

# A Comprehensive Review On Steroid Abuse And Its Impact On Various Body Systems

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## **Abstract:**

*This comprehensive review delves into the multifaceted implications of steroid abuse on diverse physiological systems within the human body. Steroid abuse is a widespread issue that transcends boundaries, affecting individuals across different age groups, genders, and athletic backgrounds. This in-depth review article explores the complex network of repercussions that users may experience by exploring the diverse impacts of steroid usage on different physiological systems. We explore the mechanisms of action, short-term and long-term health repercussions, and potential interventions for addressing this growing concern. Lastly, the article addresses the treatment Options for Steroid Abuse.*

**Keyword:** *Steroid abuse; Anabolic-androgenic steroids, widespread issue Athletic background; Long term health repercussion.*

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## **I. Introduction**

Steroids, a class of compounds encompassing both anabolic and corticosteroids, have garnered immense popularity in various fields, from professional sports to aesthetic enhancement<sup>[1]</sup>. However, the misuse and abuse of these substances have direct consequences for the human body, affecting not only the musculoskeletal system but also the cardiovascular, hepatic, renal, endocrine, and psychiatric systems. Steroids are intricate, four-ringed chemical compounds that are vital to multicellular organisms. They act as endocrine hormones and structural elements similar to cholesterol. With hydrophilic hormones acting on cell surfaces and hydrophobic hormones diffusing across membranes to activate intracellular receptors, these hormones help regulate and enable communication between cells and tissues. The biology of lipophilic steroid hormones and synthetic steroid compounds created to cure illnesses will be the main topic of this review. In particular, their functions in foetal development and resolving postnatal issues from preterm birth will be discussed<sup>[2][3]</sup>. This review aims to provide a comprehensive overview of the far-reaching effects of steroid abuse on these body systems. Anabolic-androgenic steroids (AAS), which are generated from testosterone, have been related to mood disorders like depression and mania and can cause a range of behavioural abnormalities, especially when used in large dosages. AAS affect the central nervous system by both direct receptor modification and indirect neurotransmitter influence, while low doses may have no effect and even be helpful for some mood disorders. In addition to interacting with the hypothalamus and amygdala, two brain regions linked to aggression and anxiety, they can also be turned into oestrogen and impact a variety of hormone receptors. The intricate interaction between these processes, especially the lateral hypothalamus and expanded amygdala, is responsible for the aggression and anxiety seen in AAS abusers<sup>[4]</sup>. Abuse of steroids has a wide range of short- and long-term effects and poses serious health concerns to several body systems. While using anabolic-androgenic steroids excessively can result in major problems such muscle dysmorphia, cardiovascular troubles, liver damage, hormone imbalances, and psychological impacts, they can also provide momentary improvements in strength and muscle mass. Notable concerns are also issues related to reproductive health and greater susceptibility to infections. A multifaceted strategy including medical supervision, psychosocial assistance, education, and rehabilitation is needed to address steroid abuse. In order to lessen the negative impacts of steroid usage and encourage healthier choices, awareness and intervention are essential<sup>[5]</sup>. The adrenal cortex, more especially the zona glomerulosa (ZG) and zona fasciculata (ZF), is the primary source of mineralocorticoid hormones. The ZG secretes 18-hydroxycorticosterone (18OHB) and aldosterone (Aldo), and angiotensin II primarily controls this process. On the other hand, the ZF is regulated by adrenocorticotrophic hormone (ACTH) to create deoxycorticosterone (DOC), 18-hydroxy-deoxycorticosterone (18OH-DOC), and corticosterone(B). Excessive levels of the glucocorticoid cortisol (F) can have mineralocorticoid activity, which may also contribute to hypertension via gglucocorticoid and mineralocorticoid pathways. Mineralocorticoid hypertension can be caused by elevated levels of Aldo, DOC, or B, caused by adrenal tumours genetic abnormalities orexogenoussources (suchliquoriceand9αfluorinated

steroids). Low potassium levels high blood pressure, and renin-angiotensin system suppression are the hallmarks of mineralocorticoid hypertension<sup>[6]</sup>.

**Table no 1 :** Shows general dosage ranges of anabolic steroids with Route of Administration

General dosage ranges of anabolic steroids		
Medication	Route	Dosage range
Danazol	Oral	100–800 mg/day
Drostanolone propionate	Injection	100 mg 3 times/week
Ethylestrenol	Oral	2–8 mg/day
Fluoxymesterone	Oral	2–40 mg/day
Mesterolone	Oral	25–150 mg/day
Metandienone	Oral	2.5–15 mg/day
Metenolone acetate	Oral	10–150 mg/day
Metenolone enanthate	Injection	25–100 mg/week
Methyltestosterone	Oral	1.5–200 mg/day
Nandrolone decanoate	Injection	12.5–200 mg/week
Nandrolone phenylpropionate	Injection	6.25–200 mg/week
Norethandrolone	Oral	20–30 mg/day
Oxandrolone	Oral	2.5–20 mg/day
Oxymetholone	Oral	1–5 mg/kg/day or 50–150 mg/day
Stanozolol	Oral	2–6 mg/day
	Injection	50 mg up to every two weeks
Testosterone	Oral	400–800 mg/day
	Injection	25–100 mg up to three times weekly
Testosterone cypionate	Injection	50–400 mg up to every four weeks
Testosterone enanthate	Injection	50–400 mg up to every four weeks
Testosterone propionate	Injection	25–50 mg up to three times weekly
Testosterone undecanoate	Oral	80–240 mg/day
	Injection	750–1000 mg up to every 10 weeks
Trenbolone HBC	Injection	75 mg every 10 days

**Table no.2:** Various types of steroids that effects the body system

Formulations	Examples
Tablets & Liquids	Prednisolone, Danazol, Methyltestosterone, Dexamethasone, Stanazolol, Oxymetholone, Mesterolone, Fluoxymesterone
Inhalers and nasal sprays	Beclomethasone, Fluticasone
Injection (to joints, muscles, blood vessels)	Methylprednisolone, Dexamethasone, Stanazolol, Mesterolone, Methylprednisolone
Creams, lotions, gels	Hydrocortisone
Suspensions	Stanazolol

## II. Effect Of Steroids On Different Systems

### Musculoskeletal System:

Anabolic-androgenic steroids like Danazol and methyltestosterone are often abused for their muscle-building properties, leading to short-term gains in muscle mass and strength. Long-term use of steroids like Danazol and methyltestosterone results in muscle dysmorphia, tendon and ligament injuries, and reduced bone density. The high or low dose of corticosteroids used often determines their impact on the musculoskeletal system. Regarding aseptic osteonecrosis, myopathy, and excessive dosages appear to have a lot of power yet are unpredictable, while Growth inhibition and osteoporosis relate to low doses of therapy as well as more typical incidental events. But the understanding of a threshold dosage that defines the negative. Because there are so many diseases, the impacts are still minimal addressed and the necessary mix of medication therapy<sup>[7]</sup>.

### **Cardiovascular System:**

Abuse of steroids like Methylprednisolone and dexamethasone can elevate blood pressure, increasing the risk of stroke and heart attack.<sup>[5]</sup> There are several subtypes of primary aldosteronism, including unilateral adenoma and idiopathic aldosteronism, which are disorders of the adrenal glands' zona glomerulosa. Although these subsets have reliable diagnostic criteria, little is known about their pathophysiology, which calls for more research. Adrenalin may be a major factor in the elevated sensitivity to angiotensin II seen in patients with idiopathic aldosteronism. Glucocorticoid-remediable aldosteronism is a subtype that is characterized by a substantial overproduction of cortisol pathway products and aberrant aldosterone sensitivity to ACTH. A syndrome of apparent mineralocorticoid excess connected to hypertension can also result from poor cortisol metabolism, whereas rare disorders such as congenital adrenal hyperplasia syndromes and deoxycorticosterone-secreting tumours are linked to excess mineralocorticoid secretion.

<sup>[6]</sup>. Long-term use of steroids like methyltestosterone use may contribute to the development of atherosclerotic plaques<sup>[8]</sup>.

### **Hepatic System:**

Oral anabolic steroids like methyltestosterone can strain the liver, potentially leading to liver damage or tumors. Many hepatotoxic consequences, including as cholestasis, peliosis hepatis, and liver cancers, have been related to anabolic androgen steroids (AAS). This is mostly because the liver is the primary site of clearance for these drugs. The mechanisms of liver injury include excessive collagen deposition brought on by activated Kupffer cells and hepatic stellate cells, oxidative stress from mitochondrial dysfunction, and inflammatory cell infiltration. In contrast to peliosis hepatis, which is characterized by blood-filled cysts and hyper vascular alterations in the liver, AASs can interfere with bile transport and result in cholestasis, especially when combined with 17 $\alpha$ -alkylated steroids. Long-term use of AAS is linked to the development of hepatic malignancies, especially in men. This is probably because of increased hepatocyte proliferation and changes that resemble stem cells. Even though both oral and injectable AAS forms have the potential to cause hepatotoxicity, the majority of cases of AAS-induced liver damage recover completely after stopping the medication. To comprehend the biology of AAS-related liver damage and its possible long-term effects, there has to be a greater awareness of and reporting of this condition<sup>[9]</sup>. Steroids like stanozolol and methyltestosterone abuse may disrupt bile flow, causing cholestasis<sup>[9]</sup>.

### **Renal System:**

Anabolic steroids like stanozolol, and carbon -17 alkylated anabolic steroids effects the renal system By enhancing sodium reabsorption and encouraging potassium and hydrogen excretion, mineralocorticoids—most notably aldosterone—control electrolyte balance in the kidneys, producing effects including positive sodium balance and hypokalaemia when present in excess. Hypocorticism, on the other hand, causes hyperkalaemia, hyponatremia, and sodium loss, which may lead to serious cardiovascular problems. While prolonged exposure may result in salt "escape," indicating receptor downregulation, aldosterone operates through mineralocorticoid receptors in the kidney, increasing sodium permeability and boosting the sodium/potassium-ATPase. Glucocorticoids vary in that they cause nephrocalcinosis, increase water diuresis, and renal plasma flow without substantially altering hydrogen excretion. They also affect how well electrolytes are absorbed in other tissues; for example, glucocorticoids improve the gastrointestinal tract's absorption of water and sodium, which may cause problems in the stomach.

Steroid abuse can lead to sodium and water retention, potentially impacting kidney function<sup>[11]</sup>. Long-term use may contribute to glomerular and tubular dysfunction.

### **Endocrine System:**

Corticosteroids have a major effect on how many hormones are secreted and function. While cortisol suppresses spontaneous growth hormone release in hypercorticism, it increases growth hormone production in acromegaly patients, which causes growth failure in children by slowing the maturity of the epiphyseal plate. Additionally, they reduce the efficiency of thyroxine in individuals with myxoedema and suppress the secretion of thyroid-stimulating hormones. By promoting the synthesis of adrenaline, high doses of corticosteroids enhance adrenergic effects and inhibit the release of luteinizing hormones. Adrenocortical insufficiency after withdrawal, steroid-induced hyperglycaemia, hyperlipidaemia, increased glucagon levels, and hypocalcaemia are further systemic consequences of glucocorticoids. Overall, the effects of corticosteroids on metabolic and hormonal control are complex<sup>[10]</sup>. Abuse of steroids like cortisol disrupts the hypothalamic-pituitary-gonadal axis, resulting in reduced testosterone production in males and menstrual irregularities in females<sup>[12]</sup>. Abuse of steroids like cortisol can lead to hypogonadism, infertility, and gynecomastia in males.

### **Psychiatric System:**

Nandrolone and other anabolic-androgenic steroids (AAS) have a major effect on the neuronal circuits associated with aggression and anxiety, especially in the extended amygdala, which contains the bed nucleus of the stria terminalis (BNST) and the central amygdala (CeA). Changes in corticotropin-releasing factor (CRH) and GABA neurotransmission, where imbalances between these systems contribute to heightened anxiety, are the main mechanisms via which prolonged AAS usage might result in anxiety-like behaviours. Decreased 5-HT<sub>1A</sub> receptor levels and increased 5-HT<sub>2A</sub> receptor activity are two other ways that AAS change receptor expression in the hypothalamus, which impacts serotonin and dopamine pathways that are important for aggression. AAS users' aggressive behaviour is further aggravated by altered NMDA receptor function and increased dopamine receptor activity, especially D<sub>2</sub> and D<sub>5</sub>. Altogether, the interaction of different neurotransmitters and oxidative stress in different parts of the brain highlights the intricate neurobiological processes underlying the anxiety and aggression brought on by AAS<sup>[4][11][12]</sup>.

### **Reproductive System:**

Prolonged use of anabolic steroids like methyltestosterone, mesterolone, and fluoxymesterone leads to abnormalities in the reproductive system. The abuse of anabolic steroids can lead to testicular atrophy, which is the shrinking of the testicles. This occurs because the body senses the presence of exogenous (external) testosterone and reduces its natural production of testosterone. This can result in a decrease in testicle size and impaired sperm production. Anabolic steroids can significantly reduce sperm production, leading to a low sperm count. This can result in infertility, making it difficult or impossible for the individual to father children<sup>[14]</sup>. Steroid abuse can cause or worsen erectile dysfunction, making it difficult to achieve or maintain an erection. Steroid abuse in females can disrupt the normal menstrual cycle, leading to irregular periods or even complete cessation of menstruation (amenorrhea). Virilization refers to the development of male secondary sexual characteristics in females. It can include the deepening of the voice, growth of facial and body hair (hirsutism), and enlargement of the clitoris (clitoromegaly). These changes are often irreversible<sup>[15]</sup>. Steroid abuse can lead to changes in breast tissue, including breast size reduction<sup>[16]</sup>.

### **Integumentary System:**

Abuse of steroids like methyltestosterone can cause or exacerbate acne, leading to the development of pimples, blackheads, and whiteheads. This occurs because steroids increase the production of sebum (skin oil) and can clog pores, promoting the growth of acne-causing bacteria.<sup>[17]</sup> Prolonged use of steroids like prednisone can lead to thinning of the skin, making it more fragile and prone to tearing or bruising. This is especially apparent in regions where topical steroids are used<sup>[18]</sup>. In both men and women, abuse of steroids like prednisone can lead to hirsutism, which is the abnormal growth of facial and body hair. This may be distressing and hard to reverse.

### **Skeletal System:**

Prolonged use of steroids like prednisone, and methyl prednisone can lead to a decrease in bone density, a condition known as osteoporosis. This happens because steroids can interfere with the normal balance of bone remodeling, leading to bone loss. Weakened bones are more prone to fractures<sup>[19]</sup>. The risk of fractures, especially compression fractures of the spine, rises with a reduction in bone density. Steroid misuse such as prednisone and methyl prednisone usage can result in severe pain and impairment from these fractures. In adolescents who misuse anabolic steroids, like methyltestosterone, the premature closure of growth plates (epiphyseal closure) can occur. This can result in stunted growth, as the long bones of the body no longer grow in length. Abuse of steroids like prednisone, and methyl prednisone can weaken tendons and ligaments, making them more susceptible to injuries such as tears and ruptures. Joint function and stability may be impacted by this.

### **Respiratory and Immune systems:**

Steroids, particularly systemic corticosteroids like prednisone, can suppress the immune system. This immunosuppression can increase an individual's susceptibility to respiratory infections, such as pneumonia, bronchitis, and tuberculosis<sup>[20]</sup>. Steroids like prednisone can mask the symptoms of respiratory infections, making it more challenging to detect and diagnose these conditions on time. This delay in diagnosis and treatment can lead to more severe illness. In individuals with pre-existing respiratory conditions such as asthma or chronic obstructive pulmonary disease (COPD), steroid abuse can have complex effects. While steroids like prednisone can provide short-term relief from symptoms like wheezing and shortness of breath, long-term or excessive use can lead to complications, including steroid-induced asthma, which can be difficult to manage. Steroids like beclomethasone can cause throat irritation, leading to a persistent cough and hoarseness, which can be bothersome.

### **Immune System:**

The immune system is impacted by long-term usage of steroids such as cortisone, methyl prednisone, and prednisone. Steroids are known for their immunosuppressive properties. They significantly weaken the immune system's ability to fight off infections, when abused. This can lead to an increased risk of various infections, including bacterial, viral, and fungal infections<sup>[21]</sup>. Steroid abuse can delay the body's natural healing processes and impair the immune system's response to injuries or infections. This can prolong recovery times and increase the severity of illnesses. In some cases, steroid abuse can trigger or exacerbate autoimmune disorders, where the immune system attacks the body's tissues and organs. This can lead to conditions such as lupus or rheumatoid arthritis<sup>[22]</sup>. Steroid abuse can make individuals more susceptible to opportunistic infections, which are typically harmless in healthy individuals but can cause severe illness in people with weakened immune systems. Examples include fungal infections like candidiasis and viral infections like herpes zoster (shingles).

### **III. Treatment Options For Steroid Abuse**

Combining medical, psychological, and social interventions necessitates a comprehensive and customized approach to address steroid misuse<sup>[23]</sup>. The goals of treatment plans are to address underlying problems, encourage long-term recovery, and manage the psychological and physical effects of steroid abuse. Important therapy choices are listed below:

#### **Medical Management:**

Individuals who overuse steroids must collaborate closely with medical professionals to develop a customized treatment plan. The individual's unique requirements and circumstances should be taken into consideration in this plan. Seek immediate professional medical assistance if you or someone you know is experiencing steroid abuse.

**HRT (Hormone Replacement Therapy):** Under medical supervision, HRT may be administered for patients whose endogenous testosterone production has been inhibited to restore hormonal balance<sup>[24]</sup>. Long-term steroid misuse can interfere with the body's ability to produce hormones naturally. Hormone replacement treatment could be recommended in some circumstances to assist in reestablishing hormonal balance. A healthcare provider's supervision is required for this.

**Management of Health Consequences:** Medical specialists deal with certain health conditions such as liver damage, musculoskeletal disorders, and cardiovascular problems that arise from abusing steroids.

#### **Psychological Support:**

**Therapy and Counseling:** Behavioral therapies, such as motivational enhancement therapy (MET) and cognitive-behavioral therapy (CBT), can assist people in addressing the underlying causes of their steroid addiction, enhancing their coping mechanisms, and cultivating positive body image beliefs.

**Group and Individual Counseling:** People can examine the psychological components of their substance use in individual counseling sessions and group therapy, and they can share their experiences with others going through similar difficulties.

**Psychological Assessment:** To treat co-occurring mental health conditions including mood disorders or body dysmorphic disorder, people may benefit from receiving a psychiatric assessment.

#### **Education and Awareness:**

**Psychoeducation Programs:** Educating people about the dangers and repercussions of abusing steroids can increase awareness and encourage wise decision-making.

**Education of the Family:** Including the family in the therapeutic process helps strengthen support systems and leads to a more thorough comprehension of the patient's requirements.

#### **Rehabilitation Programs:**

**Inpatient Rehabilitation:** A structured setting with round-the-clock care is offered by inpatient rehabilitation facilities in cases of extreme severity or when medical monitoring is required.

**Outpatient Rehabilitation:** Programs for outpatient therapy are flexible, enabling patients to get treatment while going about their everyday lives<sup>[25]</sup>.

**Follow-Up Care:**

**Long-Term Monitoring:** To address potential relapse triggers and manage ongoing health concerns, people in recovery require regular medical and psychological monitoring.

**Aftercare Programs:** Following extensive treatment, individuals can transition to a less regimented setting with the help of aftercare programs, which offer resources and continuing support.

## **IV. Discussion**

**Musculoskeletal System:**

**Long-Term Consequences vs. Anabolic Effects** While anabolic-androgenic steroids (AAS) like methyltestosterone and Danazol might assist increase muscle building and strength quickly, long-term usage of these drugs can lead to major issues. For example, there is a problem known as muscular dysmorphia, which is a type of eating disorder linked to steroid addiction and has additional health consequences for users. In addition, abusing steroids puts the user at risk for bone loss and alterations to tendons and ligaments, which raises concerns about both the user's long-term health and the muscle gains they have accomplished.

**Cardiovascular System:**

**Atherosclerosis and Hypertension** Steroid use has the potential to significantly raise blood pressure and cause a number of cardiovascular conditions, including heart attacks and strokes. The possibility of atherosclerosis development especially long term use of methyltestosterone makes one mindful of the risks that come with the innocent looking desire for muscles. These cardiovascular risks highlight the need of prioritizing overall health over achieving the ideal performance or appearance.

**Hepatic System:**

**liver damage and cholestasis** Methyltestosterone and its analogs are frequently linked to cholestasis, hepatotoxicity, and other types of severe liver injury. The repercussions are dire; there may be hazardous health hazards involved. As a result, the person may want to think twice before utilizing music augmentation devices to quickly strain their muscles.

**Renal System:**

**Fluid Imbalance and Renal Injury** Also, the impact of steroids on the renal system is substantial. Steroids cause excessive fluid retention which can result in electrolyte imbalances, further aggravating the renal function. Long term use is associated with an increased risk of glomerular and tubular injury, triggering concern over the renal health profile of the users.

**Endocrine System:**

**Endocrine disorders and hormonal disruption** Steroids have a significant impact on the endocrine system. Abuse of steroids throws off the hormonal balance, resulting in irregular menstruation in women and diseases such as hypogonadism and infertility in men. The reproductive system and general health may be negatively impacted for some time by such endocrine repercussions.

**Psychiatric System:**

**Anxiety and Depressive States** The psychological repercussions of steroid use are problematic, with documented mood swings, hostility, and even steroid-induced insanity. The possibility of psychological dependence makes matters more complicated because users may experience withdrawal symptoms, which can result in an abusive cycle that is hard to stop.

**Reproductive System:**

**Abnormalities and their effects on conception** Particularly concerning are the effects of steroid misuse on reproduction. Testicular shrinkage, poor sperm counts, and erectile dysfunction in males, and virilization consequences in females, emphasize the extensive and often irreversible changes that steroid usage can impose on reproductive health. These changes may have a significant impact on quality of life and family planning.

**Integumentary System:**

**Skin Disorders and Sexual Assault** The integumentary side effects, like thinning skin and pimples, can be quite upsetting for users. Increased body and facial hair, or hirsutism, highlights the physical changes that might be unwanted or detrimental to one's mental health in both men and women.

### **Skeletal System:**

Loss of Bone Density and Risk of Fractures The use of steroids carries a substantial danger to bone health, since it can lead to osteoporosis and increased risk of fractures due to decreasing bone density. In teenagers, the possibility of early epiphyseal closure underlines the hazards of steroid use during important growth phases, potentially impeding development.

### **Respiratory and Immune Systems:**

Immunodeficiency and Issues with the Respiratory System Lastly, it is impossible to ignore the impact on the immunological and respiratory systems. Immunosuppression brought on by steroids can exacerbate pre-existing respiratory disorders and make people more vulnerable to infections, creating a vicious cycle of health issues that can be challenging to break.

## **V. Conclusion**

Steroid abuse exerts a profound impact on various body systems, leading to a plethora of short-term and long-term health consequences. This review highlights the importance of addressing this issue through education, prevention, and intervention strategies to safeguard against the overuse of steroids. A variety of psychological, medical, and legal repercussions get complicated when steroids are used. Healthcare providers, legislators, and anybody else considering the use of these drugs need to be informed on the possible benefits and drawbacks of doing so. While some people may find that steroids improve their performance, when used improperly, they can have several undesirable side effects that frequently outweigh the benefits. It's a two-edged weapon that calls for awareness, education, and compassion for those who become snared in its jagged edges.

## **References**

- [1]. Ericson-Neilsen W, Kaye Ad. Steroids: Pharmacology, Complications, And Practice Delivery Issues. Ochsner J. 2014 Summer;14(2):203-7. Pmid: 24940130; Pmcid: Pmc4052587.
- [2]. Cole Tj, Short Kl, Hooper Sb. The Science Of Steroids. Semin Fetal Neonatal Med. 2019 Jun;24(3):170-175. Doi: 10.1016/J.Siny.2019.05.005. Epub 2019 May 23. Pmid: 31147162.
- [3]. Hodgens A, Sharman T. Corticosteroids. [Updated 2023 May 1]. In: Statpearls [Internet]. Treasure Island (FL): Statpearls Publishing; 2023 Jan-. Available From: <https://www.ncbi.nlm.nih.gov/books/nbk554612/>
- [4]. Bertozzi, G., Sessa, F., Albano, G.D., Sani, G., Maglietta, F., Roshan, M.H., Volti, G.L., Bernardini, R., Avola, R., Pomara, C. And Salerno, M., The Role Of Anabolic Androgenic Steroids In Disruption Of The Physiological Function In Discrete Areas Of The Central Nervous System. *Molecular Neurobiology*, 2018, 55, Pp.5548-5556.
- [5]. Vanberg, P. And Atar, D., Androgenic Anabolic Steroid Abuse And The Cardiovascular System. *Doping In Sports: Biochemical Principles, Effects And Analysis*, 2010, Pp.411-457.
- [6]. Mantero, F., Armanini, D., Boscaro, M., Carpenè, G., Fallo, F., Opocher, G., Rocco, S., Scaroni, C. And Sonino, N., Steroids And Hypertension. *The Journal Of Steroid Biochemistry And Molecular Biology*, 1991, 40(1-3), Pp.35-44.
- [7]. Theodorakidou, M., Nikola, O.A. And Lambrou, G.I., 2018. The Multiple Roles Of Steroids And Anabolic Steroids And Its Relations To Cardiovascular And Musculoskeletal Pathology: A Brief Review. *J. Res. Pract. Musculoskelet. Syst*, 2, Pp.22-30.
- [8]. Liu Jd, Wu Yq. Anabolic-Androgenic Steroids And Cardiovascular Risk. *Chin Med J (Engl)*. 2019 Sep 20;132(18):2229-2236. Doi: 10.1097/Cm9.0000000000000407. Pmid: 31478927; Pmcid: Pmc6797160.
- [9]. Petrovic A, Vukadin S, Sikora R, Bojanic K, Smolic R, Plavec D, Wu Gy, Smolic M. Anabolic Androgenic Steroid-Induced Liver Injury: An Update. *World J Gastroenterol*. 2022;28(26):3071-3080.
- [10]. Ravisankar P, Pravalika D, Anjali D, Sree Vidya V, Sai Anvith P, Pragna P. Fatty Liver Disease In Depth Analysis. *Indo American Journal Of Pharmaceutical Research*. 2015; 5(11):3622-3642.
- [11]. McKay Li, Cidowski Ja. Physiologic And Pharmacologic Effects Of Corticosteroids. In: Kufe Dw, Pollock Re, Weichselbaum Rr, Et Al., Editors. *Holland-Frei Cancer Medicine*. 6th Edition. Hamilton (On): Bc Decker; 2003. Available From: <https://www.ncbi.nlm.nih.gov/books/nbk13780/>
- [12]. Bertozzi, G., Sessa, F., Albano, G.D., Sani, G., Maglietta, F., Roshan, M.H., Volti, G.L., Bernardini, R., Avola, R., Pomara, C. And Salerno, M., The Role Of Anabolic Androgenic Steroids In Disruption Of The Physiological Function In Discrete Areas Of The Central Nervous System. *Molecular Neurobiology*, 2018, 55, Pp.5548-5556.
- [13]. Trenton, A.J. And Currier, G.W., Behavioural Manifestations Of Anabolic Steroid Use. *Cns Drugs*, 2005, 19, Pp.571-595.
- [14]. El Osta R, Almont T, Diligent C, Hubert N, Eschwège P, Hubert J. Anabolic Steroids Abuse And Male Infertility. *Basic Clin Androl*. 2016 Feb 6;26:2. Doi: 10.1186/S12610-016-0029-4. Pmid: 26855782; Pmcid: Pmc4744441.
- [15]. Thiago Gagliano-Jucá, Shehzad Basaria, Abuse Of Anabolic Steroids: A Dangerous Indulgence, *Current Opinion In Endocrine And Metabolic Research*, Volume 9, 2019, Pages 96-101, Issn 2451-9650, <https://doi.org/10.1016/J.Coemr.2019.10.002>.
- [16]. Casavant, M.J., Blake, K., Griffith, J., Yates, A. And Copley, L.M., Consequences Of Use Of Anabolic Androgenic Steroids. *Pediatric Clinics Of North America*, 2007, 54(4), Pp.677-690.
- [17]. Meena S, Gupta Lk, Khare Ak, Balai M, Mittal A, Mehta S, Bhatni G. Topical Corticosteroids Abuse: A Clinical Study Of Cutaneous Adverse Effects. *Indian J Dermatol*. 2017 Nov-Dec;62(6):675. Doi: 10.4103/Ijd.Ijd\_110\_17. Pmid: 29263550; Pmcid: Pmc5724325.
- [18]. Inakanti, Y., Thimmasarathi, V.N.R., Kumar, S., Nagaraj, A., Peddireddy, S. And Rayapati, A., Topical Corticosteroids: Abuse And Misuse. *Our Dermatology Online/Nasza Dermatologia Online*, 6(2). 2015.
- [19]. Hall, G.M., Spector, T.D., Jane Griffin, A., Sm Jawad, A., Hall, M.L. And Doyle, D.V., The Effect Of Rheumatoid Arthritis And Steroid Therapy On Bone Density In Postmenopausal Women. *Arthritis & Rheumatism: Official Journal Of The American College Of Rheumatology*, 1993, 36(11), Pp.1510-1516.
- [20]. Janahi, I.A., Rehman, A. And Baloch, N.U.A., Corticosteroids And Their Use In Respiratory Disorders. *Corticosteroids*. London: Intech Open, 2018, Pp.47-57

- [21]. Bertozzi, G., Sessa, F., Maglietta, F., Cipolloni, L., Salerno, M., Fiore, C., Fortarezza, P., Ricci, P., Turillazzi, E. And Pomara, C., Immunodeficiency As A Side Effect Of Anabolic Androgenic Steroid Abuse: A Case Of Necrotizing Myofasciitis. *Forensic Science, Medicine And Pathology*, 2019, 15, Pp.616-621.
- [22]. Benagiano M, Bianchi P, D'elios Mm, Brosens I, Benagiano G. Autoimmune Diseases: Role Of Steroid Hormones. *Best Pract Res Clin Obstet Gynaecol*. 2019 Oct;60:24-34. Doi: 10.1016/J.Bpobgyn.2019.03.001. Epub 2019 Mar 16. Pmid: 31047850.
- [23]. Hoseini, M., Yousefi, B. And Khazaei, A.A., The Prevalence Of Anabolic-Androgenic Steroids Abuse, Knowledge And Attitude Of Their Side Effects, And Attitude Toward Them Among The Female Bodybuilding Athletes In Kermanshah. *Journal Of Advanced Biomedical Sciences*, 2020, 10(3), Pp.2439-2447.
- [24]. Zou My, Cohen Re, Ursomanno Bl, Yerke Lm. Use Of Systemic Steroids, Hormone Replacement Therapy, Or Oral Contraceptives Is Associated With Decreased Implant Survival In Women. *Dent J (Basel)*. 2023 Jun 29;11(7):163. Doi: 10.3390/Dj11070163. Pmid: 37504229; Pmcid: Pmc10377784.
- [25]. Nedeljkovic U, Dackovic J, Tepavcevic Dk, Basuroski Id, Mesaros S, Pekmezovic T, Drulovic J. Multidisciplinary Rehabilitation And Steroids In The Management Of Multiple Sclerosis Relapses: A Randomized Controlled Trial. *Arch Med Sci*. 2016 Apr 1;12(2):380-9. Doi: 10.5114/Aoms.2015.47289. Epub 2015 Mar 18. Pmid: 27186184; Pmcid: Pmc4848347.