SALMOPAST

Authorised

- Mannheimia haemolytica, Inactivated
- Pasteurella multocida, serotype D4, antigens
- Pasteurella multocida, serotype A3, antigens
- Salmonella enterica, subsp. enterica, serovar Typhimurium, antigens O and H
- Salmonella enterica, subsp. enterica, serovar Dublin, antigens O and H

Product identification

Medicine name:
SALMOPAST

Active substance:
Mannheimia haemolytica, Inactivated
Pasteurella multocida, serotype D4, antigens
Pasteurella multocida, serotype A3, antigens
Salmonella enterica, subsp. enterica, serovar Typhimurium, antigens O and H
Salmonella enterica, subsp. enterica, serovar Dublin, antigens O and H

Target species:
Cattle
Sheep
Goat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Mannheimia haemolytica, Inactivated
0.60 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Pasteurella multocida, serotype D4, antigens
0.60 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Pasteurella multocida, serotype A3, antigens
0.60 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Typhimurium, antigens O and H
1.50 slow agglutination test unit(s) / 1.00 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Dublin, antigens O and H
1.50 slow agglutination test unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - All relevant tissues. 0 day
- **Sheep**
  - All relevant tissues. 0 day
- **Goat**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AB
QI03AB
QI04AB

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
7/01/1981

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/8588182 8/1981

Date of authorisation status change:
7/01/2011

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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