

# Stullmisan vet. Pulver

Authorised

- Abies sibirica extract

## Product identification

**Medicine name:**

Stullmisan vet. Pulver

---

**Active substance:**

Abies sibirica extract

---

**Target species:**

Quail

Turkey

Cattle

Chicken

Chicken (for reproduction)

Cattle (pre-ruminant)

Dog

Sheep

Horse

Cat

Guinea pig

Hamster

Pig (suckling piglet)

Horse (foal)

Rabbit (non food-producing)

Pig (piglet)

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Abies sibirica extract

30.56 milligram(s) / 1.00 gram(s)

---

**Pharmaceutical form:**

Oral powder

---

**Withdrawal period by route of administration:****Oral use:**

- 

**Quail**

- Egg. 0 day
- Meat and offal. 1 day

- 

**Turkey**

- Egg. 0 day
- Meat and offal. 1 day

- 

**Cattle**

- Meat and offal. 1 day
- Milk. 1 day

- 

**Chicken (for reproduction)**

- Meat and offal. 1 day
- Egg. 0 day

- 

**Cattle (pre-ruminant)**

- Meat and offal. 1 day

- 

#### **Sheep**

- Meat and offal. 1 day

- Milk. 1 day

- 

#### **Horse**

- Meat and offal. 1 day

- Milk. 1 day

- 

#### **Pig (suckling piglet)**

- Meat and offal. 1 day

- 

#### **Horse (foal)**

- Meat and offal. 1 day

- 

#### **Pig (piglet)**

- Meat and offal. 1 day

---

### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA07

---

### **Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

---

### **Authorisation status:**

Valid

---

### **Authorised in:**

Germany

---

### **Package description:**

Available only in German

Available only in German

Available only in German

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Pharma Stulln GmbH

---

**Marketing authorisation date:**

21/06/2010

---

**Manufacturing sites for batch release:**

Pharma Stulln GmbH

---

**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

---

**Authorisation number:**

401262.00.00

---

**Date of authorisation status change:**

29/09/2015

---

To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics
------------------------------------

This document does not exist in this language (English). You can find it in another language below.

2401262-parde-20100614.pdf