

# INMEVA, SUSPENSION FOR INJECTION

Suspended

- Chlamydia abortus, strain A22, Inactivated
- Salmonella enterica, subsp. enterica, serovar Abortusovis, strain Sao, Inactivated

## Product identification

**Medicine name:**

INMEVA, SUSPENSION FOR INJECTION

---

**Active substance:**

Chlamydia abortus, strain A22, Inactivated

Salmonella enterica, subsp. enterica, serovar Abortusovis, strain Sao, Inactivated

---

**Target species:**

Sheep

---

**Route of administration:**

Subcutaneous use

---

## Product details

**Active substance and strength:**

Chlamydia abortus, strain A22, Inactivated

1.00 relative potency / 2.00 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Abortusovis, strain Sao, Inactivated  
1.00 relative potency / 2.00 millilitre(s)

---

**Pharmaceutical form:**

Suspension for injection

---

**Withdrawal period by route of administration:**

**Subcutaneous use:**

- 

**Sheep**

- All relevant tissues. 0 day

---

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

QI04AB

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Suspended

---

**Authorised in:**

Denmark

---

**Package description:**

Polyethylene (PET) vials of 100ml

Polyethylene (PET) vials of 50ml

Polyethylene (PET) vials of 250ml

Polyethylene (PET) vials of 10ml

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Laboratorios Hipra S.A.

---

**Marketing authorisation date:**

29/05/2019

---

**Manufacturing sites for batch release:**

Laboratorios Hipra, S.A.

---

**Responsible authority:**

Danish Medicines Agency

---

**Authorisation number:**

61285

---

**Date of authorisation status change:**

29/05/2019

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0350/001

---

**Concerned member states:**

Austria Belgium Denmark Germany Greece Hungary Ireland Italy  
Luxembourg Netherlands Poland Portugal Romania Spain Sweden  
United Kingdom (Northern Ireland)

---

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.