

Supravet, 4µg/ml, Solution for injection

Authorised

- Buserelin

Product identification

Medicine name:

Supravet, 4µg/ml, Solution for injection

Active substance:

Buserelin

Target species:

Cattle (cow)
Horse (mare)
Rabbit (female for reproduction)
Pig (sow)
Pig (sow, nullipar)

Route of administration:

Intravenous use
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Buserelin

4.00 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01CA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Package description:

Glass Vial 1 x 10.0 millilitre(s)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

P.H. Farmaceutici S.r.l.

Marketing authorisation date:

23/07/2025

Manufacturing sites for batch release:

Fundacio Privada Dau

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/027/25-C

Date of authorisation status change:

23/07/2025

Reference member state:

Czechia

Procedure number:

CZ/V/0197/001

Concerned member states:

Italy

Generic of:

600000065231

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 13/03/2026

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Package Leaflet

English (PDF)

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Labelling

English (PDF)

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