

Mesenchym comp. N PlantaVet

Authorised

- Mesenchym suis GI Dil. D5
- TEXTUS CONNECTIVUS BOVIS GL DIL. D5 (HAB, VS. 41B)
- DUCTUS THORACICUS BOVIS GL DIL. D5 (HAB, VS. 41C)
- RENES BOVIS GL DIL. D5 (HAB, VS. 41A)
- FUNICULUS UMBILICALIS BOVIS GL DIL. D5 (HAB, VS. 41B)
- Borago officinalis D5
- Cuprum aceticum D5

Product identification

Medicine name:

Mesenchym comp. N PlantaVet

Active substance:

Mesenchym suis GI Dil. D5

TEXTUS CONNECTIVUS BOVIS GL DIL. D5 (HAB, VS. 41B)

DUCTUS THORACICUS BOVIS GL DIL. D5 (HAB, VS. 41C)

RENES BOVIS GL DIL. D5 (HAB, VS. 41A)

FUNICULUS UMBILICALIS BOVIS GL DIL. D5 (HAB, VS. 41B)

Borago officinalis D5

Cuprum aceticum D5

Target species:

Cattle

Horse

Route of administration:

Subcutaneous use

Product details**Active substance and strength:**

Mesenchym suis Gl Dil. D5

1.00 gram(s) / 10.00 millilitre(s)

TEXTUS CONNECTIVUS BOVIS GL DIL. D5 (HAB, VS. 41B)

1.00 gram(s) / 10.00 millilitre(s)

DUCTUS THORACICUS BOVIS GL DIL. D5 (HAB, VS. 41C)

1.00 gram(s) / 10.00 millilitre(s)

RENES BOVIS GL DIL. D5 (HAB, VS. 41A)

1.00 gram(s) / 10.00 millilitre(s)

FUNICULUS UMBILICALIS BOVIS GL DIL. D5 (HAB, VS. 41B)

1.00 gram(s) / 10.00 millilitre(s)

Borago officinalis D5

1.00 gram(s) / 10.00 millilitre(s)

Cuprum aceticum D5

1.00 gram(s) / 10.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

-

Cattle

- Meat and offal. 0 day

- Milk. 0 day

-

Horse

- Meat and offal. 0 day

- Milk. 0 day

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

Available only in German

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

SaluVet GmbH

Marketing authorisation date:

27/12/2005

Manufacturing sites for batch release:

Wala-Heilmittel GmbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

6500259.00.00

Date of authorisation status change:

27/12/2005

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