

KEFAMAST DC

*Anhydrous Cefalexin
Dihydrostreptomycin*

INDICATIONS

- For dairy cow treatment at drying off
- For the treatment of subclinical mastitis infections present at drying off and to assist in preventing new infections during the dry cow period



BENEFITS

- A long acting dry cow treatment combining a cephalosporin and an aminoglycoside giving broad spectrum antimicrobial activity
- Short withdrawals milk 40 days + 60 hours, resulting in a faster return to milk tank
- Reduces high SCC and significantly aids in the prevention of new infections during the dry period
- Excellent when combined with Boviseal
- A unique antibiotic combination which provides outstanding synergistic activity against sub-clinical mastitis

PACKAGING

| LIST NO. | UNIT PACKAGE | CASE SIZE |
|----------|--------------|-----------|
| 1KEF002 | 120 SYRINGES | 1 |

Bimeda®
Broomhill Road
Tallaght
Dublin 24, Ireland
Tel: 1850-515253

See reverse side for full indications, administration and dosage.



www.bimeda.ie

KEFAMAST DRY COW

Anhydrous Cefalexin Dihydrostreptomycin

PRESENTATION

A sterile intramammary injection.
Each 9g contains:
Cefalexin (anhydrous) 500mg

Dihydrostreptomycin 500mg
(as Dihydrostreptomycin Sulphate)

USES

For the treatment of sub-clinical mastitis infection present at drying off of cows and to assist in preventing new infections occurring during the dry period.

DOSAGE AND ADMINISTRATION

The contents of one injector should be infused into the teat canal of each quarter immediately after the last milking of the lactation. Before infusion, the teat should be thoroughly cleaned and disinfected. Care should be taken to avoid contamination of the injector nozzle after the protective cap has been removed.

CONTRA-INDICATIONS, WARNINGS, ETC.

For animal treatment only.
Do not use within 40 days of the estimated calving date.
Not to be used in animals known to be hypersensitive to any of the active ingredients. Not for use in lactating cows.

OPERATOR WARNINGS

Operators should avoid contact with this preparation as occasional skin allergy may occur. Penicillins and cephalosporins may cause hypersensitivity following injection, inhalation, ingestion or skin contact. Hypersensitivity to penicillins may lead to cross

reactions to cephalosporins and vice versa.

Allergic reactions to these substances may occasionally be serious. Do not handle this product if you know that you are sensitised, or if you have been advised not to work with such preparations. Handle this product with care to avoid exposure. If you develop symptoms such as a skin rash following exposure, seek medical advice and show this warning to the doctor. Swelling of the face, lips or eyes, or difficulty with breathing are more serious symptoms and require urgent medical attention.

WITHHOLDING PERIODS

With cows milked twice daily, milk for human consumption may only be taken from 60 hours (five milkings) after calving. Foodstuffs for human consumption must not be taken during the treatment period. If calving occurs within 40 days of treatment, milk for human consumption may only be taken from 40 days + 60 hours after treatment. Animals intended for human consumption may only be slaughtered from 28 days after the last treatment.

PHARMACEUTICAL PRECAUTIONS

Do not store above 25°C. Keep out of reach and sight of children.

LEGAL CATEGORY POM

PACKAGE QUANTITIES

120 injectors

VPA 10126/32/1

