

# Nafpenzal T Salbe zur intramammären Anwendung für Rinder (trockenstehende Milchkühe)

Authorised

- Nafcillin sodium monohydrate
- Benzylpenicillin procaine monohydrate
- Dihydrostreptomycin sulfate

## Product identification

### **Medicine name:**

Nafpenzal T Salbe zur intramammären Anwendung für Rinder (trockenstehende Milchkühe)

---

### **Active substance:**

Nafcillin sodium monohydrate  
Benzylpenicillin procaine monohydrate  
Dihydrostreptomycin sulfate

---

### **Target species:**

Cattle

---

### **Route of administration:**

Intramammary use

---

## Product details

### **Active substance and strength:**

Nafcillin sodium monohydrate  
121.95 milligram(s) / 1.00 Applicator

Benzylpenicillin procaine monohydrate  
300.00 milligram(s) / 1.00 Applicator

Dihydrostreptomycin sulfate  
134.23 milligram(s) / 1.00 Applicator

---

### **Pharmaceutical form:**

Intramammary ointment

---

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

#### **Intramammary use:**

- 

#### **Cattle**

- Milk. 5 day

bei einer Trockenstehzeit von > 35 Tagen. Bei abweichender Trockenstehzeit und für weitere Hinweise s. SPC.

- Meat and offal. 17 day

---

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

QJ51RC22

---

### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

### **Authorisation status:**

Valid

---

### **Authorised in:**

Germany

---

### **Available in:**

Germany

---

**Package description:**

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

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:**

Intervet Deutschland GmbH

---

**Marketing authorisation date:**

25/10/2004

---

**Manufacturing sites for batch release:**

INTERVET INTERNATIONAL B.V.

---

**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

---

**Authorisation number:**

6830405.00.00

---

**Date of authorisation status change:**

25/10/2004

---

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

This document does not exist in this language (English). You can find it in another language below.