

Brucellin Aquilon (--)- Solution for injection

Authorised

- Brucella abortus, strain AQ1302, protein extract

Product identification

Medicine name:

Brucellin Aquilon (--)- Solution for injection

Active substance:

Brucella abortus, strain AQ1302, protein extract

Target species:

Pig

Route of administration:

Intradermal use

Product details

Active substance and strength:

Brucella abortus, strain AQ1302, protein extract

Presentation_strength:≥ 1 RP Reference:HSE Index:2

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intradermal use:**

-

Pig

- Not applicable. 0 day 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AR

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:vial (glass), Package_size:1 vial, Content:2.5 ml

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:

CZ Vaccines S.A.U.

Marketing authorisation date:

26/01/2023

Manufacturing sites for batch release:

CZ Vaccines S.A.U.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

26/01/2023

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

Documents

Combined File of all Documents

English (PDF)

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