17α-METHYLTESTOSTERONE (ANABOLIC ANDROGENIC STEROIDS) ALTERS ACETYLCHOLINESTERASE ENZYME ACTIVITY IN DIFFERENT PARTS OF THE BRAIN IN FEMALE MICE, MUS MUSCULUS

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ABSTRACT

Aim: Anabolic androgenic steroids (AAS) are synthetic derivatives of the male sex hormone testosterone. Androgens and anabolic steroids have been used for therapeutic purpose with few exceptions. However, the abuse of AAS is a remarkably prevalent problem, particularly among athletes and adolescents. Supraphysiological doses of AAS exert profound effects on mental state and behaviors such as depression, anxiety, aggressiveness, and cognitive deterioration.

Objective: In the present investigation, we studied the impact of one of the AAS compounds, i.e., 17α-methyltestosterone on acetylcholinesterase (AChE) enzyme activity in different brain parts of mice, namely, forebrain, hippocampus, midbrain, and hindbrain.

Methods: The adult female mice were assigned to four experimental groups to which different doses of 17α-MT (0.5, 5.0 and 7.5 mg/kg bwt, respectively) were administrated s.c. for 30 days. A significant increase in AChE activity in forebrain and midbrain (low and medium dose treatment) suggests a reduction of cholinergic neurotransmission efficiency due to decrease in acetylcholine levels in trans-synaptic cleft. Further, concurrent reduction in AChE activity was observed in whole brain, hippocampus, and hindbrain of 17α-MT-treated mice suggests the impairment in neuronal transmission. Since the regulation of cholinergic system through acetylcholine hydrolysis has been largely attributed to AChE activity, a significant reduction in its activity may lead to stress-related anxiety, memory loss with some cognitive and behavioral aspects in the mice.

Conclusion: Based on the observed results, we propose that 17α-MT, an alkylated steroid compound, has a negative impact on AChE enzyme activity in different parts of mice brain, leading to impairment in neuronal transmission.

Keywords: Anabolic-androgenic steroids, Acetylcholinesterase, Forebrain, Hippocampus, Midbrain, Hindbrain.

INTRODUCTION

Anabolic androgenic steroids (AAS) are, more often than not, widely abused by professional athletes, bodybuilders, and non-athletes to enhance athletic performance, muscle mass, promote self-confidence, and relieve fatigue. Indeed, AAS was originally designed for the clinical purpose for the treatment of endometriosis, aplastic anemia, male hypogonadism, and inoperable breast cancer [1,2]. However, doping with these AAS results in a broad spectrum of secondary adverse effects such as metabolic suppression, especially lipoprotein metabolism, hepatotoxic effects, antagonism of glucocorticoid effects, impairment in cardiovascular system, gonadal dysfunction, gynecomastia, and behavioral and psychiatric side effects [2-5]. Further, of late, the use of AAS is becoming increasingly widespread among the adolescent girls and woman athletes, to improve their muscle mass, physique, appearance, and euphoria [6-9]. The actual influence of these compounds on overall physiology, central nervous system (CNS), and neurobehavioral alteration is yet to be established clearly.

A previous study from our laboratory reported that administration of a AAS compound Stanozolol leads to acceleration of granulopoiesis and stimulates an immune response (at physiological level only), though it alters the lipoprotein profile in mice [10]. Androgenic neurotransmission in the mammalian CNS is regulated predominantly by the enzyme acetylcholinesterase (AChE, EC 3.1.1.7).

AChE is one of the efficient primary cholinesterases which terminates acetylcholine-mediated neurotransmission, by rapidly hydrolyzing acetylcholine (ACh) into acetate and choline [12,13]. Further, ACh is the major neurotransmitter involved in cortical activation, attention, memory, learning, pain, control of motor tone, locomotion, and control of autonomic functions [14]. It is known that androgen receptors are present in most of the body parts including CNS and influence many neuronal functions through both classical genomic effects and also rapid membrane effects. Sustained use of AAS has an impact on those areas of the CNS that are involved in emotional and cognitive responses such as sexuality, anxiety, aggression, memory, and development of drug dependence [15-19]. The androgen 17α-methyltestosterone (17α-MT), an alkylated steroid compound at 17α position of steroid nucleus and the most frequently abused AAS compound, has been selected for the present study to investigate its impact on different areas of the brain which are associated with wide range of functions including emotional and cognitive regulation.

METHODS

2½-month-old female mice exhibiting normal estrous cycle (25-30 g) were obtained from the mice colony, maintained in the Department of Zoology, Karnatak University, Dharwad. They were housed in individual cages at room temperature (27±1°C) with natural-cum-artificial light for 12–14 h and fed with a pelleted diet (Gold Mohur, Lipton, India) and water ad libitum.
Hormone treatment
17α-MT was obtained from Sigma-Aldrich Chemical Co., USA. The mice were weighed and allocated to four experimental groups comprising five animals in each (n=5). Doses of 17α-MT administered in the experiment were so chosen as to stimulate the range of doses taken by human female users [20]. The first experimental group that received 1% of alcohol acted as a baseline (control). The 2nd, 3rd, and 4th group received 17α-MT daily, for 30 days, by s.c. injection in 1%, respectively (LD - low dose-0.5; medium dose (MD) -5.0; and High dose (HD)-7.5 mg/kg bwt). On the 31st day, the mice were sacrificed by the inhalation of diethyl ether.

Separation of different parts of brain
After sacrificing the animal, brain was extracted out from the skull and rinsed in cold phosphate-buffered saline to remove the excess blood. Before the separation of brain parts, it was kept on the cold metal plate placed on the crushed ice in a tray. After confirming its hardness precisely separated the different brain parts, namely, forebrain, midbrain, hindbrain, and hippocampus regions. Each brain part was homogenized and stored at −80°C until assay.

Quantification of AChE enzyme activity
Fresh brain from the treated as well as the control mice, immediately after autopsy, were homogenized in phosphate buffer (pH 7.6) and centrifuged for 10 min at 8000 rpm. The supernatant was used as the source of the AChE activity. The enzyme activity was estimated as per the procedure described elsewhere [21,22]. The incubation mixture comprised of 0.2 ml of 20 mM phosphate buffer (pH 7.6), 0.1 ml of 8 mM acetylcholine iodide, and 40 µl of tissue supernatant. The incubation was carried out at room temperature for 30 min. The reaction was terminated by adding 1.8 ml of (5,5-Dithiobis(2-nitrobenzoic acid). The absorbance was measured immediately at 412 nm using ultraviolet-VISIBLE spectrophotometer (Hitachi, U-2800).

Statistical analysis
Data were expressed as a mean±standard error comparison of normally distributed variables across groups were made using one-way analysis of variance followed by Tukey’s post hoc test. The level of statistical significance was set at p<0.01. Data were analyzed using SPSS (version 16.0) and the recorded values are summarized in figures.

RESULTS
The observed results are depicted in Figs. 1 and 2. The whole brain homogenates of all the 17α-MT treated mice revealed a significant depletion in AChE activity (F_0,32 = 6.80, P < 0.05; Fig. 1a) as compared with control. As shown in Fig. 1b, the activity levels of AChE in forebrain region were found to be elevated significantly (F_{1,32} = 138.48, p<0.01). In contrast, a significant depletion in AChE activity was observed in hippocampus (F_{1,32} = 128.16, p<0.01; Fig. 1c) as well as hindbrain regions (F_{1,32} = 185.8, p<0.01; Fig. 2b) in the entire treatment group as compared to control. In case of midbrain, an increased AChE activity in LD and MD treatment groups and drastic reduction in HD (p<0.01)-treated mice was noticed when compared to control (Fig. 2a).

DISCUSSION
Steroid hormones are known to affect neurotransmission through direct effects on the cellular membrane, modulation of the synthesis, and degradation of neurotransmitters by altering their metabolism [11]. ACh is an essential player in the formation, maintenance, and evocation of memory processes. In the present investigation, we evaluated the impact of one of the AAS compounds 17α-MT on the AChE enzyme activity in different brain regions. A significant increase in AChE activity in forebrain and midbrain (low and medium dose treatment) suggests a reduction of cholinergic neurotransmission efficiency due to a decrease in acetylcholine levels in the synaptic cleft which may be related to neurotoxicity and neuronal degeneration [23]. Parallel to this, subchronic treatment of nandrolone decanoate (ND) resulted in the augmented levels of AChE activity in cerebellum and striatum of rat brain affecting the cholinergic system and consequently the CNS [24]. Likewise, ND treatment induced behavioral changes including oxidative damage, inflammation, and imbalance in brain neurotransmitter systems, modulation of nerve growth factor, and neuronal apoptosis [25-27]. Furthermore, some studies have demonstrated that the prolonged overexpression of AChE activity in the CNS implies cognitive and neuroanatomic pathologies/disorders [28,29].

Further, in the present investigation, concurrent reduction in AChE activity was observed in hippocampus, hindbrain as well as whole brain of 17α-MT treated mice as compared to control suggests the accumulation of ACh at synaptic junctions which in turn would result in the impairment of neuronal transmission. Based on the observed results, we propose that the inhibitory effect of 17α-MT may dependant on its binding capacity to the AChE enzyme activity site. A previous study from our laboratory reported that Stanozolol, a potent AAS compound, induced alteration in local antioxidant cascade besides altering AChE activity in brain and kidney in female mice [30]. Since the regulation of cholinergic system through acetylcholine hydrolysis has been largely attributed to AChE activity, a significant reduction in its activity may lead to stress-related anxiety and memory loss, thus interfering with some cognitive and behavioral aspects in the mice. An excess level of ACh was shown to be neurotoxic, while the opposite is also true since low levels of ACh in the synaptic junction may influence memory negatively [31,32]. Reduction in AChE activity...
Steroid hormones: Humoral signals which alter brain cell function, but excessive AChE activity leads to disruption of ACh neurotransmission and cholinergic modulation [24]. There are also reports that show that the ingestion of AAS affects neuronal activity in the hypothalamus and forebrain [37] by promoting neurodegenerative apoptotic effect [38]. Likewise, the supraphysiological levels of testosterone initiate apoptotic cascade in neuronal cells of human neuroblastoma cell lines [39].

Based on results of the present study, the following inferences are drawn:

1. Increase in AChE enzyme activity in forebrain and midbrain indicates a reduction in cholinergic neurotransmission efficiency due to a decrease in acetylcholine levels in the trans-synaptic cleft.

2. Decreased activity of AChE in hippocampus, hindbrain, as well as whole brain suggest impairment in neuronal transmission, in turn may lead to anxiety and depression in mice.

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CONFLICTS OF INTEREST

Authors have none to declare.

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