

# Withdrawn VMP

Geautoriseerd

- IRON OXIDE RED (E172)

## Product identification

**Naam van het geneesmiddel:**

Withdrawn VMP

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**Werkzame stof:**

Alleen beschikbaar in [English](#)

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**Doeldiersoort(en):**

Hond

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**Toedieningsweg:**

Oraal gebruik

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## Product details

**Werkzame stof / Sterkte:**

Alleen beschikbaar in [English](#)  
1.00 milligram(s)/millilitre / 1.00 Blisterverpakking

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**Farmaceutische vorm:**

Kauwtablet

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**Withdrawal period by route of administration:**

**Oraal gebruik:**

- Hond
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**Anatomisch therapeutisch chemische veterinaire classificatie (ATCvet code)**

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QA01AA01

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**Afleverstatus:**Alleen beschikbaar in [German](#) [Estonian](#) [Greek](#) [English](#) [Italian](#) [Portuguese](#) [Norwegian](#)

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**Status toelating:**Valid

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**Authorised in:**Alleen beschikbaar in [Estonian](#) [English](#) [French](#) [Italian](#) [Latvian](#) [Portuguese](#) [Slovak](#) [Icelandic](#) [Norwegian](#)

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**Package description:**Alleen beschikbaar in [English](#)

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## Additional information

**Entitlement type:**Alleen beschikbaar in [English](#) [French](#) [Croatian](#) [Italian](#) [Latvian](#) [Finnish](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

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**Rechtsgrondslag productvergunning:**Alleen beschikbaar in [English](#) [Italian](#) [Latvian](#) [Norwegian](#)

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**Handelsvergunninghouder:**European Medicines Agency

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**Marketing authorisation date:**28/07/2021

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**Verantwoordelijke instantie:**European Medicines Agency

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**Toelatingsnummer:**Not Applicable

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**Wijzigingsdatum status toelating:**

27/07/2021

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

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**Source URL:** <https://medicines.health.europa.eu/veterinary/600000004400>