

The detection of age-, gender-, and region-specific changes in mouse brain tocopherol levels via the application of different validated HPLC methods

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Abstract

The aging process clearly increases the demand for antioxidant protection, especially in the brain, involving that provided by α -tocopherol (α T). However, little is known about the age-related changes in brain α T levels and the influencing effect of gender on it, in human or murine samples as well. Accordingly, the aim of the current study was to detect age-, gender- and region-specific changes in α T concentrations in mouse brain tissue and to assess the influencing effect of plasma α T levels on it.

Female and male C57BL/6 mice at the ages of 6, 16 and 66 weeks (n = 9 in each group) were applied. α T levels were determined with high performance liquid chromatography (HPLC) from the striatum, cortex, hippocampus, cerebellum, brainstem and from plasma samples. A detailed validation process was carried out for the applied HPLC method as well.

The results demonstrated that brain α T levels significantly increased in the striatum, cortex, and hippocampus with aging in both genders, but in a more pronounced way in females with an increasing magnitude of this difference. In case of the cerebellum, a moderate elevation could be detected only in females, whereas in case of the brainstem there was no significant change in α T level. With regard to plasma samples, no clear trend could be identified.

The current study is the first to present age-dependent gender-specific changes in α T level in certain brain regions of the C57Bl/6 mouse strain, and may provide meaningful information for future therapeutic studies targeting aging-related processes.

Keywords: tocopherol, mouse, brain, gender, HPLC, aging

1. Introduction

Tocopherols and tocotrienols, also known as the fat-soluble vitamin E, have an outstanding role in antioxidant mechanisms [1, 2]. This antioxidant protection may have a special relevance with regard to the brain, which organ is especially prone to oxidative injuries due to its high energy demand and elevated lipid content [3]. Accordingly, the deficiency of vitamin E often causes neurological symptoms and studies have proved significantly lower serum or plasma α -tocopherol levels in some neurological disorders, such as in Alzheimer's disease (AD) [4, 5] and in Parkinson's disease (PD) [6], which have an increasing prevalence in the elderly, and in some other neurological conditions [7–11].

With regard to the effects of normal aging on serum tocopherol levels, studies have shown a trend of similarly increasing concentrations in both gender [12, 13] which may be the consequence of the age-related alteration in lipid homeostasis [14]. Studies investigated α -tocopherol (α T) level in cerebrospinal fluid (CSF) in healthy subjects [15] and in patients with Alzheimer-type dementia (ATD) [16] as well. In addition to the delineation of positive correlations between serum and CSF α T and γ -tocopherol (γ T) levels in the healthy ones, significantly decreased CSF α T level was demonstrated in ATD compared to controls. In relation to human brain tocopherol levels, studies provided contradictory results. Craft et al. [17] carried out an examination on the regional distribution of the level of α T and γ T isomers, demonstrating significant age-related (67-90 years, n = 5 – 2 females and 3 males) decline in α T level, more pronounced in females, but due to low case number, statistical comparison could not be carried out. With regard to the regional distribution, there were no significant differences between the grey and white matter of frontal and occipital cortices. Later, Johnson et al. [18] demonstrated a significant relationship between cognitive functions and tocopherol levels in the serum, cerebellar, frontal, occipital, and temporal cortices of healthy octogenarians and centenarians. In case of serum samples, dramatically lower concentrations were measured, whereas for given brain regions, significantly higher levels were determined with aging. There was a significantly negative correlation between serum and brain α T levels, except the cerebellum, although cerebellar α T level, similarly to that of the other brain regions, also positively correlated with the scores of Mini-Mental State Examination (MMSE) and Severe Impairment Battery (SIB), both of which are measures of global cognition.

With regard to rodent studies on tocopherol homeostasis, only limited data are available about the effect of aging and gender on plasma or serum and brain tocopherol levels. Most of the available rodent tocopherol studies did not separate animals to the necessary groups for later statistical comparisons [19–26]. Although Gohil et al. [27] determined α T level in several brain regions

(cerebral cortex, hippocampus, cerebellum, midbrain and the remaining part of the brainstem) of 5 months old C57Bl/6 female (n = 5) and male (n = 3) mice and found significantly higher α T concentrations in all the five regions in females than in males, moreover significantly lower α T level was observed in the cerebellum than those of other examined brain regions, no information was obtained about the effect of aging. With regard to cholesterol levels, there was not any significant difference between genders. There is only one study [28] which assessed the effect of aging on tocopherol levels of rodents in details and reported a not significant decrease in plasma α T level (from ~ 8 to ~ 6 μ M) with aging in C57Bl/6NCr male mice (3, 6, 12, 18 and 24 months old, n = 5) and significantly increasing values only in some brain regions (cerebrum (from 10 to 20 nmol/g tissue), hippocampus (from 12 to 25 nmol/g tissue) and cerebellum (from 11 to 15 nmol/g tissue)). In addition, it was also demonstrated that triglyceride levels were not influenced by aging. However, female mice were not utilized in this study, so the effect of gender cannot be assessed.

There are many available detection methods for the measurement of tocopherol concentrations from plasma or serum and from brain samples [29]. Accordingly, Vitamin E compounds are mainly assessed by diode-array detector (DAD) [30], electrochemical detector (ECD) [31], fluorescence detector (FLD) [32] or mass spectrometry [22, 33]. In case of plasma or serum and brain samples of rodents, especially those of mice, the measurements of other tocopherols are challenging because of their small concentration levels. For the determination of their levels, more sensitive methods are necessary, because with the application of ECD, FLD and DAD, usually only α T can be measured reliably [29].

The aim of the current study was the expansion of the research of Takahashi et. al. [28] and Gohil et. al [27] to be able to quickly and reliably determine α T concentrations in mouse plasma and brain tissues searching for the influencing effects of gender and aging via the application of robust, reproducible and validated HPLC methods. As a part of this process, the applicability of FLD and ECD detectors for the measurement of α T from brain samples was investigated as well.

2. Materials and methods

2.1. Materials

α T, ammonium acetate, ascorbic acid, butylated hydroxytoluene (BHT), dioxane, n-hexane, methanol (MeOH), ethylenediaminetetraacetic acid (EDTA) and sodium-perchlorate (NaClO₄) were purchased from Sigma-Aldrich (Saint Louis, MO, USA). Rac-tocol (rT) was acquired from Matreya LLC (Pleasant Gap, PA, USA).

Acetonitrile (ACN), tetrahydrofuran (THF), absolute ethanol (EtOH), isopropanol (IPA), sodium-chloride (NaCl), disodium-hydrogen-phosphate hydrate ($\text{Na}_2\text{HPO}_4 \cdot 2 \text{H}_2\text{O}$) and sodium-hydrogen-carbonate (NaHCO_3) were purchased from VWR International (Radnar, PA, USA).

Potassium-dihydrogen-phosphate (KH_2PO_4), sodium-dihydrogen-phosphate (NaH_2PO_4), D-glucose hydrate ($\text{D-glucose} \cdot 1 \text{H}_2\text{O}$) and calcium-chloride hydrate ($\text{CaCl}_2 \cdot 2 \text{H}_2\text{O}$) were obtained from Reanal (Budapest, Hungary).

Potassium-chloride (KCl) and Triton-X were purchased from Spektrum 3D (Budapest, Hungary, from 2008: part of VWR), disodium-sulfate (Na_2SO_4) from Fine Chemical Co. (Budapest, Hungary) and magnesium-chloride hydrate ($\text{MgCl}_2 \cdot 6 \text{H}_2\text{O}$) from Scharlau (Barcelona, Spain).

2.2. Animals

For this study C57Bl/6 female and male mice were used. The animals were housed under standard laboratory conditions ($50\% \pm 2\%$ humidity, $22^\circ\text{C} \pm 1^\circ\text{C}$ temperature range and 12h - 12h light - dark cycle) in cages (max 4 per cage) with free access to food (standard rodent diet) and drinking water. We examined 6 groups of animals consisting of 6, 16 and 66 weeks old male and female mice ($n = 9$ in each group). All animal experiments were carried out in accordance with the Scientific Ethics Committee for Animal Research of the Protection of Animals Advisory Board (XXIV./352/2012.).

2.3. Sample preparation

At the age of 6, 16 and 66 weeks, the animals were deeply anesthetized with isoflurane (Forane[®]; Abott Laboratories Hungary Ltd., Budapest, Hungary). After thoracotomy, venous blood was collected from the right ventricle by intracardial puncture into Eppendorf tubes containing EDTA, followed by perfusion with artificial cerebrospinal fluid (pH = 7.4, composition in mM: 122 NaCl, 3 KCl, 1 Na_2SO_4 , 1.25 KH_2PO_4 , 10 $\text{D-glucose} \cdot 1 \text{H}_2\text{O}$, 1 $\text{MgCl}_2 \cdot 6 \text{H}_2\text{O}$, 2 $\text{CaCl}_2 \cdot 2 \text{H}_2\text{O}$, 6 NaHCO_3) for 5 min by an automatic peristaltic pump. After centrifugation of blood samples at 4°C for 5 min at 3500 RPM, the supernatant plasma (200 μL) were mixed immediately with 200 μL 85 mM ascorbic acid and 400 μL 1.14 mM BHT-EtOH solution and the samples were stored at -80°C until further use. Before measurements, 600 μL n-hexane containing 1.14 mM BHT and rT, as internal standard (IS), was added to the stabilized and freshly thawed plasma samples. After an intensive 1 minute long vortex, the samples were centrifuged at 4°C for 10 min at 12000 RPM. In the next step, 450 μL of the hexane layer was evaporated under nitrogen flow. The residue was resolved with 75 μL ACN and 50 μL EtOH-dioxane (1:1), then placed into amber-coloured HPLC vials for measurements.

The anatomical borders of 5 different brain regions (striatum, cortex, hippocampus, cerebellum and brainstem) were determined with the aid of the online-available Allen Brain Atlas: Mouse Brain (Allen Institute for Brain Science, Seattle, WA, USA; <http://mouse.brain-map.org/static/atlas>), and they were rapidly removed on ice and stored at -80°C until further use. Before measurements, the samples were weighed and sonicated (UP100H, Hielscher Ultrasound Technology, Germany; amplitude: 100%, cycle: 0.5) in 1020 μL ice-cold solution (composition in mM: 75 $\text{Na}_2\text{HPO}_4 \cdot 2\text{H}_2\text{O}$, 17 NaH_2PO_4 , 121 NaCl , 2 KCl , 1 EDTA, 67 ascorbic acid, 2 BHT and 0.2 v/v% Triton-X, 9.8 v/v% EtOH). The samples were centrifuged next at 4°C for 10 min at 12000 RPM, and the supernatants were collected and stabilized applying the same method as described in case of the plasma.

2.4. Chromatographic conditions

The concentrations of the $\beta/\gamma\text{T}$ and δT were under the limit of detection in cases of all measurements with all the applied detectors, only αT levels could be quantified, so we focus on this compound in the following part of the methodological description. For the quantification of αT and the IS (rT) from the mouse plasma samples, a previously published method of the authors was utilized [12], with the slight modification that the chromatographic separations were performed on a Kinetex C18 column, 150×4.6 mm i.d., 5 μm particle size (Phenomenex Inc., Torrance, CA, USA).

Two independent methods, utilizing ECD and FLD, were developed for the quantification of αT from certain different brain regions, also with the application of rT as an IS. Both methods involved the utilization of an Agilent 1100 HPLC system (Agilent Technologies, Santa Clara, CA, USA) equipped with a Model 105 ECD (Precision Instruments, Marseille, France) and a FLD (Agilent Technologies, Santa Clara, CA, USA). Measurements were carried out under isocratic conditions.

The first step of our ECD method development was the determination of the optimal working potential for αT and rT. As a result of our measurements (Fig. 1) the working potential was set at +700 mV, using a glassy carbon electrode and an Ag/AgCl reference electrode.

The mobile phase consisted of 91.25 v/v% MeOH, 4.25 v/v % distilled water, 4.5 v/v % IPA and 2.81 w/v% NaClO_4 and it was delivered at a rate of 1.2 ml/min at 25°C onto the reversed-phase column (Luna C18, 75×4.6 mm, 3 μm particle size, Phenomenex Inc., Torrance, CA, USA) after passage through a pre-column (SecurityGuard, 4×3.0 mm i.d., Phenomenex Inc., Torrance, CA, USA). 10 μL of aliquots were injected by the autosampler with the cooling module set at 4°C .

For the method, using FLD detection, the mobile phase consisted of pure methanol, applying the slightly modified method of Yuan et al. [34]. The flow rate was 1.8 ml/min at 25°C , using the same column as in the ECD method. The excitation and emission wavelengths were set at 292 nm and

330 nm, respectively, for the determination of both α T and rT based on spectral analysis. The injection volume was 10 μ L, and the samples were thermostated at 4°C. For ECD and FLD, 2 parallel measurements were carried out for each sample.

2.5. Validation of the applied methods

2.5.1. Calibration curve and linearity

In case of DAD measurements, calibrators were prepared in 1.14 mM BHT-EtOH, then they were arranged in six different concentration levels with concentration ranges of 1 – 50 μ M and 0.25 – 12.5 μ M for α T and rT, respectively. In case of ECD measurements, these ranges were the following: 0.05 – 2.0 μ M and 0.30 – 6.0 μ M for α T and rT, respectively, whereas in case of FLD measurements, the concentration levels were similarly between 0.10 – 2.0 μ M and 0.30 – 6.0 μ M for α T and rT, respectively. The peak area responses were plotted as a function of the corresponding concentration and linear regression computations were evaluated by the least square method with the freely available R software [35]. Good linearity ($R^2 \geq 0.99$) was detected in each method throughout the concentration ranges for all compounds.

2.5.2. Selectivity

To analyze the selectivity of the methods, we matched the chromatograms of α T and rT for a blank plasma or central nervous system (CNS) sample and those for a spiked sample. As shown in Fig. 2, both compounds can be detected without interference from other compounds.

2.5.3. Precision

With regard to the within-run precision, the coefficients of variation of the concentrations were 0.73%, 1.49% and 3.46% for α T and 0.62%, 1.84% and 0.75% for rT in case of DAD, ECD and FLD, respectively.

2.5.4. Recovery

The relative recoveries were estimated by measuring spiked samples of α T at two different concentration levels with three replicates of each. No significant differences were observed for the lower and higher concentrations. The recoveries for the plasma samples ranged from 67% to 71% with DAD, whereas the recoveries for brain samples ranged from 104% to 106% in case of ECD and 87% to 104% with FLD.

2.5.5. Limit of detection (LOD) and limit of quantification (LOQ)

Regarding the quantitative analysis, both LOD and LOQ are important parameters, LOD showing the smallest concentration that can be detected, but not necessarily quantified, whereas LOQ is the lowest analyte concentration in a sample that can be measured with an acceptable level of accuracy and precision [36, 37]. In the current study, LOD and LOQ were calculated as shown in Equation 1. The LOD values for α T were the following: 397 nM, 26 nM and 41 nM for DAD, ECD and FLD, respectively. The LOQ values for α T were 1202 nM, 77 nM and 123 nM for DAD, ECD and FLD, respectively.

$$\text{LOD} = 3.3 * \frac{\sigma}{S'} \text{ and } \text{LOQ} = 10 * \frac{\sigma}{S'}$$

Equation 1

The calculation of LOD and LOQ values by formula, where σ is the standard error of the intercept and S' is the slope of the calibration curve of the analyte. *LOD* limit of detection; *LOQ* limit of quantification.

2.5.6. Statistical analysis

All statistical calculations were performed with the use of the freely available R software [35]. We first checked the distribution of data populations with the Shapiro-Wilk test, and we also performed the Levene test to confirm the homogeneity of variances. As the distribution proved to be Gaussian and the variances were equal, two-way ANOVA was applied with Tukey HSD *post hoc* test for pairwise comparison. We decided *a priori* that the comparisons of opposite gender *and* age groups may not yield meaningful information, so they were not implemented, therefore only 9 comparisons were applied between the 6 groups. We rejected the null hypothesis when the *p*-values were ≤ 0.05 , and in such cases the differences were considered significant. If any significant change was observed, the effect size was calculated (omega-squared (ω^2) for two-way ANOVA and *Cohen's d* for Tukey HSD) [38, 39]. Data were plotted as means (\pm S.D.).

3. Results

3.1. Plasma

The results of the measurements of α T concentration from plasma samples are presented in Table 1, 2 and 3, and in Fig. 3a. The applied two-way ANOVA demonstrated significant difference for age ($F = 10.55$, $df = 2$, $p < 0.001$, $\omega^2 = 0.0276$) but no differences for gender ($F = 0.64$, $df = 1$, $p = 0.427$) or age *vs.* gender ($F = 1.11$, $df = 2$, $p = 0.339$). *Post hoc* analysis with Tukey HSD test yielded a significant increase in α T concentrations with aging only between 16 and 66 weeks old male mice ($p < 0.01$; Table 2), with a 2.234 effect size (Table 3).

3.2. Brain regions

The measurements of α T concentration from brain samples are demonstrated in Table 1, 2 and 3, and Fig. 3b-f. In light of the validation parameters, both ECD and FLD measurement are applicable for the determination of α T from brain samples and accordingly, the results of FLD and ECD measurements were averaged for each individual CNS sample. In the next step, the implementation of two-way ANOVA with Tukey HSD *post hoc* test yielded the following results.

In the striatum (Fig. 3b), there was a significant difference for age ($F = 120.019$, $df = 2$, $p < 0.001$, $\omega^2 = 0.0752$) and gender ($F = 23.062$, $df = 1$, $p < 0.001$, $\omega^2 = 0.0070$), and for age vs. gender ($F = 3.588$, $df = 2$, $p = 0.0353$, $\omega^2 = 0.0016$) as well. The Tukey HSD *post hoc* test revealed significantly elevated α T concentrations in the latter groups in the following pairwise comparisons from those of *a priori* decided: $p < 0.001$ for 6 vs. 16 weeks old females and $p < 0.01$ for males; $p < 0.001$ for both 16 vs. 66 weeks old females and males; $p < 0.001$ for both 6 vs. 66 weeks old females and males; $p < 0.05$ for 16 weeks old males vs. females and $p < 0.001$ for 66 weeks old males vs. females (Table 1 and 2).

In the cortex (Fig. 3c), there was also a significant difference for age ($F = 159.589$, $df = 2$, $p < 0.001$, $\omega^2 = 0.1042$), gender ($F = 17.377$, $df = 1$, $p < 0.001$, $\omega^2 = 0.0054$), and for age vs. gender ($F = 5.465$, $df = 2$, $p = 0.0073$, $\omega^2 = 0.0029$). The Tukey HSD *post hoc* test revealed significantly elevated α T concentrations in the latter groups in the following pairwise comparisons from those of *a priori* decided: $p < 0.001$ for 6 vs. 16 weeks old females and $p < 0.05$ for males; $p < 0.001$ for both 16 vs. 66 weeks old females and males; $p < 0.001$ for both 6 vs. 66 weeks old females and males and $p < 0.001$ for 66 weeks old males vs. females (Table 1 and 2).

Furthermore, in the hippocampus (Fig. 3d), similar results were demonstrated for age ($F = 195.500$, $df = 2$, $p < 0.001$, $\omega^2 = 0.1056$), gender ($F = 24.343$, $df = 1$, $p < 0.001$, $\omega^2 = 0.0063$), and for age vs. gender ($F = 7.045$, $df = 2$, $p = 0.0021$, $\omega^2 = 0.0033$). The Tukey HSD *post hoc* test revealed significantly elevated α T concentrations in the latter groups in the following pairwise comparisons from those of *a priori* decided: $p < 0.001$ for both 6 vs. 16 weeks old females and males; $p < 0.001$ for both 16 vs. 66 weeks old females and males; $p < 0.001$ for both 6 vs. 66 weeks old females and males and $p < 0.001$ for 66 weeks old males vs. females (Table 1 and 2).

With regard to the cerebellum (Fig. 3e) and the brainstem (Fig. 3f), there was a significant difference for age ($F = 17.091$, $df = 2$, $p < 0.001$, $\omega^2 = 0.0134$ and $F = 3.491$, $df = 2$, $p = 0.0384$, $\omega^2 = 0.0021$, respectively) and gender ($F = 10.66$, $df = 1$, $p = 0.0020$, $\omega^2 = 0.0040$ and $F = 13.295$, $df = 1$, $p < 0.001$, $\omega^2 = 0.0051$, respectively), but not for age vs. gender ($F = 2.897$, $df = 2$, $p = 0.0649$ and $F = 0.820$, $df = 2$, $p = 0.4464$, respectively). The Tukey HSD *post hoc* test revealed significantly elevated α T concentrations only in case of the cerebellum in the latter groups in the following

pairwise comparisons from those of *a priori* decided: $p < 0.01$ for 6 vs. 16 weeks old females and $p < 0.001$ for 6 vs. 66 weeks old females (Table 1 and 2).

The corresponding effect size values for the significant differences in brain α T concentrations are presented in Table 3.

4. Discussion

Evidence suggests that tocopherols may have a special role in antioxidant protection in lipid-rich structures, such as the CNS. As there is a clear worsening of brain functioning with aging, there were several approaches which aimed at the achievement of neuroprotection via the administration of exogenous α T [40–44]. However, only limited data are available on the changes of endogenous tocopherol levels either in human or murine brain samples with aging [12, 13, 17, 18, 28]. Therefore, there is a special need for the fine assessment of age-related changes in α T levels. The identification of clear trends with regard to either certain brain regions or genders may help to understand the differences in the sensitivity to oxidative damage. Although the main focus may be paid on human studies, the assessment of rodents from this point of view may also yield relevant information in light of the fact that most preclinical research on neurodegeneration is carried out in animals belonging to this subfamily.

Accordingly, the aim of the current study was to determine region-, age- and gender-specific changes in brain α T level in the C57Bl/6 mouse strain, which is one of the most commonly applied strains in the research on neurodegeneration. Furthermore, our study was supplemented with the assessment of plasma samples as well to be able to judge the possible influence of peripheral changes on brain α T levels. The results demonstrated that brain α T levels significantly increased in the striatum, cortex, and hippocampus with aging in both genders. This increase was more pronounced in females and the magnitude of this difference also rose with aging in case of all the above-mentioned brain regions. However, in case of the cerebellum, a moderate elevation could be detected only in females, whereas in case of the brainstem there was no significant change in α T level. With regard to plasma samples, no clear trend could be identified, a significant difference was found only between 16 and 66 weeks old males. These findings are in line with those found by Takahashi et al. [28] and Gohil et al. [27], i.e., there is a clear elevation of cortical and hippocampal α T levels, but only a moderate increase in cerebellar α T level with aging and significantly higher values in females. The novelty of the current study is the presentation of such a pronounced elevation in striatal α T level, while no change in brainstem α T level, and furthermore, the first delineation that the difference between genders significantly increases with aging in case of the striatum, cortex and hippocampus. Similar to that found by Takahashi et al., plasma α T level

seemingly does not influence the characteristic change in striatal, cortical and hippocampal α T levels. In addition to the clear and unbiased demonstration of region-, age- and gender-specific changes in mouse brain α T level via the application of different validated HPLC methods, this article also aims at highlighting the possible underlying factors behind these findings and their hypothetical significance as well.

For better understanding of these results, we looked further into the transport and metabolism of tocopherols. The transport of vitamin E in the mammalian body is well documented [45–47]. Due to the chemical properties of vitamin E compounds, their circulation is associated with lipoproteins, but their transcellular transport is mediated by carrier proteins [45]. The major forms of vitamin E, α T and γ T are taken up by diet and absorbed from the small intestine, followed by their transportation to the liver by chylomicrons. In hepatocytes, a special binding protein, α T transport protein (α -TTP), is expressed and has a major role in α T uptake. Studies have shown that α -TTP recognizes preferably α T by its chemical characters [45]. From liver cells, α T is transferred to the serum by very-low density lipoproteins (VLDL). Following the hydrolysis of VLDL by plasma enzymes to low-density lipoprotein (LDL), it delivers α T to extrahepatic tissues via the LDL receptor-mediated pathway [47, 48].

In the brain the CSF has an important role in micronutrient distribution, including vitamins as well [46]. The transportation of vitamins B₁, B₃ and E from blood to CSF with simple diffusion cannot provide enough concentrations for tissue cells, therefore several specific systems (facilitated diffusion, active sodium-dependent or independent systems, receptor related systems and many other mechanisms) are included as well [46, 49]. Accordingly, it was proposed that brain α T homeostasis has a strict regulation, but the precise description of the pharmacokinetics of vitamin E transport into the CSF and brain is not available yet, although some possible mechanisms were previously described in the literature [46].

One of these mechanisms may be connected to the scavenger receptor class B type 1 (SR-B1) [50–52], which is responsible for α T transport across the blood-brain barrier (BBB) by promoting the uptake of HDL-associated α T in brain capillary endothelial cells (BCEC). Balázs et al. [50] proposed that SR-B1 is located at the caveolae of the apical membrane of BCEC and facilitates the selective uptake of HDL-associated α T by transcytosis across the BBB. In the study of Srivastava et al. [51], C57Bl/6 mice were fed with high-cholesterol and high-fat containing diet and their brain SR-B1 expressions consequently increased, although hepatic SR-B1 expression was not influenced. Furthermore, Mardones et al. [52] demonstrated that SR-B1 knock-out (KO) female and male (n = 3-6, 2 - 4 months old) mice have significantly higher plasma and lower CNS α T levels, consistent with the postulated role of SR-B1 at the BBB. Surprisingly, the whole brain levels were not

decreased enough to cause deficiency symptoms. There are no available data on the influence of aging and gender on SR-B1 expression.

The second possible mechanism may be connected to phospholipid transfer protein (PLTP) [53]. PLTP is highly expressed in the choroid plexus (CP) which raises the possibility that PLTP is involved in the transfer of α T from plasma into the CSF [54, 55]. In PLTP KO female mice (n = 6) Desrumaux et al. [54] investigated the brain concentration of α T and found it decreased by 30% compared to controls and found considerably higher (by 450%) lipofuscin level, which is the final product of the reaction of lipid peroxide with proteins during oxidative stress. In the substantia nigra. PLTP deficiency also resulted in the accumulation of brain cholesterol derivatives oxidized at position 7 and besides this, Yokota et al. [56] proved that lipofuscin accumulations could disappear with the application of vitamin E dietary supplementation. Similar to SR-B1, there are also no available data on the influence of aging and gender on PLTP expression.

α -TTP serves as a third mechanism of α T transport in the mammalian brain [56]. The presence of this protein is demonstrated in rodent and human brains as well, although the expression level in normal brain tissue is low [57]. In α -TTP KO female and male mice (n = 2 – 2), the plasma and tissue concentrations were less than 0.46 μ M and 0.23 nmol/g ww, respectively. Moreover, supplementation with oral vitamin E raised the plasma concentration of these mice close to normal (\sim 3.4 μ M), but the brain levels increased only with \sim 20%, however, this small increase in the brain could eliminate the neurological signs [56]. Gohil et al. [27] also investigated α -TTP expression in several CNS and liver tissue samples of 5 months old α -TTP KO and control mice with the finding that α -TTP protein or its mRNA could only be detected from the liver of controls. Later, Takahashi et al. [28] successfully determined α -TTP expressions from the cerebrum, hippocampus and cerebellum. However, they utilized only 12-month-old male mice for that purpose, therefore the effect of aging and gender could not be analysed.

Another investigated pathway is in relation with afamin, a member of the albumin super family and may serve as a transport protein involved in α T homeostasis as well [58]. Afamin is synthesized endogenously by BCEC and assists α T transport to astrocytoma cells, but to a lesser extent than HDL-mediated transport. The expression of afamin was detected by immunohistochemistry in porcine, mouse and *post mortem* human brains. It was also demonstrated that afamin level correlates with the concentration of α T in the CSF, but not with α T level in the serum [58, 59]. However, there are no available data on the influence of aging and gender on afamin expression.

Although the above-mentioned findings may support the role of SR-B1, PLTP, α -TTP and afamin in the transport and regulation of the α T levels in the CNS, the assessment of the effect of aging and gender on these transport processes warrants further studies.

Probably the major significance of the current findings is that the elevations of α T concentrations by aging may serve as to compensate the burden of oxidative stress which increases with aging [12, 43, 60]. Furthermore, the demonstrated gender-related differences, i.e., significantly higher α T levels in female striatal, cortical and hippocampal brain regions, may at least partially explain the decreased vulnerability of e.g., C57Bl/6 female mice to certain neurotoxins, e.g., 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine [61–64] compared to the male ones.

In light of the fact that there are only limited and contradictory data are available on the changes of human brain tocopherol levels with aging, the human relevance of the current findings cannot be determined at present. However, the results may draw attention to the importance of antioxidant protection against oxidative damage, the extent of which clearly increases with aging. With regard to the effect of gender on age-related pathological alterations, several human studies were published [65–68]. Király et al. [68] demonstrated that male brain ages faster, following the assessment of the volume of cortical and subcortical grey matter, including that of the hippocampus, putamen and caudate nucleus, within the age-range of 21 to 58 years. Although the age range is presumably different, this male preponderance was not reassured in the most common neurodegenerative disorder, AD [69], but was proposed in the second most common neurodegenerative disorder, PD [70]. Nevertheless, the fact that there is not enough human data to be able to determine the extent of the influence of the alterations in tocopherol homeostasis on the pathogenetic process in major age-related neurodegenerative disorders, warrants further studies.

In conclusion the current study is the first to demonstrate age-dependent gender-specific changes in α T level in certain brain regions of the C57Bl/6 mouse strain which finding may explain the increased vulnerability of male C57Bl/6 mice to certain neurotoxins targeting these regions, but the delineation of possible underlying mechanisms and the demonstration of their human relevance may require future studies. Nevertheless, a deeper insight into this aspect of antioxidant protection may help the development of therapeutic strategies against age-related pathogenetic processes.

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6. Conflict of interest

The authors declare no conflict of interest.

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8. Figure captions

Fig. 1

Plotted peak areas vs. voltage diagram of αT and rT . αT α -tocopherol; rT rac-tocol.

Fig. 2

Representative chromatograms of rT and αT . The chromatograms of blank and spiked (with rT and αT) mouse plasma samples measured with DAD (a) and those of blank and spiked (with rT and αT) mouse brain sample homogenizations measured with ECD (b) and FLD (c) are demonstrated. αT α -tocopherol; DAD diode-array detector; ECD electrochemical detector; FLD fluorescence detector; rT rac-tocol.

Fig. 3

αT concentration levels in plasma and in different brain regions of mice. We observed significant elevation of αT levels in the plasma (a), striatum (b), cortex (c), hippocampus (d) and cerebellum (e), but not in the brainstem (f) with aging. To avoid the overcomplication of this figure, the results of statistical comparisons are presented elsewhere (Table 1, 2 and 3), except that for the interaction which may help in the interpretation of the presence or absence of increasing difference between males and females with aging. Data are presented as mean (\pm S.D.); $n = 9$; αT α -tocopherol; *N.S.* not significant.