Because Congo red is known to inhibit oligomerization of A β by stabilizing A β monomer (Podlisny et al., 1995, 1998), we next examined whether A β -mediated lipid release is inhibited after concurrent treatment with Congo red. A β was incubated at high concentration for 24 hr at 37°C, filtered, and added into neuronal cultures. As shown in Figure 2c, iA β promoted lipid release from neurons, whereas A β incubated with 10 μ M Congo red for 24 hr at 37°C lost its function as a lipid acceptor. In addition, freshly dissolved A β did not induce lipid release from neurons (Fig. 2c), suggesting that the aggregated form of A β is necessary for acquiring a lipid acceptor function.

Because the concentrations of A β used in this study were high,

Figure 3. Density gradient ultracentrifugation analysis of the culture medium of neurons in the presence of $iA\beta$ or H_2O_2 . Neuronal cultures plated in six-well plastic plates were incubated with $A\beta$ 1–40 ($10 \mu M$) in serum-free N_2 medium for 24 hr. The culture medium was collected, filtered through a $0.45 \mu m$ filter, and subjected to an initial discontinuous density gradient prepared using KBr solution as described in Materials and Methods. *a*, After ultracentrifugation, fractions were obtained, and the density, cholesterol, and phospholipids content in each fraction were determined. *b*, Aliquots of $10 \mu l$ from each fraction were mixed with the same volume of SDS buffer, subjected to SDS gel electrophoresis, and immunoblotted with antibodies against $A\beta$ (BA27), apoE (AB947), and apoJ. GM1 ganglioside in each fraction was detected with HRP-conjugated chorea toxin-B. *c*, Forty-eight hours after plating in serum-free N_2 medium, the neuronal cultures plated in six-well plastic plates were washed in DMEM and incubated with $A\beta$ 1–40 ($10 \mu M$) or 5 mM H_2O_2 in serum-free N_2 medium for indicated periods. The percentage of LDH released from the cultures was determined as described in Materials and Methods. The data are mean $\pm$ SE of triplicate. *d*, Forty-eight hours after plating in serum-free N_2 medium, the neuronal cultures plated in six-well plastic plates were washed in DMEM and incubated with 5 mM H_2O_2 in serum-free N_2 medium for 24 hr. The culture medium was collected, filtered through a $0.45 \mu m$ filter, and subjected to an initial discontinuous density gradient prepared using KBr solution as described in Materials and Methods. After ultracentrifugation, fractions were obtained, and the density and concentrations of cholesterol and phospholipids were determined.

they may have induced oxidation of cell membranes and thereby could be cytotoxic (Schubert et al., 1995; Mark et al., 1996), leading to lipid leakage from cells damaged by $A\beta$. Therefore, we next performed experiments to determine whether free radicals are involved in $iA\beta$ -mediated lipid release in the neuronal cultures. The neuronal cultures were subjected to the following treatments: none, $iA\beta$, $iA\beta$ + *N*-acetyl-L-cysteine (NAC), NAC, H_2O_2 , and H_2O_2 + NAC for 8 hr. The amount of lipids released into the culture media in each culture was determined. NAC at a concentration of 1 mM had no effect on $iA\beta$ -mediated lipid release, whereas lipid leakage caused by H_2O_2 was significantly inhibited by 1 mM NAC (Fig. 2*d*).

It is known that lipid release is an active cellular process and that intracellular signaling molecules such as PKC are involved in cellular cholesterol release (Theret et al., 1990; Mendez et al., 1991; Li and Yokoyama, 1995; Mendez, 1997). To confirm that the $iA\beta$ -mediated lipid release is an active cellular process and not a nonspecific physicochemical phenomenon, we next examined the effect of a PKC inhibitor, H7, on the $iA\beta$ -mediated lipid release from neurons. As shown in the Figure 2*e*, H7 completely inhibited lipid release mediated by $iA\beta$.

Density gradient analysis of lipid particles produced by neurons in the presence of $iA\beta$

The characteristics of the released cholesterol and phospholipids and the localization of $iA\beta$ added exogenously to the serum-free media in cultured neurons were examined. The conditioned media of the neuronal cultures treated with $iA\beta$ ($10 \mu M$) were filtered with a $0.45 \mu m$ Millipore filter and centrifuged at 34,500 rpm for 48 hr at $4^\circ C$ in a tube using a Beckman SW 41-Ti rotor. Figure 3, *a* and *b*, show the results of density gradient ultracentrifugation of the conditioned medium containing $iA\beta$. They show that most of the cholesterol and phospholipids are distrib-

uted similarly across the gradient, with both having densities of ~ 1.08 – 1.18 gm/ml (fractions 5, 6, 7, and 8), which correspond to the densities of HDL (Fig. 3*a*). The cholesterol to phospholipid ratio (w/w) at peak density was 0.35. Western blotting analysis using anti- $A\beta$ antibody, BA27, and anti-apoE antibody, AB947, was performed with the samples from the fractions. Exogenously added $A\beta$ was recovered mainly from fractions 4–8 (Fig. 3*b*). $A\beta$ in these fractions was detected as monomers and dimers under denatured conditions. Endogenous GM1 ganglioside was identified in fractions 5–8 (Fig. 3*b*). However, apoE was not detected in these fractions, and apoJ, which is secreted by neurons, was localized in fractions 10–12 (Fig. 3*b*). GM1 is known to have a strong positive curvature-imposing potency, which is required for the formation of small particles such as HDL (Epan, 1998), suggesting that GM1 may contribute to the formation of $A\beta$ -lipid particles by bending of lipid membranes in a convex manner.

These results indicate that $iA\beta$ -mediated lipid release results in the formation of $A\beta$ -lipid particles. To confirm that $iA\beta$ -mediated lipid release is not the nonspecific lipid leakage from neuronal cultures by free radicals generated by $iA\beta$, we determined whether HDL-like particles were generated in the conditioned medium of the neuronal cultures damaged by H_2O_2 . As shown in Figure 3*c*, $iA\beta$ at $20 \mu M$ has no toxic effect on neuronal cultures until 144 hr of treatment, whereas H_2O_2 at 5 mM exhibits a toxic effect on neuronal cultures at 24 hr of treatment assayed by LDH release (Fig. 3*c*). The conditioned media of these cultures were collected, and lipids, the release of which from neurons was caused by H_2O_2 , were analyzed by density-gradient ultracentrifugation. As shown in Figure 3*d*, the distribution pattern of lipids shows no peak of HDL and was quite different from that mediated by $iA\beta$ (Fig. 3*a*). These lines of evidence together with the finding shown in Figure 2*d*

the solution was fractionated. The density and cholesterol and phospholipid content in each fraction were determined (*c*). Aliquots of 10 μ l from each fraction were mixed with the same volume of SDS buffer, subjected to SDS gel electrophoresis, and immunoblotted with antibodies against A β (BA27), apoE (AB947), and apoJ. GM1 ganglioside in each fraction was detected with HRP-conjugated chorea toxin-B (*d*).

clearly indicate that lipid release mediated by iA β is not nonspecific lipid leakage from damaged cells by a cytotoxic effect of iA β .

Oligomeric A β promotes lipid release from astrocytes

We further examined the effects of iA β on the release of cholesterol and phosphatidylcholine from cultured astrocytes. The astrocyte cultures were labeled with [^{14} C]acetate for 48 hr, washed in DMEM, and incubated with iA β at various concentrations for 4 hr. Lipid concentration released into the media and cellular lipid content were determined. The dose-response curves for the release of cholesterol and phosphatidylcholine at 4 hr of incubation with iA β are shown in Figure 4, *a* and *b*, respectively. Incubation with iA β promoted the release of cholesterol and phosphatidylcholine from astrocytes in a dose-dependent manner. Saturation of iA β -mediated release of these lipids was observed at iA β concentrations higher than 1 μ M. In contrast to iA β , freshly solubilized A β at 10 and 30 μ M did not promote lipid release from astrocytes (Fig. 4, *a* and *b*, respectively).

Density gradient analysis of lipid particles produced by astrocytes in the presence of iA β

The conditioned media of the astrocyte cultures treated with iA β (10 μ M) were filtered with a 0.45 μ M Millipore filter and centrifuged at 34,500 rpm for 48 hr at 4°C in a tube using a Beckman SW 41-Ti rotor. Density gradient analysis of lipid released from astrocytes showed that a major part of cholesterol and phospholipids distributed similarly across the gradient, with both having densities at ~1.08–1.17 gm/ml (fractions 5, 6, 7, and 8), which corresponded to the densities of HDL (Fig. 4*c*). The cholesterol to phospholipids ratio (w/w) at peak density was 2.22. Western blotting analysis using anti-A β antibody, BA27, and anti-apoE antibody, AB947, was performed with the samples from the fractions. Exogenously added A β was recovered mainly in the

fractions between 4 and 8 (Fig. 4*d*). A β was detected as monomer and dimer under denaturing conditions. Endogenous GM1 ganglioside and apoE were identified in the fractions between 5 and 8 (Fig. 4*d*). However, apoJ, which is secreted from neurons, was localized in fractions 10, 11, and 12 (Fig. 4*d*).

Morphological and biochemical analysis of lipid particles associated with A β

To confirm the conformation of A β -lipid particles isolated by density gradient analysis of filtered culture medium of neurons in the presence of iA β , fractions containing A β -lipid particles were diluted with distilled water to a density of 1.006 gm/ml and centrifuged at 34,500 rpm for 48 hr at 4°C in a tube using a Beckman SW 41-Ti rotor. The conditioned media of the neuronal cultures in the presence of 0.25 μ M of apolipoprotein E3 (apoE3) were also analyzed. The lower part of each solution was subjected to immunoelectron microscopic study. Analysis of samples from neuronal conditioned media in the presence of iA β revealed that lipoprotein particles were spherical, with a mean diameter of 29.4 ± 1.1 nm, similar to the appearance of, but larger than, HDL-like particles formed via the apoE-mediated manner, the mean diameter of which is 11.4 ± 0.5 nm (Fig. 5*a,b*). In addition, A β was found to be associated with lipid particles as demonstrated by the immunoreactivity of lipid particles against the anti-A β antibody, 6E10, using electron microscopy (Fig. 5*c*). In contrast, A β immunoreactive lipid particles were not detected in negative control samples (Fig. 5*d*), suggesting that iA β is directly associated with lipid particles.

We further examined whether A β -lipid particles contain other acceptors such as apoE and apoJ. The filtered conditioned medium of neuronal cultures was subjected to immunoprecipitation using antibodies including anti-apoE antibody, AB947, anti-A β antibodies, 6E10, anti-apoJ antibody, or normal mouse IgG as a

Figure 4. iA β promotes lipid release from astrocytes in culture. Astrocyte-rich cultures were labeled with [^{14} C]acetate for 48 hr as described in Materials and Methods. Cells were then washed three times with DMEM and incubated with iA β 1–40 or fresh A β at various concentrations for 4 hr. The conditioned media were collected and then filtered. The lipids that were released into the medium and the intracellular lipids were extracted and analyzed as described in Materials and Methods. iA β (○) promoted the release of cholesterol (*a*) and phosphatidylcholine (PC) (*b*) in a dose-dependent manner; fresh A β (●) did not. Data are mean $\pm$ SE for four samples. The scale bars are smaller than the symbol size at 0 μ M (*a* and *b*). Density gradient ultracentrifugation analysis was performed with the conditioned medium of astrocytes in the presence of iA β . Astrocytes plated in six-well plastic plates were incubated with A β 1–40 (10 μ M) in DMEM for 24 hr. The culture medium was collected, filtered through a 0.45 μ M filter, and subjected to an initial discontinuous density gradient prepared using KBr solution as described in Materials and Methods. After ultracentrifugation,

Figure 5. Electron micrographs of lipid particles associated with iA β and apoE3. Neuronal cultures were incubated with 10 μ M iA β or 0.25 μ M apoE3 for 24 hr. The conditioned media of these cultures were collected, filtered with a 0.45 μ m Millipore filter, subjected to an initial discontinuous density gradient prepared using KBr solution, and centrifuged at 34,500 rpm for 48 hr using a SW 41-Ti rotor. After centrifugation, 12 fractions were isolated, and lipid concentration and the density in each fraction were determined. HDL fractions were then diluted with 10-fold volumes of distilled water, followed by centrifugation at 34,500 rpm for 48 hr using a SW 41-Ti rotor. Electron microscopic examination of the lower part of each solution was performed. Negative staining of electron micrographs of lipid particles in the presence of apoE and iA β is shown (*a* and *b*, respectively). Results of immunoelectron microscopy show that exogenously added iA β 1–40 forms complexes with lipid particles as demonstrated using the antibody directed against human A β 1–17, 6E10 (Kim et al., 1990). *c*, Gold labeling is considered to be associated with lipid particles. *d*, In contrast, lipid particles were not labeled with gold without 6E10. Scale bar, 50 nm.

negative control. The ratio of cholesterol and phospholipids associated with protein G-Sepharose to those in the conditioned medium is shown in Figure 6. Approximately 12% of total cholesterol and phospholipids in the conditioned medium was immunoprecipitated with 6E10, whereas the percentages of both lipids immunoprecipitated with AB947, anti-apoJ antibody, and normal mouse IgG were significantly lower than those with 6E10 in neuronal culture (Fig. 6*Aa,b*). In the conditioned medium of astrocytes incubated with 10 μ M iA β , the percentages of cholesterol and phospholipids immunoprecipitated with 6E10 were 4.7 and 3.1%, respectively, of those in the conditioned medium, and those immunoprecipitated with AB947 were 4.7 and 3.2% of those in the conditioned medium, respectively (Fig. 6*Ba,b*). However, as seen in neuronal cultures, the percentages of either lipid immunoprecipitated with anti-apoJ antibody and normal mouse IgG were significantly lower than those with 6E10 or AB947 in astrocyte cultures.

Figure 6. Binding of A β by immobilized anti-A β antibody, 6E10, results in capture of lipids. Filtered conditioned medium of neuronal and astrocyte cultures incubated with iA β for 48 hr was subjected to immunoprecipitation using anti-A β antibody (6E10), anti-apoE antibody (AB947), anti-apoJ antibody, and normal mouse IgG. *A*, The protein-G-Sepharose-associated lipids were determined using the kits described in Materials and Methods. The quantity of cholesterol (*a*) and phospholipids (*b*) immunoprecipitated with 6E10 from the conditioned medium of neuronal culture incubated with iA β was ~12% of the total cholesterol and phospholipids in the initial conditioned medium. However, those immunoprecipitated with anti-apoE antibody, anti-apoJ antibody, or normal mouse IgG were significantly low. *B*, In astrocyte culture medium, the quantity of cholesterol (*a*) and phospholipids (*b*) immunoprecipitated with 6E10 and AB947 from the conditioned medium incubated with iA β was significantly higher than those with anti-apoJ antibody or normal mouse IgG. Western blot analysis using anti-apoE antibody (AB947) and anti-A β antibody (BA27) was performed with the immunoprecipitates. The bands corresponding to A β monomers and dimers is labeled only in immunoprecipitates by 6E10, whereas no band or a faint band was detected in those by AB947, anti-apoJ antibody, and normal mouse IgG in neurons (*Ac*). The bands corresponding to A β monomers and dimers and apoE are labeled in both immunoprecipitates by 6E10 and AB947, respectively, whereas no band or a faint band was detected in those by anti-apoJ and normal mouse IgG for astrocytes (*Bc*). Data are mean $\pm$ SE for four samples. * p < 0.01 versus 6E10, anti-apoJ, and normal IgG (*Ac*); * p < 0.01 versus anti-apoJ and normal IgG (*Bc*).

Western blotting analysis using anti-A β antibody, BA27, and anti-apoE antibody, AB947, was performed with the immunoprecipitates. The pellet fractions from the immunoprecipitation of neuronal conditioned medium with the anti-A β antibody contained exogenously added A β , being compatible with lipid content in association with immunoprecipitates (Fig. 6*Ac*). In contrast, those from the immunoprecipitation of astrocyte-conditioned medium with the anti-A β antibody contained both exogenously added A β and endogenous apoE (Fig. 6*Bc*). In addition, those of astrocyte-conditioned medium with the anti-apoE antibody contained both exogenously added A β and endogenous apoE (Fig. 6*Bc*).

Binding and internalization of A β -lipid particles into neurons

Because iA β 1–40 was found to be an acceptor of lipids from neurons and astrocytes to form A β -lipid particles, it was considered appropriate to perform further study on binding and internalization of A β -lipid particles produced by iA β . The conditioned media of [^{14} C]acetate-labeled astrocytes in the presence of iA β at 10 μ M or apoE3 at 0.25 μ M in DMEM after 24 hr incubation were collected. A β -lipid particles were recovered from HDL fractions by density gradient centrifugation and dialyzed. The fractions containing A β -lipid particles were collected and dialyzed, and the radioactivity contained was normalized by a scintillation counter. Neuronal cultures were then incubated with the equal amount of [^{14} C]labeled-A β -lipid particles in DMEM for 20 min at 4 or 37°C. The cultures were washed with cold PBS three times and dried under the fresh airflow. The lipids were extracted with hexane/isopropanol (3:2 v/v), separated by HPTLC, and quantified with BAS2500. Figure 7 demonstrates the binding activity and internalization efficacy of A β -lipid particles and HDL-like particles formed by exogenously added iA β and apoE3, respectively. The ratio of the labeled-cholesterol activity associated with neurons per total cholesterol activity in the added medium was significantly lower at both 4 and 37°C in the cultures incubated with conditioned medium treated with iA β 1–40 (Fig. 7*a*). In contrast, it was significantly higher both at 4 and 37°C in the cultures incubated with apoE. Similar results were observed for the ratio of the labeled phosphatidylcholine in association with the cells (Fig. 7*b*).

DISCUSSION

In the present study, we found a novel action of A β : oligomeric A β can promote lipid release from astrocytes and neurons to form A β -lipid particles consisting of cholesterol, phospholipids, GM1 ganglioside, and A β . A β -lipid particles produced by oligomeric A β have very low binding affinity to neurons and therefore are not internalized into neurons, suggesting that oligomeric A β may affect intracellular lipid metabolism. Because high concentrations of A β are known to induce oxidation and can be cytotoxic (Schubert et al., 1995; Mark et al., 1996), we have examined the toxicity of A β used in this study and found that iA β has no cytotoxic effect on neurons until 144 hr of treatment, as demonstrated by LDH assay. We have also found that NAC, a potent antioxidant molecule, has no effect on iA β -mediated lipid release, and lipids released from the cells after the addition of H $_2$ O $_2$ do not form lipid particles, which were recovered in HDL fractions. These lines of evidence clearly indicate that lipid release mediated by iA β is not nonspecific lipid leakage from damaged cells by cytotoxic effect of iA β . Because Congo red is a well known dye that not only binds to A β fibrils and A β oligomers to inhibit fibril formation but also inhibits A β oligomerization by stabilizing A β monomer (Podlisny et al., 1995, 1998), we examined the effect of Congo red on iA β -mediated lipid release. We also performed experiments to determine whether freshly dissolved A β can serve as a lipid acceptor. We found that neither freshly prepared A β nor iA β incubated with 10 μ M Congo red removes lipids from neurons. These findings suggest that oligomerized A β is required for acquiring the ability to promote lipid release from the cells. In addition, iA β -filtered does not contain any fibrils or ADDLs (Fig. 1*c*). Moreover, all conditioned media of cultures treated with iA β were examined after filtration, and electron microscopic study did not detect any fibrils in the samples after 24 hr incubation (data not shown). These lines of evidence exclude the possibility that

Figure 7. Binding affinity and internalized efficacy of A β -lipid particles into neurons. Astrocyte-rich cultures were labeled with [^{14} C]acetate for 48 hr as described in Materials and Methods. Cells were then washed three times with DMEM and incubated with 10 μ M iA β 1–40 or 0.25 μ M human recombinant apoE3 for 24 hr. The conditioned media were obtained, filtered using a 0.45 μ m filter, and subjected to density-gradient ultracentrifugation at 34,000 rpm for 48 hr in a Beckman SW 41-Ti rotor. The HDL fractions were then collected and dialyzed. The radioactivity in each sample was determined by a scintillation counter and normalized with DMEM. The normalized conditioned medium containing iA β or apoE3 was added to neuronal cultures at 4 or 37°C. Twenty minutes after the addition, the cultures were washed three times with cold PBS and dried under air flow at room temperature. The lipids in each culture were extracted by incubation with hexane/isopropanol (3:2 v/v) solution for 1 hr. Then the solution was moved into tubes and dried under N $_2$ gas. The extracted lipids were then dissolved in chloroform, developed in HPTLC, and quantified by BAS2500 (Fuji Film, Tokyo, Japan). *a*, The ratio of the labeled cholesterol associated with neurons was significantly lower at both 4 and 37°C in the cultures incubated with conditioned medium treated with iA β 1–40. However, it was significantly higher at both 4 and 37°C in the cultures incubated with apoE. Similar results were observed for the ratio of the labeled phosphatidylcholine in association with the cells (*b*). Data are mean $\pm$ SE for six samples. * p < 0.003.

A β fibrils, but not A β oligomers, act as lipid acceptors. Taken together, oligomerization to dimers, trimers, tetramers, and possibly larger assemblies is indispensable for A β to acquire the ability to promote lipid release.

The molecular mechanism by which oligomeric A β , but not monomeric A β , promotes cholesterol and phospholipid release remains obscure. However, one can explain the mechanism based on the differential binding affinity of A β to lipids. It has been reported that the binding efficacy of lipids, including cholesterol, phosphatidylcholine, and free fatty acids, to A β is increased when the added A β forms polymers (Avdulov et al., 1997). Aggregated A β exhibits strong electrostatic interactions with the surface of model membranes, which appear to mediate its neurotoxicity (Hertel et al., 1997). In addition, a synthetic peptide in an amphipathic β -sheet structure was found to associate with lipids, including cholesterol and phosphatidylcholine, and is being pro-

posed as a model for apolipoprotein B (Osterman et al., 1984), suggesting that cholesterol and other lipids bind to hydrophobic areas of amphipathic β -sheet. These lines of evidence suggest that cholesterol and other lipids may bind to hydrophobic areas of aggregated $A\beta$ to form $A\beta$ -lipid particles.

Another interesting point is that $A\beta$ -lipid particles generated in the cultured astrocytes are cholesterol rich compared with HDL-like particles produced by apolipoproteins, such as apoE or apoAI (Ito et al., 1999; Michikawa et al., 2000). In addition, our finding that the average size of $A\beta$ -lipid particles produced by $iA\beta$ (29.4 ± 1.1 nm) is larger than that of HDL-like particles produced by apoE (11.4 ± 0.5 nm) in neuronal cultures suggests that there could be two species of lipid particles, one produced by $iA\beta$ and the other produced by apoE. The size of HDL-like particles produced by apoE is consistent with that previously reported in astrocyte conditioned medium or in CSF (LaDu et al., 1998; Fagan et al., 1999). The mechanisms underlying these cell type- and acceptor type-specific discrepancies are not clear, and further studies are needed to address these issues.

Interestingly, $A\beta$ -lipid particles produced by $iA\beta$ cannot bind to neurons and thus are not internalized into the cells, whereas HDL-like particles produced by apoE can do so. This may be because neurons have no receptors to bind $A\beta$. It has been shown that under physiological conditions, a significant amount of soluble $A\beta$ is associated with apoE-containing HDL particles in CSF, AD brain, and culture medium (Koudinov et al., 1994, 1996; Biere et al., 1996). In contrast to oligomeric $A\beta$, when monomeric $A\beta$ forms complexes with such particles, apoE receptors bind and internalize these lipid complexes into the cells, thereby modulating the amount of $A\beta$ in the brain parenchyma through cellular clearance mechanisms (Holtzman et al., 1999). This may be because the protein component of HDL under physiological conditions is predominantly apoE and not $A\beta$, and thus apoE can function as a ligand to its receptors. These lines of evidence raise the question of whether $A\beta$ directly binds to apoE and not to lipids, or directly binds to lipids to form apoE-lipid- $A\beta$ complexes. It is difficult to answer this question using astrocyte cultures, because astrocytes synthesize and secrete both apoE and $A\beta$ (Figs. 4d, 6Bc). However, our data on $iA\beta$ -mediated lipid release from neurons can exclude the former possibility, because lipid particles associated with $iA\beta$ contained neither apoE nor apoJ (Figs. 3b, 6Ac). These data indicate that $iA\beta$ interacts directly with lipid particles. Previous studies have shown that $A\beta$ oligomers (dimers, trimers, tetramers, and possibly larger assemblies) are formed in the conditioned media of certain cell lines that constitutively secrete $A\beta$ and that endogenous and synthetic $A\beta$ can assemble into stable oligomers at physiological concentrations in culture (Podlisny et al., 1995, 1998; Xia et al., 1997). Recently, $A\beta$ oligomers have been identified in CSF of AD patients (Pitschke et al., 1998). These lines of evidence together with our present results may allow us to assume that oligomeric $A\beta$ accumulates extracellularly under pathophysiological conditions such as the AD brain, and this oligomeric $A\beta$, in turn, may stimulate lipid release from neurons, leading to disruption of cholesterol homeostasis in the CNS.

We have recently found that the deficiency in intracellular cholesterol content causes tau phosphorylation (Fan et al., 2001). We have also demonstrated that tau is hyperphosphorylated in brains of mice with Niemann-Pick disease type C (Sawamura et al., 2001), which is known as a cholesterol storage disease involving late endosomes and lysosomes with defective intracellular sterol trafficking (Scriver et al., 1995). On the basis of these lines

of evidence, together with the findings reported here, we hypothesize that oligomeric $A\beta$ promotes lipid release, which in turn may reduce cellular cholesterol levels, thereby promoting tau phosphorylation and neurodegeneration, as observed in AD brain. Our observations in the present study also provide new insight into the central issue concerning the pathogenesis of AD, that is, the relationship between amyloid plaque formation and the development of neurofibrillary tangles in neurons.

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