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:
Medicinal product subject to medical 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:
Aquilon Cyl S.L.

Marketing authorisation date:
26/01/2023

Manufacturing sites for batch release:
CZ VETERINARIA, S.A.

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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