

Suibiovac ART Inaktywowane bakterie Bordetella bronchiseptica (BB 4-78) nie mniej niż  $8 \times 10^9$ , Inaktywowane bakterie Pasteurella multocida typ D (CECT 4325) nie mniej niż  $8 \times 10^9$ , Letalna dermonekrotoksyna Pasteurella multocida (DNT) nie mniej niż  $1 \mu\text{g}$ , 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  $8 \times 10^9$ , Inaktywowane bakterie Pasteurella multocida typ D (CECT 4325) nie

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

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**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

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**Target species:**

Pig

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## 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)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Pig**

- Meat and offal. 0 day

**Subcutaneous use:**

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**Pig**

- Meat and offal. 0 day

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

QI09AB04

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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:**

Poland

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**Package description:**

Available only in [Polish](#)

Available only in [Polish](#)

Available only in [Polish](#)

Available only in [Polish](#)

Available only in [Polish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Informed consent application (Article 13c of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Biowet Drwalew Sp. z o.o.

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**Marketing authorisation date:**

12/05/2016

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**Manufacturing sites for batch release:**

Laboratorios Ovejero S.A.U.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2540

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**Date of authorisation status change:**

12/05/2016

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