



# Voluntary Anti-Doping Association Official Prohibited List

This document contains the Official Prohibited List (Substances and Methods) of the Voluntary Anti-Doping Association (VADA). VADA guidelines concerning these specific substances and groups are intended to closely track internationally recognized standards for substances prohibited by sport, such as the World Anti-Doping Agency (WADA) Official Prohibited List of 2021. Therefore, nomenclature for these substances, classification groups, and other uses by the WADA Prohibited List is largely preserved.

## VADA Prohibited Substance List

The following classification groups and substances are **prohibited at all times** during participation in the VADA program. The following classification groups and substances listed herein are not restricted to the specifically-listed common or chemical names, nor are they restricted to the specific compounds or isomers listed below. Moreover, VADA has the right, at any time, to modify, edit, and add any substance or method according to any new laws, guidelines, VADA policies, or anti-doping ideals.

### **S0. NON-APPROVED SUBSTANCES**

Any pharmacological substance which is not addressed by any of the subsequent sections of the List and with no current approval by any governmental regulatory health authority for human therapeutic use (e.g. drugs under pre-clinical or clinical development or discontinued, designer drugs, substances approved only for veterinary use) is prohibited at all times.

### **S1. ANABOLIC AGENTS**

**Anabolic agents are prohibited.**

#### **1. Anabolic Androgenic Steroids (AAS)**

when administered exogenously, including but not limited to:

- 1-Androstenediol (5 $\alpha$ -androst-1-ene-3 $\beta$ ,17 $\beta$ -diol)
- 1-Androstenedione (5 $\alpha$ -androst-1-ene-3,17-dione)
- 1-Androsterone (3 $\alpha$ -hydroxy-5 $\alpha$ -androst-1-ene-17-one)
- 1-Epiandrosterone (3 $\beta$ -hydroxy-5 $\alpha$ -androst-1-ene-17-one)
- 1-Testosterone (17 $\beta$ -hydroxy-5 $\alpha$ -androst-1-ene-3-one)
- 4-Androstenediol (androst-4-ene-3 $\beta$ , 17 $\beta$ -diol)

4-Hydroxytestosterone (4,17 $\beta$ -dihydroxyandrost-4-en-3-one)  
5-Androstenedione (androst-5-ene-3,17-dione)  
7 $\alpha$ -hydroxy-DHEA  
7 $\beta$ -hydroxy-DHEA  
7-Keto-DHEA  
19-Norandrostenediol (estr-4-ene-3,17-diol)  
19-Norandrostenedione (estr-4-ene-3,17-dione)  
Androstanolone (5 $\alpha$ -dihydrotestosterone, 17 $\beta$ -hydroxy-5 $\alpha$ -androstan-3-one)  
Androstenediol (androst-5-ene-3 $\beta$ ,17 $\beta$ -diol)  
Androstenedione (androst-4-ene-3,17-dione)  
Bolasterone  
Boldenone  
Boldione (androsta-1,4-diene-3,17-dione)  
Calusterone  
Clostebol  
Danazol ([1,2]oxazolo[4',5':2,3]pregna-4-en-20-yn-17 $\alpha$ -ol)  
Dehydrochlormethyltestosterone (4-chloro-17 $\beta$ -hydroxy-17 $\alpha$ -methylandrosta-1,4-dien-3-one)  
Desoxymethyltestosterone (17 $\alpha$ -methyl-5 $\alpha$ -androst-2-en-17 $\beta$ -ol and 17 $\alpha$ -methyl-5 $\alpha$ -androst-3-en-17 $\beta$ -ol)  
Drostanolone  
Epiandrosterone (3 $\beta$ -hydroxy-5 $\alpha$ -androst-17-one)  
Epi-dihydrosterone (17 $\beta$ -hydroxy-5 $\beta$ -androstan-3-one)  
Epitestosterone  
Ethylestrenol (19-norpregna-4-en-17 $\alpha$ -ol)  
Fluoxymesterone  
Formebolone  
Furazabol (17 $\alpha$ -methyl [1,2,5]oxadiazolo(3',4':2,3)-5 $\alpha$ -androstan-17 $\beta$ -ol)  
Gestrinone  
Mestanolone  
Mesterolone  
Metandienone (17 $\beta$ -hydroxy-17 $\alpha$ -methylandrosta-1,4-dien-3-one)  
Metenolone  
Methandriol  
Methasterone (17 $\beta$ -hydroxy-2 $\alpha$ ,17 $\alpha$ -dimethyl-5 $\alpha$ -androstan-3-one)  
Methyl-1-testosterone (17 $\beta$ -hydroxy-17 $\alpha$ -methyl-5 $\alpha$ -androst-1-en-3-one)  
Methylclostebol  
Methyldienolone (17 $\beta$ -hydroxy-17 $\alpha$ -methylestra-4,9-dien-3-one)  
Methylnortestosterone (17 $\beta$ -hydroxy-17 $\alpha$ -methylestr-5 $\alpha$ -4-en-3-one)  
Methyltestosterone  
Metribolone (methyltrienolone, 17 $\beta$ -hydroxy-17 $\alpha$ -methylestra-4,9,11-trien-3-one)  
Mibolerone  
Nandrolone (19-nortestosterone)  
Norboletone  
Norclostebol (4-chloro-17 $\beta$ -ol-estr-4-en-3-one)  
Norethandrolone  
Oxabolone  
Oxandrolone  
Oxymesterone

Oxymetholone  
Prasterone (dehydroepiandrosterone, DHEA, 3 $\beta$ -hydroxyandrost-5-en-17-one)  
Prostanazol (17 $\beta$ -((tetrahydropyran-2-yl)oxy)-1'H-pyrazolo(3,4:2,3)-5 $\alpha$ -androstane)  
Quinbolone  
Stanozolol  
Stenbolone  
Testosterone  
Tetrahydrogestrinone (17-hydroxy-18 $\alpha$ -homo-19-nor-17 $\alpha$ -pregna-4,9,11-trien-3-one)  
Trenbolone (17  $\beta$ -hydroxyestr-4,9,11-trien-3-one)

and other substances with a similar chemical structure or similar biological effect(s).

## 2. Other Anabolic Agents

**Including, but not limited to:**

- Clenbuterol, selective androgen receptor modulators (SARMs, e.g. andarine, LGD-4033 (ligandrol), enobosarm (ostarine) and RADA 140), tibolone, zeranol and zilpatrol

## **S2. PEPTIDE HORMONES, GROWTH FACTORS, RELATED SUBSTANCES, AND MIMETICS**

**The following substances, and other substances with similar chemical structure or similar biological effects(s), are prohibited:**

1. Erythropoietins (EPO) and agents affecting erythropoiesis, including, but not limited to:

1.1 Erythropoietin-receptor agonists, e.g.

Darbepoetins (dEPO);  
Erythropoietins (EPO);  
EPO based constructs (e.g. EPO-Fc, methoxy polyethylene glycol-epoetin beta (CERA))  
EPO-mimetic agents and their constructs (e.g. CNTO 530 and peginesatide).

1.2 Hypoxia-inducible factor (HIF) activating agents, e.g.

Cobalt;  
Daprodustat (GSK1278863);  
IOX2;  
Molidustat (BAY 85-3934);  
Roxadustat (FG-4592);  
Vadadustat (AKB-6548);  
Xenon.

1.3 GATA inhibitors, e.g.

K-11706

1.4 Transforming growth factor beta (TGF- $\beta$ ) signaling inhibitors, e.g.

Luspatercept;  
Sotatercept.

1.5 Innate repair receptor agonists, e.g.

Asialo EPO;  
Carbamylated EPO (CEPO).

**2. Peptide Hormones and their Releasing Factors,**

**2.1** Chorionic Gonadotrophin (CG) and Luteinizing Hormone (LH) and their releasing factors in males, e.g. busserelin, deslorelin, gonadorelin, goserelin, leuprorelin, nafarelin and triptorelin;

**2.2** Corticotrophins and their releasing factors, e.g. corticorelin;

**2.3** Growth hormone (GH), its fragments and releasing factors, including, but not limited to: Growth hormone fragments, e.g. ADD-9604 and hGH 176-191;

Growth hormone-releasing (GHRH) and its analogues, e.g. CJC-1293, CJC-1295, sermorelin and tesamorelin; growth hormone secretagogues

(GHS), e.g. lenomorelin (ghrelin) and its mimetics, e.g. anamorelin, ipamorelin, macimorelin and tabimorelin; GH-releasing peptides (GHRPs), e.g. alexamorelin, GHRP-1,

GHRP-2 (pralmorelin), GHRP-3, GHRP-4, GHRP-5, GHRP-6, and examorelin (hexarelin).

**3. Growth Factors and Growth Factor Modulators, including, but not limited to:**

Fibroblast growth factors (FGFs)

Hepatocyte growth factor (HGF)

Insulin-like growth factor-1 (IGF-1) and its analogues

Mechano growth factors (MGFs)

Platelet-derived growth factor (PDGF)

Thymosin-  $\beta$ 4 and its derivatives e.g. TB-500

Vascular endothelial growth factor (VEGF)

and other growth factors or growth factor modulators affecting muscle, tendon or ligament protein synthesis/degradation, vascularisation, energy utilization, regenerative capacity or fibre type switching.

**S3. BETA-2 AGONISTS**

**All selective and non-selective beta-2 agonists, including all optical isomers, are prohibited.**

**Including, but not limited to:**

Arformoterol;

Fenoterol;

Formoterol;

Higenamine;

Indacaterol;

Levosalbutamol;

Olodaterol;

Procaterol;

Reproterol;

Salbutamol;

Salmeterol;

Terbutaline;

Tretoquinol (trimetoquinol)  
Tulobuterol;  
Vilanterol.

**Exceptions:**

- Inhaled salbutamol: maximum 1600 micrograms over 24 hours in divided doses not to exceed 800 micrograms over 12 hours starting from any dose;
- Inhaled formoterol: maximum delivered dose of 54 micrograms over 24 hours;
- Inhaled salmeterol: maximum 200 micrograms over 24 hours;
- Inhaled vilanterol: maximum 25 micrograms over 24 hours.

**Note:**

The presence in urine of salbutamol in excess of 1000 ng/ml or formoterol in excess of 40 ng/ml is not consistent with therapeutic use of the substance and will be considered as an *Adverse Analytical Finding (AAF)* unless the *Athlete* proves, through a controlled pharmacokinetic study, that the abnormal result was the consequence of a therapeutic dose (by inhalation) up to the maximum dose indicated above.

**S4. HORMONE AND METABOLIC MODULATORS**

**The following hormone and metabolic modulators are prohibited:**

1. **Aromatase inhibitors** including, but not limited to:
  - 2-Androstenol (5 $\alpha$ -androst-2-en-17-ol)
  - 2-Androstenone (5 $\alpha$ -androst-2-en-17-one)
  - 3-Androstenol (5 $\alpha$ -androst-3-en-17-ol)
  - 3-Androstenone (5 $\alpha$ -androst-3-en-17-one)
  - 4-Androstene-3,6,17 trione (6-oxo)
  - Aminoglutethimide
  - Anastrozole
  - Androsta-1,4,6-triene-3,17-dione (androstatrienedione)
  - Androsta-3,5-diene-7,17-dione (arimistane)
  - Exemestane
  - Formestane
  - Letrozole
  - Testolactone
2. Anti-estrogenic substances (anti-estrogens and selective estrogen receptor modulators (SERMS)). Including, but not limited to:
  - Bazedoxifene
  - Clomifene
  - Cyclofenil
  - Fulvestrant
  
  - Ospemifene
  - Raloxifene
  - Tamoxifen
  - Toremifene

3. Agents preventing activin receptor IIB activation including, but not limited, to:

Activin A-neutralizing antibodies;

Activin receptor IIB competitors such as:

Decoy activin receptors (e.g. ACE-031);

Anti-activin receptor IIB antibodies (e.g. bimagrumab);

Myostatin inhibitors such as:

Agents reducing or ablating myostatin expression;

Myostatin-binding proteins (e.g. Follistatin, myostatin, propeptide);

Myostatin-neutralizing antibodies (e.g. Domagrozumab, landogrozumab, stamulumab).

4. Metabolic modulators:

4.1 Activators of the AMP-activated protein kinase (AMPK), e.g. AICAR, SR9009; and Peroxisome Proliferator Activated Receptor  $\delta$  (PPAR $\delta$ ) agonists, e.g. 2-(2-methyl-4-((4-methyl-2-(4-(trifluoromethyl) phenyl)thiazol-5-yl)methylthio)phenoxy) acetic acid (GW 1516, GW501516);

4.2 Insulins and insulin-mimetics

4.3 Meldonium

4.4 Trimetazidine

## **S5. DIURETICS AND OTHER MASKING AGENTS**

The following diuretics and masking agents are prohibited, as are other substances with a similar chemical structure or similar biological effect(s).

**Including, but not limited to:**

- Desmopressin; probenecid; plasma expanders, e.g. intravenous administration of albumin, dextran, hydroxyethyl starch and mannitol;
- Acetazolamide; amiloride; bumetanide; canrenone; chlortalidone; etacrynic acid; furosemide; indapamide; metolazone; spironolactone; thiazides, e.g. Bendroflumethiazide, chlorothiazide and hydrochlorothiazide; triamterene and vaptans, e.g. tolvaptan

**Exceptions:**

- Drospirenone; pamabrom; and topical ophthalmic administration of carbonic anhydrase inhibitors (e.g. dorzolamide, brinzolamide);
- Local administration of felypressin in dental anaesthesia

**Note:**

The detection in an *Athlete's Sample* at all times, as applicable, of any quantity of the following substances subject to threshold limits: formoterol, salbutamol, cathine, ephedrine, methylephedrine and pseudoephedrine, in conjunction with a diuretic or masking agent, will be considered as an *Adverse Analytical Finding (AAF)* unless the *Athlete* has an approved *Therapeutic Use Exemption (TUE)* for that substance in addition to the one granted for the diuretic or masking agent.

## **S6. STIMULANTS**

**All stimulants, including all optical isomers, e.g. d- and l- where relevant, are prohibited.**

**Stimulants include, but are not limited to:**

Adrafinil  
Amfepramone  
Amfetamine  
Amfetaminil  
Amiphenazole  
Benfluorex  
Benzylpiperazine  
Bromantan  
Clobenzorex  
Cocaine  
Cropropamide  
Crotetamide  
Fencamine  
Fenetylline  
Fenfluramine  
Fenproporex  
Fonturancetam (4-phenylpiracetam (carphedon))  
Furfenorex  
Lisdexamfetamine  
Mefenorex  
Mephentermine  
Mesocarb  
Methamphetamine (*d*-)  
p-methylamphetamine  
Modafinil  
Norfenfluramine  
Phendimetrazine  
Phentermine  
Prenylamine  
Prolintane

3-Methylhexan-2-amine (1,2-dimethylpentylamine)  
4-Methylhexan-2-amine (methylhexaneamine)  
4-Methylpentan-2-amine (1,3 dimethylbutylamine)  
5-Methylhexan-2-amine (1,4-dimethylpentylamine)  
Benzfetamine  
Cathine\*\*  
Cathinone and its analogues, e.g. mephedrone, methedrone, and  $\alpha$ -pyrrolidinovalerophenone  
Dimetamfetamine (dimethylamphetamine)  
Ephedrine\*\*\*

Epinephrine\*\*\*\* (adrenaline)  
Etamivan  
Etilamfetamine  
Etilefrine  
Famprofazone  
Fenbutrazate  
Fencamfamin  
Heptaminol  
Hydroxyamfetamine (parahydroxyamphetamine)  
Isometheptene  
Levmetamfetamine  
Meclofenoxate  
Methylenedioxyamphetamine  
Methyephedrine\*\*\*  
Methylphenidate  
Nikethaminde  
Norfenefrine  
Octodrine (1,5-dimethylhexylamine)  
Octopamine;  
Oxilofrine (methylnephrine)  
Pemoline  
Penetrazol  
Phenethylamine and its derivatives  
Phenmetrazine  
Phenpromethamine  
Propylhexedrine  
Pseudoephedrine\*\*\*\*\*  
Selegiline  
Sibutramine  
Strychnine  
Tenamfetamine (methylenedioxyamphetamine)  
Tuaminoheptane

and other substances with a similar chemical structure or similar biological effect(s).

**Exceptions:**

- Clonidine
- Imidazole derivatives for dermatological, nasal or ophthalmic use (e.g. brimondine, clonazoline, fenoxazoline, indanazoline, naphazoline, oxymetazoline, xylometazoline) and those stimulants included in WADA's 2021 Monitoring Program\*.

\*Bupropion, caffeine, nicotine, phenylephrine, phenylpropanolamine, pipradrol, and synephrine: These substances are included in WADA's 2021 Monitoring Program, and are not considered Prohibited Substances.

\*\*Cathine: Prohibited when its concentration in urine is greater than 5 micrograms per milliliter.

\*\*\*Ephedrine and methylephedrine: Prohibited when the concentration of either in urine is greater 10 micrograms per milliliter.

\*\*\*\*Epinephrine (adrenaline): Not prohibited in local administration, e.g. nasal, ophthalmologic, or co-administration with local anaesthetic agents.

\*\*\*\*\*Pseudoephedrine: Prohibited when its concentration in urine is greater than 150 micrograms per milliliter.



## **S7. NARCOTICS**

*Substance of Abuse* in this section: diamorphine (heroin)

**The following narcotics, including all optical isomers, e.g. d- and l- where relevant are prohibited:**

Buprenorphine  
Dextromoramide  
Diamorphine (heroin)  
Fentanyl and its derivatives  
Hydromorphone  
Methadone  
Morphine  
Nicomorphine  
Oxycodone  
Oxymorphone  
Pentazocine  
Pethidine

## **S9. GLUCOCORTICOIDS**

All glucocorticoids are prohibited when administered by oral, intravenous, intramuscular or rectal routes.

**Including but not limited to:**

Beclometasone  
Bethamethesone  
Budesonide  
Ciclesonide  
Cortisone  
Deflazacort  
Dexamethasone  
Flucortolone  
Flunisolide  
Fluticasone  
Hydrocortisone  
Methylprednisolone  
Mometasone  
Prednisolone  
Prednisone  
Triamcinolone acetonide

## **PROHIBITED METHODS**

### **M1. MANIPULATION OF BLOOD AND BLOOD COMPONENTS**

**The following are prohibited:**

1. The *Administration* or reintroduction of any quantity of autologous, allogenic (homologous) or heterologous blood, or red blood cell products of any origin into the circulatory system.

2. Artificially enhancing the uptake, transport or delivery of oxygen.

Including, but not limited to:

Perfluorochemicals; efaproxiral (RSR13) and modified haemoglobin products, e.g. haemoglobin-based blood substitutes, and microencapsulated haemoglobin products, excluding supplemental oxygen by inhalation.

3. Any form of intravascular manipulation of the blood or blood components by physical or chemical means.

### **M2. CHEMICAL AND PHYSICAL MANIPULATION**

**The following are prohibited:**

1. *Tampering, or Attempting to Tamper*, to alter the integrity and validity of *Samples* collected during *Doping Control*.

Including, but not limited to:

*Sample* substitution and/or adulteration, e.g. addition of proteases to *Sample*.

2. Intravenous infusions and/or injections of more than a total of 100ml per 12-hour period except for those legitimately received in the course of hospital treatments, surgical procedures or clinical diagnostic investigations.

### **M3. GENE AND CELL DOPING**

**The following, with the potential to enhance sport performance, are prohibited:**

1. The use of nucleic acids or nucleic acid analogues that may alter genome sequences and/or alter gene expression by any mechanism. This includes but is not limited to gene editing, gene silencing and gene transfer technologies.

2. The use of normal or genetically modified cells.