Behaviour and Degenerative Changes in the Basal Forebrain Systems of Aged Rats (12 Months Old) after Levo-Acetyl-Carnitine Treatments

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ABSTRACT

One group of six male control rats [12 months old] and one group of six male rats of the same age, singularly maintained in a cage, and treated with acetyl-L-carnitine-HCl [γ-trimethyl-beta-acetyl-butyrobetaine-HCl: Sigma-Tau code ST200 or ALCAR: 60 mg/kg/day [7]/po] for six months were tested in the spatial learning/memory Morris maze-water task and for atrophy and cell loss in seven myelo- and cytostructurally defined basal forebrain (BF) cholinergic regions [Freddi et al., 2009]. Coronal sections 25 μm thick were cut through the BF regions and processed every 200 μm for choline acetyltransferase (ChAT) immunohistochemistry. The ALCAR-treated rats had significantly shorter exit times on the Morris maze-water task test than the control rats (average ± SD 28.3 ± 12.4 s vs. 61.16 ± 4.67 s; t = 6.07, DOF = 10, P = 0.0001). Degenerative morphological changes in the BF ChAT-positive cells were observed in the substantia innominata pars anterior of the control rats but not in the treated animals (P < 0.05). In the BF, the counted and estimated average number of ChAT + cells in the 12-month-old ALCAR-treated rats (ChAT-ALCAR-12+ [Nos. 2,3,4]) was higher but not significantly (15.288 ± 3281) than that counted and estimated in the 12-month-old control rats (ChAT-CT-12 [Nos. 1,2,3]) (11.508 ± 3868), t = 1.82, DOF = 10, P = 0.319). In the substantia innominata pars posterior, the ChAT+ cells were significantly more numerous (P < 0.05) in the 12-month-old ALCAR-treated rats (ChAT-ALCAR-12 + [Nos. 2,3,4]) than in the control rats (ChAT-CT-12 [Nos. 1,2,3]). Above all, these results demonstrate that treatment with ALCAR from the age of 6 up to 12 months significantly attenuated spatial learning/memory impairment on the Morris maze-water behavioral task (P < 0.001) and also importantly reduced degeneration in size and number of cholinergic cells in the nucleus basalis magnocellularis of the BF. Accordingly, the surviving cholinergic neurons found in the BF of the ALCAR-treated rats might play an important role in modulating cortical activity and facilitating processes of attention, learning and memory.

Keywords: Acetyl-L-Carnitine-HCl; Ageing; Choline Acetyltransferase Neurons; Substantia Innominata; Morris Maze-Water Task

1. Introduction

Acetyl-L-carnitine ([ALCAR] gamma-trimethyl-beta-acetyl-butyrobetaine-hydrochloride, Sigma-Tau code: ST200) is the metabolically active form of acetylcarnitine present in the central nervous system (CNS). ALCAR acts as a shuttle for long-chain fatty acids between the cytoplasm and the mitochondria, is involved in the production of cellular energy and the removal of toxic accumulation of fatty acids from the mitochondria. It is actively transported across the blood-brain barrier and displays affinities for GABAergic and L-glutamate carriers [1]. Acetyl-L-carnitine is a source of acetyl groups and acts as a partial cholinergic agonist synergistically with ACh and L-glutamate, increasing the firing rate of rat neocortical neurons [2], thus working in the CNS [3-9].

Rodents present an age-related decline in nerve growth factor (NGF) and basal forebrain (BF) cholinergic neurons, as well as in learning and memory ability [10,11]. Acetyl-L-carnitine protects the CNS and NGF [8,9,12-13], improves cognitive deficits, and very importantly reduces rat mortality [14,15]. Rats and mice show an age-dependent decline in ACh and cell loss in the septo-hippocampal and BF regions [16-19]. Contrasting data about the age-dependent cholinergic cell BF decline have been reported by Horneberger et al. [20], who noted that aging causes a progressive shrinkage of AChE positive neurons in the BF nuclei, which was most pronounced in the nucleus basalis magnocellularis; however, the number of
cells remained unchanged.

To date, no studies have investigated in 12-month-old rats, untreated and treated with ALCAR, for the severity and the extension of cholinergic cell loss in the seven myelo- and cytoarchitecturally defined BF regions which comprise [15]: the medial septum (MS) and diagonal band of Broca (DBB), the magnocellular preoptic nucleus (MCPO), the olfactory tubercle (Otu), the substantia innominata pars anterior (SiA, also called the ventral pallidum), the substantia innominata pars posterior (SiP, also called the extended amygdala), and the globus pallidus (GP), where basal-cortical neurons projecting to the meso- and isocortex [21,22] intermingle with the BF cells projecting into the caudal hypothalamus [23].

Although cholinergic cells account for only 5% - 6% of the total population of the BF, this study focused on: 1) the topographical location and severity of cholinergic deficit in 12-month-old rats with impaired performance on the spatial/learning Morris water-maze test; and 2) the effect of acetyl-L-carnitine treatment on rats aged from 6 to 12 months to see whether this neuropsychopharmacological compound can significantly attenuate spatial learning/memory impairment on the Morris maze-water task [13] and significantly reduce degeneration of the largest cholinergic neurons in the seven myelo- and cytoarchitecturally defined BF regions [15,24-26].

2. Methods
2.1. Animals

All experiments were carried out in accordance with Directive 86/609/EEC regulating experimental animal care and with the aim to maximally reduce stress, suffering and the number of animals used [15,25].

Twelve male Sprague-Dawley Ico rats (OFA-SD-IOPS Caw) (IFFACredo, Italy) (age, 6 months; approximate body weight 545 ± 20 g) bought by Sigma-Tau (Rome, Italy) were singularly maintained in cages at the Sigma-Tau laboratory, given food and water ad libitum and kept at the same position in the pool at all times during the experiment. The pool was placed in a corner of a room containing several extra-maze cues (objects) which the animals could use as reference points to locate the platform. Dried milk powder was added to the water to make it opaque. The day before each experiment, the platform was removed, the water was changed, and the rat was allowed to swim freely for 3 minutes so as to become familiar with the surroundings. On test days the animals were placed in the tank at one of the four different equidistant starting points along the edge and timed to see how long they took to reach the platform. After the rats had found the platform, they were left on it for 30 seconds before beginning a new trial. Four trials were carried out per session(s) [S1 + S2 + S3 + S4] [15]. The experiment was divided into two phases: an acquisition phase consisting of eight sessions, and a retention phase, presented in this study, consisting of four sessions.

We calculated the average time (in s) that each animal took to climb onto the platform during the retention phase of four sessions [S1(Sa,b,c,d) + S2(Sa,b,c,d) + S3(Sa,b,c,d) + S4(Sa,b,c,d)]: the last test was performed before death by aortic perfusion. The average time needed to escape onto the platform (in seconds (s)) was statistically analyzed using two- and one-way ANOVA and post-hoc Student’s t-test multiple comparison of the means.

2.2. Behavioral Morris Water-Maze Test

After six months of ALCAR treatment (or not), all twelve 12-month-old rats were tested, during daylight, for spatial learning/memory on the Morris water-maze task [27]. The test apparatus consisted of a large round plastic tank (130 cm in diameter and 45 cm high) filled to a depth of 40 cm with 770 litres of water kept at 24°C. The test apparatus was divided into four quadrants by four points positioned at equal distances along the edge and arbitrarily labelled North, South, East and West. A white escape platform (10 cm in diameter) was located 1 cm beneath the water surface in the middle of the NW quadrant, and kept at the same position in the pool at all times during the experiment. The pool was placed in a corner of a room containing several extra-maze cues (objects) which the animals could use as reference points to locate the platform. Dried milk powder was added to the water to make it opaque. The day before each experiment, the platform was removed, the water was changed, and the rat was allowed to swim freely for 3 minutes so as to become familiar with the surroundings. On test days the animals were placed in the tank at one of the four different equidistant starting points along the edge and timed to see how long they took to reach the platform. After the rats had found the platform, they were left on it for 30 seconds before beginning a new trial. Four trials were carried out per session(s) [S1 + S2 + S3 + S4] [15]. The experiment was divided into two phases: an acquisition phase consisting of eight sessions, and a retention phase, presented in this study, consisting of four sessions.
2.3. Surgery and Perfusion

After completing the behavioral tests at the Sigma-Tau laboratory, the five control (ChAT-CT-12: Nos. 1,2,3,4,5 (body weight, 713 ± 36.5 g) and the four ALCAR-treated (ChAT-ALCAR-12: Nos. 1, 2, 3, 4 (body weight, 730 ± 73.5 g) male Sprague-Dawley Ico rats were anaesthetized with chloral hydrate (278 ± 15 mg/ip) for surgery and perfusion at the Medical School of the University of Milan (Milan, Italy). One control rat (ChAT-CT-12: No. 6) and 2 ALCAR-treated rats (ChAT-ALCAR-12: Nos. 5, 6) were delivered to the Pharmacy School of the University of Milan (Milan, Italy) for another parallel study.

As previously reported [15], the brains of the five control (ChAT-CT-12: Nos. 1,2,3,4,5 (body weight, 713 ± 36.5 g) and the four ALCAR-treated (ChAT-ALCAR-12: Nos. 1, 2, 3, 4 (body weight, 730 ± 73.5 g) male Sprague-Dawley Ico rats were fixed by perfusion through the ascending aorta, first with a phosphate buffered solution [(PBS; 0.1 M, pH 7.4, containing 20 mM MgCl2) (<100 ml within 30 s)], then with a fixative solution containing 3% paraformaldehyde, 0.2% picric acid and 0.05% glutaraldehyde (pH 7.4) in 0.1 M PBS) (approximately 1000 ml over 30 min), and finally with 10% sucrose-PBS (pH 7.4) (approximately 250 ml over 15 min). All solutions were perfused at a temperature between 10°C and 18°C. The brains were removed, cut into two coronal blocks, immersed overnight, and shaken at 4°C in 30% sucrose/PBS, then frozen at −50°C in liquid freon and stored at −80°C.

2.4. Immunocytochemistry

As previously published [15,21-24], coronal brain sections 25 μm thick of the ChAT-CT-12+: Nos. 1,2,3 and of the ChAT-ALCAR-12+: Nos. 2,3,4 rats were cut on a freezing microtome and collected every 200 μm in PBS (pH 7.3). The sections were processed for visualization of choline acetyltransferase (ChAT+) labeling using the peroxidase antiperoxidase (PAP) technique with a ChAT monoclonal antibody (1:4) prepared in rats (Boehringer Mannheim, Germany). The ChAT antibody was obtained by fusing mouse myeloma cells (X63Ag8.653) with spleen cells of Wistar rats after immunization with highly purified choline acetyltransferase from pig brains. The sections were incubated overnight, first with the primary antibody, then with a rabbit anti-rat antibody (1:50, ICN, Italy) for 45 min, and finally with rat PAP (1:50 - 1:75) (Sternberger Monoclonals Inc., USA) for 45 min. The last two steps were repeated in a double-bridge procedure. The primary and secondary antibodies were diluted in Tris solution (TS, 0.1 M) containing 1% normal rabbit serum (NRS) which was also used for rinses between incubations. Before incubation, the sections were placed in TS containing a higher concentration of bovine serum albumin (BSA 10%) for blocking. The peroxidase (PAP) was revealed by 6 min incubation with 0.05% 3,3′diaminobenzidine (DAB) and 0.01% hydrogen peroxidase (H2O2) in Tris water (TW). Control sections were incubated without the primary antibody and processed with the secondary antiserum, PAP, and DAB.

2.5. Estimation and Morphological Image Analysis of the ChAT+ Cells in the BF

As previously reported [15,21-24], to map and estimate the BF ChAT-immunoreactive cells in the BF of the 3 control (ChAT-CT-12+: Nos. 1, 2, 3) and of the 3 ALCAR-treated (ChAT-ALCAR-12+: Nos. 2, 3, 4) rats—where the perfusion and immunohistochemistry were best developed—we used a Biocom image-analysis system (Biocom, Paris, France) with a computer-based atlas of cholinergic BF cell groups. The image-analysis system consisted of a Leitz Dialux microscope equipped with an x/y movement-sensitive stage and a video camera (COHU) connected to a Compaq Deskpro/25 M computer. The atlas consisted of 6 templates (800 μm apart) extending from AP levels A10.9 - A6.9 through the BF regions, divided into seven contours distributed over more than one level. Labeled ChAT+ cells were mapped [ChAT-CT-12+: Nos. 1,2,3 and ChAT-ALCAR-12+: Nos. 2, 3, 4] or drawn [ChAT-CT-12+: No. 1 and ChAT-ALCAR-12+: No. 2] in the seven subregions of the BF by fitting each section to the appropriate atlas template at low power (2.5 × objective) and tracing or indicating the ChAT-CT-12+: Nos.1, 2, 3 + cells and the ChAT-ALCAR-12+: Nos. 2, 3, 4 + cells of the rats using a mouse-driven cursor at high power (40 × objective).

2.6. Statistical Analysis

As previously described [15,21-24], the morphometric measurements of the ChAT-CT-12+ and the ChAT-ALCAR-12+ cell groups under study were averaged for each BF nucleus and are presented here, as previously described for 2 rats: one control (ChAT-CT-12+: No. 1) and one ALCAR-treated (ChAT-ALCAR-12+ No. 2) rat. Data were analyzed with Student’s t-test. Abercrombie’s estimation of cholinergic cells has recently been demonstrated not to differ significantly from stereological estimation of cholinergic cells [24,25]. Therefore, we performed average cell counts per 3 rats from each group [ChAT-CT-12+: Nos. 1, 2, 3 and ChAT-ALCAR-12+: Nos. 2, 3, 4] in the seven cyto- and myelos-structurally defined BF regions [25] and computed the total numbers by using Abercrombie’s formula (Abercrombie 1946), viz. using the average morphometric measurements of the large diameters of the ChAT-CT-12+ (No. 1) and ChAT-ALCAR-12+ (No. 2) cells calculated from the longest diameter of the nuclei.
The computer-based maps through the BF were completed for publication using the Corel Draw 8 software program.

3. Results

3.1. Morris Behavioral Water-Maze Test

The average exit time in the last four sessions (S1(a,b,c,d) + S2(a,b,c,d) + S3(a,b,c,d) + S4(a,b,c,d)) of the final trials during the retention phase of spatial learning and memory consolidation of the Morris water-maze test [13], i.e., the time taken to reach the platform, was significantly greater on average in the 12-month-old control rats [ChAT-CT-12+ (Nos. 1, 2, 3)] (61.16 ± 4.67 (s)) than in the 12-month-old ALCAR-treated rats [ChAT-ALCAR-12+ (Nos. 2, 3, 4)] (28.30 ± 12.40 (s)) (Table 1). The control animals had consistently longer exit times. Prolonged ALCAR treatment of the rats, from the age of 6 to 12 months, very significantly attenuated the impairment in spatial learning (P < 0.0001). Moreover, the exit times in the last four sessions of the final trial indicate that the aged ALCAR-treated rats showed the most significant drop in exit time in the last session [13] (see Table 1 and Figure 2).

3.2. Localization of ChAT Basal Forebrain Neurons

Stained BF sections: in the 12-month old controls [ChAT-CT-12+ (Nos. 1, 2, 3)] and in the ALCAR-treated [ChAT-ALCAR-12+ (Nos. 2, 3, 4)] rats, the cells containing ChAT were evident by a dark floccular brown staining produced by DAB. As previously reported [24], ChAT+ cells were examined only if they were visible as whole cells containing nuclei with at least 3 dendrites.

In the BF, the ChAT+ cells in the control rats (ChAT-CT-12+: Nos. 1, 2, 4) (Figure 1) and the ChAT+ cells in the ALCAR-treated rats (ChAT-ALCAR-12+: Nos. 2, 3, 4) (Figure 2) were distributed throughout the myelo-cyto-architectonic subregions: the medial septum (MS), the diagonal band of Broca (DBB), the magnocellular preoptic nucleus (MCPO), the olfactory tubercle (OTu), the substantia innominata pars anterior (Slp), also called the ventral pallidum, the substantia innominata pars posterior (Slp), also called the sublenticular substantia innominata

Table 1. Time taken by 12-month-old control rats (ChAT-CT-12; Nos. 1, 2, 3) and ALCAR-treated 12-month-old rats (ChAT-ALCAR-12; Nos. 2, 3, 4) to find the platform (seconds) on the Morris water-maze test.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Time taken to find the platform (seconds)</th>
<th>Trial Exit time (s)</th>
<th>Exit time (s)</th>
<th>Trial S1-S4 Mean ± SD</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>S1(a,b,c,d)</td>
<td>S2(a,b,c,d)</td>
<td>S3(a,b,c,d)</td>
</tr>
<tr>
<td>Control</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ChAT-CT-12+ (Nos. 1, 2, 3)</td>
<td>Average</td>
<td>67.0</td>
<td>62.3</td>
<td>59.3</td>
</tr>
<tr>
<td></td>
<td>SD</td>
<td>1.73</td>
<td>2.51</td>
<td>4.72</td>
</tr>
<tr>
<td>ALCAR</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ChAT-ALCAR-12+ (Nos. 2, 3, 4)</td>
<td>Average</td>
<td>42.3</td>
<td>33.3</td>
<td>24.3</td>
</tr>
<tr>
<td></td>
<td>SD</td>
<td>2.08</td>
<td>1.15</td>
<td>1.52</td>
</tr>
</tbody>
</table>

1) Two-way ANOVA showed that the ALCAR-treated rats (ChAT-ALCAR-12; Nos. 2, 3, 4) had significantly shorter exit times than the untreated control rats (ChAT-CT-12; Nos. 1, 2, 3); across groups (P = 0.0001), sections (P = 0.0001) and between group and sections (P = 0.0001); 2) One-way ANOVA for each group showed that all ALCAR-treated rats (ChAT-ALCAR-12; Nos. 2, 3, 4) had significantly shorter exit times than the untreated control rats (ChAT-CT-12; Nos. 1, 2, 3) (P = 0.0001); 3) Rats in the same group (ChAT-CT-12; Nos. 1,2,3 and ChAT-ALCAR-12; Nos. 2, 3, 4) took approximately the same time to reach the platform.

Figure 1. Computer atlas figure showing six levels of the BF. The contours of the major subregions were drawn with the aid of a computer image-analysis system (Biocom, Paris, France). Distribution of ChAT+ cells in the ChAT-CT-12+ No. 1 rat was directly mapped at high magnification (40×) on the atlas template. Each symbol represents one labeled cell.
or extended amygdala, and the globus pallidus (GP). Throughout the BF, the ChAT+ cells of the 12-month-old control and ALCAR-treated rats were found to be oriented according to the principal direction of the passing fibers within the seven BF subregions.

The ChAT+ cells in the BF of the control rat [ChAT-CT-12+ (No. 1)] varied from small to medium in size, ranging from 11.9 to 21.9 µm in large diameter; they formed a population with one mode. Indeed, the majority of the ChAT+ cells had medium to large-sized neurons with a long diameter of 17.9 ± 1.8 µm. In summary, the average large diameter of the ChAT+ cells measured in the ALCAR-treated rat [ChAT-ALCAR-12+ (No. 2)] (Table 2). Although the cell population of the ChAT-CT-12+ (No. 1) and of the ChAT-ALCAR-12+ (No. 2) overlapped in surface area and diameter, the population of the ChAT-CT-12+ (No. 1) was significantly (P < 0.05) smaller than that of the ChAT+ cells of the ALCAR-treated rat [ChAT-ALCAR-12+ (No. 2)] (Table 2).

Throughout the BF, the ChAT+ cells were more numerous in the ALCAR-treated rats [(ChAT-ALCAR-12+ (Nos. 2, 3, 4)] than in the control rats [ChAT-CT-12+ (Nos. 1, 2, 3)] (15.288 ± 3281 vs. 11.508 ± 3868, respectively) (Table 3).

3.3. Regional Characterization and Distribution of ChAT+ Cells

Located rostrally in the deeper inner ring of the BF where the transverse fiber systems sweep beneath the accum-bens-striatum and pars anterior of the anterior commissure, ChAT+ cells are located within the anterior portion of the substantia innominata (Sla). The ChAT+ cells in the subcommissural Sla are frequently elongated, multipolar neurons that commonly lie with their long axes parallel to the transverse fibers that travel throughout this

Table 2. Morphometry of choline acetyltransferase-positive (ChAT+) neurons in the basal forebrain of one aged 12-month-old control rat (ChAT-CT-12+ # 1) and one ALCAR-treated rat (ChAT-ALCAR-12+ # 2).

<table>
<thead>
<tr>
<th>Structure</th>
<th>No. of Cells</th>
<th>Small diameter</th>
<th>Large diameter</th>
<th>Surface area</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sla</td>
<td>5</td>
<td>8.1 ± 2.2</td>
<td>14.0 ± 0.9*</td>
<td>99.2 ± 16.4</td>
</tr>
<tr>
<td>BF</td>
<td>35</td>
<td>10.0 ± 2.4</td>
<td>16.8 ± 2.0*</td>
<td>134.5 ± 37.1</td>
</tr>
<tr>
<td>Sla</td>
<td>5</td>
<td>8.5 ± 2.4</td>
<td>15.3 ± 0.4*</td>
<td>100.8 ± 14.7</td>
</tr>
<tr>
<td>BF</td>
<td>36</td>
<td>10.3 ± 2.3</td>
<td>17.9 ± 1.8*</td>
<td>140.2 ± 40.7</td>
</tr>
</tbody>
</table>

1) Measurements in µm and µm² (Mean ± SD) in the basal forebrain (right site of the brain) for ChAT+ cells in one animal from each group: aged control rat (ChAT-CT-12+ No. 1) and one ALCAR-treated rat (ChAT-ALCAR-12+ No. 2) (16, 22, 23,24,25).
ever, the similarity between the Morris water-maze escape latency times for the aged control and treated rats in this study belong to the category of the so-called “intermediate performer rats” as classified by Taglialetela et al. [13,28] (see Figures 1 and 2).

The behavioral findings of this study strongly indicate and confirm that prolonged treatment with ALCAR can very significantly ($P < 0.0001$) improve spatial learning and memory on the Morris water-maze test [Freddi et al., 2009]. This study confirmed that the 12-month-old control rats with impaired spatial learning and memory performance on the Morris water-maze test [13,28] had significantly smaller and fewer BF cholinergic neurons than the ALCAR-treated rats [16]. It has been claimed that ALCAR affects the cholinergic central systems through its manifold properties, namely, by increasing the catalyst of L-carnitine acylation to acetyl-L-carnitine, serving as a precursor for acetylcholine, maximizing energy production, promoting cellular membrane stability, counteracting membrane changes, particularly those related to age, and by supporting physiological cellular functioning at the mitochondrial level [15,13,28,29]. We suggest that ALCAR treatments from an age of 6 to 12 months can, with various different degrees of impact, counteract age-dependent neurodegenerative central and cognitive function changes—and perhaps peripheral motor pathologies which are considerably accelerated during aging. ALCAR treatment during ageing seems to strongly influence the activity of the cholinergic basal complex, so as to facilitate cortical activation, attention, learning and mnemonic processes.

4.2. Morphology and Estimation of BF ChAT+ Cells

Our morphological analysis focused on the myelo- and cytostructural BF regions where cholinergic cells are located [25,26]. Our results suggest that the average values of the morphological parameters of the ChAT-immunoreactive neurons in the BF of 12-month-old control rats (ChAT-CT-12) were lower than those of the cholinergic cells observed in the BF of young rats [24] and lower than the cholinergic neurons observed projecting into the meso- and isocortex [22] but similar to the cholinergic cells projecting into the caudal hypothalamus [23]. Interestingly, the large diameter of the ChAT+ cells in the BF of the ALCAR-treated 12-month-old rats was similar to the large diameter of the ChAT+ cells calculated in 518 cholinergic cells in the young control rats [24].

Accurate morphological analysis revealed that the average size of ChAT+ neurons in the anterior portion of the substantia innominata of the spatial learning/memory-impaired control rats was significantly smaller than that in the 12-month-old ALCAR-treated rats. The average of the large diameter of the ChAT+ neurons of the substantia innominata of the spatial learning/memory-impaired control rats was significantly smaller than that in the 12-month-old ALCAR-treated rats [16].

region. The ChAT+ neurons in the Sla of the 12-month-old control rats were significantly smaller than those in the ALCAR-treated rats, measuring 14.0 ± 0.9 μm vs. 15.3 ± 0.4 μm in long diameter on average ($P < 0.05$) [24].

Caudally, behind the crossing of the anterior commissure, and dorsally, deep in the magnocellular preoptic nucleus where transverse fibers sweep under the globus pallidus and caudate putamen, ChAT+ cells are distributed through the posterior portion of the substantia innominata. Rostrally, the cluster of ChAT cells are oriented in parallel with passing fibers of the stria medullaris and inferior thalamic peduncle, which turn dorsomedially into the rostral pole of the thalamus, and also with the perforating blood vessels which traverse the magnocellular preoptic nucleus to continue through the substantia innominata into the lentiform nucleus. In such aggregates the ChAT+ cells are multipolar neurons that form vertically oriented clusters by parallel alignment of soma and dendrites. We observed more ChAT+ cells in the Sla of the ALCAR-treated rats [(ChAT-ALCAR-12+ (Nos. 2, 3, 4)] than in the control rats [ChAT-CT-12+ (Nos. 1, 2, 3)] (on average 1894 vs. 1387, respectively) ($P < 0.05$) [24].

4. Discussion

4.1. Morris Behavioral Water-Maze Task

A possible limitation to this study is that we did not evaluate or record the motor performance or the distance traveled in cm or the time spent in the correct quadrant or the representative trajectories of the two rat groups during swimming performance to find the platform.

Table 3. Average crude cell counts and average estimated total cholinergic viz. choline acetyltransferase-positive (ChAT+) neurons in the basal forebrain of 12-month-old control rats (ChAT-CT-12+ Nos. 1, 2, 3) and of ALCAR-treated rats (ChAT-ALCAR-12+ Nos. 2, 3, 4).

<table>
<thead>
<tr>
<th>Structure</th>
<th>Sites sampled</th>
<th>Crude Counts ± SD</th>
<th>Estimated Totals ± SD</th>
<th>Probability of t-test</th>
</tr>
</thead>
<tbody>
<tr>
<td>ChAT-CT-12 + Nos. 1, 2, 3</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sia</td>
<td>2</td>
<td>73 ± SD 5*</td>
<td>1387 ± 95*</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>Total BF</td>
<td>6</td>
<td>601 ± 202</td>
<td>11,508 ± 3868</td>
<td>n.s.</td>
</tr>
<tr>
<td>ChAT-ALCAR-12+ Nos. 2, 3, 4</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sia</td>
<td>2</td>
<td>103 ± 19*</td>
<td>1894 ± 349*</td>
<td></td>
</tr>
<tr>
<td>BF</td>
<td>6</td>
<td>820 ± 176</td>
<td>15,288 ± 3281</td>
<td></td>
</tr>
</tbody>
</table>

1. Labeled ChAT+ cells on the right side of the brain were plotted from A10.9 to A6.9 (16, 22, 23, 24, 25) and their number tabulated for each structure in 25-μm thick sections at intervals of 800 μm. The average crude cell count (±SD) and the estimated total number for each structure are presented;
2. The averages of the estimated totals were obtained by multiplying the cell count by the number of sections per interval (32), corrected for the average BF cell size of each structure and 25-μm thick sections at intervals of 800 μm, using Abercrombie’s formula (16, 22, 23, 24, 25); 3) (*) Student’s t-test showed that there was a significant number of ChAT+ neurons in the substantia innominata pars posterior of the ALCAR-treated rats (ChAT-ALCAR-12+ Nos. 2, 3, 4) in comparison with the control rats (ChAT-CT-12+ Nos. 1, 2, 3).
innominata pars anterior found in the 12-monthold ALCAR-treated rats was similar to the average large diameter of the ChAT+ cells in the substantia innominata pars anterior of the young rats [24].

Accurate morphological analysis and estimation of the ChAT cells number revealed that the average estimated number of ChAT+ cells throughout the substantia innominata pars posterior of the spatial learning/memory-impaired control rats was significantly lower than that estimated in the 12-month-old ALCAR treated rats (1387 vs. 1894, respectively; P < 0.05). The number of cholinergic cells estimated in the ALCAR-treated rats was nearly the same as that estimated in the substantia innominata pars posterior in the younger rats (approximately 2182) [24].

Interestingly, ALCAR treatment from the age of 6 to 12 months was found to preserve a large number of cholinergic neurons in the BF of the rats (15, 288). This estimated value is similar to that in the young rats (18, 236, [24]) and higher than that reported in 21-month-old ALCAR-treated rats [8,15].

Despite differences in immunocytochemical techniques and rat strain and sex, the results of this study do partly agree with previous reports that: 1) there is a reduction in NGF+/putative-cholinergic neurons in the BF of rats with longer latency on the spatial water-maze task [10,11] aging causes progressive shrinkage of acetyl cholinesterase (AChE)-positive neurons in the BF nuclei [20]; 3) there was a 30% - 50% reduction in ChAT and AChE activities in the BF [29]; 4) there was a significant reduction in AChE-positive cells in the BF of 24- and 31-month old Wistar rats [16].

To summarize, treatment with ALCAR in rats aged between 6 and 12 months can significantly boost performance on the spatial learning/memory Morris water-maze task and may importantly and significantly prevent the loss of the mango cellular cholinergic neurons and the change in size within some cholinergic subgroups. The largest diameter of BF cholinergic cells in the 12-month old ALCAR-treated rats was similar to that of the ChAT+ cells in the BF [24], yet similar to those of the cholinergic cells projecting into the prefrontal and parietal cortices [21].

In conclusion, this study confirms that early chronic ALCAR treatment, from the age of 6 months, may better rescue cholinergic neurotransmission which is known to decline with age [15,22,24,25].

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