

# HorStem 15 x 10<sup>6</sup> cells/ml - Suspension for injection

Authorised

- Equine allogeneic umbilical cord-derived mesenchymal stem cells

## Product identification

**Medicine name:**

HorStem 15 x 10<sup>6</sup> cells/ml - Suspension for injection

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**Active substance:**

Equine allogeneic umbilical cord-derived mesenchymal stem cells

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**Target species:**

Horse

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**Route of administration:**

Intraarticular use

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## Product details

**Active substance and strength:**

Equine allogeneic umbilical cord-derived mesenchymal stem cells

Presentation\_strength:15,000,000 cells Comments:range from 12,000,000 to 18,000,000 cells/ml Index:0

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Intraarticular use:**

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**Horse**

- Not applicable. 0 day Zero days

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

QM09AX

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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:**

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)

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**Available in:**

Austria , Belgium , Denmark , France , Germany , Ireland , Netherlands , Norway , Poland , Spain , Sweden

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**Package description:**

Packaging:Vial (cyclic polyolefin), Package\_size:1 vial, Content:1 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

EquiCord S.L.

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**Marketing authorisation date:**

19/06/2019

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**Manufacturing sites for batch release:**

EquiCord S.L.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

19/06/2019

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

## Documents

Combined File of all Documents

English (PDF)

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