PARTICULARS TO APPEAR ON <THE OUTER PACKAGE> <AND> <THE IMMEDIATE PACKAGE> {NATURE/TYPE}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Large Animal Diprevon Solution for Injection

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Active Ingredient:	mg/ml
Diprenorphine	3.0
(as Diprenorphine hydrochloride	3.26)
Preservative:	
Chlorocresol	1.0
Colouring Agent:	
Methylthioninium chloride	0.1

3. PHARMACEUTICAL FORM

Solution for injection.

4. PACKAGE SIZE

10.5 ml

5. TARGET SPECIES

Horses and Deer

6. INDICATION(S)

7. METHOD AND ROUTE(S) OF ADMINISTRATION

Horse: Intravenous injection

Deer: Intramuscular injection

8. WITHDRAWAL PERIOD

Must not be administered to animals intended for human or animal consumption.
Not to be used in horses 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.

9. SPECIAL WARNING(S), IF NECESSARY

<User Warnings>

10. EXPIRY DATE

11. SPECIAL STORAGE CONDITIONS

Keep the container in the outer carton.

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

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

13. THE WORDS "FOR ANIMAL TREATMENT ONLY" AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE *[Distribution category]*

POM-V

For animal treatment only

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

Abbeyvet Export LLP
Sherburn Enterprise Park
Aviation Way
Sherburn-in-Elmet
Leeds
LS25 6NB
United Kingdom

16. MARKETING AUTHORISATION NUMBER(S)

Vm 21757/4001

17. MANUFACTURER'S BATCH NUMBER

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS {NATURE/TYPE}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Large Animal Diprevon Solution for Injection

2. QUANTITY OF THE ACTIVE SUBSTANCE(S)

3. CONTENTS BY WEIGHT, BY VOLUME OR BY NUMBER OF DOSES

10.5 ml

4. ROUTE(S) OF ADMINISTRATION

Horse: Intravenous injection

Deer: Intramuscular injection

5. WITHDRAWAL PERIOD

Must not be administered to animals intended for human or animal consumption.
Not to be used in horses 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.

6. BATCH NUMBER

7. EXPIRY DATE

8. THE WORDS “FOR ANIMAL TREATMENT ONLY”

PACKAGE LEAFLET FOR:

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

Abbeyvet Export LLP
Sherburn Enterprise Park
Aviation Way
Sherburn-in-Elmet
Leeds
LS25 6NB
United Kingdom

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

Large Animal Etorphilon Solution for Injection

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

Active Ingredient:	mg/ml
Diprenorphine	3.0
(as Diprenorphine hydrochloride	3.26)
Preservative:	
Chlorocresol	1.0
Colouring Agent:	
Methylthioninium chloride	0.1

4. INDICATION(S)

To reverse neuroleptanalgesia induced by etorphine hydrochloride.

5. CONTRAINDICATIONS

6. ADVERSE REACTIONS

7. TARGET SPECIES

Horses and Deer

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

9. ADVICE ON CORRECT ADMINISTRATION

10. WITHDRAWAL PERIOD(S)

Must not be administered to animals intended for human or animal consumption.
Not to be used in horses 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.

11. SPECIAL STORAGE PRECAUTIONS

12. SPECIAL WARNING(S)

<User Warnings>

For Animal Treatment Only

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

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED

15. OTHER INFORMATION

POM - V

Vm 21757/4001

Approved: 03/01/2018