Implication of JAK1/STAT3/SOCS3 Pathway in Aging of Cerebellum of Male Rat: Histological and Molecular study

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Aging causes morphological and functional changes in the cerebellum. This work aimed to demonstrate the implication of JAK1/STAT3/SOCS3 on aging–induced changes of rat cerebellum. Thirty male rats were divided into: adult (12 months), early senile (24 months) and late senile (32 months) groups. Immunohistochemical reaction of the cerebellum to GFAP and caspase-3 was assessed and the expression of JAK1, STAT3, SOCS3 proteins was also evaluated. TNFα as well as the activities of malondialdehyde (MDA) and reduced glutathione (GSH) in cerebellar tissue were also measured. The cerebellum of late senile rats revealed more degenerative changes than early senile rats in the form of increase in GFAP and caspase-3 immunoreaction. Additionally, there was decrease in JAK1 and STAT3 expression in early and late senile rats and increase in SOCS3 when compare early and late senile groups with adult one. Enhancement of TNFα was noticed with aging as well as significant decrease in GSH and increase in MDA in early senile group. Moreover, late senile group revealed significant decrease in GSH and increase in MDA. It could be concluded that aging resulting in variable changes of the cerebellum as detected by morphological changes, immunohistochemical reactions of caspase-3 and GFAP and expression of JAK1/STAT3/SOCS3 proteins. Additionally, inflammatory marker TNFα and the activity of oxidative/antioxidative stress markers; malondialdehyde (MDA) and reduced glutathione (GSH) were also affected with aging.

The cerebellum is a fundamental part of the brain that is responsible for movement and awareness activities, such as training of motor activity, recognition of time and fine movements1. The cerebellar cortex acts for designing motor functions, storing memories and adapting some behaviors, so it is the focus of the different research aspects including structural morphology, components of nerve cell and connections of fibers2.

Astroglial cells play a crucial role in aging-induced changes of the cerebellar cortex of rat; astrocyte is a kind of glial cells that is largely affected by aging more than other glial cells. GFAP is extensively used as an indicator of astrogliosis as it is mostly present in fibrous astrocytes as one of their filamentous content3. Astrogliosis or gliosis is an astroglial proliferation that is accompanied by several brain injuries including aging and the regulation of gliosis is related to changes in GFAP expression4.

Caspase-3 is one of cysteine proteinases family that mediates apoptosis in a successive manner by destructing the related components within the cell. Additionally, caspase-3 is mediated by death receptor on the cell surface and mitochondrial apoptosis pathways5,6.

Signal transducers-Janus kinase (JAK) and transcription activators (STAT) are newly discovered proteins that their intracellular signaling pathway affects the replication of DNA and presentation of genetic material that are included in cellular stimulation, multiplication, immunological reactions and programmed cell death. The JAK-STAT link controls the transport of knowledge and chemical signals from the cytoplasm to the nucleus through the plasma membrane mediated by the STAT proteins7.

The signaling pathway of JAK-STAT in the brain is primarily associated with regulation of gene during development, release of hormone, inflammatory processes or carcinogenesis. The different types of JAK and STAT

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are presented in many regions of the central nervous system, like cerebrum, hippocampal tissue and cerebellar cortex.

The pathway of JAK-STAT could be controlled by many factors as SOCS protein; also known as suppressor of cytokine signaling that can negatively affect the duration of action and severity of cytokine-induced JAK/STAT signaling. Oxidative stress has a crucial effect in the pathology of normal aging. It was found that STAT3 protects from a variety of stresses that induce the formation of reactive oxygen species (ROS)9.

Aging is associated with decreased immunity and chronic inflammation10. Senescent cells act as a source of chronic inflammation due to senescence-associated secretory phenotype (SASP) secreted by those cells such as inflammatory cytokines and chemokines, which are mediators of inflammation and intracellular signaling11.

Aging causes morphological and functional changes in the cerebellum12. The cerebellar ultrastructure, immunohistochemical and molecular roles and signaling that explain the pathological effects of aging on cerebellar cortex are not yet completely studied.

The present study aimed to demonstrate the impact of JAK1/STAT3/SOCS3 signaling link on the age-related histological and ultrastructural changes of the cerebellar cortex of male rat. In addition, the study analyzed the age-related cerebellar immunohistochemical expression of GFAP and caspase-3, inflammatory process by measuring TNFα as well as the activity of oxidative/antioxidative stress markers; malondialdehyde (MDA) and reduced glutathione (GSH).

**Material and methods**

**Animals and experimental design.** This study was carried out on thirty Wistar male albino rats; the sample size was calculated by G*Power 3.1.9.4. The rats were obtained from the Animal and Experimental house of Faculty of Medicine, Cairo University, Egypt. The animals were environmentally acclimatized for 5 days before euthanization and the international rules for handling, care and use of rats in the experimental study were followed with allowance of free access to drink water and rat pellet diet. The rats were housed in metal cages measured 41 × 29 × 16 cm under standard diurnal conditions of light/dark cycle at 14 h: 10 h with temperature at 22 ± 2 °C. The approval for the experimental design was taken from Cairo University Institutional Animal Care and Use Committee, Egypt (CU/III/F/68/19). This included the method of animal handling, and euthanasia. All the precautions were followed to minimize animal suffering. The guidelines for the experimental processing were in accordance with the standards of National Guide for Care and Use of Laboratory Animals (NIH Publications No. 8023, revised 1978).

The rats were divided into three age groups as follows:

**Group I (adult):** this group consisted of 10 rats of 12 months age.

**Group II (early senile):** this group consisted of 10 rats of 24 months age.

**Group III (late senile):** Consisted of 10 rats of 32 months of age.

The rats of 12, 24, 32 months age were parallel to thirty, sixty, eighty years old in humans, respectively13.

**Tissue sampling.** **Light microscopic examination.** The rats were euthanized at 12 pm by single intraperitoneal injection of sodium pentobarbital at dosage of 60 mg/kg bw. The skulls were opened and the cerebellas from 5 rats of each group were put immediately in a fixative solution of ten percent of formol saline and kept for one day at the standard atmospheric temperature of the laboratory center. After that, the cerebellar tissues were rinsed in ascending concentrations of alcohol (70%, 90%, 100%) for dehydration followed by immersion in xylene lasting 2 hrs. Tissues were embedded in paraffin blocks; at first left in loose wax followed by hard wax at 60°C for further 1 hr, then, 4-5 µm thickness sections were cut for the following.

Hematoxylin and eosin staining. To study the histological structure of cerebellum in all groups by using light microscopic examination.

Immunohistochemical study. Immunostaining was done by dehydration and de-waxing of paraffin blocks then cerebellar sections of 5 µm thick were cut and buffered with 10-mM sodium citrate for antigen retrieval, 0.9% hydrogen peroxide in absolute methanol was used for blocking activity of internal peroxidase. Not related antibodies were blocked by addition of protein for 1 h (ab64226, Abcam, Cambridge, UK). Incubation of sections was done with mouse monoclonal GFAP antibodies (Abcam, Cambridge, UK) 1:20 dilutions for 1 h. Sections were rinsed with mouse monoclonal GFAP antibodies (Abcam, Cambridge, UK) 1:200 dilutions and rabbit monoclonal caspase-3 antibodies (Abcam, Cambridge, UK) 1:200 dilutions for 1 h. Sections were rinsed in phosphate buffered saline (PBS) then heated with Rat IgG antibody (ab150160, Abcam, Cambridge, UK), followed by washing in PBS and incubation with Novolink polymer (Novo-castra). Deletion of the primary antibody in automated staining design and tonsillar human tissue were done for the 2 negative controls for GFAP and caspase-3.

Histomorphometric measurement of GFAP and caspase-3 immunoreacted sections. The area percent of GFAP-immunostained astrocytes as well as caspase-3 cytoplasmic immunoreaction was measured in 10 non-overlapping microscopic fields picked randomly from each slide. Four sections were analyzed for one cerebellum and this technique was applied to the cerebellum of each rat in a given group, i.e. 40 measurements were taken from each rat group. This histomorphometric measurement was assessed by using image analyzer (Leica Qwin 500 software Image computer system Ltd., Cambridge, England). Images were captured live on to the screen from sections under a light microscope (Olympus Bx-40, Olympus Optical Co. Ltd., Japan) then the video images were digitalized. The cells stained with dark brown colour were considered positive.
Electron microscopic examination. From the other 5 cerebellar cortices from each group, 1mm$^3$ in thickness was immediately fixed in 3% glutaralddehyde, buffered with phosphate buffered saline (PBS) and processed for preparation of semithin slices. Ultrathin sections were prepared and mixed with uranyl acetate and lead citrate. Transmission electron microscopic (TEM) analysis was carried out by a JEM-1400 (JEOL, Tokyo, Japan).

Detection of JAK1, STAT3 & SOCS3 by Western Blotting method (using V3 Western Workflow™ Complete System, Bio-Rad®, Hercules, CA, USA). Extraction of proteins from cerebellar tissue homogenates was done by using ice-cold radioimmunoprecipitation assay (RIPA) buffer then centrifuged at 12,000 rpm for twenty minutes. Similar concentrations of protein (20-30 µg of total protein) were separated by 10% acrylamide gel electrophoresis and transferred to polyvinylidene difluoride membranes (Pierce, Rockford, IL, USA). The membranes were rinsed with PBS after transfer and were blocked with 5% skimmed milk powder in PBS for one hour at room temperature. The blots were formed using antibodies for JAK1, STAT3, SOCS3 and beta actin gained by (Thermoscientific, Rockford, Illinois, USA) and heated at 4 °C overnight with slight shaking. After washing, secondary antibodies with marked peroxidase were added, and the membranes were heated for one hour. Band intensity was analyzed by ChemiDoc™ imaging system with Image Lab™ software version 5.1 (Bio-Rad Laboratories Inc., Hercules, CA, USA). The results were expressed following normalization for β-actin protein expression.

Measurement of TNF alpha by ELISA. The kit was supplied by MyBiosource, USA.

Measurement of oxidative marker (malondialdehyde-MDA). One hundred milligrams of the cerebellar tissue were homogenized in 1 mL phosphate buffered saline, pH 7.0 with the micropestle. 20% Trichloroacetic acid (TCA) was added to cerebellar tissue homogenate precipitating the protein and then centrifuged. Thiobarbituric acid (TBA) solution was added to the supernatants after their collection then was boiled for 10 minutes in a water bath then the absorbance was measured. The concentration of MDA in supernatants of cerebellar homogenate was calculated using the standard curve.

Measurement of antioxidant marker (reduced Glutathione-GSH). It depends on the reduction of 5,5dithiobis (2-nitrobenzoic acid) (DTNB) with reduced glutathione (GSH) producing a yellow compound. The reduced chromogen is directly proportional to GSH concentration and its absorbance can be measured at 405 nm by using its kit.

Estimation of the thickness of the layers of cerebellar cortex. The thickness of the different layers of the cerebellar cortex (the molecular and granular layers) was measured using Leica Qwin 500 LTD image analyzer computer system (software Qwin 500, Cambridge, UK). In each animal of the different age groups, the used sections were stained with hematoxylin and eosin under magnification X 400.

Data analysis. The data obtained from the image analyzer were expressed as mean ± SD and analyzed using the SPSS (Statistical Package for Social Sciences), version 21, Chicago, USA. The data were subjected to one-way ANOVA test for comparison between the different groups. Results were considered statistically significant with the value of ($p \leq 0.05$) and highly significant with the $p$ value $\leq 0.001$. Data was tabulated and represented.
graphically. The difference; either increase or decrease in all studied parameters were expressed as a percent by using special statistical equation.

**Ethical approval.** All applicable international, national, and/or institutional guidelines for the care and use of animals were followed.

**Results**

**Aging–induced histological changes in the cerebellum.** Microscopic examination of H&E-stained cerebellar tissue showed that early senile rats (group II) exhibited degenerative changes including damaged purkinje cells which were surrounded with vacuolated neuropil, areas of complete disappearance of purkinje cells, and vacuolated areas in the molecular layer. The hematoxylin and eosin – stained sections of the late senile

|                | Adult Mean ± SD | Early senile Mean ± SD | Late senile Mean ± SD | P-value                                                                 |
|----------------|-----------------|------------------------|-----------------------|-------------------------------------------------------------------------|
| GFAP           | 25.0500 ± 2.55  | 29.7133 ± 1.90         | 41.3167 ± 6.46        | Early senile versus adult p = 0.21 NS                                   |
|                |                 |                        |                       | Late senile versus adult p < 0.001 **                                   |
|                |                 |                        |                       | Late senile versus early senile p = 0.001 **                            |
| Caspase-3      | 23.7533 ± 1.44  | 29.71 ± 2.08           | 42.50 ± 5.11          | Early senile versus adult p = 0.021†                                    |
|                |                 |                        |                       | Late senile versus adult p < 0.001 **                                   |
|                |                 |                        |                       | Late senile versus early senile p < 0.001 **                            |

Table 1. Effect of aging on GFAP and caspase-3 immunoreaction in all groups. SD: standered deviation; NS: non-significant (p > 0.05); †statistically significant (p ≤ 0.05); **highly significant (p ≤ 0.001).
rats of group III revealed areas of complete loss of the Purkinje cell layer. The few remaining Purkinje cells appeared destroyed and surrounded with vacuolated neuropil. Vacuolated areas in the molecular and the granular layers were noticed. Additionally, interstitial hemorrhage in the medulla was detected (Fig. 1a–c).

Effect of aging on density of glial fibrillary acidic protein (GFAP) immunoreaction in the granular layer of the cerebellar cortex. Immunohistochemical study of GFAP in the cerebellar cortex sections revealed astrocytes that appeared dark brown in the granular layer. There was slight GFAP immunoreaction in adult group (Fig. 2a), while in early senile group; there was mild increase in GFAP (Fig. 2b). Late senile group revealed extensive increase in density of GFAP in the granular layer (Fig. 2c). There was high significant increase (p ≤ 0.001) in GFAP immunoreaction in late senile rats as compared to either adult rats or early senile rats by 65% and 39%, respectively. However, there was statistically non-significant difference (p = 0.21) regarding GFAP immunostaining in early senile rats as compared to adult one (Fig. 2d, Table 1).

Assessment of aging–induced cerebellar apoptosis by immunohistochemical study of caspase-3. The sections from the cerebellum reacted immunohistochemically to caspase-3 as follows: group I (adult male rat) revealed mild caspase-3 cytoplasmic immunoreaction (Fig. 3a). However, the sections from group II (early senile rat) demonstrated an increase in caspase-3 immunoreactivity localized in the cytoplasm of the cerebellar cells (Fig. 3b). In addition, the examination of the specimens from groups III (late senile) showed extensively caspase-3 cytoplasmic immunoreactivity (Fig. 3c). There was statistically significant moderate increase (p = 0.021) by 25% in caspase-3 immunoreaction in early senile rats as compared to adult rats, also there was highly statistically significant increase (p < 0.001) regarding caspase-3 immunostaining in late senile rats as compared to either adult or early senile rats by 79% and 43%, respectively (Fig. 3d, Table 1).
Figure 4. (a, b) Electron micrographs from rat cerebellum of group I (adult) showing purkinjie cell (P) with euchromatic nucleus (N). The perikaryon displaying healthy mitochondria (M) and Golgi apparatus (GA). Granular cell (G) and myelinated nerve axons with regular compact myelin sheath (arrow) are also seen. (TEM; a X 5000, b X 6000). (c, d) Electron micrographs from rat cerebellum of group II (early senile) revealing purkinjie cell (P) containing mitochondria with disrupted cristae (M). Note the lipid droplet (L) in the surrounding neuropil. Granular cells (G) with shrunken nucleus (N), mitochondria with disrupted cristae (m), indented nuclear membrane (thick arrow) and many glial cells (GL) are also seen. Note the disrupted myelin sheath (thin arrows) and the vacuolated axoplasm (a) and longitudinally-cut ramified dendrite of purkinjie cell (asterisk). (TEM; c X 8000, d X 6000, n value = 10 rats for each age group). (e–h) Electron micrographs from rat cerebellum of group III (late senile). (e) Demonstrating purkinjie cell (P) containing shrunken nucleus (N) with irregular membrane (wavy arrow), irregular nucleolus and numerous cytoplasmic lysosomes (double splitted arrow). Many astrocytes (AS) with shrunken nucleus (n) and markedly vacuolated cytoplasm (V) are also seen. Note the swollen astrocytic process (AP). (f) Revealing many ballooned mitochondria (M)
within the purkinjie cell as well as granular cells (G) with irregular nucleus (n), vacuolated cytoplasm (V) and ballooned mitochondria with disrupted cristae (m) are also observed. g) Displaying granular cells (G) with vacuolated cytoplasm (V) and congested blood capillary (BC). The endothelial cell (ED) showing irregular nucleus. Intersticial edema (E) and many glial cells (GL) are noticed. h) Showing disrupted (D) and splitted (arrow) axonal myelin sheath, vacuolated axoplasms (a), mitochondria with disrupted cristae (M) and edema surrounding the axons (E). (TEM; e X 2000, f X 6000, g X 5000, h X 12000, n value = 10 rats for this age group).

Aging–induced ultrastructural changes in the cerebellum. Electron microscopic examination of adult rat cerebellar cortex revealed purkinjie cell with euchromatic nucleus. The perikaryon demonstrated healthy mitochondria and Golgi apparatus. There were normal granular cell and regular myelin sheath of the myelinated nerve axons (Fig. 4a,b).

Cerebellar sections from early senile rats using electron microscope revealed purkinjie cell had mitochondria with disturbed cristae. The granular cells demonstrated shrunken nuclei, indented nuclear envelope and disturbed mitochondrial cristae as well as many glial cells. The axons showed disrupted myelin sheath and vacuolated axoplasms. Lipid droplets surrounding neuropil were noticed (Fig. 4c,d).

Ultrastructural examinations of cerebellar sections of the late senile rats demonstrated purkinjie cell with shrunken nuclei and irregular nuclear envelope, ballooned mitochondria and numerous cytoplasmic lysosomes. The granular cells showed markedly vacuolated cytoplasm with shrunken irregular nuclei and damaged mitochondria. Many glial cells as well as congested blood capillaries with damaged endothelial cell nuclei and interstitial edema were also found. The axons demonstrated disrupted and splitted myelin sheaths, vacuolated axoplasms, mitochondria with disrupted cristae and edema surrounding the axons (Fig. 4e–h).

Modulation of the thickness of the different layers of cerebellum with aging. Morphometric measurement of the different layers of the cerebellum revealed statistically highly significant decrease (*p < 0.001*) in the whole thickness of the cerebellar cortex with early senile and late senile groups by 28% and 58%, respectively as compared to adult group. Moreover, high significant decrease (*p < 0.001*) by 41% was observed in late senile group as compared with early senile one. Additionally, there was highly significant decrease (*p < 0.001*) in the molecular layer of the cerebellar cortex of rat with aging (early senile and late senile) by 31% and 66%, respectively as compared to adult group. Furthermore, significant decrease by 51% was noticed high (*p < 0.001*) in the molecular layer of late senile group when compared with early senile group. There was statistically significant increase (*p = 0.012*) by 78% in the granular layer in early senile group as compared to adult one and highly significant increase (*p < 0.001*) in this layer in late senile group as compared to either adult or early senile groups by 4 and 2 folds, respectively. In addition, moderate significant decrease (*p = 0.027*) by 13% was noticed in the purkinjie layer of early senile group in comparison with adult one and high decrease (*p < 0.001*) was observed in the purkinjie layer in late senile group as compared to either adult or early senile groups by 58% and 52%, respectively (Fig. 5, Table 2).

Changes of JAK1, STAT3 and SOCS3 signaling pathway with aging: (Figs. 6, 7 and Table 3).

Western blotting technique was used to identify the activities of JAK1, STAT3 and SOCS3 proteins in the cerebellar homogenates. There was statistically significant decrease (*p = 0.001*) in the expression of JAK1 by 6% in the early senile group as compared with adult group. Additionally, the decrease in JAK1 was high (*p < 0.001*) when compare late senile group with either adult or early senile groups by 79% and 78%, respectively.

Regarding STAT3, there was statistically highly significant decrease (*p < 0.001*) in late senile group as compared with either adult or early senile groups by 80% and 79%, respectively. However, statistically non-significant difference was observed between early senile and adult groups regarding STAT3.

The increase in SOCS3 was statistically high (*p < 0.001*) when compare the late senile group with either adult group by one and half fold or early senile group by 88%. However, there was no-difference between early senile and adult groups.

Assay of TNFα as a marker of inflammatory induction by aging. ELISA displayed an enhancement of TNFα with aging which was confirmed statistically; early senile and late senile groups revealed highly statistically significant increase (*p < 0.001*) as compared with adult one by 75% and three-fold increase, respectively. In addition, late senile group demonstrated an increase in TNFα by 39% when compared with early senile group (Fig. 8, Table 3).

Modulation of antioxidant marker (GSH) and oxidant marker (MDA) with aging. There was statistically significant (*p ≤ 0.001*) decrease in GSH and increase in MDA in early senile group as compared with adult group by 16.7% and 12%, respectively. Moreover, late senile group revealed statistically significant decrease in GSH and increase in MDA as compared to either adult one by 55% and 5 folds or early senile group by 46% and 4 folds, respectively (Fig. 9, Table 3).

Discussion
The present research assessed aging-induced histological, immunohistochemical and ultrastructural changes of cerebellum. The study emphasized on the age modulation of the cerebellar JAK1/STAT3/SOCS3 as measured by western blotting technique. In addition, the detection of oxidant and antioxidant markers as well as inflammatory marker induced by normal aging was also evaluated.

Histological examination of cerebellar tissue revealed moderate degenerative changes in early senile rats in the form of damaged purkinjie cells which confirmed by disruption of their mitochondrial cristae as evaluated by the electron microscope. Vacuolated neuropil with scattered areas of complete disappearance of purkinjie
The functional decrease was attributed to continuous degenerative changes of the cerebellum in the form of complete loss of the purkinjie cell layer that was confirmed by shrinkage of their nuclei with irregular nuclear envelope as assessed by electron microscope. The few remaining purkinjie cells appeared destroyed and surrounded with vacuolated neuropil. Additionally, ballooned mitochondria and numerous cytoplasmic lysosomes were noticed in ultrathin sections of late senile group. Extensive vacuolated areas in the molecular and granular layers were observed, additionally, interstitial hemorrhage was detected. The granular cells appeared in the ultrathin sections of the late senile group with markedly vacuolated cytoplasm, shrunken irregular nuclei and damaged mitochondria. Henrique et al.16 reported no age-related mitochondrial changes, however regression in the mitochondrial physiology with aging was found by several studies17,18. This functional decrease was attributed to continuous cytoplasmatic collection of Ca^{2+} that resulting in dangerous effects such as apoptosis19. Squier and Bigelow20 added that the mitochondria of senile mice reveal activation of permeability transition pore (PTP) that resulting in enhanced formation of mitochondrial oxidant agents with increased intracellular Ca^{2+} and release of pro-apoptotic components. Luo et al.21 emphasized on the reduction of mitochondria in aged synapses and this decrease may be due to abnormal secretion of neurotransmitter which in turn causing suppression of the electrical impulses from neurons. Earlier, Louis et al.22 attributed the aging declining in the activity of the purkinjie cells to the insufficient amount of noradrenaline in the synapses, particularly in the cerebellum. Furthermore, levels of neurotransmitter hydrolases decrease, which weakens the activity of glutamate dehydrogenase, causing glutamate accumulation which in turn harm the neurons and can cause further neuronal apoptosis23,24. Morphometric measurement of the different layers of the cerebellum revealed statistically significant decrease of the whole thickness of the cerebellar cortex and its molecular layer with aging (early senile and late senile) as compared to adult group. In addition, statistically significant decrease was noticed in the thickness of the purkinjie layer with aging (early senile and late senile groups) in comparison with adult group. Hara et al.25 demonstrated significant decrease of purkinjie cell count in aged rats as the purkinjie cells are the most vulnerable and

| Thickness         | Adult Mean (µm) ± SD | Early senile Mean (µm) ± SD | Late senile Mean (µm) ± SD | P value                                  |
|-------------------|----------------------|-----------------------------|---------------------------|-----------------------------------------|
| Whole thickness   | 3316.77 ± 187.53     | 2395.8 ± 35.79              | 1400.43 ± 210.2           | Early senile versus adult p < 0.001**  |
|                   |                      |                             |                           | Late senile versus adult p < 0.001**   |
|                   |                      |                             |                           | Late senile versus early senile p < 0.001**  |
| Molecular layer   | 2395.60 ± 35.79      | 1643.74 ± 82.22             | 807.75 ± 46.009           | Early senile versus adult p < 0.001**  |
|                   |                      |                             |                           | Late senile versus adult p < 0.001**   |
|                   |                      |                             |                           | Late senile versus early senile p < 0.001**  |
| Granular layer    | 144.04 ± 54.90       | 257.20 ± 24.24              | 994.78 ± 82.27            | Early senile versus adult p = 0.012*   |
|                   |                      |                             |                           | Late senile versus adult p < 0.001**   |
|                   |                      |                             |                           | Late senile versus early senile p < 0.001**  |
| Purkinjie layer   | 251.36 ± 32.14       | 219.36 ± 5.71               | 105.18 ± 0.399            | Early senile versus adult p = 0.027*   |
|                   |                      |                             |                           | Late senile versus adult p < 0.001**   |
|                   |                      |                             |                           | Late senile versus early senile p < 0.001**  |

Table 2. Effect of aging on the thickness (µm) of the different layers of the cerebellar cortex in the different groups. SD: standered deviation; NS: Non significant (p > 0.05); *statistically significant (p ≤ 0.05); **highly significant (p ≤ 0.001).
sensitive to the changes of aging. Nearer findings were found by Zhang et al. who reported reduction in total thickness of the cerebellar cortex secondary to reduction in the molecular layer thickness as a result of decrease in its number of neurons (basket and stellate cells). The later neurons act as inhibitory synapses with purkinjie cells and their destruction could suppress the inhibitory impulse to the purkinjie cells. The decrease in cerebellar cortex thickness is considered the most important factor in aging deterioration of the normal physiologic mechanisms of the brain. Hadj-Sahraoui et al. attributed the reduction in molecular layer thickness to the destructive effect of aging on the foliation of the purkinjie cells dendrites.

The current research work revealed significant increase in the granular layer thickness which was moderate in early senile group and severe in late senile group. Also, many glial cells were noticed in electron microscopic examination of early and late senile cerebellar sections. Moreover, GFAP immunoreaction proved the detection of astrocytes in the granular layer of the cerebellar cortex as there was strong positive GFAP immunoreaction in early and late aged rats in comparison with adult rats as well as increase in GFAP immunostaining in late senile rats as compared to early senile one. The increase in the granular layer thickness with aging is due to astrocytes proliferation in this layer. Jalenques et al.; Sabbatini et al. detected astrocytes by GFAP immunoreaction in variable mammalian species and demonstrated proliferation of these cells in response to aging. The changes in the astrocytes could be attributed to the neurodegenerative changes of aging; however, there was no clear mechanism explains the age-induced GFAP in astrocytic cells. Zhang et al. suggested that the enhancement in the size and count of astrocytes may inhibit the damaging mechanism of aged nerve cells and degeneration of the cerebellar cortex. The authors attributed the astrocytic proliferation to their compensatory reaction to the reduction of the cerebellar cortex thickness by occupying the empty spaces produced by the loss of neurons. Earlier, Hilber and Caston and Caston et al. stated that there is no clear link between reductions in the thickness of the different layers of the cerebellar cortex; however, the suppression in the motor activity with aging is confirmed by the decrease in the count of nerve cells.

Deverman and Patterson demonstrated that cytokines' stimulation of JAK-STAT signaling pathway resulting in transformation of the cortical progenitors into mature astrocytes that in turn express the marker glial fibrillary acidic protein (GFAP). Levy and Darnell revealed that the stimulation of JAK resulting in phosphorylation of
between JAK/STAT pathway with normal as well as abnormal cellular growth. It was found by Guo et al. Furthermore, Smith et al. showed gradual decrease during growth and into adulthood. Schwaiger et al. described inflammation of CNS as a complex mechanism that occurs between different types of brain cells such as microglial, astrocytic, neuronal and endothelial cells. Degos et al. demonstrated that JAK/STAT linkage is activated by release of stimulatory items in response to inflammation. Earlier, Schwaiger et al. reported that the regeneration of axons following optic nerve injury. Moreover, Schindler et al. emphasized on the crucial role of apoptosis in the brain insults that secondary lead to neuronal loss. Sierra et al. reported that the loss of granular cells could affect the motor activity in aged ones as they are confined to conservation of memories along the different durations, regulation of movement mechanisms and control the double directional amplitude of movements.

Immunohistochemical study of adult cerebellar sections of this work revealed high statistically significant increase in caspase-3 immunoreaction in late senile animals in comparison to either adult or early senile groups. Also, there was moderate statistically significant increase regarding caspase-3 immunostaining in late aged rats in comparison to adult. Also, Sun et al. found an increase in the brain apoptotic cells with aging mainly in the granular layer. Sastre et al. attributed the aging-induced apoptosis to the increase in the formation of reactive oxygen radicals with age. Dayer et al. emphasized on the crucial role of apoptosis in the brain insults that secondary lead to neuronal loss. Sierra et al. pointed out that apoptotic cells are scavenged by microglial cells that are primarily located in the brain tissue.

In the present work, activities of JAK1, STAT3 and SOCS3 proteins in cerebellar homogenates were detected by using blotted method. There was reduction in JAK1 and STAT3 expression and enhancement in SOCS3 expression in late aged group in comparison with either adult or early aged groups. However, there was statistically non-significant difference was detected between early senile group and adult regarding STAT3 and SOCS3. De-Fraja et al. pointed out that the expression of JAK/STAT proteins differed throughout the developmental period; their level showed an increase during embryonic period, a particular JAK1 and STAT3 and their level showed gradual decrease during growth and into adulthood. Schwaiger et al. reported that the regeneration of neurons in response to lesion would include the pathway JAK-STAT as the activation and expression of STAT3 is high following axonal injury specifically in regenerating neurons. Moreover, Schindler et al. revealed that stimulation of STAT can induce non-apoptotic proteins transcription. The latter information was coinciding with the result of the present work. Schindler et al. demonstrated that the activation of the ratios of STAT1/STAT3 or STAT1/STAT5 ratio would add to the mechanism of programmed cell death, as STAT3 and STAT5 are more antagonists to apoptosis than STAT1.

In present study, there was an increase in the cerebellar inflammation induced by aging either in early senile or late senile rats as measured by TNFα; a marker of inflammation. Regis et al. confirmed this finding and demonstrated that JAK/STAT link is activated by release of stimulatory items in response to inflammation. Earlier, Schweiger et al. stated that the pathway of JAK/STAT is included in the regeneration of the neurons and limitation of spreading of inflammation in case of injury by forming scars around it. Yu et al. pointed out relation between JAK/STAT pathway with normal as well as abnormal cellular growth. It was found by Guo et al. that STAT3 is closely related to neuronal survival and that the decrease of STAT3 function could be directly related with neuronal death. Herrmann et al. demonstrated that activation of STAT3 would result in formation of the glial scars (astrogliosis) following injury of CNS, an important step for decrease propagation of inflammation. Furthermore, Smith et al. revealed that removal of SOCS3 enhances regeneration of axons following optic nerve injury. This removal in turn, would stimulate gp130-mediated signaling pathway (involving JAK-STAT).

Planas et al. described inflammation of CNS as a complex mechanism that occurs between different types of brain cells such as microglial, astrocytic, neuronal and endothelial cells. Degos et al. added that inflammation follows stimulation of microglial and astrocytic cells that mediated by cytokines, chemokines or growth factors. Moreover, Vezzani et al. mentioned the crucial role of CNS inflammation in saving integrity of the brain against

|                | Adult        | Early senile | Late senile | p value         |
|----------------|--------------|--------------|-------------|----------------|
| JAK1           | 1.02±0.014   | 0.96±0.033   | 0.21±0.011  | Early senile versus adult p=0.001** |
|                |              |              |             | Late senile versus adult p<0.001 **|
|                |              |              |             | Late senile versus early senile p<0.001** |
| STAT3          | 0.86±0.018   | 0.84±0.032   | 0.17±0.014  | Early senile versus adult p>0.05NS |
|                |              |              |             | Late senile versus adult p<0.001 **|
|                |              |              |             | Late senile versus early senile p<0.001** |
| SOCS3          | 0.47±0.018   | 0.51±0.054   | 0.96±0.018  | Early senile versus adult p>0.05NS |
|                |              |              |             | Late senile versus adult p<0.001 **|
|                |              |              |             | Late senile versus early senile p<0.001** |
| TNFα           | 45.23±0.16   | 79.08±15     | 110.28±0.23 | Early senile versus adult p<0.001** |
|                |              |              |             | Late senile versus adult p<0.001 **|
|                |              |              |             | Late senile versus early senile p<0.001** |
| GSH            | 38.25±0.18   | 31.85±0.52   | 17.25±0.18  | Early senile versus adult p<0.001** |
|                |              |              |             | Late senile versus adult p<0.001 **|
|                |              |              |             | Late senile versus early senile p<0.001** |
| MDA            | 29.25±0.18   | 32.71±0.76   | 76.04±0.91  | Early senile versus adult p<0.001** |
|                |              |              |             | Late senile versus adult p<0.001 **|
|                |              |              |             | Late senile versus early senile p<0.001** |

Table 3. Effect of aging on the JAK1, STAT3, SOCS3, TNFα, GSH and MDA in all groups. SD: standard deviation; NS: Non significant (p>0.05); *statistically significant (p≤0.05); **statistically highly significant (p<0.001).
the injurious items but could be worsens several disorders of the brain; involving cerebral ischemia and neuro-degenerative pathologies.

Earlier research by Planas et al.48, emphasized on the effect of reactive oxygen species and interleukins in activation of STAT1 and STAT3 by JAK1-induced effect. This pathway stimulation aimed to enhance the expression of inflammatory genes. In the present work, there was statistically significant increase of GSH (anti-oxidant) and decrease in MDA (oxidant) in early senile and late senile groups in comparison with adult group. Planas et al.48 stated that activation of STAT isoform resulting in variable effects on inflammatory processes, proliferation or differentiation as they may be included in nerve cell survival.

The attachment of cytokines to the receptors of on the surface of the cell membrane could stimulate kinase components of JAKs resulting in their activation. The stimulated JAKs phosphorylate tyrosine particles on the receptors resulting in formation of binding locations for stimulator of STAT proteins. Homo- or hetero-dimers of STAT3 were formed as a result of release of stimulated STATS from the receptors. These dimers move to the nucleus and attached to transcription elements to regulate expression of proliferative or anti-inflammatory genes51.

The present research concluded that aging resulting in variable changes of the cerebellum; apoptosis was confirmed by caspase-3 marker and aging-induced astrogliosis was detected by GFAP. Additionally, inflammatory marker TNFα and the activity of oxidative/antioxidative stress markers; malondialdehyde (MDA) and reduced glutathione (GSH) were also affected with aging. The expression of JAK1/STAT3/SOCS3 proteins was affected by aging and enhancement of JAK1/STAT3 activity with inhibiting of SOCS3 can act as a promising protective line against age-related cerebellar changes.

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**Figure 8.** Bar chart revealing the mean of TNFα in adult, early senile and late senile groups; (***) a highly statistically significant increase in late senile versus either early senile or adult groups. (#) highly statistically significant increase in early senile group versus adult group (p < 0.001). (n value = 10 rats for each age group).

**Figure 9.** Bar chart demonstrating the mean of GSH and MDA in adult, early senile and late senile groups; (#) a highly statistically significant (p ≤ 0.001) decrease in GSH and increase in MDA in early senile group versus adult group. (***) a highly statistically significant decrease in GSH and increase in MDA in late senile group versus either adult or early senile groups. (n value = 10 rats for each age group).
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Competing interests
The authors declare no competing interests.

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