PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE – 500ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

KILLITCH 25% w/v CUTANEOUS EMULSION

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Active ingredient: Benzyl Benzoate 25% w/v

3. PHARMACEUTICAL FORM

See under section 7. Method and Route of Administration

4. PACKAGE SIZE

500 ml

5. TARGET SPECIES

See under section 7. Method and Route of Administration

6. INDICATION(S)

• Proven to prevent sweet itch when used before symptoms appear
•Treats and soothes irritated areas
&
See under section 7. Method and Route of Administration

7. METHOD AND ROUTE(S) OF ADMINISTRATION

DIRECTIONS FOR USE
Cutaneous liquid for treatment of the condition known as sweet itch in horses and ponies. Apply to all affected areas by rubbing in with fingers or soft cloth as soon as sweet itch is diagnosed and for the remaining part of the sweet itch season (March-September). Initial application of Killitch (75ml) should be made twice daily, reducing to once daily and then 3-4 times a week as the irritation lessens. All affected areas should be kept clean by shampooing and rinsing well at least once a week. After shampooing or heavy exercise, re-apply to ensure adequate protection. With horses which are known to suffer from sweet itch, it is recommended that Killitch is applied in the early spring before symptoms become evident, so that the condition may never take a hold. Avoid contact with eyes and mucous membranes. If signs of disease persist or appear, consult a veterinary surgeon.

8. WITHDRAWAL PERIOD

Not to be used on horses intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

Keep out of reach of children. Avoid contact with eyes. Dispose of empty packaging and any remaining product in the household refuse. For animal treatment only.

9. SPECIAL WARNING(S), IF NECESSARY

SHAKE BOTTLE BEFORE USE. Protective rubber gloves should be worn when using this cutaneous solution. Using this product may cause skin irritation in sensitive individuals. Wash hands thoroughly after use. In the case of eye or skin contact, wash the affected area immediately with plenty of water. If irritation occurs, seek medical advice showing the product label. FOR EXTERNAL USE ONLY.HARMFUL IF SWALLOWED. DO NOT INDUCE VOMITING - seek urgent medical attention showing the product label to the doctor or nurse. Do not eat, drink or smoke while using this product. Store in tightly closed, original container. Do not store above 25ºC. Protect from frost. Protect from direct sunlight.

10. EXPIRY DATE

• Batch no
• Expiry date

11. SPECIAL STORAGE CONDITIONS

See text under section 9. Special Warnings

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

See text under section 8. Withdrawal Period

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

See text under section 8. Withdrawal Period for "For Animal treatment only".

AVM-GSL **UK authorised veterinary medicinal product**

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

See text "Keep out of reach of children" under section 8. Withdrawal Period for "For Animal treatment only".

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

Carr & Day & Martin Limited, Woodland Granaries, Narrow Lane, Wymeswold, Loughborough, LE12 6SD

16. MARKETING AUTHORISATION NUMBER(S)

Vm 01974/4003

17. MANUFACTURER'S BATCH NUMBER

Please see base of bottle for:
• Batch no
• Expiry date

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE – 1000 ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

KILLITCH 25% w/v CUTANEOUS EMULSION

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Active ingredient: Benzyl Benzoate 25% w/v

3. PHARMACEUTICAL FORM

See under section 7. Method and Route of Administration

4. PACKAGE SIZE

1 litre

5. TARGET SPECIES

See under section 7. Method and Route of Administration

6. INDICATION(S)

• Proven to prevent sweet itch when used before symptoms appear
•Treats and soothes irritated areas
&
See under section 7. Method and Route of Administration

7. METHOD AND ROUTE(S) OF ADMINISTRATION

DIRECTIONS FOR USE
Cutaneous liquid for treatment of the condition known as sweet itch in horses and ponies. Apply to all affected areas by rubbing in with fingers or soft cloth as soon as sweet itch is diagnosed and for the remaining part of the sweet itch season (March-September). Initial application of Killitch (75ml) should be made twice daily, reducing to once daily and then 3-4 times a week as the irritation lessens. All affected areas should be kept clean by shampooing and rinsing well at least once a week. After shampooing or heavy exercise, re-apply to ensure adequate protection. With horses which are known to suffer from sweet itch, it is recommended that Killitch is applied in the early spring before symptoms become evident, so that the condition may never take a hold. Avoid contact with eyes and mucous membranes. If signs of disease persist or appear, consult a veterinary surgeon.

8. WITHDRAWAL PERIOD

Not to be used on horses intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

Keep out of reach of children. Avoid contact with eyes. Dispose of empty packaging and any remaining product in the household refuse. For animal treatment only.

9. SPECIAL WARNING(S), IF NECESSARY

SHAKE BOTTLE BEFORE USE. Protective rubber gloves should be worn when using this cutaneous solution. Using this product may cause skin irritation in sensitive individuals. Wash hands thoroughly after use. In the case of eye or skin contact, wash the affected area immediately with plenty of water. If irritation occurs, seek medical advice showing the product label. FOR EXTERNAL USE ONLY.HARMFUL IF SWALLOWED. DO NOT INDUCE VOMITING - seek urgent medical attention showing the product label to the doctor or nurse. Do not eat, drink or smoke while using this product. Store in tightly closed, original container. Do not store above 25ºC. Protect from frost. Protect from direct sunlight.

10. EXPIRY DATE

- Batch no
- Expiry date

11. SPECIAL STORAGE CONDITIONS

See text under section 9. Special Warnings

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

See text under section 8. Withdrawal Period

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

See text under section 8. Withdrawal Period for "For Animal treatment only".

AVM-GSL **UK authorised veterinary medicinal product**

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

See text "Keep out of reach of children" under section 8. Withdrawal Period for "For Animal treatment only".

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

Carr & Day & Martin Limited, Woodland Granaries, Narrow Lane, Wymeswold, Loughborough, LE12 6SD

16. MARKETING AUTHORISATION NUMBER(S)

Vm 01974/4003

17. MANUFACTURER'S BATCH NUMBER

- Batch no
- Expiry date

Approved 09 February 2022