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Impact of sphingosine and acetylsphingosines on the aggregation and toxicity of metal-free and metal-treated amyloid- β

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Pathophysiological shifts in the cerebral levels of sphingolipids in Alzheimer's disease (AD) patients suggest a link between sphingolipid metabolism and the disease pathology. Sphingosine (SP), a structural backbone of sphingolipids, is an amphiphilic molecule that is able to undergo aggregation into micelles and micellar aggregates. Considering its structural properties and cellular localization, we hypothesized that SP potentially interacts with amyloid- β (A β) and metal ions that are found as pathological components in AD-affected brains, with manifesting its reactivity towards metal-free A β and metal-bound A β (metal-A β). Herein, we report, *for the first time*, that SP is capable of interacting with both A β and metal ions and consequently affects the aggregation of metal-free A β and metal-A β . Moreover, incubation of SP with A β in the absence and presence of metal ions results in the aggravation of toxicity induced by metal-free A β and metal-A β in living cells. As the simplest acyl derivatives of SP, *N*-acetylsphingosine and 3-*O*-acetylsphingosine also influence metal-free A β and metal-A β aggregation to different degrees, compared to SP. Such slight structural modifications of SP neutralize its ability to exacerbate the cytotoxicity triggered by metal-free A β and metal-A β . Notably, the reactivity of SP and the acetylsphingosines towards metal-free A β and metal-A β is determined to be dependent on their formation of micelles and micellar aggregates. Our overall studies demonstrate that SP and its derivatives could directly interact with pathological factors in AD and modify their pathogenic properties at concentrations below and above critical aggregation concentrations.

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Introduction

Accounting for approximately 50% of the brain's dry weight, lipids serve as signaling molecules, energy storage, and the building blocks of cellular membranes in neurons.^{1,2} Sphingolipids, among the different types of lipids, are essential for the

structural integrity of neuronal membranes with implications in cellular recognition, neurotransmission, myelin sheath formation, and apoptotic regulation.^{3,4} Previous research has examined the pertinence of sphingolipid metabolism in brain homeostasis through the concept of sphingolipid rheostat, defined as the balance in the salvage pathway of sphingolipids.^{5,6} For these reasons, multiple aspects of sphingolipids have been investigated with respect to the development, function, and homeostasis of the brain as well as neurodegeneration.^{3,5,7-10}

As illustrated in Fig. 1a, sphingosine (SP) is an amino alcohol with an unsaturated hydrocarbon chain presenting the structural backbone of sphingolipids.^{5,11,12} Amphiphilic compounds, including SP, are known to form micelles and micellar aggregates at concentrations above their critical aggregation concentrations (CACs).¹³⁻¹⁷ Several reports suggest the subcellular localization of SP in the plasma membrane, cytosol, lysosome, mitochondria, Golgi apparatus, and endoplasmic reticulum.^{7,18,19} As a component of cellular regulation circuits, SP is engaged in regulating the release of neurotransmitters by controlling pre-synaptic vesicle fusion under normal conditions.^{20,21} In addition, pathological shifts in the sphingolipid rheostat manifesting increased levels of SP are associated with

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Fig. 1 SP and acetyl sphingosines studied in this work. (a) Chemical structures of SP, N-Ac-SP, and 3-O-Ac-SP. An acetyl group was incorporated into the structure of SP at the N- or 3-O-position yielding N-Ac-SP and 3-O-Ac-SP, respectively. SP, (2S,3R)-2-amino-octadec-4-trans-ene-1,3-diol; N-Ac-SP, (2S,3R,4E)-2-(acetylamino)-4-octadecene-1,3-diol; 3-O-Ac-SP, (2S,3S,E)-2-amino-1-hydroxy-octadec-4-en-3-yl acetate. Potential donor atoms for metal binding are highlighted in blue. (b) Amino acid sequences of Aβ₄₀ and Aβ₄₂ (top) as well as hydrophobicity and metal binding of Aβ (bottom). Top: the amino acid residues involved in metal coordination and the self-recognition site are highlighted in blue and red, respectively. Bottom: (left) structure of Aβ₄₀ (PDB ID: 2LFM)³⁵ exhibiting a degree of hydrophobicity; (right) examples of Zn(II) and Cu(II) coordination to Aβ.^{30,31} Possible fifth ligands on the metal centers are not shown in this figure. (c) Synthetic routes to 3-O-Ac-SP. Reagents and conditions: (i) (1) vinylmagnesium bromide, THF, -30 °C (30 min) to 0 °C (1 h); (ii) (2) sat. NH₄Cl (aq); (iii) 1-pentadecene, second-generation Grubbs catalyst; (iv) CH₂Cl₂, room temperature, 16 h; (v) acetyl chloride, 4-dimethylaminopyridine; (vi) pyridine : CH₂Cl₂ (2 : 1), room temperature, overnight; (vii) trifluoroacetic acid; (viii) CH₂Cl₂ : H₂O (2 : 1), room temperature, overnight.

pro-apoptotic conditions contributing towards neurodegeneration.^{6,21,22} Recently, elevated concentrations of SP have been detected in the post-mortem brains of Alzheimer's disease

(AD) patients.²² Despite such notions, a connection between SP and AD is only beginning to be revealed.

AD is the most common neurodegenerative disease exhibiting significant intricacy in its pathology involving several pathogenic features, such as proteopathy and metal ion dys-homeostasis.^{23–34} The proteopathic implications of AD include the pathology of amyloid-β (Aβ), while the metal ion hypothesis focuses on the dualistic pathogenic qualities of loss-of-function and gain-of-toxicity through metal ion dyshomeostasis and miscompartmentalization.^{23–34} Increasing evidence indicates the independent and synergistic contributions of Aβ and metal ions towards neurodegeneration.^{23–34} Aβ, the main component of senile plaques, is an aggregation-prone peptide that forms neurotoxic species.^{24–35} Metal ions [e.g., Zn(II) and Cu(II)] colocalized with Aβ aggregates in senile plaques can coordinate to Aβ peptides, as displayed in Fig. 1b, and affect the aggregation of Aβ.^{24–34} The contemporary viewpoint on the complex pathology of AD considers the inter-relationships among different pathological elements found in the disease.^{24,25,32}

In this study, SP was hypothesized to interact with Aβ and metal ions and subsequently alter the aggregation and cytotoxicity of metal-free Aβ and metal-bound Aβ (metal-Aβ). Such interactions under pathological conditions are supported by four fundamental aspects: (i) potential points of convergence at the subcellular level for SP, Aβ, and metal ions,^{7,18,19,36} (ii) enhanced concentrations of SP, Aβ, and metal ions in AD-affected brains,^{22,24–26} (iii) amphiphilic properties of both SP and Aβ, and (iv) inclusion of potential metal-binding sites on the structure of SP. Three sphingolipids, including SP, N-acetylsphingosine (N-Ac-SP), and 3-O-acetylsphingosine (3-O-Ac-SP) shown in Fig. 1a, were utilized to experimentally confirm these interactions at the molecular level in association with their ability to form micelles and micellar aggregates and evaluate their impact on the aggregation and cytotoxicity of both metal-free Aβ and metal-Aβ. Overall, we provide a multidisciplinary perspective on the interactions and pathogenic connections between our molecular subjects.

Results and discussion

Preparation and properties of SP and acetyl sphingosines

SP, a skeletal structure of sphingolipids, is subject to a diverse range of structural variations such as acylation, phosphorylation, and glycosylation, often serving as a measure to control cellular signaling.^{5,12,37} In particular, acylation is a universal modification of biomolecules with nitrogen (N) and oxygen (O) atoms generating amide and ester groups, respectively.³⁸ In this work, we incorporated the acetyl functionality, as the simplest acyl group, onto the structure of SP to construct N-Ac-SP and 3-O-Ac-SP, as presented in Fig. 1a, and employed them as representative sphingolipids, along with SP. SP and N-Ac-SP were obtained commercially, while 3-O-Ac-SP was synthesized following previously reported procedures with slight modifications, as summarized in Fig. 1c.^{39–43} The Grignard reaction of Garner's aldehyde with vinylmagnesium bromide produced 1. Next, 2 was generated via the cross-metathesis reaction of 1 with 1-pentadecene using the second-generation Grubbs' catalyst. The subsequent



acetylation of **2** with acetyl chloride under basic conditions afforded **3**. Lastly, after acid-catalyzed deprotection of the cyclic *N,O*-aminal and *tert*-butoxycarbonyl group in **3**, the final product, **3-O-Ac-SP**, was obtained in a modest yield (Fig. S1†).

SP and the two acetyl sphingosines are expected to interact with A β and metal ions based on their molecular structures depicted in Fig. 1a. For A β interaction, the alkenyl chains and the amino alcohol moieties with or without acetyl substitution of these molecules may interact with the hydrophobic and hydrophilic regions of A β , respectively, as presented in Fig. 1b.²⁶ All three compounds embody potential metal-binding sites. **SP** has two bidentate sites for metal ions, *i.e.*, the moieties of 2-aminoethanol and 1,3-propanediol. The two acetyl sphingosines also have possible metal-binding sites composed of N and O donor atoms. Moreover, the amphiphilicity of these compounds

drives their self-aggregation into micelles and micellar aggregates at concentrations above their CACs.^{13–17} Their corresponding CACs were measured by dynamic light scattering in appropriate buffered solutions. The CACs of **SP**, **N-Ac-SP**, and **3-O-Ac-SP** were determined to be *ca.* 3–81 μ M, 7–27 μ M, and 29–68 μ M, respectively, depending on pH and the presence of a salt (*e.g.*, NaCl) (Fig. S2†). The aggregation is initiated at their corresponding CACs. The concentration ratio of micelles and micellar aggregates to monomers theoretically increases in a manner proportional to their concentrations above the CACs.^{44,45}

Interactions of **SP** and acetyl sphingosines with A β

To analyze the molecular-level interactions between **SP**, **N-Ac-SP**, or **3-O-Ac-SP** and A β , binding studies were conducted by

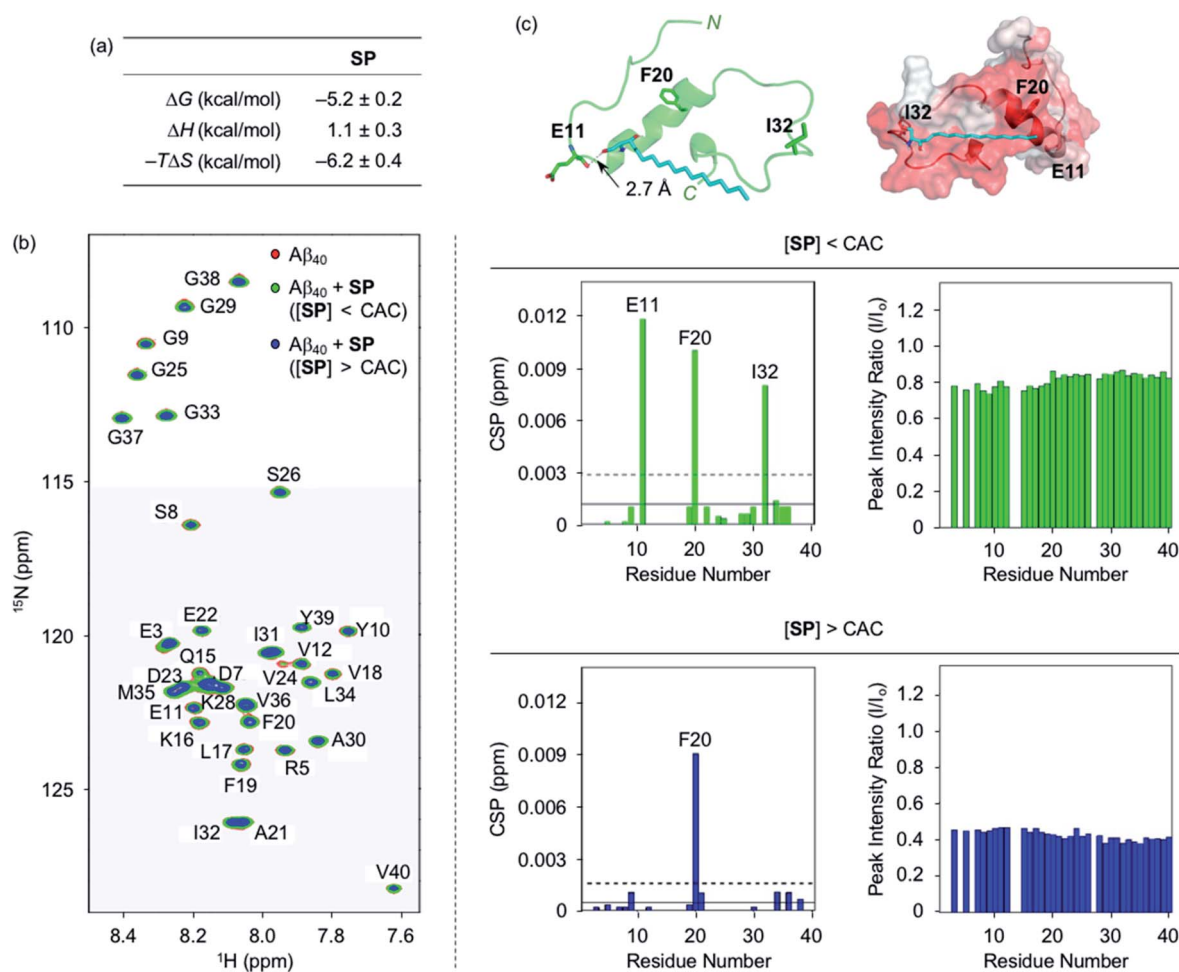


Fig. 2 Interactions of **SP** with A β . (a) Thermodynamic parameters for binding of **SP** to A β_{40} determined by ITC. ITC data are shown in Fig. S3.† Conditions: [A β_{40}] = 500 μ M; [SP] = 200 μ M (1% v/v DMSO); 20 mM HEPES, pH 7.4; 10 $^\circ$ C. (b) 2D ^1H - ^{15}N SOFAST-HMQC NMR analysis of the samples containing **SP** and ^{15}N -labeled A β_{40} . The spectra (left), CSPs (middle), and peak intensity ratios (right) of ^{15}N -labeled A β_{40} were acquired upon addition of **SP** at the concentrations below or above the CAC. Two horizontal lines represent the average chemical shift (solid line) plus one standard deviation (dashed line). Conditions: [^{15}N -labeled A β_{40}] = 40 μ M; [SP] = 2 μ M (below the CAC) or 100 μ M (above the CAC) (1% v/v DMSO); 20 mM HEPES, pH 7.4; 10% v/v D $_2\text{O}$; 10 $^\circ$ C. (c) Possible interactions between **SP** (cyan) and A β_{40} monomer (PDB ID: 2LFM)³⁵ visualized by docking studies [cartoon (left) and surface (right) versions]. The amino acid residues in A β_{40} with relatively large CSPs observed from the NMR studies are depicted in stick representation (left). The dashed line indicates possible hydrogen bonding between **SP** and A β_{40} monomer within 3.0 Å. Hydrophobic contacts of **SP** onto the self-recognition site and the C-terminal region of A β_{40} monomer are observed (right). Hydrophilic to hydrophobic amino acid residues are indicated in a gradient from white to red.



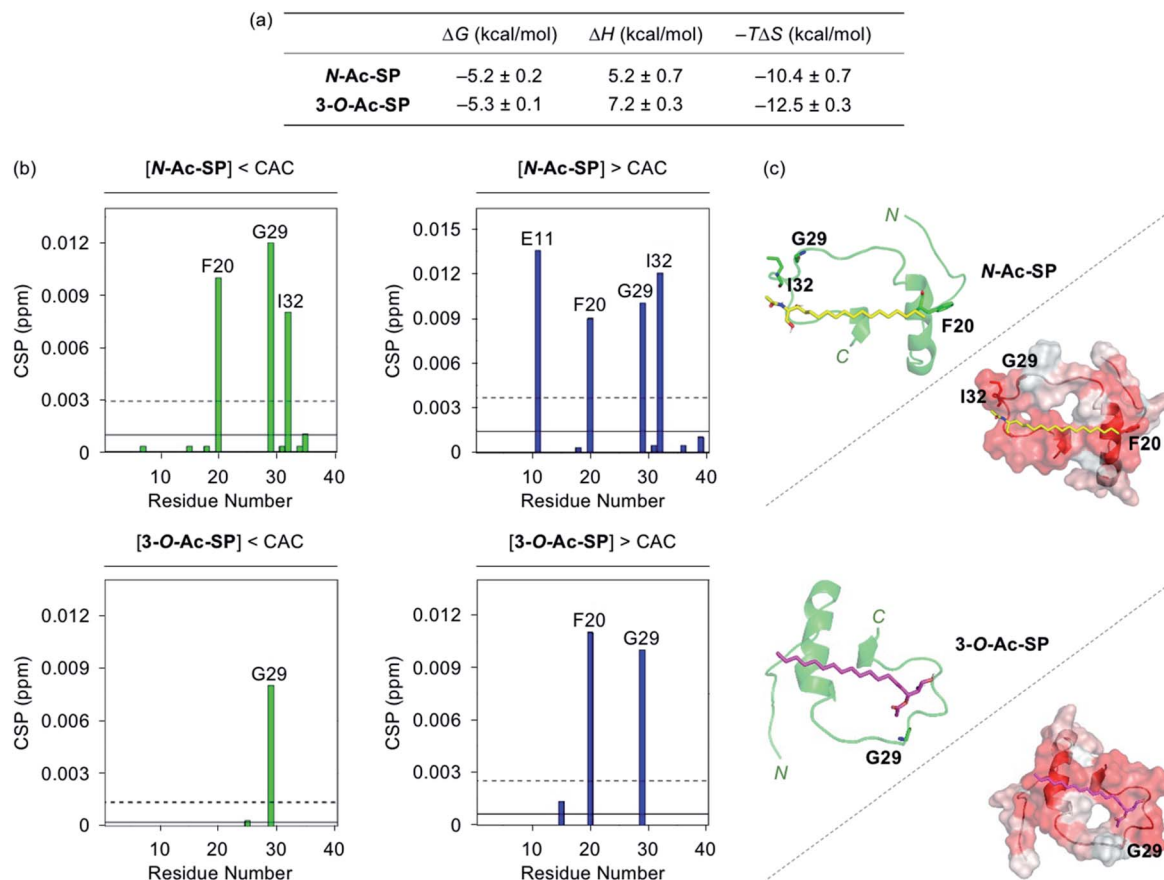


Fig. 3 Interactions of *N*-Ac-SP and 3-*O*-Ac-SP with A β monomer. (a) Thermodynamic parameters for binding of *N*-Ac-SP and 3-*O*-Ac-SP to A β ₄₀ measured by ITC. ITC data are shown in Fig. S4.† Conditions: [A β ₄₀] = 500 μ M; [acetylsphingosine] = 200 μ M (1% v/v DMSO); 20 mM HEPES, pH 7.4; 10 °C. (b) CSPs in A β ₄₀ upon addition of *N*-Ac-SP and 3-*O*-Ac-SP at the concentrations below or above their CACs obtained from 2D ¹H–¹⁵N SOFAST-HMQC NMR spectra (Fig. S5†). Average CSPs (solid line) are indicated with standard deviations (dashed line). Conditions: [¹⁵N-labeled A β ₄₀] = 40 μ M; [compound] = 2 μ M (below the CAC) or 100 μ M (above the CAC) (1% v/v DMSO); 20 mM HEPES, pH 7.4; 10% v/v D₂O; 10 °C. (c) Possible conformations of *N*-Ac-SP (yellow) and 3-*O*-Ac-SP (pink) docked with A β ₄₀ monomer (PDB ID: 2LFM)³⁵ [cartoon (left) and surface (right) versions]. The amino acid residues in A β ₄₀ with relatively large CSPs are presented in stick representation (left). Hydrophobic contacts of the acetylsphingosines onto the self-recognition site and the C-terminal region of A β ₄₀ monomer are displayed (right). Hydrophilic to hydrophobic amino acid residues are indicated in a gradient from white to red.

preferentially *via* the 2-aminoethanol moiety. The metal-binding properties of the acetylsphingosines were also considered with their potential metal-binding sites (*i.e.*, 1,3-propanediol for *N*-Ac-SP; 2-aminoethanol for 3-*O*-Ac-SP shown in Fig. 1a). *N*-Ac-SP includes the *N*-(2-hydroxyethyl)acetamide group as an additional site for possible metal binding. Unfortunately, the K_d value of this moiety for Zn(II) could not be obtained even with excess concentrations of Zn(II) (data not shown). Therefore, SP and 3-*O*-Ac-SP exhibit a greater Zn(II)-binding ability than *N*-Ac-SP through the 2-aminoethanol group.

Cu(II)-binding properties of 2-aminoethanol were analyzed *via* solution speciation studies with UV-vis variable-pH titrations. Spectrophotometric titrations of 2-aminoethanol (L) in the absence of Cu(II) were first performed to estimate its acidity constant (pK_a), as shown in Fig. S6.† The solution speciation diagram depicts the presence of neutral (L) and monoprotonated (LH) forms of 2-aminoethanol in the pH range from 3 to 11. At pH 6.8, a condition plausibly representing

a physiologically acidotic environment where A β aggregation is suggested to be facilitated in the presence of Cu(II),⁵¹ and at physiological pH (pH 7.4), the LH form of the ligand was predicted to be major. Employing the pK_a value [9.79(0)] of the ligand, solution speciation experiments in the presence of Cu(II) were further carried out to determine the Cu(II)-to-ligand stoichiometry and its binding affinity for Cu(II). As shown in Fig. 4b, Cu(II)-ligand complexes with 1 : 1 and/or 1 : 2 Cu(II)-to-ligand stoichiometry were indicated in the solution speciation diagram at pH 6.8 and pH 7.4. Moreover, the binding affinity of the ligand for Cu(II) was predicted based on the values of stability constants ($\log \beta$) and pCu [$pCu = -\log[Cu_{free}]$, where $[Cu_{free}]$ indicates the concentration of unchelated Cu(II)]. Considering the protonation and complexation for 2-aminoethanol at the given pH and concentrations of the ligand and Cu(II), the value of pCu was obtained as indicative of the relative Cu(II)-binding ability of the ligand under experimental conditions.^{52,53} 2-Aminoethanol exhibited the pCu values as 7.96 and 9.08 at pH 6.8 and pH 7.4, respectively, that are expected to be





Fig. 4 Metal-binding properties of 2-aminoethanol. (a) ^1H NMR spectra (left) and changes in the chemical shifts ($\Delta\delta$; right) of 2-aminoethanol with various concentrations of $\text{Zn}(\text{II})$. Red curves were fitted to the $\Delta\delta$ values of the peaks assigned to the protons at the A and B positions to estimate a binding affinity of the compound for $\text{Zn}(\text{II})$. Error bars indicate the standard error from three independent experiments. Conditions: [2-aminoethanol] = 100 μM ; [Zn(II)] = 0.005, 0.01, 0.05, 0.10, 0.15, 0.32, 0.82, and 1.81 mM; [DSS (2,2-dimethyl-2-silapentane-5-sulfonate sodium salt)] = 10 μM (internal reference); $\text{H}_2\text{O} : \text{D}_2\text{O}$ (9 : 1). (b) Solution speciation studies of $\text{Cu}(\text{II})$ -2-aminoethanol (L) complexes. Top: UV-vis variable-pH titration spectra (left) and solution speciation diagram (right) were obtained by spectrophotometric titrations of $\text{Cu}(\text{II})$ -ligand complexes (F_{Cu} = fraction of species at given pH). Bottom: the values of $\log \beta$ and pCu of $\text{Cu}(\text{II})$ -ligand complexes were summarized in the table. The errors in the last digit are shown in parentheses. Charges are omitted for clarity. Conditions: [2-aminoethanol] = 400 μM ; [Cu(II)] = 200 μM ; room temperature; $I = 0.10$ M NaCl.

higher than pZn at each pH based on Irving-Williams series.⁵⁴ According to previously reported studies, $\text{Cu}(\text{II})$ in the solution of aminosugars and aminoglycosides containing aminoalcohol moieties has been observed to lower the pK_a of hydroxyl groups in the ligands, corroborating the higher binding affinity of 2-aminoethanol for $\text{Cu}(\text{II})$ than $\text{Zn}(\text{II})$.⁵⁵⁻⁵⁷ It should be noted that the values of pCu for the 1,3-propanediol and *N*-(2-hydroxyethyl) acetamide moieties could not be obtained due to their insignificant optical changes upon addition of $\text{Cu}(\text{II})$ (data not shown). Overall, **SP** and **3-O-Ac-SP** can coordinate to $\text{Cu}(\text{II})$ through the 2-aminoethanol functionality. The comparable metal-binding affinities of 2-aminoethanol in **SP** and **3-O-Ac-SP** with those of $\text{A}\beta$ [$K_d = 10^{-9}$ to 10^{-6} M for $\text{Zn}(\text{II})$ - $\text{A}\beta$ at pH 7.4; $K_d = 10^{-6}$ to 10^{-12} M and pCu = ca. 6-12 for $\text{Cu}(\text{II})$ - $\text{A}\beta$ in the pH range from 6.5 to 7.4]^{24,58} support their ability to competitively bind to metal ions with $\text{A}\beta$.

To evaluate whether **SP** containing the 2-aminoethanol group can chelate out the metal ion from metal- $\text{A}\beta$, its interaction with metal- $\text{A}\beta$ was analyzed by 2D SOFAST-HMQC NMR and electron paramagnetic resonance (EPR) spectroscopy (Fig. 5, S7, and S8[†]). As depicted in Fig. 5b, the addition of $\text{Zn}(\text{II})$ into the solution of ^{15}N -labeled $\text{A}\beta_{40}$ led to remarkable changes in the NMR signals of the amino acid residues in the *N*-terminal region. Particularly, the peaks of the amino acid residues close to the $\text{Zn}(\text{II})$ -binding site in $\text{A}\beta$ (Fig. 1b), e.g., R5, D7, S8, G9, E11, and V12, disappeared, indicative of $\text{Zn}(\text{II})$ binding to $\text{A}\beta$.^{59,60} When **SP** at the concentration above its CAC was introduced into the solution of ^{15}N -labeled $\text{A}\beta_{40}$ in the presence of $\text{Zn}(\text{II})$, the peak signals corresponding to most of the aforementioned amino acid residues were restored to certain extents. This suggests that $\text{Zn}(\text{II})$ binding to $\text{A}\beta$ could be partially disrupted by **SP**. *N*-Ac-**SP**, which does not include the 2-aminoethanol moiety, was not able to affect binding of $\text{Zn}(\text{II})$ to $\text{A}\beta$ at the concentration

above its CAC (Fig. S9[†]), as expected from its relatively poor metal-binding ability. It should be noted that the ability of the compounds at concentrations below their CACs to chelate out $\text{Zn}(\text{II})$ from $\text{Zn}(\text{II})$ - $\text{A}\beta$ could not be distinguishably detected under our experimental conditions (*i.e.*, [^{15}N -labeled $\text{A}\beta_{40}$] = 40 μM ; [compound] = 2 μM), as presented in Fig. S10 and S11.[†] Overall, **SP** with its micellar species potentially sequesters $\text{Zn}(\text{II})$ from $\text{Zn}(\text{II})$ - $\text{A}\beta$, which is expected based on the $\text{Zn}(\text{II})$ -binding affinity of 2-aminoethanol in **SP** (Fig. 4a).

The interaction of **SP** with $\text{Cu}(\text{II})$ - $\text{A}\beta$ at the concentration above its CAC was examined by EPR spectroscopy. For the EPR studies, $\text{A}\beta_{16}$, an *N*-terminal fragment of full-length $\text{A}\beta$ that binds to metal ions and shows no significant aggregation,²⁴ was used. Upon treatment of **SP** into the solution of $\text{Cu}(\text{II})$, the shifts in symmetry of the *g*- and *A*-values were observed from axial to isotropic, as depicted in Fig. 5c and S8,[†] suggesting its binding to $\text{Cu}(\text{II})$. This is expected from the $\text{Cu}(\text{II})$ -binding affinity of 2-aminoethanol (Fig. 4b). The sample of $\text{Cu}(\text{II})$ with $\text{A}\beta_{16}$ produced an axial EPR signal at $g_{\perp} = 2.05$ and $g_{\parallel} = 2.28$ with the hyperfine splitting of $A_{\perp} = 6.96$ G and $A_{\parallel} = 167.75$ G (Fig. 5c and S8[†]), indicative of the complexation between $\text{Cu}(\text{II})$ and $\text{A}\beta_{16}$.⁶¹ When **SP** was added into the solution containing $\text{Cu}(\text{II})$ and $\text{A}\beta_{16}$, the EPR spectrum was different from that of the $\text{Cu}(\text{II})$ samples with either **SP** or $\text{A}\beta_{16}$. $\text{Cu}(\text{II})$ binding was further quantified by weighted summation of the normalized spectra of the $\text{Cu}(\text{II})$ samples containing either $\text{A}\beta_{16}$ or **SP** following the previously reported procedure.⁶²⁻⁶⁴ As a result, the EPR spectrum of the $\text{Cu}(\text{II})$ sample treated with both $\text{A}\beta_{16}$ and **SP** exhibited a mixture of $\text{Cu}(\text{II})$ bound to **SP** (10%) and $\text{A}\beta_{16}$ (90%). Thus, the EPR studies demonstrate the ability of **SP** to potentially chelate out $\text{Cu}(\text{II})$ from $\text{Cu}(\text{II})$ - $\text{A}\beta$. In contrast, the EPR spectra for the $\text{Cu}(\text{II})$ samples with *N*-Ac-**SP** in the absence and presence of $\text{A}\beta_{16}$ were almost identical with those of compound-free $\text{Cu}(\text{II})$ samples





Fig. 5 Interactions of SP with metal- $\text{A}\beta$ analyzed by 2D ^1H - ^{15}N SOFAST-HMQC NMR and EPR spectroscopy. (a) 2D NMR spectra of ^{15}N -labeled $\text{A}\beta_{40}$ with and without Zn(II) and SP at the concentration above the CAC. (b) CSPs of the Zn(II) samples in the absence (left) and presence (right) of SP at the concentration above the CAC. Upon addition of Zn(II) to the solution of ^{15}N -labeled $\text{A}\beta_{40}$ with and without SP, the NMR signals of the amino acid residues indicated in asterisks (*) disappeared. Conditions: [^{15}N -labeled $\text{A}\beta_{40}$] = 40 μM ; [Zn(II)] = 40 μM ; [SP] = 100 μM (above the CAC) (1% v/v DMSO); 20 mM HEPES, pH 7.4; 10% v/v D_2O ; 10 $^\circ\text{C}$. (c) EPR spectra of the Cu(II) samples with and without SP and $\text{A}\beta_{16}$. The simulated spectrum (blue), similar to the normalized experimental spectrum of the Cu(II) sample with $\text{A}\beta_{16}$ and SP (black), was obtained by weighted summation of the normalized spectra of the Cu(II) samples with either SP (10%; pink) or $\text{A}\beta_{16}$ (90%; dark yellow). Conditions: [$\text{A}\beta_{16}$] = 100 μM ; [Cu(II)] = 100 μM ; [SP] = 400 μM (above the CAC) (1% v/v DMSO); 20 mM HEPES, pH 6.8, 150 mM NaCl; 37 $^\circ\text{C}$; 24 h; constant agitation. The EPR spectra were collected with the following experimental parameters: microwave frequency, 9.41 GHz; microwave power, 2 mW; modulation frequency, 100 kHz; modulation amplitude, 10 G; time constant, 0.01 ms; conversion time, 60 ms; sweep time, 120 s; number of scan, 4; temperature, 100 K.

with and without $\text{A}\beta_{16}$ (Fig. S8 and S12[†]), indicating that this compound cannot significantly affect Cu(II) coordination to $\text{A}\beta$. These distinct behaviors of the compounds against metal- $\text{A}\beta$ at the concentrations above their CACs are also supported by the previous studies displaying that ionic micelles can induce structural changes of peptides and proteins in a manner affecting their Cu(II)-binding properties.^{65–67} Taken together, as expected from the compounds' metal-binding affinities, the spectroscopic studies manifest that SP potentially sequesters the metal ion from metal- $\text{A}\beta$.

Influence of SP and acetylsphingosines on the aggregation of metal-free and metal-associated $\text{A}\beta$

The effects of SP, *N*-Ac-SP, and 3-*O*-Ac-SP, at concentrations below and above the CACs, on the aggregation of $\text{A}\beta_{40}$ and $\text{A}\beta_{42}$ with and without Zn(II) and Cu(II) were evaluated. Upon incubation of SP with freshly prepared metal-free and metal-treated $\text{A}\beta$, as displayed in Fig. 6, S14, and S15,[†] the size distribution and the morphology of the resultant $\text{A}\beta$ species were analyzed

by gel electrophoresis with western blotting (gel/Western blot) and transmission electron microscopy (TEM), respectively. $\text{A}\beta$ species with molecular weights (MWs) in a range from *ca.* 4 kDa (monomer) to *ca.* 240 kDa (aggregate) can be probed as visible bands in gel/Western blot.⁶⁸ In particular, the smearing bands from compound-treated $\text{A}\beta$ samples, distinct from compound-free $\text{A}\beta$ samples, indicate that the compound can vary $\text{A}\beta$ aggregation. $\text{A}\beta$ aggregates that are too large to penetrate the gel matrix can be visualized by TEM.

At the concentration below the CAC, SP did not significantly affect the aggregation of metal-free $\text{A}\beta_{42}$, as shown in Fig. 6b. In the case of Zn(II)- $\text{A}\beta_{42}$ incubated with SP, however, more intense bands were detected at *ca.* 25 kDa with the MW range larger than *ca.* 50 kDa, relative to compound-untreated Zn(II)- $\text{A}\beta_{42}$. The sample of Cu(II)- $\text{A}\beta_{42}$ treated with SP indicated an increase in the levels of dodecameric or larger $\text{A}\beta_{42}$ species (*ca.* ≥ 50 kDa), compared to that of compound-free Cu(II)- $\text{A}\beta_{42}$. Upon increasing the concentration of SP beyond the CAC, changes in the MW distributions of $\text{A}\beta_{42}$ became more dramatic, showing





Fig. 6 Impact of SP on metal-free and metal-induced $A\beta_{42}$ aggregation. (a) Scheme of $A\beta_{42}$ aggregation experiments. (b) Analysis of the size distribution of the resultant $A\beta_{42}$ species by gel/Western blot using an anti- $A\beta$ antibody (6E10). Lanes: (C) $A\beta_{42} \pm Zn(II)$ or $Cu(II)$; (SP) C + SP. The original gel images are shown in Fig. S13.† (c) Morphologies of the resultant $A\beta_{42}$ aggregates from (b) detected by TEM. Conditions: $[A\beta_{42}] = 25 \mu M$; $[Zn(II)$ or $Cu(II)] = 25 \mu M$; [compound] = $10 \mu M$ (below the CAC) or $100 \mu M$ (above the CAC) (1% v/v DMSO); 20 mM HEPES, pH 7.4 [for metal-free and $Zn(II)$ -containing samples] or pH 6.8 [for $Cu(II)$ -containing samples], 150 mM NaCl; $37^\circ C$; 24 h; constant agitation. Scale bars = 200 nm.

smearing bands at *ca.* 4–240 kDa for metal-free $A\beta_{42}$ and $Cu(II)$ - $A\beta_{42}$. The intensity of the bands from the sample containing $Zn(II)$ - $A\beta_{42}$ and SP was enhanced at *ca.* 4–20 kDa and above *ca.* 50 kDa.

In the TEM studies, as displayed in Fig. 6c, the incubation of SP with metal-free and metal-treated $A\beta_{42}$ resulted in the generation of $A\beta_{42}$ aggregates indicating a mixed morphology with both fibrillary and amorphous characteristics. The overall size of the aggregate species was reduced with the treatment of SP, compared to those of the compound-free samples. At the concentration above the CAC, the SP-induced changes in the sizes of $A\beta_{42}$ aggregates became more prominent, forming remarkably smaller $A\beta$ species. Based on these observations, SP was demonstrated to modulate the aggregation of both metal-free and metal-treated $A\beta_{42}$ and such reactivities were further intensified with its micellar species. The results of gel/Western blot and TEM studies with metal-free $A\beta_{40}$ and metal- $A\beta_{40}$ showed lacking reactivity from the samples incubated with SP at the concentration below the CAC (Fig. S14†). At the concentration above the CAC, SP altered the aggregation of metal-free $A\beta_{40}$ and $Cu(II)$ - $A\beta_{40}$, but it did not significantly affect $Zn(II)$ - $A\beta_{40}$ aggregation. As shown in the ITC and 2D NMR studies (*vide supra*), hydrophobic interactions between SP and $A\beta$ may promote the effect of the compound on the formation of $A\beta$ aggregates. The metal-binding properties of SP could further

support its ability to change the aggregation of metal- $A\beta$. In particular, the capability of SP at concentrations above the CAC to partially sequester the metal ion from metal- $A\beta$ is expected to contribute to its reactivity towards the aggregation of metal- $A\beta$. It is noteworthy that multiple interactions of SP with metal- $A\beta$ at concentrations above the CAC are possible based on the observations and findings through the NMR and EPR studies. For example, the transient ternary complexation between metal- $A\beta$ and SP and the interactions of metal-free and/or metal-bound SP with the hydrophobic core and C-terminal region of $A\beta$ (Fig. S7†) could simultaneously occur, along with metal displacement by SP. These could result in the modulative impact of SP with its micellar species on the aggregation of metal- $A\beta$.

The generation of $A\beta$ aggregates in the presence of *N*-Ac-SP and 3-*O*-Ac-SP was further analyzed (Fig. 7). At the concentrations under the CACs, *N*-Ac-SP produced new bands larger than *ca.* 25 kDa or 50 kDa for the samples of metal-free $A\beta_{42}$ and metal- $A\beta_{42}$, as depicted in Fig. 7b. When 3-*O*-Ac-SP was treated to metal-free $A\beta_{42}$ and metal- $A\beta_{42}$, the amount of aggregates with MW larger than *ca.* 50 kDa was increased, along with that of hexameric species at *ca.* 25 kDa, under metal-free and $Zn(II)$ -present conditions. At the concentrations above the CACs, *N*-Ac-SP noticeably varied the MW distributions larger than *ca.* 25 kDa for metal-free $A\beta_{42}$, 7–25 kDa and above *ca.* 100 kDa for $Zn(II)$ - $A\beta_{42}$, and 7–240 kDa for $Cu(II)$ - $A\beta_{42}$. In the case of 3-*O*-Ac-SP, a new smearing band in the MW range larger than *ca.* 25 kDa was detected from metal-free $A\beta_{42}$, while that with MW larger than *ca.* 50 kDa was observed for metal- $A\beta_{42}$. In particular, the treatment of 3-*O*-Ac-SP with $Cu(II)$ - $A\beta_{42}$ led to the detection of a band at *ca.* 25 kDa. The TEM studies revealed that both *N*-Ac-SP and 3-*O*-Ac-SP at the concentrations below their CACs fostered the production of smaller $A\beta_{42}$ aggregates, compared to compound-untreated metal-free $A\beta_{42}$ and metal- $A\beta_{42}$, as presented in Fig. 7c. Upon addition of the acetyl-sphingosines at the concentrations above their CACs, relative to compound-free peptide samples, smaller aggregates of $A\beta_{42}$ were exhibited with a mixture of morphologies with filamentous or amorphous qualities in the absence and presence of metal ions. As shown in Fig. S15,† the acetylsphingosines did not discernably influence the aggregation of metal-free and metal-treated $A\beta_{40}$. Overall, SP, *N*-Ac-SP, and 3-*O*-Ac-SP are capable of modifying $A\beta$ aggregation in the absence and presence of metal ions to distinct degrees at concentrations below and above their CACs.

Impact of SP and acetylsphingosines on the toxicity induced by metal-free $A\beta$ and metal- $A\beta$ in living cells

Prior to cell studies with metal-free and metal-treated $A\beta$, the cytotoxicity of SP, *N*-Ac-SP, and 3-*O*-Ac-SP was investigated by the MTT assay employing human neuroblastoma SH-SY5Y (5Y) cells [MTT = 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide]. The half-maximal inhibitory concentration (IC_{50}) values for SP and *N*-Ac-SP were determined to be in the micromolar range, as summarized in Fig. S16a.† The IC_{50} value for 3-*O*-Ac-SP could not be obtained under our





Fig. 7 Influence of *N*-Ac-SP and *3-O*-Ac-SP on the aggregation of metal-free $A\beta_{42}$ and metal- $A\beta_{42}$. (a) Scheme of $A\beta_{42}$ aggregation experiments. (b) Analysis of the size distribution of the resultant $A\beta_{42}$ species by gel/Western blot using an anti- $A\beta$ antibody (6E10). Lanes: (C) $A\beta_{42} \pm$ Zn(II) or Cu(II); (1) C + *N*-Ac-SP; (2) C + *3-O*-Ac-SP. The original gel images are shown in Fig. S13.† (c) Morphologies of the resultant $A\beta_{42}$ aggregates from (b) detected by TEM. Conditions: $[A\beta_{42}] = 25 \mu\text{M}$; $[Zn(II)$ or $Cu(II)] = 25 \mu\text{M}$; [acetylsphingosine] = $10 \mu\text{M}$ (below the CAC) or $100 \mu\text{M}$ (above the CAC) (1% v/v DMSO); 20 mM HEPES, pH 7.4 [for metal-free and Zn(II)-containing samples] or pH 6.8 [for Cu(II)-containing samples], 150 mM NaCl; 37 °C; 24 h; constant agitation. Scale bars = 200 nm.



Fig. 8 Effect of SP, *N*-Ac-SP, and *3-O*-Ac-SP on the toxicity induced by metal-free $A\beta_{42}$ or metal- $A\beta_{42}$ in living cells. Cell viability, determined by the MTT assay, was calculated in comparison to that of the cells treated with an equivalent amount of the buffered solution with 0.2% v/v DMSO (20 mM HEPES, pH 7.4 or pH 6.8, 150 mM NaCl). Error bars indicate the standard error from three independent experiments. Conditions: $[A\beta_{42}] = 10 \mu\text{M}$; $[Zn(II)$ or $Cu(II)] = 10 \mu\text{M}$; [compound] = $10 \mu\text{M}$ (0.2% v/v DMSO); 20 mM HEPES, pH 7.4 [for metal-free and Zn(II)-containing samples] or pH 6.8 [for Cu(II)-containing samples], 150 mM NaCl.

experimental conditions due to its relatively less toxicity (*ca.* >55% cell survival up to $100 \mu\text{M}$). Moving forward, to examine the effects of SP and the acetylsphingosines on the cytotoxicity induced by $A\beta_{42}$ in the absence and presence of metal ions, the samples containing $A\beta$ with either Zn(II) or Cu(II), a compound, or both were treated to 5Y cells, as shown in Fig. 8. The treatment of SP with metal-free and metal-treated $A\beta_{42}$ resulted in a further reduction in cell viability by *ca.* 15%. In contrast, when the cells were incubated with *N*-Ac-SP or *3-O*-Ac-SP with metal-free and metal- $A\beta_{42}$, the cell viabilities were not significantly affected. It should be noted that incubation of the compounds with 5Y cells at their concentrations used for cell studies with $A\beta$ presented cell survival rates greater than *ca.* 80% in the absence and presence of Zn(II) or Cu(II) (Fig. S16b†). These results suggest that SP, but not *N*-Ac-SP and *3-O*-Ac-SP, may have a deteriorative impact on the toxicity induced by metal-free and metal-treated $A\beta_{42}$.

Conclusions

As the lipid membrane-mediated perturbations in the properties of $A\beta$ come to the fore, lipidomics is emerging as an aspect of AD prompting the advancement of our knowledge towards the relevancy of lipids in its pathology.^{26,69-74} AD-associated disturbances in the cerebral composition of sphingolipids, along with their neurobiological implications, suggest their potential roles in the progression of the disease.^{21,22} In this



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