STANOZOLOL

PRODUCT OVERVIEW

Specification	Details
Generic Name	**Stanozolol**
Drug Class	**Anabolic Androgenic Steroid (AAS)**
Active Ingredient	**Stanozolol**
Chemical Formula	$C_{21}H_{32}N_2O$
Molecular Weight	**328.49 g/mol**
Administration Route	**Oral Administration / Intramuscular Injection (Aqueous Suspension)**
Ester Type	**Non-Esterified Pyrazole-Derived Dihydrotestosterone (DHT) Derivative**
Approximate Half-Life	**Oral: Approximately 8–9 Hours / Injectable Suspension: Approximately 24 Hours**
Release Profile	**Rapid Systemic Availability**
Bioavailability	**High, Following Oral Administration**
Primary Medical Use	**Approved in selected countries for hereditary angioedema and historically investigated for selected catabolic conditions**

INTRODUCTION

Stanozolol is a synthetic anabolic androgenic steroid structurally derived from dihydrotestosterone (DHT). Unlike esterified injectable anabolic steroids, stanozolol is a non-esterified compound available in both oral tablets and aqueous intramuscular suspension.Its unique pyrazole ring modification differentiates it from most other anabolic steroids and contributes to its distinctive pharmacokinetic profile. Due to its high oral bioavailability and relatively short elimination half-life, stanozolol and dihydrotestosterone physiological properties that differ from testosterone.derived anabolic steroids.Stanozolol has been extensively investigated in clinical medicine and sports pharmacology due to its oral bioavailability, injectable formulation, and well-characterized anabolic profile.

HISTORY

Stanozolol was developed during pharmaceutical research aimed at producing an orally active anabolic steroid with improved anabolic properties and reduced androgenic activity.

Following its introduction into clinical medicine, stanozolol was investigated for hereditary angioedema, selected catabolic disorders, tissue wasting, and disorders affecting connective tissue metabolism.

Its extensive scientific evaluation has contributed significantly to research involving:

- Endocrine pharmacology
- Connective tissue physiology
- Protein metabolism
- Sports medicine

CHEMICAL STRUCTURE

Stanozolol is a synthetic derivative of dihydrotestosterone containing a fused pyrazole ring and a 17-alpha-alkyl modification that enhances bioavailability.

Unlike esterified anabolic steroids, stanozolol does not require enzymatic ester hydrolysis before becoming biologically active.

Its molecular structure primarily influences:

Oral bioavailability
Metabolic stability
Androgen receptor affinity
Pharmacokinetic characteristics

Following oral administration or intramuscular injection, biologically active stanozolol becomes available to interact with androgen receptors throughout the body.

CHEMICAL PROFILE

Parameter	Information
Chemical Name	**Stanozolol**
Molecular Formula	$C_{21}H_{32}N_2O$
Molecular Weight	**328.49 g/mol**
Hormone Classification	**Anabolic Androgenic Steroid**
Ester Group	**None (17α-Alkylated DHT Derivative)**
Active Hormone	**Stanozolol**

PHARMACOLOGY

Following oral administration or intramuscular injection, stanozolol is absorbed into systemic circulation without requiring ester hydrolysis.

Once absorbed, stanozolol binds to androgen receptors located throughout multiple tissues including:

* Skeletal muscle
* Bone
* Bone marrow
* Connective tissue
* Liver
* Sebaceous tissue

As a dihydrotestosterone derivative, stanozolol does not undergo aromatization into estradiol.

Conversion Pathway	Enzyme	Product
Stanozolol → Hepatic Metabolites	**Hepatic Enzymes**	**Inactive Metabolites**
Aromatization	**Aromatase**	**No Significant Conversion**

MECHANISM OF ACTION

Stanozolol produces its biological effects primarily through activation of intracellular androgen receptors.

The biological process includes:

* Stanozolol enters target cells.
* Stanozolol binds to androgen receptors.
* The receptor-hormone complex becomes activated.
* The complex translocates into cellular nuclei.
* It interacts with androgen response elements on DNA.
* Gene transcription is regulated.
* Androgen protein synthesis and tissue-specific physiological responses are influenced.

Activation of androgen receptors contributes to:

* Protein synthesis
* Nitrogen retention
* Lean tissue maintenance
* Bone remodeling
* Red blood cell production
* Male reproductive physiology
* Endocrine regulation

PHARMACODYNAMICS

Stanozolol affects multiple physiological systems through androgen receptor signaling.

SKELETAL MUSCLE SYSTEM

Stanozolol contributes to normal muscle physiology through:

* Regulation of protein metabolism
* Support of muscle tissue maintenance
* Influence on nitrogen balance
* Regulation of muscle fiber function

Stanozolol interacts with androgen receptors, contributing to anabolic activity in androgen-responsive tissues.

SKELETAL SYSTEM

Stanozolol contributes to skeletal physiology by influencing:

* Bone remodeling
* Osteoblast activity
* Osteoclast activity
* Bone mineral metabolism

CONNECTIVE TISSUE

Stanozolol contributes to skeletal physiology by influencing:

* Structural protein metabolism.
* Connective tissue maintenance
* Tissue remodeling
* Collagen metabolism

REPRODUCTIVE SYSTEM

Stanozolol interacts with androgen-sensitive tissues and contributes to:

* Hormonal regulation
* Maintenance of androgen-responsive tissues
* Endocrine signaling
* Reproductive physiology

PHARMACOKINETICS

Following oral administration or intramuscular injection, stanozolol is rapidly absorbed into systemic circulation without requiring esterification.

The oral formulation exhibits rapid gastrointestinal absorption, while the aqueous injectable suspension provides sustained absorption.Both the intramuscular hormone.It produces a prolonged release profile.

Both formulations deliver biologically active stanozolol directly into systemic circulation.

PHARMACOKINETIC PROFILE

Parameter	Description
Ester Type	**None (17α-Alkylated DHT Derivative)**
Absorption	**Rapid gastrointestinal absorption**
Release Pattern	**Gradual release from aqueous suspension**
Half-Life	**Approximately 8–9 Hours**
Duration of Activity	**Approximately 24 Hours**
Plasma Stability	**Moderate**
Protein Binding	**High**
Metabolism	**Primarily hepatic**
Active Hormone	**Stanozolol**
Elimination	**Urinary excretion of metabolites**

PHYSIOLOGICAL EFFECTS

Normal stanozolol activity contributes to:

Muscular Function
* Protein synthesis
* Nitrogen retention
* Maintenance of lean body tissue
* Muscle physiology

Bone Health
* Bone remodeling
* Skeletal maintenance
* Bone metabolism

Connective Tissue
* Protein metabolism
* Connective tissue maintenance
* Collagen metabolism
* Tissue remodeling

Hematological Function
* Red blood cell production
* Hormonal transport support
* Normal erythropoiesis

CLINICAL APPLICATIONS

Stanozolol has been investigated and utilized for selected medical conditions involving hereditary angioedema, impaired protein metabolism, and selected catabolic disorders.

Clinical evaluation may include:

* Medical history
* Physical examination
* Laboratory assessment
* Hormonal and hematological evaluation

Historical and investigational applications include:

* Hereditary angioedema
* Protein catabolic disorders
* Selected catabolic disorders
* Endocrine pharmacology research

SAFETY CONSIDERATIONS

Stanozolol therapy requires individualized medical evaluation and regular monitoring.

Important considerations include:

* Cardiovascular health
* Blood parameters
* Liver function tests
* Liver and kidney function
* Lipid profile
* Individual physiological response

POTENTIAL ADVERSE EFFECTS

Possible adverse effects may include:

Endocrine Effects
* Suppression of endogenous testosterone production
* Hormonal imbalance
* Reproductive system function

Dermatological Effects
* Acne
* Increased skin oil production
* Hair loss in genetically predisposed individuals

Cardiovascular Effects
* Changes in lipid profile
* Cardiovascular load pressure

Hepatic Effects
* Elevation of liver enzymes
* Hepatic stress associated with 17α-alkylated oral administration

Reproductive Effects
* Reduced fertility
* Suppression of spermatogenesis
* Hormonal suppression

CONTRAINDICATIONS

Condition	Reason
Prostate Cancer	**Potential androgen sensitivity**
Male Breast Cancer	**Hormone-sensitive condition**
Pregnancy	**Risk of fetal virilization**
Severe Cardiac, Hepatic or Renal Disease	**Requires specialist evaluation**
Known Hypersensitivity	**Risk of allergic reaction**

DRUG INTERACTIONS

Potential interactions may occur with:

Anticoagulants
Stanozolol may influence anticoagulant response and require monitoring.

Antidiabetic Medications
Changes in glucose metabolism may alter therapeutic requirements.

Corticosteroids
Combined administration may influence fluid balance and metabolic parameters.

Other Hormonal Therapies
Combined hormonal therapy requires professional medical supervision.

LABORATORY MONITORING

Laboratory Test	Purpose
CBC	**Assess blood parameters**
Hematocrit	**Monitor red blood cell concentration**
Hemoglobin	**Evaluate erythropoiesis**
Liver Function Tests	**Assess hepatic health**
Kidney Function Tests	**Evaluate renal function**
Lipid Profile	**Monitor cardiovascular risk markers**
Total Testosterone	**Evaluate endocrine status**
Estradiol (E2)	**Assess hormonal balance**
PSA (When Appropriate)	**Monitor prostate health**

STORAGE & STABILITY

* Store according to official product labeling.
* Protect from excessive heat and direct sunlight.
* Avoid freezing.
* Maintain original packaging integrity.
* Store away from children.
* Inspect packaging before use.

PRODUCT AUTHENTICITY

Every genuine NEXON product should include appropriate quality verification features.

Authentication may include:

* Batch number tracking
* Manufacturing information
* Optional code verification
* QR authentication system
* Authorized distributor verification

NEXON emphasizes transparency, traceability, and product quality control.

FREQUENTLY ASKED QUESTIONS

What is Stanozolol?
Stanozolol is a synthetic anabolic androgenic steroid derived from dihydrotestosterone (DHT), available in both oral and injectable formulations.

Does Stanozolol convert to estrogen?
No. Stanozolol is a DHT-derived anabolic steroid and is not a substrate for aromatase; therefore, it does not undergo meaningful conversion to estradiol.

Is this Stanozolol available as both an oral and injectable product?
Unlike most injectable anabolic steroids, injectable stanozolol is formulated as an aqueous suspension rather than an esterified solution, allowing administration without esterification.

What medical conditions has Stanozolol been used to treat?
Stanozolol has been approved in selected countries for hereditary angioedema and has also been studied in selected catabolic conditions and connective tissue disorders.

Is Stanozolol hepatotoxic?
The oral formulation contains a 17α-alkyl modification that may increase hepatic stress. Liver function monitoring is recommended during therapy.

NEXON QUALITY COMMITMENT

NEXON is committed to maintaining high standards of manufacturing quality, product consistency, traceability, and scientific transparency.

Through quality control systems, authentication procedures, and responsible information sharing, NEXON aims to provide reliable pharmaceutical products and educational resources for healthcare professionals and informed users.

SCIENTIFIC REFERENCES

*Kicman AT, Pharmacology of Anabolic Steroids.

*Basaria S. Androgen Abuse in Athletes: Detection and Consequences.

*Handelsman DJ. Androgen Physiology and Pharmacology.

*Shahidi NT. Chemistry, Biological Action and Clinical Applications of Anabolic-Androgenic Steroids.

*Llewellyn W. Anabolics.

*Evans NA. Current Concepts in Anabolic-Androgenic Steroids.

*Kouhkalei CD. Anabolic-Androgenic Steroids.

*Nieschlag E, Behre HM. Andrology: Male Reproductive Health and Dysfunction.

*Kicman AT, Gower DB. Anabolic Steroids in Sport and Exercise.

*Wilson JD. Androgen Physiology and Pharmacology.