PARTICULARS TO APPEAR ON THE OUTER PACKAGE

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Multiject IMM Intramammary Suspension

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Multiject IMM is an intramammary suspension, lactating cow, containing in each 5 g syringe:

Procaine Penicillin	100 mg
Streptomycin Sulfate	100 mg
Neomycin Sulfate	100 mg
Prednisolone	10 mg

In a milk dispersible mineral oil base

3. PHARMACEUTICAL FORM

Intramammary suspension

4. PACKAGE SIZE

Packs of 24 single dose syringes

5. TARGET SPECIES

Cattle (milking cows)

6. INDICATION(S)

Multiject IMM is indicated in the treatment of acute and subacute bovine mastitis in milking cows, accompanied by pain and inflammation caused by bacterial infection sensitive to penicillin, streptomycin and neomycin therapy.

7. METHOD AND ROUTE(S) OF ADMINISTRATION

The contents of one syringe should be infused gently into each infected quarter via the teat canal immediately after milking once daily for three consecutive days. Aseptic precautions should be observed at all times.

8. WITHDRAWAL PERIOD

Milk for human consumption must not be taken from a cow during treatment. Milk for human consumption may only be taken from 108 hours from the last treatment. Where any other milking routine is followed consult your veterinary surgeon.

With other milking routines, the basis of the veterinary surgeon's advice should be that milk may be taken for human consumption only after the same period from the last treatment.

Animals must not be slaughtered for human consumption during treatment. Cattle may be slaughtered for human consumption only after 7 days from the last treatment.

During a course of treatment the situation should be reviewed frequently by close veterinary supervision

9. SPECIAL WARNING(S), IF NECESSARY

Protective gloves should be worn when infusing heifers, to avoid skin contact with the product.

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.

1. Do not handle this product if you know you are sensitised, or if you have been advised not to work with such preparations.
2. Handle this product with great care to avoid exposure, taking all recommended precautions.
3. If you develop symptoms following exposure such as skin rash, you should seek medical advice and show the doctor this warning. Swelling of the face, lips or eyes or difficulty with breathing are more serious symptoms and require urgent medical attention.

Wash hands after use.

10. EXPIRY DATE

EXP: DD/MM/YY

11. SPECIAL STORAGE CONDITIONS

Do not store above 25ºC.

The syringe must only be used once.

Part used syringes must be discarded.

12. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Dispose of any unused product and empty syringes in accordance with guidance from your local waste regulation authority.

13. THE WORDS "FOR ANIMAL TREATMENT ONLY" AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE

For animal treatment only.

POM-V

To be supplied only on veterinary prescription

14. THE WORDS "KEEP OUT OF THE SIGHT AND REACH OF CHILDREN"

KEEP OUT OF THE SIGHT AND REACH OF CHILDREN

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

MANUFACTURED BY:

Norbrook Laboratories Limited

Newry

Co. Down

Northern Ireland

DISTRIBUTED BY:

Norbrook Laboratories Limited

Carnbane Industrial Estate

Newry

Co. Down

BT35 6QQ

Northern Ireland

16. MARKETING AUTHORISATION NUMBER(S)

ManA 2000

VM: 02000/4062

17. MANUFACTURER'S BATCH NUMBER

B.N.:

UK AUTHORISED VETERINARY MEDICINAL PRODUCT

LOGO

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Multiject IMM Intramammary Suspension

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Contents:

Procaine Penicillin	100 mg
Streptomycin Sulfate	100 mg
Neomycin Sulfate	100 mg
Prednisolone	10 mg

In a milk dispersible mineral oil base

3. PHARMACEUTICAL FORM

Intramammary suspension

4. PACKAGE SIZE

5g

5. TARGET SPECIES

Cattle (milking cows)

6. INDICATION(S)

See carton text for indications

7. METHOD AND ROUTE(S) OF ADMINISTRATION

See carton text for method and route of administration

8. WITHDRAWAL PERIOD

Milk: 108 hours.

Meat: 7 days

9. SPECIAL WARNING(S), IF NECESSARY

Penicillins/cephalosporins may occasionally cause severe allergic reactions. See carton text for user warning.

10. EXPIRY DATE

EXP: DD/MM/YY

11. SPECIAL STORAGE CONDITIONS

Do not store above 25ºC.

The syringe must only be used once.

Part used syringes must be discarded.

12. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

13. THE WORDS "FOR ANIMAL TREATMENT ONLY" AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE

For animal treatment only.

POM-V

To be supplied only on veterinary prescription

14. THE WORDS "KEEP OUT OF THE SIGHT AND REACH OF CHILDREN"

KEEP OUT OF THE SIGHT AND REACH OF CHILDREN

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

MANUFACTURED BY:

Norbrook Laboratories Limited

Newry

Co. Down

Northern Ireland

DISTRIBUTED BY:

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Carnbane Industrial Estate

Newry

Co. Down

BT35 6QQ

Northern Ireland

16. MARKETING AUTHORISATION NUMBER(S)

ManA 2000

VM: 02000/4062

17. MANUFACTURER'S BATCH NUMBER

B.N.:

DOM:

UK AUTHORISED VETERINARY MEDICINAL PRODUCT

PACKAGE LEAFLET FOR MULTIJECT INTRAMAMMARY SUSPENSION

1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT

Norbrook Laboratories Limited

Newry, Co. Down

Northern Ireland

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

MULTIJECT INTRAMAMMARY SUSPENSION

3. STATEMENT OF THE ACTIVE SUBSTANCE (S) AND OTHER INGREDIENTS

Multiject IMM is an intramammary suspension, lactating cow, containing in each 5 g syringe:

Procaine Penicillin	100 mg
Streptomycin Sulfate	100 mg
Neomycin Sulfate	100 mg
Prednisolone	10 mg

In a milk dispersible mineral oil base

4. INDICATION(S)

Multiject IMM is indicated in the treatment of acute and subacute bovine mastitis in milking cows, accompanied by pain and inflammation caused by bacterial infection sensitive to penicillin, streptomycin and neomycin therapy.

5. CONTRAINDICATIONS

6. ADVERSE REACTIONS

If you notice any serious effects or other effects not mentioned in this leaflet, please inform your veterinary surgeon.

7. TARGET SPECIES

Cattle (milking cows)

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

The contents of one syringe should be infused gently into each infected quarter via the teat canal immediately after milking, once daily for three consecutive days.

9. ADVICE ON CORRECT ADMINISTRATION

Aseptic precautions should be observed at all times.

10. WITHDRAWAL PERIOD(S)

Milk for human consumption must not be taken from a cow during treatment. Milk for human consumption may only be taken from 108 hours from the last treatment. Where any other milking routine is followed consult your veterinary surgeon.

With other milking routines, the basis of the veterinary surgeon's advice should be that milk may be taken for human consumption only after the same period from the last treatment.

Animals must not be slaughtered for human consumption during treatment. Cattle may be slaughtered for human consumption only after 7 days from the last treatment.

During a course of treatment the situation should be reviewed frequently by close veterinary supervision.

11. SPECIAL STORAGE PRECAUTIONS

Do not store above 25ºC.

The syringe must only be used once.

Part used syringes must be discarded.

12. SPECIAL WARNING(S)

Protective gloves should be worn when infusing heifers, to avoid skin contact with the product.

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.

1. Do not handle this product if you know you are sensitised, or if you have been advised not to work with such preparations.
2. Handle this product with great care to avoid exposure, taking all recommended precautions.
3. If you develop symptoms following exposure such as skin rash, you should seek medical advice and show the doctor this warning. Swelling of the face, lips or eyes or difficulty with breathing are more serious symptoms and require urgent medical attention.

Wash hands after use

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY

Dispose of any unused product and empty syringes in accordance with guidance from your local waste regulation authority

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED

21st March 2012

15. OTHER INFORMATION

DISTRIBUTED BY:

Norbrook Laboratories Limited

Carnbane Industrial Estate

Newry

Co. Down

BT35 6QQ

Northern Ireland

Package Quantities

Packs of 24 single dose syringes.

BN:

D.O.M:

Exp:

ManA 2000

Vm 02000/4062

POM-V

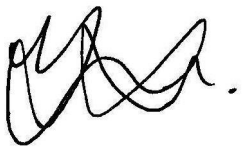
To be supplied only on veterinary prescription

KEEP OUT OF REACH AND SIGHT OF CHILDREN

FOR ANIMAL TREATMENT ONLY

LOGO

UK AUTHORISED VETERINARY MEDICINAL PRODUCT



Approved: 28 October 2022