

## 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

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## Product details

**Active substance and strength:**

Abies sibirica extract

30.56 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Oral powder

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

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**Quail**

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

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**Turkey**

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

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**Cattle**

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

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**Chicken (for reproduction)**

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

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**Cattle (pre-ruminant)**

- Meat and offal. 1 day

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### **Sheep**

- Meat and offal. 1 day

- Milk. 1 day

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### **Horse**

- Meat and offal. 1 day

- Milk. 1 day

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

- Meat and offal. 1 day

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### **Horse (foal)**

- Meat and offal. 1 day

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### **Pig (piglet)**

- Meat and offal. 1 day

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

QA07

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

Veterinary medicinal product not subject to veterinary prescription

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### **Authorisation status:**

Valid

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### **Authorised in:**

Germany

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### **Package description:**

Available only in German

Available only in German

Available only in German

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Pharma Stulln GmbH

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**Marketing authorisation date:**

21/06/2010

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**Manufacturing sites for batch release:**

Pharma Stulln GmbH

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

401262.00.00

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**Date of authorisation status change:**

29/09/2015

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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.

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