Original article

Decreased plasmatic spermidine and increased spermine in mild cognitive impairment and Alzheimer’s disease patients

Helena P. G. Joaquim*1,2
Alana C. Costa1,2
Orestes V. Forlenza1,2
Wagner F. Gattaz1,2
Leda L. Talib1,2

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Abstract

Background: Current evidence suggests that upregulation of polyamines system plays a role both in cognitive deficit and synaptic loss observed in Alzheimer’s disease (AD). Objective: The aim of this study was to determine the plasmatic concentration of polyamines in mild cognitive impairment (MCI) and AD patients in comparison with healthy controls (HC). Methods: Plasmatic polyamines were quantified using the AbsoluteIDQ® p180 and liquid chromatography coupled to tandem mass spectrometry (LC/MS-MS). Results: The study group comprised 34 AD patients, 20 MCI and 25 HC. All individuals were followed for 4 years. During this period 8 amnestic MCI patients (40% of the MCI sample at baseline) converted to AD. Spermidine level was lower in both patient groups (AD; MCI) compared to HC (p = 0.007). Plasma levels of spermine were higher in the MCI group (p < 0.001), but decreased in the sub-sample of MCI patients who converted to AD (p = 0.043). No statistically significant differences were found in ornithine and putrescine levels (p = 0.056 and p = 0.126, respectively). Discussion: Our results suggest dynamic changes in the expression of polyamines in the MCI-AD continuum.

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Keywords: Biomarkers, Alzheimer’s, mild cognitive impairment, polyamines, plasma.

Introduction

The clinical diagnosis of Alzheimer’s disease (AD) is a probabilistic definition that takes into account the patient’s complaints, the objective characterization of cognitive impairment, family history of dementia, neuropsychiatric examination, laboratory tests and brain imaging1-2. Brain imaging and cerebrospinal fluid (CSF) levels of beta-amyloid peptide 42 (Aβ42), Tau and phosphorylated Tau can help in differential diagnosis of AD and can be useful for predicting AD in individuals with mild cognitive impairment (MCI)3-5. The search for biological markers in AD and related disorders is important for a better understanding of the pathophysiological process, and the clinical translation of this knowledge can provide support to the diagnostic workup and the prediction of dementia in patients with mild cognitive symptoms. New technologies have raised the possibility of using biological markers to reinforce the diagnosis of probable AD at early stages of the disease process5-6, addressing in vivo molecules related to core pathogenic mechanisms of AD and to surrogate markers of the neurodegenerative process, i.e., related to the formation of neuritic plaques and neurofibrillary tangles. Neuritic plaques are extracellular lesions and their main constituent is the amyloid-β42 peptide (Aβ42). Neurofibrillary tangles are intracellular lesions and are mostly composed of hyperphosphorylated Tau protein7-9. Both are commonly referred to as ‘molecular’ markers of the disease process9-13,14,15.

Polyamines (PAs) are low-molecular-weight organic compounds that perform important functions in cell growth and proliferation, gene transcription, protein synthesis, conformation of nucleic acid and apoptosis15-17. The biosynthesis of natural PA initiates from two amino acids, arginine (Arg) and ornithine (Orn). Orn is decarboxylated by ornithine decarboxylase (ODC), the limiting enzyme in the PA synthesis, first to putrescine, which is the immediate precursor for the synthesis of the spermidine and spermine. In mammalian cells, the main natural PAs are putrescine, spermidine and spermine17, that are localized inside and outside of cells, as well as in the plasma membrane (Figure 1)18-20.

The effects of abnormal increases in the concentration of neural PAs are not clear. Nevertheless, current evidence suggests that upregulation of PA system plays a role in both cognitive deficit and synaptic loss observed in AD21. It appears that PAs and Aβ have several potential interaction sites in vivo, particularly in cellular organelles, in the cytoplasm and inside neuronal cells22. Aβ peptides have been shown to be responsible for upregulated PA metabolism, specifically, increased PA uptake and high ODC activity23-25. Also, PAs are critically involved in microtubule assembly and stabilization26-28. Recently, Vemula et al.24 determined the time course of the arginine metabolic profile changes in the frontal cortex and hippocampus (among others) when TAU synthesis is increased. They speculate a shift of L-arginine metabolism to favor the polyamine as a protective mechanism. In addition, spermidine and spermine have the ability to modulate several ion channels in the brain specifically related to the N-methyl-D-aspartate (NMDA) receptor29-30. Dysfunction of the NMDA receptor, established as neuronal excitotoxicity is hypothesized to be involved in the etiology of AD27,28.
Exploring the role of polyamines in the pathophysiology of AD and its prodromal phase MCI represents an alternative to developing biomarkers. Thus, the aim of this study was to determine the plasmatic concentration of PAs in MCI and AD patients compared to healthy controls.

Material and methods

Subjects

Eligible subjects were recruited from a cohort of community-dwelling older adults attending a psychogeriatric outpatient facility at the Institute of Psychiatry, Faculty of Medicine, University of São Paulo, Brazil. Participants, mostly from the hospital catchment area, were either outpatient undergoing treatment for cognitive disorders or healthy volunteers in a longitudinal follow-up study conducted at the memory clinic. Seventy nine subjects were enrolled in this study: 34 AD, 20 amnestic MCI and 25 physically and cognitively healthy controls (Table 1). All results were controlled for years of education and gender because of statistical differences.

Table 1. Socio-demographic characteristics of patients and controls

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>AD (n=34)</th>
<th>MCI (n=20)</th>
<th>HC (n=25)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender (M/F)</td>
<td>10/24</td>
<td>3/17</td>
<td>7/18</td>
<td>0.47</td>
</tr>
<tr>
<td>Age (mean ± sd)</td>
<td>75.0 ± 6.7</td>
<td>73.9 ± 6.2</td>
<td>74.4 ± 5.8</td>
<td>0.81</td>
</tr>
<tr>
<td>Years of education (mean ± sd)</td>
<td>6.2 ± 3.8</td>
<td>7.9 ± 4.4</td>
<td>13.2 ± 5.7</td>
<td>0.01</td>
</tr>
<tr>
<td>MMSE (mean ± sd)</td>
<td>19.0 ± 5.0</td>
<td>27.0 ± 2.0</td>
<td>29.0 ± 1.0</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>CAMCOG (mean ± sd)</td>
<td>57.0 ± 18.0</td>
<td>86.0 ± 9.0</td>
<td>95.0 ± 6.0</td>
<td>&lt;0.01</td>
</tr>
</tbody>
</table>

M: male; F: female; sd: standard deviation; AD: Alzheimer’s disease; MCI: mild cognitive impairment; HC: healthy control; MMSE: Mini-Mental State Examination; CAMCOG: Cambridge Cognitive Test. Bold values indicate significant p-values.

All individuals were followed during 4 years. During this period, 8 amnestic MCI patients converted to AD (converters MCI-AD) and 12 did not (MCI-NC). These eight MCI patients performed new blood collection after conversion.

The study was approved by the Local Ethics Committee of University of Sao Paulo (CAPPesq nº 943.883) and performed in accordance with the Helsinki declaration. All subjects provided written informed consent prior to inclusion in the study. The sociodemographic characteristics of the patients and controls are summarized in Table 1. All results were controlled for years of education and gender because of statistical differences.

Sample preparation

Blood samples of all subjects were collected in EDTA-coated tubes (Vacutainer, Bencton Dickison) for plasma metabolite determination after 8 hours of fasting. Samples were centrifuged at 20 °C and 1,800g for 15 min and stored at -80 °C until analysis.

Polyamines quantification

We analyzed endogenous metabolites with an AbsoluteIDQ® p180 kit (Biocrates Life Science AG, Innsbruck, Austria), and performed a targeted quantitative and quality controlled assay using liquid chromatography coupled to tandem mass spectrometry (LC-MS/MS). In brief, after the addition of 10 μl if the supplied internal standard solution to each well on a filterspot of the 96-well extraction plate, 10 μl of each plasma sample, quality control samples, blank, zero sample, or calibration standard were added to the appropriate wells. The plate was then dried under a gentle stream of nitrogen. The samples were derivatized with phenyl isothiocyanate (PITC) for the biogenic amines, and dried again. Sample extracts were diluted with either 40% methanol in water for the LC-MS/MS analysis. This technique uses isotope-labelled internal standards and provides quantitative results based on calibration curves and quality control analyses. Low/mid/high level QC samples provided by Biocrates Life Science AG were prepared and analyzed on each plate as recommended by the manufacturer. These QC samples were used for a technical validation of each kit plate. To allow appropriate inter-plate abundance scaling based specifically on this cohort of samples, we generated a Study Pool QC by combining approximately 10 μl from the first 20 samples for analysis. This sample was frozen and analyzed on each plate. This analysis was performed on a triple-quadrupole mass spectrometer (Xevo TQ-S, Waters Corporation, USA), using positive electrospray ionization operating in the Multiple Reaction Monitoring (MRM) mode. MRM transitions (compound-specific precursor to product ion transitions) for each analyte and internal standard were collected over a scheduled retention time window using tune files and acquisition methods provided in the AbsoluteIDQ® p180 kit. The plasma samples were processed according manufacturer instructions. Chromatographic separation of biogenic amines was performed using an ACQUITY UPLC System (Waters Corporation) using an ACQUITY 2.1 mm × 50 mm 1.7 μm BEH C18 column fitted with an ACQUITY BEH C18 1.7 μm VanGuard guard
column, and quantified by calibration curve plotting ratio of analyte to internal standard versus standard concentration, fitted using a linear regression with 1/x weighting. All biogenic amines utilize either deuterated or 13C stable-isotope labeled internal standard of the exact analyte or closely-eluting compound of similar class. MassLynx v4.1 software (Waters Technologies) was used to calculate the concentrations of metabolites, peak integration and calibration. The data from MassLynx were analyzed using Biocrates’ MetIDQ v5.4.8 software.

Statistical analysis
We performed the statistical analysis using the software SPSS v.22 (Statistical Package for Social Science, Chicago, IL). For gender distribution analysis, we used Chi-square test. We checked normality assumptions with QQ plots. For variables with normal distribution we analyzed by ANOVA test; Kruskal-Wallis test for variables with non-normal distribution; Dunn-Bonferroni for pairwise comparison. We used Wilcoxon test to compare longitudinally MCI patients that have converted to AD. The statistical significance adopted for all analyses was p ≤ 0.05 (α = 95%).

Results
Plasmatic concentrations of spermidine were lower in patients with any degree of cognitive impairment (AD and MCI) as compared to healthy controls (p = 0.007) (Figure 2).

Spermine levels were higher in the MCI group compared to the other two groups (p = <0.001).

The MCI patients grouped in MCI who subsequently converted in AD (MCI_AD) and MCI no converters presented no statistical differences in polyamines levels, however, the sub-sample of MCI patients who progressed to dementia upon follow-up also had lower plasma concentrations of spermine at baseline (Figure 3).

No statistically significant differences were found in plasmatic levels of arginine, ornithine and putrescine between the three groups (Table 2). The difference in frequency of participant’s gender was the same in all groups (p = 0.47; Table 1). There was also no association with any polyamines’ levels and gender. Either way, the results were controlled by gender and years of education.

Discussion
In this study, we determined plasmatic levels of the amino acids arginine and ornithine, and the polyamines putrescine, spermidine and spermine, searching for differences in the concentrations of these amine-containing molecules in patients with mild and severe cognitive impairment (i.e., MCI and AD) compared to healthy, age-matched controls. We found no alteration either on arginine, ornithine or putrescine. Mean spermine concentrations were three times higher in the MCI group as compared to controls and AD but, interestingly, these levels were actually decreased in the sub-sample of MCI patients who converted to AD. Spermidine levels were also lower in AD and MCI groups when compared to healthy controls. The above findings are in agreement with the emerging evidence from studies supporting changes in PAs in cognitive/neurodegenerative disorders such as AD and its prodromal phase MCI. Brain metabolic profiling studies also indicate that changes in PAs pathway do occur in AD, along with an increment of spermidine and spermine in localized brain regions as frontal lobe and parietal

<table>
<thead>
<tr>
<th>Metabolite</th>
<th>AD (Mean ± SD)</th>
<th>MCI (Mean ± SD)</th>
<th>HC (Mean ± SD)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Arginine</td>
<td>63.02 ± 18.71</td>
<td>63.02 ± 18.71</td>
<td>69.62 ± 18.04</td>
<td>0.111</td>
</tr>
<tr>
<td>Ornithine</td>
<td>59.34 ± 18.47</td>
<td>64.07 ± 19.42</td>
<td>64.07 ± 19.24</td>
<td>0.056</td>
</tr>
<tr>
<td>Putrescine</td>
<td>0.14 ± 0.05</td>
<td>0.14 ± 0.05</td>
<td>0.08 ± 0.07</td>
<td>0.126</td>
</tr>
<tr>
<td>Spermidine</td>
<td>7.47 ± 4.18</td>
<td>1.54 ± 1.51</td>
<td>0.90 ± 0.48</td>
<td>0.007</td>
</tr>
<tr>
<td>Spermine</td>
<td>13.13 ± 11.28</td>
<td>41.71 ± 18.33</td>
<td>14.34 ± 15.73</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

AD: Alzheimer's disease; MCI: mild cognitive impairment; HC: healthy controls. Data are presented as mean ± standard deviation. Bold values indicate significant p-values.
lobe; putrescine levels were increased in frontal lobe11. Further, changes in concentrations of plasmatic PAs have also been described. Graham et al. proposed that these changes might help predict the conversion to dementia in MCI patients up to 2 years before the onset of functional deficits11. Their study showed that MCI subjects who subsequently developed dementia (AD) present higher plasmatic PAs levels as compared to stable cases of MCI1. Our study corroborates Graham et al., since we found an increased PAs level 4 years before the onset in a small subset of MCI AD patients.

Luo et al. showed that PAs might interact with amyloid-beta (Aβ) promoting its fibrillation and reducing the neurotoxicity of Aβ peptides13. Hence, PAs, especially spermine, may interact with toxic Aβ forms promoting the structural transition of Aβ toward less toxic fibrillar conformations. This increase of spermine level might be a physiological attempt to decrease neurotoxicity of Aβ2-14. In accordance with these findings, our results suggest that pathological process in AD is associated with depletion of protective PAs spermine and spermidine, given that the plasmatic concentrations of these two molecules are reduced both at pre-dementic and dementic stages of the MCI-AD continuum. Notwithstanding, it is possible that adaptive responses may upregulate the expression of the former, leading to a temporary, but significant three-fold increase in the expression of spermine in MCI. We hypothesize that, in the presence of continuous/progressive Aβ toxicity, the exhaustion of this adaptive response may ultimately lead decrease in the availability of this PA31.

There are few metabolomics studies with plasma reporting polyamine changes in AD patients. The findings regarding putrescine and spermidine plasmatic levels are still inconclusive in AD and MCI31. Our results do not show any statistically significant differences in putrescine levels between the three groups. However, it has been reported that stable MCI subjects present higher plasmatic putrescine levels31. Although putrescine is increased in certain brain regions of AD patients, in plasma it seems not to accumulate because it is preferentially uptaken as a precursor for the synthesis of the spermidine and, subsequentially, the spermine31,32. Spermidine in plasma has been reported to be decreased in stable MCI patients, but not in MCI who afterwards developed to AD, which was interpreted as possible predictor of conversion to AD31.

The small sample size from each diagnostic group is an important limitation of the study; therefore the results should be replicated in an independent and larger sample to reinforce these findings. In addition, it is necessary to validate these findings in other sample cohorts.

Conclusions
Interestingly, our results suggest dynamic changes in the expression of spermine in the MCI-AD continuum, which reinforce the need for additional studies in this field aiming at a better understanding of the role of these important mediators in the pathophysiology of AD.

Author contributions
Authors ACC, HPG and LLT managed the literature searches. ACC and HPG undertook the statistical analysis. ACC wrote the first draft of the manuscript. Authors ACC and HPG wrote the protocol and performed the laboratory analyses. Authors OFV was responsible for clinical assessment. Authors LLT and WFG designed the study. All authors contributed and approved the final manuscript.

Conflict of interest
The authors declare no conflict of interest.

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References


