

# Cefa-Cure Vet. tabletter 50 mg

Geautoriseerd

- Cefadroxil monohydrate

## Product identification

### **Naam van het geneesmiddel:**

Cefa-Cure Vet. 50 mg tabletter

Cefa-Cure Vet. tabletter 50 mg

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### **Werkzame stof:**

Alleen beschikbaar in [English](#)

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### **Doeldiersoort(en):**

Kat

Hond

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### **Toedieningsweg:**

Oraal gebruik

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## Product details

### **Werkzame stof / Sterkte:**

Alleen beschikbaar in [English](#)

52.50 milligram(s) / 1.00 Tablet

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### **Farmaceutische vorm:**

Tablet

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### **Withdrawal period by route of administration:**

**Oraal gebruik:**

- **Kat**
- **Hond**

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

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QJ01DB05

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**Afleverstatus:**

Alleen beschikbaar in [Czech](#) [Estonian](#) [English](#) [French](#) [Italian](#) [Latvian](#) [Portuguese](#) [Slovenian](#) [Finnish](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

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**Status toelating:**

Valid

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**Authorised in:**

Denemarken

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**Package description:**

Alleen beschikbaar in [Danish](#)

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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](#) [French](#) [Italian](#) [Latvian](#) [Norwegian](#)

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**Handelsvergunninghouder:**

Intervet International B.V.

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**Marketing authorisation date:**

27/10/1988

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**Productielocaties partijvrijgifte:**

Intervet Productions S.r.l.

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**Verantwoordelijke instantie:**

DKMA

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**Toelatingsnummer:**

13185

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**Wijzigingsdatum status toelating:**

27/10/1988

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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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