

# CHELIDONIUM -HOMACCORD VETERINARIO

Authorised

- ATROPA BELLA-DONNA D10
- ATROPA BELLA-DONNA D1000
- ATROPA BELLA-DONNA D200
- ATROPA BELLA-DONNA D30
- ATROPA BELLA-DONNA D4
- CHELIDONIUM MAJUS D10
- CHELIDONIUM MAJUS D200
- FEL TAURI D10
- FEL TAURI D200

## Product identification

**Medicine name:**

CHELIDONIUM -HOMACCORD VETERINARIO

**Active substance:**

ATROPA BELLA-DONNA D10  
ATROPA BELLA-DONNA D1000  
ATROPA BELLA-DONNA D200  
ATROPA BELLA-DONNA D30  
ATROPA BELLA-DONNA D4  
CHELIDONIUM MAJUS D10  
CHELIDONIUM MAJUS D200  
FEL TAURI D10

FEL TAURI D200

---

**Target species:**

Cattle (cow)  
Cattle  
Ornamental bird  
Dog  
Goat  
Sheep (ewe)  
Sheep  
Horse  
Cat  
Rabbit  
Fish  
Rodents  
Pig

---

**Route of administration:**

Oral use  
Intramuscular use  
Intravenous use  
Subcutaneous use

---

## Product details

**Active substance and strength:**

ATROPA BELLA-DONNA D10  
15.00 milligram(s) / 5.00 millilitre(s)  
ATROPA BELLA-DONNA D1000  
15.00 milligram(s) / 5.00 millilitre(s)  
ATROPA BELLA-DONNA D200  
15.00 milligram(s) / 5.00 millilitre(s)  
ATROPA BELLA-DONNA D30  
15.00 milligram(s) / 5.00 millilitre(s)  
ATROPA BELLA-DONNA D4  
15.00 milligram(s) / 5.00 millilitre(s)

CHELIDONIUM MAJUS D10  
30.00 milligram(s) / 5.00 millilitre(s)

CHELIDONIUM MAJUS D200  
30.00 milligram(s) / 5.00 millilitre(s)

FEL TAURI D10  
5.00 milligram(s) / 5.00 millilitre(s)

FEL TAURI D200  
5.00 milligram(s) / 5.00 millilitre(s)

---

**Pharmaceutical form:**

Solution for injection

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Italy

---

**Package description:**

Available only in [Italian](#)

Available only in [Italian](#)

---

## Additional information

**Entitlement type:**

Homeopathic Registration

---

**Marketing authorisation holder:**

Biologische Heilmittel Heel GmbH

---

**Marketing authorisation date:**

30/10/2012

---

**Manufacturing sites for batch release:**

Biologische Heilmittel Heel GmbH

---

**Responsible authority:**

Ministry Of Health

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

4/10/2012

---

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

## Documents

Package Leaflet and Labelling

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