Metabolic profiling of Alzheimer’s disease brains

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Alzheimer’s disease (AD) is an irreversible, progressive brain disease and can be definitively diagnosed after death through an examination of senile plaques and neurofibrillary tangles in several brain regions. It is to be expected that changes in the concentration and/or localization of low-molecular-weight molecules are linked to the pathological changes that occur in AD, and determining their identity would provide valuable information regarding AD processes. Here, we propose definitive brain metabolic profiling using ultra-performance liquid chromatography coupled with electrospray time-of-flight mass spectrometry analysis. The acquired data were subjected to principal components analysis to differentiate the frontal and parietal lobes of the AD/Control groups. Significant differences in the levels of spermine and spermidine were identified using S-plot, mass spectra, databases and standards. Based on the investigation of the polyamine metabolite pathway, these data establish that the downstream metabolites of ornithine are increased, potentially implicating ornithine decarboxylase activity in AD pathology.

Alzheimer’s disease (AD) is a type of dementia that causes problems with memory, thinking and behavior among older people. In the AD brain, two abnormal structures called plaques and tangles have been the prime suspects in the severe depression of metabolic mechanisms, killing important nerve cells and impairing higher brain functions. Recently, various researchers compared a wide range of pathophysiological markers between mutation carriers and non-carriers as a function of the parental age at onset in order to evaluate correlates of events such as clinical, cognitive, imaging, and biochemical changes in the large international cohort of AD patients. In this result, the AD process begins more than 30 years before the clinical onset of dementia and is associated with a series of pathophysiological changes that occur over decades in cerebrospinal fluid (CSF) biomarkers related to plaques of amyloid beta (Aβ), tangles of tau protein, brain Aβ deposition and brain metabolism as well as progressive cognitive impairment. Glucose metabolism in particular declines from the first stage of the disease to the onset of the expected symptoms (−30 years). Although we still do not know how the AD process begins and/or continues, it appears likely that a simple amyloid cascade hypothesis does not fit with the reality of the neurogenic etiology. Moreover, because the underlying cause of a diagnosis of dementia varies, it is not easy to identify the mechanism and brain regions involved in causing AD. So far as known, it is the key of AD process that this pathology and metabolic mechanisms may be contributed to the distinct cognitive profile in brain. Therefore, it is critical to identify the link between metabolic processes and the disease pathology that causes the distinct cognitive profile and brain lesions of AD patients. AD can be definitively diagnosed after death through definitive examination of senile plaques and fibrillary tangles in several brain regions using staining procedures such as Gallyas-Braak (GB). It would be very useful to be able to correlate metabolic changes in specific regions of the brain with the actual pathological changes that occur in AD. Monitoring various metabolites in the brain will enable the creation of multilateral framework to determine the pathophysiologic features and processes of the neurogenic etiology.

Metabolic profiling of various AD patients using mass spectrometry (MS) techniques revealed that some low-molecular-weight molecules were decreased/increased in the AD versus control groups. Recently, liquid chromatography coupled with mass spectrometry (LC/MS) lipidomic analysis indicated a fundamental change in the AD brain such that the lipid changes may contribute to pathogenesis. Lipidomic MS used to identify lipid abnormalities showed that stearoyl-CoA desaturase and sphingolipids were elevated in the AD brain. The greater sensitivity and definitive brain metabolic profiling by MS allows for the identification of a larger number of metabolites compared to other biological samples, which is especially useful for fingerprinting the pathogenesis of...
AD. However, the identification of biomarkers through less-invasive techniques can often be achieved through comparison to known metabolites and/or database matching metabolites from easy-to-acquire fluids (plasma and CSF) from AD patients without using genotype, pathology and/or diagnostic data\textsuperscript{11–19}. The dramatic change in various metabolic mechanisms in the AD brain is poorly understood.

In this study, we propose to use novel and definitive brain metabolic profiling using ultra-performance liquid chromatography coupled with electrospray time-of-flight mass spectrometry (UPLC-ESI/TOF/MS) analysis to evaluate the diversity of low-molecular-weight molecules patterns from specific brain regions. The brain tissues used for metabolic profiling were therefore based on reliable backgrounds confirmed through multiple independent methods. Our study represents a major top-down use of definitive brain metabolomics using MS and multivariate data analysis regarding to AD pathology. We could identify specific metabolites such as spermidine (SPD) and spermine (SPM), and we discuss the metabolic mechanism of polyamine in specific brain region in the establishment of AD.

**Results**

**Pathological examination of brain samples.** The descriptive diagnosis of all study variables is summarized in Table 1. In addition, the apolipoprotein-E (ApoE) genotype was 3/3 in all AD patients. In the AD group, we indicated the Braak Stage of AD in the post-mortem pathological diagnosis and the duration of disease ranged from 6 to 20 years pre-mortem. In the pathological examination of the AD brain, we assessed the functional properties of the visual pathology by determining the presence of plaques/tangles in brain tissues (Fig. 1). The plaques/tangles pattern of GB staining from AD/Control patients allowed each brain to be used in our comparison. Brain regions such as the frontal lobe (FL), parietal lobe (PL) and occipital lobe (OL) showed a pronounced propensity for GB staining properties in the AD/Control groups. Based on these results, we categorized the AD and Control groups for definitive brain metabolic profiles of low-molecular-weight molecules.

**UPLC-ESI/TOF/MS analysis of metabolites in brain tissues.** We propose that MS metabolic profiling of AD pathology can be used to changes in brain metabolites related to the presence of plaques and tangles. Our previous protocol was applied for UPLC-ESI/TOF/MS molecular-weight phase, the detected m/z values in MS data collection with a scan mode (positive/negative ionizations) from m/z 100–1000 using two different columns (T3-C18 and HS-F5) for pathologically confirmed AD/Control brain tissues\textsuperscript{20,21}. Samples were prepared by centrifugal ultrafiltration to separate the low-molecular-weight phase for UPLC-ESI/TOF/MS profile analysis and to remove the protein phase (> 3,000 Da). Using these low-molecular-weight phases, the detected m/z values in MS

![Figure 1](https://www.nature.com/scientificreports)
chromatograms ranged from 1201 to 4699 compounds on positive and 385 to 1110 compounds on negative ionizations in all brain tissues (Fig. S1). These peak responses of low-molecular-weight molecules were extracted based on a 100 counts cutoff (marker intensity threshold) because of the dependability statistics. These raw data were analyzed for peak detection and alignment and then exported for a principal components analysis (PCA) and orthogonal partial least-squares-discriminant analysis (OPLS-DA).

For validation of UPLC-ESI/TOF/MS analysis, the pooled parietal lobe tissues from AD (n = 10) and Control (n = 10) were performed on the same day for determining the intra-day accuracy, replicate (n = 6) analytes of various peaks. The intra-day accuracy was expressed as the relative standard deviation (RSD, %) of retention time (0.08 – 1.63%) and peak area (1.44 – 8.93%) for typical m/z values (Table S1). For multivariate data analysis, each sample of real patients was used for single running based on this validation.

**Multivariate statistical analysis of metabolites in brain tissues.**

PCA for all samples based on the AD and Control groups revealed an interesting distribution in specific brain regions. The representative PCA score-plots of FL, PL and OL tissues after positive ionization with the T3-C18 UPLC column are shown in Fig. 2a–c. All PCA score-plots are shown in Fig. S2. The FL and PL of the AD group converged on a particular set. On the other hand, the Control groups in all regions indicated a disproportionate lack of coherence in the PCA score-plot patterns of detected m/z values. In the OL region, the distributions were rarely different between the AD and Control groups. One feature that these different metabolites share is their contribution to the PCA score-plots with integrated m/z values, which indicates that they are very important metabolites for fingerprinting the pathogenesis of AD. Thus, OPLS-DA was used to differentiate between the AD and Control groups in each tissue (n = 10 for each sample) (Fig. 2d–f). Each point represents the metabolite, m/z and retention time pair; the X-axis represents variable contribution, and the further this pair point departs from zero, the more the ion contributes to the difference between the two groups of AD and Control; the Y-axis represents variable confidence, and the further the metabolite, m/z and retention time pair departs from zero, the higher the confidence level of the ion is for the difference between the two groups of AD and Control. Hence, in this study, this pair points at the two ends of “S” represent those components contributing most to the difference between the two groups with most confidence, which could be regarded as the most differentiating components between AD and Control. For the extraction of potential markers of group separation, the cut-off of variable confidence was approximately 0.8 for the increased markers and ~0.8 for the decreased markers in S-plot. Thus, in this study, the S-plot showed that the quality differentiation is employed by the decreased and increased m/z values with a correlation of rigorous ± 0.85. In these S-plots, the increased (red-box) and decreased (blue-box) peaks show the important metabolites in the FL and PL regions. This result showed that a total of 431 peaks were analyzed to identify possible biomarker candidates based on all brain regions (Table 2). In the OPLS-DA scores plot (Fig. S3) using Pareto scaling with mean centering, each coordinate represents a brain region, and it could be observed that the determined brain regions are clearly divided into two clusters. All the observations were located within the Hotelling T2 (0.95) ellipse, which confirmed the fact the metabolite difference exists between AD and Control. In addition, univariate analysis was shown in Tables S2 and S3. From the results of the S-plot, specific peaks were extracted from the MS chromatograms of the biomarker candidates in the FL and PL brain regions. We identified five metabolites in both the FL and PL regions based on the shape of the peaks off the T3-C18 column as a common characteristic of all patients, which could be used as possible biomarkers candidates.

Based on the available metabolomics databases, a list of retention times, m/z values, elemental compositions and possible biomarkers candidates is presented in Table 3. The m/z 146.16 and m/z 203.22 peaks were speculated to represent SPD and SPM in the AD group. Additionally, these peaks were confirmed by comparison to known standards using UPLC-ESI/TOF/MS chromatograms and spectra (Fig. S4). These results indicated that the m/z 146.16 and m/z 203.22 peaks were SPD and SPM based on metabolomics databases and standard matching. SPD and SPM are very interesting polyamines with regard to memory-associated brain structures.

**UPLC-ESI/MS/MS analysis of polyamines in brain tissues.**

The analytical determination of biogenic polyamines such as SPD and SPM is not a simple task due to their structure and because they are usually present at low levels in complex matrices. Furthermore, biogenic polyamines do not exhibit a satisfactory absorption at visible, ultraviolet and fluorescent wavelengths. In previous reports, a chemical derivatization was applied for assaying polyamines in biological samples using UPLC-ESI/tandem MS (UPLC-ESI/MS/MS)14. Thus, these polyamines were analyzed using sensitive and selective UPLC-ESI/MS/MS to determine the metabolic mechanism in all brain tissues. UPLC-MS/MS with the derivatization assay was used for the analysis of biogenic polyamines such as SPD, SPM, ornithine (ORN), putrescine (PUT), acetyl-spermidine (Ac-SPD) and acetyl-spermine (Ac-SPM) in brain tissues. Selected reaction monitoring (SRM) chromatograms of the polyamines in the FL region of AD are shown in Fig. 3a. The detection levels of SPD and SPM in the FL and PL regions of the AD samples were increased compared with the OL region, and the difference was statistically significant (Fig. 3b–d). Moreover, PUT, Ac-SPD and Ac-SPM were increased in the FL region of the AD group (Fig. 3b). However, ORN remained at approximately the same levels between the AD and Control groups.

**Metabolite pathway of polyamines.** To identify the metabolic pathway of SPD and SPM, web-based databases and functional pathway analysis were used for this profiling. The primary pathway was the ornithine decarboxylation system, and the secondary was the SPD and SPM biosynthesis/metabolic pathways (Fig. 4). Location-based metabolite sets for SPD and SPM were found mainly in this system. The detailed construction of the altered polyamine metabolism pathways with a higher score was generated using KEGG as a reference map.

**Discussion**

In the present study, we focused on the definitive brain metabolic profiles of low-molecular-weight molecules in specific brain regions with regard to the GB staining properties of plaques/tangles based on samples from reliable AD backgrounds. This novel and accurate UPLC-ESI/TOF/MS of low-molecular-weight molecules allowed for the discovery of the interesting biomarkers SPD and SPM in specific regions of the AD and Control brain tissues. Thus, the use of sensitive and selective UPLC-ESI/MS/MS with a derivatization could allow us to determine the levels of polyamines such as SPD, SPM, ORN, PUT, Ac-SPD and Ac-SPM, in specific brain regions related to AD pathology. Based on our data, the polyamine metabolite pathway in a specific brain region is likely to be responsible for up-regulated polyamine metabolism and elevated ornithine decarboxylase (ODC) activity (Fig. 4). In addition, our results support the hypothesis that N-methyl-D-aspartate (NMDA) receptor excitotoxicity is caused by excess SPD and SPM, and that ODC activity was stimulated by the effects of AD processes in localized brain regions. Our approach overcomes the limitations of previous pathological examinations using antibody-targeted analytes and stained plaques and tangles in biological samples from AD patients.
Table 2 | A total of 431 peaks representing possible biomarker candidates based on all brain regions

| Ionization | Positive mode | Negative mode |
|------------|---------------|---------------|
|            | T3-C18 | HS-F5 | T3-C18 | HS-F5 |
| Column     | Increase | Decrease | Increase | Decrease |
| FL         | 65 | 62 | 57 | 33 |
| PL         | 56 | 11 | 32 | 33 |
| OL         | 0 | 0 | 16 | 3 |

Figure 2 | PCA score-plots and S-plots between AD and Control groups in each brain region (n = 10 for each sample). (a), PCA score-plot of frontal lobe region (explained variance (R2): component 1, 0.32 and component 2, 0.49, and predictive ability (Q2): component 1, 0.22 and component 0.21). (b), PCA score-plot of parietal lobe region (R2: component 1, 0.3 and component 2, 0.45, and Q2: component 1, 0.21 and component 2, 0.17). (c), PCA score-plot of occipital lobe region (R2: component 1, 0.18 and component 2, 0.28, and Q2: component 1, 0.08 and component 2, 0.03). (d), S-plot of frontal lobe region. (e), S-plot of parietal lobe region. (f), S-plot of occipital lobe region.
Metabolomics has substantially advanced our understanding of the analytical features that constitute part of AD research. Several studies have proposed or performed metabolic profiling of low-molecular-weight molecules in biological AD samples, such as CSF and plasma based on MS techniques. In these previous studies, potential biomarkers have revealed insights into the presence of sphingomyelin, ceramide and desmosterol in AD plasma. A peripheral plasma biomarker would offer significant advantages over

Table 3 | Possible biomarkers candidates for AD from increased m/z values based on the metabolomics databases

| Brain region | Possible biomarkers | Ret. Time | m/z  | Elemental Composition (mDa, i-FIT) |
|--------------|--------------------|-----------|------|-----------------------------------|
| FL Spermidine | 0.6022 146.1656 | −0.1, 0.0, C_{19}H_{32}N_{3} |
| PL Spermidine | 0.6046 146.1658 | 0.1, 0.0, C_{19}H_{32}N_{3} |
| FL Spermine  | 0.618 203.2221  | −1.5, 0.0, C_{19}H_{32}N_{4} |
| PL Spermine  | 0.5938 203.2226  | −1.0, 0.0, C_{19}H_{32}N_{4} |
| FL 3-[(R)-Dimethylamino]([1R,2R]-2-hydroxycyclohexyl)methyl]phenol | 8.5688 250.1775 | −3.2, 0.8, C_{13}H_{20}NO |
| PL 3-[(R)-Dimethylamino]([1R,2R]-2-hydroxycyclohexyl)methyl]phenol | 8.202 250.1796 | −1.1, 1.8, C_{13}H_{20}NO |
| FL Unknown   | 10.8745 343.2235 | 2.9, 1.3, C_{22}H_{31}O_{3} |
| PL 3-[(R)-(Dimethylamino)[(1R,2R)-2-hydroxycyclohexyl]methyl]phenol | 8.742 250.1796 | 2.9, 1.3, C_{22}H_{31}O_{3} |
| FL Unknown   | 5.5884 416.2824 | 3.0, 1.4, C_{17}H_{22}N_{3}O_{6}S |
| PL 10.4778 343.2286 | 1.3, 3.5, C_{22}H_{31}O_{3} |

Bottom note: Both FL and PL regions based on the shape of each peak using T3-C18 column as a common characteristic of all patients.

Figure 3 | UPLC-ESI/MS/MS with derivatization for the analysis of biogenic polyamine in the brain. (a), SRM chromatograms of polyamine in AD frontal lobe (No. 11). (b), Concentration levels of polyamine in frontal lobe (n = 10 for each sample). (c), Concentration levels of polyamine in occipital lobe (n = 10 for each sample). * P < 0.05, ** P < 0.01, *** P < 0.005.
the first in a novel class of AD medications that act on the glutamatergic system, and it has been shown to have a modest effect in moderate-to-severe disease. Dysfunction of the NMDA receptor, manifested as neuronal excitotoxicity, is hypothesized to be involved in the etiology of AD. The balance between these mechanisms within specific neuronal networks may determine whether the activity of the NMDA receptor increases or decreases Aβ in vivo. Aβ pathology may drive an abnormal conformation of the NMDA receptor or deleteriously enhance the association of the NMDA receptor with certain molecules. For example, the NMDA subunit could be stimulated by SPM in the dimeric pairs of NR1 and NR2, and NR1 and NR2A. The specific polyamine that binds to the NMDA receptor may contribute more to local variability in brain injury than the glutamate site, and it has the potential to ameliorate excitotoxicity in specific AD-related brain regions. Thus, our discovery of an abnormal increase in the localized concentration of SPM and SPM indicated that over-oxidation mediated by specific NMDA receptors might contribute to localized focal brain in AD.

The effects of abnormal increases in the concentration of neural polyamines are not clear. In this discussion, we focus on the apparent cause-and-effect chain of abnormal elevated polyamines in the AD brain region. For example, Aβ peptides have been shown to be responsible for up-regulated polyamine metabolism, specifically, increased polyamine uptake and elevated ODC activity. In addition, Yatin et al. reported that Aβ causes an increase in polyamine metabolism manifested by up-regulated SPM uptake and increased ODC activity and that up-regulation of polyamine metabolism due to Aβ is a result of enhanced neurotoxicity. These extraordinary behaviors of Aβ plaque deposition, polyamine metabolism, ODC activity and NMDA receptor excitotoxicity occurred in localized AD brain regions. Fig. 4 shows the metabolism of polyamines from a web-based database and functional pathway analysis. PUT is synthesized from ORN by ODC. SPD is then synthesized from PUT, and SPM from SPD, by transfer of the aminopropyl moiety of decarboxylated S-adenosylmethionine. The enzymes catalyzing these reactions are spermidine synthase (SPDS) and spermine synthase (SPMS). Different enzymes convert SPD to SPM, and SPM to PUT. Spermine oxidase (SMO) directly catalyzes the conversion of SPM to SPD, and both spermidine/spermine N-acetyltransferase (SSAT) and acetylpyridoxal polyamine oxidase (PAO) catalyze the conversion of AcSPM to SPD and Ac-APD to PUT. To investigate the regulation of the polyamine content, the concentrations of SPD, SPM, ORN, PUT, Ac-SPD and Ac-SPM in brain tissues were analyzed by UPLC-ESI/MS/MS. In previous reports, a chemical derivatization was applied for assaying polyamines in biological samples using sensitive and selective UPLC-ESI/MS/MS to determine the metabolic mechanism. UPLC-ESI/MS/MS with the derivatization assay was used for the analysis of biogenic polyamines such as SPD, SPM, ORN, PUT, Ac-SPD and Ac-SPM without a change of ORN are associated with ODC activity in the AD processes (Fig. 4). Thus, one theory posits that the state-dependency effect of excess SPD and SPM in focal brain would be adverse impact of NMDA receptor.

Methods

Materials. Polyamine standards such as SPD, SPM, ORN, PUT, Ac-SPD and Ac-SPM were purchased from Wako Pure Chemical Co. (Osaka, Japan), Kanto Chemical Co. (Tokyo, Japan) and Sigma-Aldrich Co. (St. Louis, MO), 1,6-Diaminohexane (DAH) and 4-N,N-Dimethylaminosulfonyl)-7-fluoro-2,1,3-benzoxadiazole (DBD-F) were obtained from Tokyo Chemical Industry Co. (Tokyo, Japan). DAH was used as the internal standard. Organic solvents, sodium tetraborate (Borax) and 1 mol/L Tris-HCl (pH 7.5) were purchased from Wako Pure Chemical Co. (Osaka, Japan). Formic acid (FA), methanol and acetonitrile of LC-MS grade were obtained from Kanto Chemical Co. (Tokyo, Japan). All other chemicals were of analytical grade and were used without further purification. De-ionized and distilled water was used throughout the study (Aquarius PWU 200 automatic water distillation apparatus, Advancec, Tokyo, Japan).

Autopsy and sampling of brain tissues. The utilized tissues from the Fukushima Brain Bank were used for the accurate, reliable and detailed pathological evaluation of
The pathological conference 62. Senile plaques were confirmed using criteria of the monoclonal clone pSyn#64. 1
plaques, an anti-human pTau antibody (INNOGENETICS, clone AT8), and amyloid antibody (Dako, mouse monoclonal, clone pSUN641). 1
We obtained written informed consent for autopsy from the patients’ guardians as well as their permission to use the results of diagnosis, research, and genetic analysis at the Choyu Medical Institute Fukushima Hospital. The brain was removed at autopsy, weighed, cut mid sagitally and examined for vascular and other macroscopically detectable lesions. Specimens for diagnostic examination were taken from the hemisphere that showed abnormalities on CT scanning or from the left hemisphere if no difference between the left and the right hemisphere was found by CT. Specimens were separated into the cerebrum, cerebellum, midbrain, pons and medulla and dissected into slices. These slices were fixed in buffered 10% formalin as a hemispheric block. The other brain regions were also separated, and stored at –80°C until use and used after gaining approval from authorities including the University of Shizuoka ethics panel.

GB staining was carried out to screen for senile plaques and neurofibrillary tangles in several brain regions. 62-64 For routine immunohistochemistry, an anti-human β-amyloid antibody (Dako, mouse monoclonal, 1:30) was used for detecting senile plaques, an anti-human pTau antibody (INNOGENETICS, clone AT8), and an anti-phosphorylated t-tau antibody (Wako, mouse monoclonal clone pSUN641) for detecting Lewy bodies and neuritis.

All of our cases were reviewed and discussed with several doctors at a clinicopathological conference. 65 Senile plaques were confirmed using criteria of the Consortium to Establish a Registry for Alzheimer’s Disease (CERAD) and Braak and Braak staging, and a grade higher than C was selected as indicative of AD. 66 The ApoE genotype was determined and correlated with neuropathological findings. DNA was extracted from brain tissues by the phenol-chloroform method. Peripheral blood of the elderly in the PBC group was collected in tubes containing EDTA, and DNA was extracted using a QIAamp DNA Blood kit (Qiagen, Valencia, CA) and stored at 4°C. ApoE genotyping was carried out by the polymerase chain reaction (PCR)-restriction fragment length polymorphism (RFLP) method. 67

Sample preparation of low-molecular-weight molecules in brain tissues. These supernatant solutions were applied to centrifugal ultrafiltration using Amicon Ultra-0.5 mL 3 K (Ultracel-3K, regenerated cellulose 3,000 M.W., Millipore Co. Ltd., Billerica, MA). In the first step, the brain tissues and two biacARB beads (3.0 mm) in the tubes were placed in the holes of an aluminum block and immediately homogenized for 3 min by a Shake Master (Bio Medical Sciences, Tokyo, Japan). In the second step, the homogenized tissues were weighed in a tube, and 500 μL of the extraction solvent, i.e., the mixture of chloroform and chloroform:HCl buffer (50 mM, pH 7.4) (4:1, v/v), was added as the reaction volume (at the ratio of 100 mg/mL). For the third step, the extraction solution and two zirconia beads (3.0 mm) in the tubes were homogenized for 3 min by a Shake Master again. Finally, these supernatant solutions were applied to centrifugal ultrafiltration using Amicon Ultra-0.5 mL 3 K. A 5 μL aliquot of the filtrate was then subjected to the UPLC-ESI/TOF/MS system.

Validation of UPLC-ESI/TOF/MS system. The quality control in the pooled parietal lobes tissues from AD (n = 10) and Control (n = 10) were performed on the same day for determining the intra-day accuracy, replicate (n = 6) analyses of various peaks. The intra-day accuracy was expressed as the relative standard error (RSD %) of retention time and peak area for typical m/z values (r). For multivariate data analysis, each sample of real patients was obtained from single running based on this validation.

Multivariate data analysis of m/z values. The UPLC-ESI/TOF/MS data were analyzed for peak detection and alignment from m/z 100 to 1000, and exported for PCA and OPLS-DA by MarkerLynxTM XS V4.1 SCN803 (Waters Co., Milford, MA).

Identification of low-molecular-weight molecules by database comparison. The elemental compositions of the low-molecular-weight molecules on the S-plot were identified based on the mass and retention time, and the values of mDa (the difference from the exact mass) and i-FIT (the correctness of isotope patterns of elemental composition; the lower i-FIT normalized values mean high) of each compound. Low-molecular-weight molecules with good mDa values and i-FIT levels were extracted from these prospective formulas and the MS spectra of the unknowns were matched to standard model compounds. The MarkerLynxTM XS V4.1 SCN803 combined lists of biomarkers candidates extracted from the following five metabolomics databases: NIST (http://www.nist.gov/pml/data/asd.cfm), Massbank (http://www.massbank.jp), KEGG (http://www.kegg.jp/), BioCyc (http://biocyc.org/), and the human metabolute database (http://www.vhmdb.org). A mass tolerance of 0.5 mDa was set as well as a maximum elemental composition of C = 500, H = 1000, N = 200, O = 200, S = 10, P = 10, and Cl = 10. The software automatically filters compounds from different libraries that have the same KEGG.

Analysis of polyamine in brain tissues. A UPLC-ESI/MS/MS was used for measuring polyamines as derivatives in brain tissues. A 20-μL extraction solution of brain tissues using the above protocol was added to 20 μL of acetonitrile containing the internal standard (DHA 500 nmol/L). Then, the mix solution was dried using a Personal Evaporator (EZ-2, Genevac Co., NY). The residues was added to 150 μL of D2O and acetonitrile (40 mMol/L) and 150 μL of 100 mmol/L Borax (pH 9.3) at 60°C for 30 min, and then UPLC-ESI/MS/MS was performed. The UPLC system was used a Waters ACQUITY UPLC (Waters, Milford, MA, USA). The UPLC separation was achieved using a Waters ACQUITY UPLC BEH C18 (2.1 × 100 mm, 1.7 μm; Waters, Milford, MA) maintained at 40°C and the mobile phase consisted of 0.1% FA in water (solvent A) and 0.1% FA in acetonitrile (solvent B). The gradient was as follows: 8%→20-60-90 (0-8-10 min), with a flow rate of 0.4 mL/min. The injection volume was 5 μL. The separated compounds were detected using a Waters Xevo™ TQ-S triple quadrupole mass spectrometer (Waters, MA). The mass spectrometer was operated with an electrospray source in the positive ionization mode. The source conditions were a capillary voltage of 3.0 kV, source offset of 0 V, collision gas and was regulated at 0.15 mL/min. The LH resolution 1, HF resolution 1, and DM resolution 1 were 78, 54, and 150, respectively. Argon was used as the collision gas and was regulated at 0.15 mL/min. The LH resolution 1, HM resolution 1, and DM resolution 1 were 75, 35, and 150, respectively. The separation, detection conditions and monitoring ions are the same as those listed in Table S4.

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