

# INMEVA, SUSPENSION FOR INJECTION

Authorised

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

## Product identification

**Medicine name:**

INMEVA, SUSPENSION FOR INJECTION

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**Active substance:**

Chlamydia abortus, strain A22, Inactivated

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

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**Target species:**

Sheep

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**Route of administration:**

Subcutaneous use

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## 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)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Sheep**

- All relevant tissues. 0 day

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

QI04AB

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Luxembourg

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**Package description:**

Polyethylene (PET) vials of 10ml

Polyethylene (PET) vials of 250ml

Polyethylene (PET) vials of 100ml

Polyethylene (PET) vials of 50ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Laboratorios Hipra S.A.

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**Marketing authorisation date:**

25/06/2019

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**Manufacturing sites for batch release:**

Laboratorios Hipra S.A.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

V 510/19/06/2124

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**Date of authorisation status change:**

25/06/2019

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**Reference member state:**

France

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**Procedure number:**

FR/V/0350/001

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**Concerned member states:**

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

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