Suibiovac ART Inaktywowane bakterie Bordetella bronchiseptica (BB 4-78) nie mniej niż 8 x 10^9, Inaktywowane bakterie Pasteurella multocida typ D (CECT 4325) nie mniej niż 8 x 10^9, Letalna dermonekrotoksyna Pasteurella multocida (DNT) nie mniej niż 1 µg, Zawiesina do wstrzykiwań

Authorised

- Bordetella bronchiseptica, strain BB4-78, Inactivated
- Pasteurella multocida, serogroup D, strain CECT 4325, Inactivated
- Pasteurella multocida, protein dO (non-toxic derivative of dermonecrotic toxin), recombinant

Product identification

Medicine name:
Suibiovac ART Inaktywowane bakterie Bordetella bronchiseptica (BB 4-78) nie mniej niż 8 x 10^9, Inaktywowane bakterie Pasteurella multocida typ D (CECT 4325) nie

mniej niż 8 x 10^9, Letalna dermonekrotoksyna Pasteurella multocida (DNT) nie mniej niż 1 µg, Zawiesina do wstrzykiwań

Active substance:
Bordetella bronchiseptica, strain BB4-78, Inactivated

Pasteurella multocida, serogroup D, strain CECT 4325, Inactivated

Pasteurella multocida, protein dO (non-toxic derivative of dermonecrotic toxin), recombinant

Target species:
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Bordetella bronchiseptica, strain BB4-78, Inactivated
8000.00 million unit(s)/millilitre / 2.00 millilitre(s)

Pasteurella multocida, serogroup D, strain CECT 4325, Inactivated
8000.00 million unit(s)/millilitre / 2.00 millilitre(s)

Pasteurella multocida, protein dO (non-toxic derivative of dermonecrotic toxin), recombinant
1.00 microgram(s) / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 0 day

Subcutaneous use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB04

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

Authorisation status:
Valid

Authorised in:
Poland

Package description:
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Biowet Drwalew Sp. z o.o.

Marketing authorisation date:
12/05/2016

Manufacturing sites for batch release:
Laboratorios Ovejero S.A.U.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2540

Date of authorisation status change:
12/05/2016

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