

BOLDENONE BASE

UNESTERIFIED BOLDENONE FORMULATION

100 mg mL

MICRONIZED INJECTABLE SUSPENSION

PRODUCT OVERVIEW

Specification	Details
Generic Name	Boldenone Base
Drug Class	Unesterified Anabolic-Androgenic Steroid
Active Ingredient	Micronized Boldenone
Administration Route	Intramuscular Injection
Dosage Form	Micronized Injectable Suspension
Chemical Class	1-Dehydrotestosterone / Androstadienolone
Primary Target	Androgen Receptor
Clinical Status	Fictional Non-Approved Formulation

INTRODUCTION

Boldenone Base is an unesterified anabolic-androgenic steroid formulation containing micronized boldenone, also known as 1-dehydrotestosterone. Unlike esterified derivatives, the active molecule does not require ester cleavage before becoming pharmacologically available.

The displayed presentation is a suspension rather than a conventional esterified oil solution. Particle size, vehicle composition, sterility, and suspension uniformity are therefore critical formulation characteristics.

HISTORY

Boldenone emerged from twentieth-century research into synthetic anabolic-androgenic steroids. Most recognized preparations used esterification to modify depot release and handling characteristics, especially the long-chain undecylenate ester.

Unesterified boldenone represents the parent active hormone without an attached fatty-acid ester. This distinguishes it from boldenone acetate, cypionate, and undecylenate.

Boldenone Base is not an established contemporary approved human medicine. The displayed 100 mg/mL suspension is a fictional NEXON catalog formulation.

CHEMICAL STRUCTURE

Boldenone has the molecular framework androst-1,4-dien-17β-ol-3-one, the molecular formula $C_{19}H_{26}O_2$, and an approximate molar mass of 286.42 g/mol.

It contains an additional double bond between carbon atoms 1 and 2 and no ester group at the 17β-hydroxyl position. Consequently, ester hydrolysis is not required for receptor-active boldenone to be present.

- 1-Dehydrotestosterone steroid nucleus
- Unesterified 17β-hydroxyl group
- Micronized suspended particles
- Directly active parent hormone
- Formulation-dependent release characteristics

CHEMICAL PROFILE

Parameter	Information
Active Hormone	Boldenone / 1-Dehydrotestosterone
Ester	None / Unesterified
Molecular Formula	C19H26O2
Molar Mass	Approximately 286.42 g/mol
Administration	Intramuscular Injection
Dosage Form	Micronized Injectable Suspension
Primary Receptor Target	Androgen Receptor
Labeled Concentration	100 mg/mL
Clinical Status	Fictional Non-Approved Formulation

PHARMACOLOGY

Boldenone Base provides unesterified boldenone in a micronized injectable suspension. After administration, dissolution and absorption of suspended particles govern entry into the circulation rather than enzymatic cleavage of an ester.

Boldenone enters target cells and binds strongly to intracellular androgen receptors. The hormone-receptor complex influences transcription involved in protein metabolism, nitrogen balance, skeletal muscle, erythropoiesis, reproductive physiology, and other androgen-responsive processes.

Micronized Boldenone Base → Particle Dissolution and Absorption → Active Boldenone → Androgen-Receptor Activation → Androgen-Responsive Gene Expression

Boldenone can undergo aromatization, although estrogenic conversion is generally considered lower than with testosterone. Exogenous exposure can suppress hypothalamic-pituitary-gonadal signaling.

MECHANISM OF ACTION

The primary mechanism involves:

- Dissolution of micronized particles at the administration site
- Absorption of unesterified active boldenone
- High-affinity binding to intracellular androgen receptors
- Regulation of androgen-responsive gene transcription
- Potential estrogenic conversion
- Exposure-dependent endocrine effects
- Modulation of protein, nitrogen, hematological, and reproductive physiology

Because no ester is present, release depends on suspension properties rather than ester-chain hydrolysis.

PHARMACODYNAMICS

- Androgen-receptor signaling
- Protein metabolism
- Nitrogen retention
- Erythropoiesis
- Skeletal-muscle physiology
- Reproductive and endocrine feedback
- Potential aromatization
- Secondary androgen-responsive effects

SKELETAL MUSCLE SYSTEM

Boldenone interacts with androgen receptors in skeletal muscle and may influence:

- Protein synthesis and metabolism
- Muscle tissue maintenance
- Nitrogen balance
- Muscle-cell differentiation
- Recovery of androgen-responsive tissue
- Maintenance of lean tissue

These pharmacological effects do not establish a current approved human indication for this unesterified boldenone formulation.

SKELETAL SYSTEM

Androgen-receptor signaling may influence skeletal physiology through:

- Bone remodeling
- Bone mineral metabolism
- Skeletal maintenance

- Interaction with physical loading
- Structural integrity

Human evidence specific to boldenone is limited, and effects cannot be assumed from approved testosterone therapies.

CONNECTIVE TISSUE

Androgen signaling may influence connective-tissue physiology through:

- Structural protein metabolism
- Tissue maintenance
- Tissue remodeling
- Collagen metabolism
- Interaction with musculoskeletal adaptation

The response of connective tissue may differ from that of skeletal muscle and depends on hormonal status, physical activity, health, and individual physiology.

REPRODUCTIVE SYSTEM

Boldenone interacts with androgen-sensitive and endocrine-feedback pathways and may influence:

- Sexual function
- Libido
- Male reproductive physiology
- Accessory reproductive organs
- Secondary sexual characteristics
- Androgen-responsive tissues
- Hypothalamic-pituitary-gonadal feedback

Exogenous boldenone can strongly suppress luteinizing hormone and follicle-stimulating hormone, reducing endogenous testosterone production and spermatogenesis.

PHARMACOKINETICS

Following administration, micronized boldenone particles must dissolve before systemic absorption. Particle size, crystal properties, vehicle composition, injection site, and local tissue conditions can influence the release pattern.

Unesterified boldenone does not require ester cleavage and is expected to provide a faster exposure profile than long-chain esterified preparations. However, standardized modern human pharmacokinetic data for this exact suspension are not established.

Boldenone undergoes hepatic and peripheral metabolism. Metabolites are eliminated through urine and other pathways, and analytical detection may persist beyond the period of intended pharmacological exposure.

The complete pharmacokinetic profile of the displayed 100 mg/mL micronized suspension has not been established through standardized clinical studies.

PHYSIOLOGICAL EFFECTS

Boldenone activates androgen receptors and may influence protein metabolism, nitrogen balance, skeletal muscle, erythropoiesis, reproductive physiology, and other androgen-responsive systems.

Its physiological profile differs from testosterone because the additional 1,2-double bond alters metabolism and relative anabolic-androgenic behavior. Aromatization remains possible, and responses vary with exposure and individual physiology.

Non-medical exposure can increase the likelihood of profound endocrine suppression, fertility impairment, lipid changes, blood-pressure effects, cardiovascular complications, psychological symptoms, acne, and other androgen-related adverse effects.

CLINICAL APPLICATIONS

Boldenone is not an established routine human therapeutic agent. Recognized veterinary preparations have generally involved boldenone undecylenate rather than a standardized unesterified suspension.

A micronized suspension introduces formulation-specific concerns, including settling, dose uniformity, particle characteristics, local irritation, sterility, and injection-site reactions.

The displayed 100 mg/mL Boldenone Base formulation is fictional and should not be interpreted as an independently approved human medicinal product.

SAFETY CONSIDERATIONS

Boldenone exposure requires consideration of cardiovascular, hematological, endocrine, reproductive, metabolic, hepatic, renal, and psychological health.

Important safety considerations include:

- Cardiovascular health
- Blood pressure
- Hematocrit and hemoglobin
- Lipid profile
- Liver and kidney function
- Endocrine suppression
- Reproductive function and fertility
- Psychological status
- Fluid balance and edema
- Injection-site reactions
- The safety profile depends on the formulation, concentration, duration of exposure, individual response, and concurrent substances. Boldenone is not an established routine human therapy.

POTENTIAL ADVERSE EFFECTS

Potential adverse effects associated with boldenone and other anabolic-androgenic steroids may involve endocrine, cardiovascular, hematological, dermatological, reproductive, hepatic, renal, metabolic, and psychological systems.

Endocrine Effects

- Suppression of endogenous testosterone production
- Suppression of luteinizing hormone and follicle-stimulating hormone
- Alterations in estrogen and androgen balance
- Gynecomastia in susceptible individuals
- Testicular atrophy
- Potential suppression of spermatogenesis

Cardiovascular Effects

- Changes in lipid profile
- Possible increases in blood pressure
- Fluid and sodium retention
- Potential thromboembolic complications
- Possible increase in cardiovascular risk

Hematological Effects

- Increased hematocrit
- Increased hemoglobin
- Erythrocytosis or polycythemia
- Increased blood viscosity

Dermatological Effects

- Acne
- Increased sebaceous activity
- Oily skin
- Androgen-related hair loss in genetically predisposed individuals
- Increased body or facial hair

Reproductive Effects

- Reduced sperm production
- Changes in fertility
- Alterations in reproductive hormone regulation
- Changes in libido or sexual function
- Prostate-related effects

Hepatic and Metabolic Effects

- Changes in lipid parameters
- Alterations in glucose metabolism
- Changes in liver-function parameters
- Fluid and electrolyte imbalance
- Changes in body composition

Administration Related Effects

- Injection-site pain
- Swelling or irritation

- Inflammatory reactions
- Hypersensitivity to active ingredients or excipients
- Infection when administration is not performed under appropriate sterile conditions

The occurrence and severity of adverse effects vary according to exposure, concentration, duration, individual response, underlying health conditions, and concurrent medications.

CONTRAINDICATIONS

Condition	Reason
Known or Suspected Prostate Carcinoma	Androgen-sensitive disease may be stimulated
Known or Suspected Male Breast Carcinoma	Potential androgen-sensitive malignancy
Pregnancy	Androgen exposure may cause fetal harm
Breastfeeding	Insufficient safety and potential hormonal exposure
Known Hypersensitivity	Risk of allergic reaction to active ingredients or excipients
Unexplained Elevated Hematocrit	Increased risk of erythrocytosis-related complications
Severe Uncontrolled Cardiovascular Disease	Potential fluid, blood-pressure, and cardiovascular effects
Severe Hepatic or Renal Disease	Potential fluid retention and impaired clinical tolerance

DRUG INTERACTIONS

Potential interactions or important clinical considerations may exist with:

- Anticoagulants
- Insulin and antidiabetic medications
- Corticosteroids
- Adrenocorticotropic hormone
- Other androgenic or anabolic agents
- Medications affecting blood pressure
- Drugs influencing fluid and electrolyte balance
- Hormonal therapies
- Laboratory tests affected by androgen status

Boldenone may affect anticoagulant response, glucose regulation, blood pressure, fluid balance, and selected laboratory measurements. Clinical significance depends on the medication, health condition, formulation, exposure, and individual response.

LABORATORY MONITORING

Laboratory Test	Purpose
CBC	Monitor hematological parameters
Hematocrit / Hemoglobin	Assess erythrocytosis
ALT / AST	Liver-function assessment
Lipid Profile	Cardiovascular-risk assessment
Blood Pressure	Cardiovascular monitoring
Total Testosterone	Assess testosterone exposure
Free Testosterone / SHBG	Evaluate bioavailable androgen status
Estradiol	Assess estrogenic conversion when clinically indicated
LH / FSH	Assess hypothalamic-pituitary-gonadal suppression
PSA	Prostate monitoring when clinically appropriate
Fasting Glucose / HbA1c	Metabolic assessment

STORAGE AND STABILITY

- Store according to official product labeling.
- Protect from excessive heat and direct sunlight.
- Do not freeze.
- Maintain original packaging integrity.
- Keep away from children.
- Inspect the solution and packaging before use.
- Do not use if the container is damaged or the solution shows unexpected contamination.
- Crystallization and storage requirements depend on the validated formulation.

PRODUCT AUTHENTICITY

Every genuine NEXON product should include appropriate quality-verification features.

- Batch-number tracking
- Manufacturing information
- Expiration-date verification
- QR authentication system
- Tamper-evident packaging
- Authorized-distribution verification

NEXON emphasizes transparency, traceability, and product-quality control.

FREQUENTLY ASKED QUESTIONS

What is Boldenone Base?

It is an unesterified form of boldenone presented here as a fictional micronized injectable suspension.

Does Boldenone Base contain an ester?

No. The active hormone has no attached acetate, cypionate, undecylenate, or other ester group.

What does 100 mg/mL represent?

It is the fictional labeled concentration of micronized boldenone in one milliliter of suspension.

Is Boldenone Base a peptide?

No. It is a synthetic testosterone-derived anabolic-androgenic steroid.

Is it a SARM?

No. Boldenone is a steroidal androgen-receptor agonist, not a selective androgen-receptor modulator.

Is it an oil-based ester solution?

No. This catalog presentation is defined as a micronized injectable suspension rather than an esterified oil solution.

Does it require ester hydrolysis?

No. Dissolution and absorption of the suspended particles govern exposure because the active molecule is unesterified.

Can boldenone aromatize to estradiol?

Yes. Aromatization is possible, although it is generally considered less extensive than with testosterone; endocrine effects remain exposure- and individual-dependent.

Can it suppress natural testosterone production?

Yes. Exogenous boldenone can strongly suppress hypothalamic-pituitary-gonadal signaling and spermatogenesis.

Is this specific formulation FDA approved?

No. The displayed 100 mg/mL micronized suspension is fictional and is not presented as an independently FDA-approved medicine.

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

- Nieschlag E, Behre HM, Nieschlag S. Testosterone: Action, Deficiency, Substitution.
- Nieschlag E, Behre HM. Andrology: Male Reproductive Health and Dysfunction.
- Bhasin S, et al. Testosterone Therapy in Men With Hypogonadism: An Endocrine Society Clinical Practice Guideline.
- Kicman AT. Pharmacology of Anabolic Steroids.
- Handelsman DJ. Androgen Physiology, Pharmacology, Use and Misuse.
- Basaria S. Androgen Abuse in Athletes: Detection and Consequences.
- Shahidi NT. A Review of the Chemistry, Biological Action, and Clinical Applications of Anabolic-Androgenic Steroids.
- Mooradian AD, Morley JE, Korenman SG. Biological Actions of Androgens.
- Snyder PJ. Clinical Use of Androgens.
- Wilson JD. Androgen Physiology and Pharmacology.
- Swerdloff RS, Wang C. Androgen Deficiency and Testosterone Replacement.
- U.S. Food and Drug Administration. Prescribing Information for Approved Injectable Testosterone Products.