Atorvastatin misuse alters histological and protein expression in the hippocampal CA3 region of male rat models

Glory B Samuel1, Moses B Ekong2* and Monday I Akpanabiatu1

1Department of Biochemistry, Faculty of Basic Medical Sciences, University of Uyo, Nigeria
2Department of Anatomy, Faculty of Basic Medical Sciences, University of Uyo, Nigeria

Abstract

Atorvastatin, a cholesterol-lowering drug blocks cholesterol production through inhibition of 3-hydroxy-3-methylglutaryl coenzyme A reductase. The therapeutic dose of the drug is regarded as safe, but does same apply if misused? This study therefore investigated the histology and structural protein expressions in hippocampal CA3 region in rat models. Thirty male albino Wistar rats weighing 170–210 g were selected into five groups (n=6), and divided into control, 10, 20, 40 and 80mg Atorvastatin (ATVS)/kg body weight of the animal by oral gavages and daily for 28 days. The animals were sacrificed 24 hours after the last drug administration by intracardial perfusion using 10% buffered formalin under ketamine hydrochloride anaesthesia, and processed for histology using haematoxylin and eosin, and Cresyl fast violet methods. Representative sections were immunolabelled for neuron specific enolase (NSE) and glial fibrillary acidic protein (GFAP). Results showed significantly (p < 0.05) less body weight increase in the test groups, with pyknosis, hypertrophy, dark nuclei, less Nissl staining and significantly (p < 0.05) lower cellular population observed in the CA3 of the 20, 40 and 80 mg/kg ATVS groups. There was also decreased NSE and increased GFAP expressions in CA3 of the 20, 40 and 80mg/kg ATVS groups compared with the control group. In conclusions, the misuse of ATVS influenced body weight increase and caused hippocampal CA3 histological modifications, as well as altering the balance of some structural protein expression. These effects were dose-dependent and may lead to loss of the hippocampal CA3 functional activity with resultant neurodegeneration.

Introduction

Atorvastatin, one of the statins groups of drugs, is cholesterol-lowering medication, which acts by blocking cholesterol production through the inhibition of 3-hydroxy-3-methylglutaryl coenzyme A (HMG-CoA) reductase. The reductase then converts the HMG-CoA to mevalonate, the rate-limiting step in cholesterol biosynthesis [1]. Atorvastatin is primarily used to treat dyslipidemia and the prevention of cardiovascular diseases [2,3]. The dose and potency of the statins used and individual responsiveness determines the magnitude of low density lipoprotein cholesterol (LDL-C) reduction and cardiovascular diseases benefit [4]. Thus, statins are the first choice agents for lowering LDL-C and non-high density lipoprotein cholesterol (HDL-C), which are important targets for pharmacotherapy, hence, are generally effective and remain the first choice treatment for high blood cholesterol levels [5,6]. Cholesterol is highly synthesized in the central nervous system during the time of active myelination which occurs in early neural development and however decreases in mature adult brains [7,8]. In adulthood, the level of cholesterol in the central nervous system (CNS) determines its normal functions, and this is independent of the systemic cholesterol level [9,10]. At this stage of life the CNS does not rely entirely on systemic cholesterol production, but on the brain’s inherent capacity to synthesise its own cholesterol because of the limited metabolic turnover during this stage of life [7]. The brain cholesterol is mostly synthesized de novo by astrocytes, as well as the reutilisation of free cholesterol arising from neuronal death [11,12]. As such, reductions in plasma cholesterol concentration following statin treatment are unlikely to cause acute disruption in CNS cholesterol homeostasis [13,14]. Nevertheless, cholesterol level is affected by statins if the neurons and glia functions are disrupted [9,15]. It is reported that loss of astrocytes cholesterol causes neurite impairment [15].

Atorvastatin is reported to have neuroprotective ability, including reduction of glial cell activation and cerebral oedema, and restoration of blood-brain barrier [16-21]. It also protects against traumatic brain injury models, and increases neurogenesis in the dentate gyrus, reduce delayed neuronal death in the hippocampal CA3 region, improve spatial learning in rat after traumatic brain injury, and is also reported to improve memory [16,22,23].

Negative impact of statins especially on the central nervous system has also been reported. Statins are reported to have negative effects on cognition, induction of neurite inhibition and proliferation, with morbidity and mortality effects in high-risk conditions also reported [24-26].

The therapeutic dose of ATVS has shown neuroprotective effects on the hippocampal function, but could same be obtained if misused? Brain cholesterol has been associated with a half-life of from 6 months to 5 years, requiring chronic statin therapy before significant effects on the CNS cholesterol is observable [27-29]. However, can there be

*Correspondence to: Moses B Ekong, Department of Anatomy, Faculty of Basic Medical Sciences, University of Uyo, Nigeria, Tel: +2348030868505; Email: mbe_flashpoint@yahoo.com

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histological changes due to overdose? The present study examines the hippocampal protective ability in situations of abuse as underscored in the expression of neuron specific enolase and glial fibrillary acidic protein.

Materials and methods

Thirty (30) male albino Wistar rats weighing between (170 - 210 g) were obtained from the Animal House facility of the College of Health Science, University of Uyo, Nigeria for this study. Prior to the experiment, ethical permission was obtained from the Faculty of Basic Medical Sciences Ethical Committee, and the animals were handled according to the guidelines of the National Institute of Health of the United States of America.

The animals were housed in wooden cages designed with wire guage roof for ventilation and with provisions for feed and water. They were acclimatized for 3 days before the start of the experiment and maintained under controlled environmental conditions (temperature: 25 ± 5°C and 12 hour dark/ light cycle). All animals were given feed twice daily and were allowed access to water ad libitum. The cages were cleaned and beddings changed twice a week.

The animals were randomly selected on the basis of average weight into five study groups of six rats each. 10 mg of Atorvastatin (ATVS) was solubilized in 40 ml of distilled water and made into a concentration of 0.25 mg/ml. After acclimatization period, the rats were subjected to different dosages of the drug with group 1 (control) receiving an equivalent volume of water as placebo. The test groups: 2-4 received oral doses of 10, 20, 40 and 80 mg ATVS/ kg body weight for 28 days respectively. The body weights were measured prior and after the experiment to determine body weight changes (if any).

At the end of drug treatment period (on the 28th day), the animals were fasted overnight (12hrs) and each animal was anaesthetized with 60 mg/kg ketamine hydrochloride (Rotex Medical, Germany) intraperitoneally. Incision was made through the thoraco-abdominal wall to the thoracic cavity and perfusion-fixed with 10% neutral buffered formalin for 24 hours. Thereafter, the hippocampal region of all the brains were excised and routinely processed for paraffin wax embedding. Serial sections were obtained at 10 μm, and representatives sections were processed for histological studies with haematoxylin and eosin, as well as Cresyl fast violet. The other sections were immunolabelled with neuron specific enolase (NSE) and glial fibrillary acidic protein (GFAP).

Briefly, serial paraffin sections on slides were brought to water and antigen retrieval was performed using citrate buffer (pH 6.0) in a microwave oven for 5 minutes, followed by protein block using 3% hydrogen peroxide for 10 minutes. Sections were thereafter pre-incubated in 2% normal goat serum for 30 minutes and incubated in monoclonal mouse anti-enolase-2 (Novocastra, Leica Biosystems, 22C9, 1:100) for neuron specific enolase and mouse monoclonal anti GFAP (Novocastra, Leica Biosystems, NCL-L-GFAP-GA5, 1:100) for GFAP, followed by 1-hour incubation in goat anti-mouse secondary antibody (1:100) for an hour. Detection of reaction was by means of the avidin-biotin complex with diaminobenzidine as the chromagen.

Cellular density was determined manually by means of ImageJ® software. Briefly, images of the whole hippocampal CA3 region were obtained for each section and randomly mapped with the ImageJ® gridlines. Counting of cell nuclei was done manually taking into consideration the nuclei on the upper and right borders of the mapped areas.

Statistical analysis

One-way analysis of variance using Graphpad Prism was carried out, followed by post hoc Tukey’s test. Data obtained were expressed as mean ± standard deviation. Probability level of p < 0.05 was considered significant.

Results

Body weight change

There were increased body weights of animals in all the groups. However, there were significantly (p < 0.05) lower body weight increases in the test groups compared with the control group. No difference was observed among the test groups (Table 1 and Figure 1).

Histomorphology of the Hippocampal CA3

The hippocampal CA3 region had normal three cortical layers; molecular, pyramidal and polymorphic. In the control group the molecular layer consisted mostly of nerve processes with sparse cellular population. The pyramidal layer was made up of dense pyramidal cells, while the polymorphic layer was mostly nerve processes with sparse cellular population (Figure 2A). The hippocampal CA3 region of

| Groups (n = 6) | Initial Body Weight (g) F = 71.18 | Final Body Weight (g) F = 4.805 | Body Weight Gain (g) F = 11.71 |
|---------------|------------------------------------|---------------------------------|-------------------------------|
| Control       | 174.33 ± 6.33                      | 224.33 ± 19.87                  | 50.00 ± 13.45                 |
| 10 mg ATVS    | 179.83 ± 5.03                      | 192.50 ± 20.75                  | 12.67 ± 15.72 ***             |
| 20 mg ATVS    | 192.5 ± 2.09                       | 205.33 ± 17.67                  | 12.83 ± 15.58 ***             |
| 40 mg ATVS    | 205.33 ± 1.75                      | 209.50 ± 8.41                   | 4.19 ± 6.66 ***               |
| 80 mg ATVS    | 210.67 ± 5.77                      | 220.33 ± 15.00                  | 19.66 ± 9.43 **               |

Data presented as Mean ± Standard Deviation

** - Significantly different from the control group at p < 0.05
*** - Significantly different from the control group at p < 0.001

Figure 1. The Percentage Body Weight Change of the Experimental Groups

Data presented as Mean ± Standard Deviation

** - Significantly different from the control group at p < 0.01
*** - Significantly different from the control group at p < 0.001

ATVS - Atorvastatin

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the 10 mg/kg ATVS group showed amorphous pyramidal cell nuclei compared with the control group (Figure 2B). The hippocampal CA3 region of the 20 mg/kg ATVS group showed the pyramidal cells having pyknotic nuclei compared with the control group (Figure 2C).

The hippocampal CA3 region of 40 mg/kg ATVS group showed hypertrophied and pyknotic pyramidal cells, as well as cells of the polymorphic layer compared with the control group (Figure 2D), while the hippocampal CA3 region of the 80 mg/kg ATVS group also showed pyknotic pyramidal cells, as well as cells of the polymorphic layer compared with the control group (Figure 2E).

There was no difference in cell population between the 10 mg/kg ATVS group, but significantly (p < 0.05) lower number of cells in the 20, 40 and 80 mg/kg ATVS groups compared with the control group. The 20, 40 and 80 mg/kg ATVS groups were significantly (p < 0.05) lower than the 10 mg/kg ATVS group, while the 20 mg/kg ATVS was significantly (p < 0.05) higher than the 40 mg/kg ATVS group. However, there was no difference between the cell population of the 20 mg/kg ATVS and 80 mg/kg ATVS groups as well as between the 40 mg/kg ATVS and 80 mg/kg ATVS groups (Figure 3).

Nissl staining of the hippocampal CA3

The hippocampal CA3 region of the control group showed prominent Nissl stained cells, which was same with that of the 10 mg/kg ATVS group. The hippocampal CA3 region of the 10 mg/kg ATVS group showed pyknotic pyramidal cell nuclei, as well as cells of the polymorphic layer with sparse cellular population. The pyramidal layer consists of dense pyramidal cells (arrows), and the polymorphic layer also has sparse cellular population. The pyramidal layer consists of dense pyramidal cells (arrows), and the polymorphic layer also has sparse cellular population. The pyramidal layer consists of dense pyramidal cells (arrows), and the polymorphic layer also has sparse cellular population.

There was no difference in cell population between the 10 and 40 mg/kg ATVS groups, but significantly (p < 0.05) lower number of cells in the 20 and 40 mg/kg ATVS groups compared with the control group. The 20 and 80 mg/kg ATVS groups were significantly (p < 0.05) lower than the 10 mg/kg ATVS group. However, there was no difference between the cell population of the 20 mg/kg ATVS and the 40 and 80 mg/kg ATVS groups, as well as between the 40 mg/kg ATVS and 80 mg/kg ATVS groups (Figure 3).

Neuron Specific Enolase (NSE) Immunolabelling

The hippocampal CA3 region of the control group showed cytosolic NSE expression throughout the pyramidal cells, with the soma clearly revealed, while the hippocampal CA3 region of the 10 mg/kg ATVS group showed reduced NSE expression in the pyramidal cells compared with the control group. The 20 and 80 mg/kg ATVS groups were significantly (p < 0.05) lower than the 10 mg/kg ATVS group. However, there was no difference between the cell population of the 20 mg/kg ATVS and the 40 and 80 mg/kg ATVS groups, as well as between the 40 mg/kg ATVS and 80 mg/kg ATVS groups (Figure 5).

Glial Fibrillary Acidic Protein (GFAP)

The control group showed GFAP expression throughout the hippocampal CA3 region layers with the soma and processes of the astrocytes clearly revealed (Figure 8A). In the 10 mg/kg ATVS group, GFAP was expressed throughout the hippocampal CA3 region layers, but mostly in the processes of the astrocytes compared with the control group (Figure 8B). In the 20, 40 and 80 mg/kg ATVS groups, GFAP was much expressed throughout the hippocampal CA3 region layers in the soma and processes of the astrocytes compared with the control group (Figure 8C-8E).

**Figure 3.** Cellular population of the Experimental Groups

Data presented as Mean ± Standard Deviation

*** Significant different from the control group at p < 0.001

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kg ATVS of group (Figure 4A,4B). The hippocampal CA3 region of the 20 mg/kg ATVS group showed hypertrophied cells with less Nissl staining (Figure 4C), while the hippocampal CA3 region of the 40 and 80 mg/kg ATVS groups showed some dark cells with some having less Nissl staining compared with the control group (Figure 4D,4E).

There was no difference in cell population between the 10 and 40 mg/kg ATVS groups, but significantly (p < 0.05) lower number of cells in the 20 and 80 mg/kg ATVS groups compared with the control group. However, there was no difference between the cell population of the 20 mg/kg ATVS and the 40 and 80 mg/kg ATVS groups, as well as between the 40 mg/kg ATVS and 80 mg/kg ATVS groups (Figure 7).
There was no difference in GFAP-labelled astrocyte population between the 10 mg/kg ATVS group, but significantly (p < 0.05) higher number of GFAP-labelled astrocytes in the 20, 40 and 80 mg/kg ATVS groups compared with the control group. The 20, 40 and 80 mg/kg ATVS groups were significantly (p < 0.05) higher than the 10 mg/kg ATVS group, while the 20 mg/kg ATVS group was significantly (P < 0.05) higher than the 80 mg/kg ATVS group. However, there was no difference between the NSE-labelled cell population of the 20 mg/kg ATVS and the 40 mg/kg ATVS groups, as well as between the 40 mg/kg ATVS and 80 mg/kg ATVS groups (Figure 9).

Figure 4. Sections of the CA3 of the experimental animals. Molecular (M); Pyramidal (Py); Polymorphic (Pm) layers. CFV, × 400
A: The CA3 of the control group showing prominent Nissl stained cells (arrows).
B: The CA3 of 10 mg/kg ATVS group showing prominent Nissl stained cells (arrows).
C: The CA3 of 20 mg/kg ATVS group showing hypertrophied cells (arrows).
D: The CA3 of 40 mg/kg ATVS group showing some dark cells (arrows).
E: The CA3 of 80 mg/kg ATVS group showing some dark cells (arrows).

Figure 6. Photomicrographs of the hippocampus of the experimental animals. molecular (M), pyramidal (Py) and polymorphic (Pm) layers. NSE, ×400
A: The hippocampus of the control group showing NSE expression (arrows) throughout the pyramidal cells of the hippocampus with the cell bodies clearly revealed.
B: The hippocampus of the 10 mg/kg ATVS group showing reduced NSE expression (arrows) in the pyramidal cells of the hippocampus.
C: The hippocampus of 20 mg/kg ATVS group showing reduced NSE expression (arrows) in the pyramidal cells of the hippocampus.
D: The hippocampus of 40 mg/kg ATVS group showing reduced NSE expression (arrows) in the pyramidal cells of the hippocampus.
E: The hippocampus of 80 mg/kg ATVS group showing reduced NSE expression (arrows) in the pyramidal cells of the hippocampus.

Figure 7. NSE-labelled Cell Population of the Experimental Groups
Data presented as Mean ± Standard Deviation
** - Significantly different from the control group at p < 0.01
*** - Significantly different from the control group at p < 0.001
ATVS - Atorvastatin
This study assessed the effects of the abuse of one of the statins, ATVS on body weight, and the histology and some protein expression of the hippocampal CA3 region of adult Wistar rats. The results showed less body weight increase in the test groups compared with the control group, with no difference among the test groups. This result indicates that the given dose of ATVS significantly alter body weights increase. Body weight is measured by the amount of fat deposits, which reflect in part the rate of caloric intake [30]. It is reported that statins lower body cholesterol levels, which may have been the results observed in the present experiment [5]. Cholesterol deposition may be a means by which body weight increase occur together with other lipid components [30,31]. Thus, this protective activity will ensure protection of the cardiovascular system, as increased adiposity usually result in damage to endothelial cells of arteries and capillary vessels, decrease in blood flow, impairment of metabolism, and the decrease in nutritive and oxygen levels in the brain [32]. As the CNS lipid deposits are not directly proportional to the systemic deposits, this may not have applied to the brain area under study [33].

As the interest of the present study was on memory consolidation, the hippocampal CA3 was paramount in this research. The histology of the hippocampal CA3 of the 10 mg/kg ATVS group showed amorphous pyramidal cells, with that of the 20 mg/kg ATVS group showing pyknosis and less Nissl staining. In the 40 mg/kg and 80 mg/kg ATVS groups, the polymorphic layer and the pyramidal cells appeared hypertrophied with pyknotic nuclei and some dark nuclei with less Nissl staining in others. There was significantly (p < 0.05) lower number of cells in the 20, 40 and 80 mg/kg ATVS groups compared with the control and the 10 mg/kg ATVS groups, while there was no difference in cell population between the 10 mg/kg ATVS and control groups. Pyknosis indicates cell death process and is the most characteristic feature of apoptosis [34]. In adults, apoptosis occurs as a homeostatic mechanism to maintain cell populations in tissues, and also as a defence mechanism against damage [35]. This may have been the case in the hippocampal CA3 of the present study.

There was no difference in cell population in the hippocampal CA3 between the 10 mg/kg ATVS group compared with the control group, an indication that at therapeutic doses adverse effects may not be established. Therapeutic level of most statins is regarded as safe [36]. However, lower cellular population were observed in the 20, 40 and 80 mg/kg ATVS groups compared with the control group, an indication that at therapeutic doses adverse effects may not be established. Therapeutic level of most statins is regarded as safe [36]. However, lower cellular population were observed in the 20, 40 and 80 mg/kg ATVS groups compared with the control group, an indication that higher dosages of this statin can cause cell death processes resulting in cellular loss. This corroborates the cell death process already observed. Ekong et al. [37] reported that rat brain tissues loss of cells is indicative of neurodegenerative process, which may be the case in the present study.

Brain tissue biomarkers are mainly derived from neurons or astrocytes [38,39]. Hence, NSE and GFAP were used in the present study. NSE is an isoenzyme in the cytoplasm of neurons which marks the integrity and activity of neurons [40–42]. In the present study, there was decreased NSE expression in the polymorphic cells of the hippocampal CA3 region of the 20, 40 and 80 mg/kg ATVS groups, with lower population of cells in the 20, 40 and 80 mg/kg ATVS groups compared with the control group. These are indications of the decreased neuronal glycolysis or neuronal degeneration. Decreased neuronal glycolysis leads to a concomitant reduction in neurotransmitter production...
and axonal transport and this is usually associated with degenerative processes [42].

In the 10 mg/kg ATVS group, GFAP was expressed throughout the hippocampal CA3 region layers, but mostly in the processes of the astrocytes, while in the 20, 40 and 80 mg/kg ATVS groups, GFAP was much expressed throughout the hippocampal CA3 region layers in the soma and processes of the astrocytes, with higher population of GFAP-labelled astrocytes in the 20, 40 and 80 mg/kg ATVS groups compared with the control group. GFAP is the hallmark intermediate filament protein in astrocytes, with increased expression indicating the up-regulation of the GFAP protein in the astrocytes, an astroglial activation or astrogliosis process, which is known to occur following CNS trauma and also in the process of neurodegeneration [43,44]. Astrocytes have a range of control and homeostatic functions in health and disease [45]. It has also been reported as a blood biomarker, released during neurodegenerating process into the surrounding interstitial fluid where it appears in the peripheral blood likely via disruptions in the blood-brain barrier [46].

The hippocampal CA3 region is an integral portion of the hippocampal formation and its pyramidal cells serve to connect the dentate gyrus and CA1 [47]. It plays an important role in the encoding of new spatial information within short-term memory and also important for encoding of spatial information requiring multiple trials including the acquisition of arbitrary and relational associations [48]. These processes may be altered with statin administration due to neuronal and glia injuries that may initiate cellular death processes.

**Conclusion**

The abuse of ATVS influences body weight increase and caused hippocampal CA3 histological modifications, as well as altering the balance of some structural protein expression. These effects were dose-dependent and may lead to loss of the hippocampal CA3 functional activity with resultant neurodegeneration.

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