

ANNEX I

SUMMARY OF PRODUCT CHARACTERISTICS

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

CORTEXONAVET 2 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each ml contains:

Active substance:

Dexamethasone 2.0 mg
(as dexamethasone sodium phosphate)

Excipient:

Qualitative composition of excipients and other constituents	Quantitative composition if that information is essential for proper administration of the veterinary medicinal product
Benzyl alcohol (E1519)	15.6 mg
Water for injections	

Clear, colourless solution, free from visible particles.

3. CLINICAL INFORMATION

3.1 Target species

Cattle, horses, pigs, dogs and cats.

3.2 Indications for use for each target species

Horses, cattle, pigs, dogs and cats:

Treatment of inflammatory or allergic conditions.

Cattle:

Treatment of primary ketosis (acetonaemia). Induction of parturition.

Horses:

Treatment of arthritis, bursitis or tenosynovitis.

3.3 Contraindications

Except in emergency situations, do not use in animals suffering from diabetes mellitus, renal insufficiency, cardiac insufficiency hyperadrenocorticism, or osteoporosis.

Do not use in viral infections during the viraemic stage or in cases of systemic mycotic infections.

Do not use in animals suffering from gastrointestinal or corneal ulcers, or demodicosis.

Do not administer by the intra-articular route where there is evidence of fractures, bacterial joint infections and aseptic bone necrosis.

Do not use in cases of hypersensitivity to the active substance, to corticosteroids or to any of the excipients.

Refer to section 3.5 *Special precautions for use*.

3.4 Special warnings

None.

3.5 Special precautions for use

Special precautions for safe use in the target species:

Anti-inflammatory corticosteroids, such as dexamethasone, are known to exert a wide range of side effects. Whilst single high doses are generally well tolerated, they may induce severe side-effects in long term use and when esters possessing a long duration of action are administered. Dosage in medium to long term use should therefore generally be kept to the minimum necessary to control symptoms. Response to long-term therapy should be monitored at regular intervals by a veterinary surgeon.

Use of corticosteroids in horses has been reported to induce laminitis. Therefore horses treated with such preparations should be monitored frequently during the treatment period.

During therapy effective doses suppress the hypothalamo-pituitary-adrenal axis. Following cessation of treatment, symptoms of adrenal insufficiency extending to adrenocortical atrophy can arise and this may render the animal unable to deal adequately with stressful situations. Consideration should therefore be given to means of minimising problems of adrenal insufficiency following the withdrawal of treatment (for further discussion see standard texts).

Because of the pharmacological properties of the active ingredient, special care should be taken when the veterinary medicinal product is used in animals with a weakened immune system.

Corticosteroids may delay wound healing and the immunosuppressant actions may weaken resistance to or exacerbate existing infections. In the presence of bacterial infection, antibacterial drug cover is usually required when steroids are used. In the presence of viral infections, steroids may worsen or hasten the progress of the disease.

Except in cases of acetonæmia and induction of parturition, corticosteroid administration is to induce an improvement in clinical signs rather than a cure.

The underlying disease should be further investigated. When treating groups of animals, use a draw-off needle to avoid excessive broaching of the stopper.

Following intra-articular administration, use of the joint should be minimised for one month and surgery on the joint should not be performed within eight weeks of use of this route of administration.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

Care should be taken to avoid accidental self-injection.

In case of accidental self-injection, seek medical advice immediately and show the package leaflet or the label to the physician.

The veterinary medicinal product should not be administered by pregnant women.

People with known hypersensitivity to dexamethasone or to any of the excipients should avoid contact with the veterinary medicinal product.

Avoid contact with skin and eyes. In the event of accidental eye or skin contact, wash the area thoroughly with clean running water.

Wash hands after use.

Special precautions for the protection of the environment:

Not applicable.

3.6 Adverse events

Cattle, horses, pigs, dogs and cats:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):	Hypersensitivity reaction; Cushingoid symptoms ¹ ; Polyuria ² ; Polydipsia ² , Polyphagia ² ; Sodium retention ³ , Water retention ³ , Hypokalaemia ³ , Hyperglycaemia ⁴ , Changes in blood biochemical and haematological parameters, Elevated liver enzymes; Cutaneous calcinosis; Gastro-intestinal ulceration ⁵ , Pancreatitis ⁶ ; Hepatomegaly ⁷ ; Laminitis ⁸ ; Milk production decrease ⁹ .
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¹ Cushingoid symptoms involving significant alteration of fat, carbohydrate, protein and mineral metabolism, e.g. redistribution of body fat, muscle weakness and wastage and osteoporosis may result.

² After systemic administration and particularly during the early stages of therapy.

³ In long term use.

⁴ Transient.

⁵ Reported in animals treated with corticosteroids and in animals with spinal cord trauma.

⁶ Increased risk of acute pancreatitis.

⁷ With increased serum hepatic enzymes.

⁸ In horses.

⁹ In cattle.

Reporting adverse events is important. It allows continuous safety monitoring of a veterinary medicinal product. Reports should be sent, preferably via a veterinarian, to either the marketing authorisation holder <or its local representative> or the national competent authority via the national reporting system. See the last section of the package leaflet for respective contact details.

3.7 Use during pregnancy, lactation or lay

Pregnancy:

Apart from the use of the veterinary medicinal product to induce parturition in cattle, corticosteroids are not recommended for use in pregnant animals. Administration in early pregnancy is known to have caused foetal abnormalities in laboratory animals. Administration in late pregnancy may cause early parturition or abortion.

If the veterinary medicinal product is used for induction of parturition in cattle, then a high incidence of retained placentae may be experienced and possible subsequent metritis and/or subfertility. Such use of dexamethasone, particularly at early time points, may be associated with reduced viability of the calf.

Lactation:

Use of the veterinary medicinal product in lactating cows may cause a reduction in milk yield.

3.8 Interaction with other medicinal products and other forms of interaction

Concurrent use with non-steroidal anti-inflammatory drugs may exacerbate gastrointestinal tract ulceration.

Because corticosteroids can reduce the immune response to vaccination, dexamethasone should not be used in combination with vaccines or within two weeks after vaccination.

Administration of dexamethasone may induce hypokalaemia and hence increase the risk of toxicity from cardiac glycosides. The risk of hypokalaemia may be increased if dexamethasone is administered together with potassium depleting diuretics.

Concurrent use with anticholinesterase may lead to increased muscle weakness in patients with myasthenia gravis.

Glucocorticoids antagonise the effects of insulin.

Concurrent use with phenobarbital, phenytoin and rifampicin can reduce the effects of dexamethasone.

3.9 Administration routes and dosage

Routes of administration:

Horses: intravenous, intramuscular or intra-articular use.

Cattle, pigs, dogs and cats: intramuscular use.

Normal aseptic technique should be observed.

To measure small volumes of less than 1 ml, a suitably graduated syringe should be used to ensure accurate administration of the correct dose.

For the treatment of inflammatory or allergic conditions: The following average doses are advised. However the advised dose used should be determined by the severity of the signs and the length of time for which they have been present.

<i>Species</i>	<i>Dosage</i>
Horses, cattle and pigs:	0.06 mg/kg body weight corresponding to 1.5 ml/50 kg
Cattle, pigs, dogs and cats:	0.1 mg/kg body weight corresponding to 0.5 ml/10 kg

For the treatment of primary ketosis in cattle (acetoanaemia): 0.02 to 0.04 mg/kg body weight corresponding to a dose of 5-10 ml per cow given by intramuscular injection is advocated dependent on the size of the cow and the duration of the signs. Care should be taken not to overdose Channel Island breeds. Larger doses will be required if the signs have been present for some time or if relapsed animals are being treated.

For the induction of parturition – to avoid foetal oversize and mammary oedema in cattle.

A single intramuscular injection of 0.04 mg/kg body weight corresponding to 10 ml per cow after day 260 of pregnancy.

Parturition will normally occur within 48-72 hours.

For the treatment of arthritis, bursitis or tenosynovitis: by intra-articular injection in the horse. Dose 1-5 ml.

These quantities are not specific and are quoted purely as a guide. Injections into joint spaces or bursae should be preceded by the removal of an equivalent volume of synovial fluid. Strict asepsis is essential.

3.10 Symptoms of overdose (and where applicable, emergency procedures and antidotes)

An overdose can induce drowsiness and lethargy in horses.

3.11 Special restrictions for use and special conditions for use, including restrictions on the use of antimicrobial and antiparasitic veterinary medicinal products in order to limit the risk of development of resistance

Not applicable.

3.12 Withdrawal periods

Cattle:

Meat and offal: 8 days.

Milk: 72 hours.

Pigs:

Meat and offal: 2 days.

Horses:

Meat and offal: 8 days.

Milk: Not authorised for use in horses producing milk for human consumption.

4. PHARMACOLOGICAL INFORMATION

4.1 ATCvet code: QH02AB02

4.2 Pharmacodynamics

This preparation contains the sodium phosphate ester of dexamethasone, a fluoro-methyl derivative of prednisolone, which is a potent glucocorticoid with minimal mineralocorticoid activity. Dexamethasone has ten to twenty times the anti-inflammatory activity of prednisolone.

Corticosteroids suppress the immunologic response by inhibition of dilatation of capillaries, migration and function of leucocytes and phagocytosis. Glucocorticoids have an effect on metabolism by increasing gluconeogenesis

4.3 Pharmacokinetics

After extravascular (intramuscular, subcutaneous, intra-articular) administration, this soluble ester of dexamethasone is rapidly resorbed from the injection site followed by immediate hydrolysis into the parent compound, dexamethasone. Absorption of dexamethasone is rapid. The time to reach maximum plasma concentrations (C_{max}) of dexamethasone in cattle, horses, pigs and dogs is within 20 min after intramuscular administration. Bioavailability following i.m. administration (compared to i.v. administration) is high in all species. Elimination half-life after intravenous administration in horses is 3.5 h. After intramuscular administration, apparent elimination half-life has been shown to range between 1 and 20 hours according to the species.

5. PHARMACEUTICAL PARTICULARS

5.1 Major incompatibilities

In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

5.2 Shelf life

Shelf life of the veterinary medicinal product as packaged for sale: 2 years.

Shelf life after first opening the immediate packaging: 28 days.

5.3 Special precautions for storage

Keep the vial in the outer carton in order to protect from light.

5.4 Nature and composition of immediate packaging

Cardboard box with 1 colourless, type I glass vial of 50 or 100 ml, which is closed with a bromobutyl type I rubber stopper and sealed with an aluminium cap.

Not all pack sizes may be marketed.

5.5 Special precautions for the disposal of unused veterinary medicinal products or waste materials derived from the use of such products

Medicines should not be disposed of via wastewater or household waste.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any national collection systems applicable to the veterinary medicinal product concerned.

6. NAME OF THE MARKETING AUTHORISATION HOLDER

Laboratorios SYVA S.A.

7. MARKETING AUTHORISATION NUMBER(S)

8. DATE OF FIRST AUTHORISATION

Date of first authorisation: DD/MM/YYYY

9. DATE OF THE LAST REVISION OF THE SUMMARY OF THE PRODUCT CHARACTERISTICS

DD/MM/YYYY

10. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS

Veterinary medicinal product subject to prescription.

Detailed information on this veterinary medicinal product is available in the [Union Product Database \(https://medicines.health.europa.eu/veterinary\)](https://medicines.health.europa.eu/veterinary).

ANNEX III
LABELLING AND PACKAGE LEAFLET

A. LABELLING

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

CARTON BOX / 50 ml vial
CARTON BOX / 100 ml vial

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

CORTEXONAVET 2 mg/ml solution for injection

2. STATEMENT OF ACTIVE SUBSTANCES

Dexamethasone (as dexamethasone sodium phosphate) 2.0 mg/ml

3. PACKAGE SIZE

50 ml
100 ml

4. TARGET SPECIES

Cattle, horses, pigs, dogs and cats.

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Horses: intravenous, intramuscular or intra-articular use.
Cattle, pigs, dogs and cats: intramuscular use.

7. WITHDRAWAL PERIODS

Withdrawal periods:

Cattle: Meat and offal: 8 days.
Milk: 72 hours.

Pigs: Meat and offal: 2 days.

Horses: Meat and offal: 8 days.

Milk: Not authorised for use in horses producing milk for human consumption.

8. EXPIRY DATE

Exp. {mm/yyyy}

Once opened, use by:

Shelf life after first opening the immediate packaging: 28 days.

9. SPECIAL STORAGE PRECAUTIONS

Keep the vial in the outer carton in order to protect from light.

10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”

Read the package leaflet before use.

11. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

Laboratorios SYVA S.A.

14. MARKETING AUTHORISATION NUMBERS

15. BATCH NUMBER

Lot {number}

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE

{Label/100 ml vial}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

CORTEXONAVET 2 mg/ml solution for injection.

2. STATEMENT OF ACTIVE SUBSTANCES

Dexamethasone (as dexamethasone sodium phosphate) 2.0 mg/ml

3. TARGET SPECIES

Cattle, horses, pigs, dogs and cats

4. ROUTES OF ADMINISTRATION

-

Horses: **i.v.**, **i.m.** or intra-articular use

Cattle, pigs, dogs and cats: **i.m.** use

Read the package leaflet before use.

5. WITHDRAWAL PERIODS

Withdrawal period:

Cattle:

Meat and offal: 8 days.

Milk: 72 hours.

Pigs:

Meat and offal: 2 days.

Horses:

Meat and offal: 8 days.

Milk: Not authorised for use in horses producing milk for human consumption.

6. EXPIRY DATE

Exp. {mm/yyyy}

Once opened use by...

Shelf life after first opening the immediate packaging: 28 days.

7. SPECIAL STORAGE PRECAUTIONS

Keep the vial in the outer carton in order to protect from light.

8. NAME OF THE MARKETING AUTHORISATION HOLDER

Laboratorios SYVA S.A.

9. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS

{Label/50 ml vial}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

CORTEXONAVET

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Dexamethasone (as dexamethasone sodium phosphate) 2.0 mg/ml

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

Once opened use by...

Shelf life after first opening the immediate packaging: 28 days.

B. PACKAGE LEAFLET

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

CORTEXONAVET 2 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

2. Composition

Each ml contains:

Active substance:

Dexamethasone 2.0 mg
(as dexamethasone sodium phosphate)

Excipients:

Benzyl alcohol (E1519) 15.6 mg

Clear, colourless solution, free from visible particles

3. Target species

Cattle, horses, pigs, dogs and cats.

4. Indications for use

Horses, cattle, pigs, dogs and cats:

Treatment of inflammatory or allergic conditions.

Cattle:

Treatment of primary ketosis (acetonaemia). Induction of parturition.

Horses:

Treatment of arthritis, bursitis or tenosynovitis.

5. Contraindications

Except in emergency situations, do not use in animals suffering from diabetes mellitus, renal insufficiency, cardiac insufficiency hyperadrenocorticism, or osteoporosis.

Do not use in viral infections during the viraemic stage or in cases of systemic mycotic infections.

Do not use in animals suffering from gastrointestinal or corneal ulcers, or demodicosis.

Do not administer by the intra-articular route where there is evidence of fractures, bacterial joint infections and aseptic bone necrosis.

Do not use in cases of hypersensitivity to the active substance, to corticosteroids or to any of the excipients.

Refer to section *Special warnings*.

6. Special warnings

Special warnings:

None.

Special precautions for safe use in the target species:

Anti-inflammatory corticosteroids, such as dexamethasone, are known to exert a wide range of side effects. Whilst single high doses are generally well tolerated, they may induce severe side-effects in long term use and when esters possessing a long duration of action are administered. Dosage in medium to long term use should therefore generally be kept to the minimum necessary to control symptoms. Response to long-term therapy should be monitored at regular intervals by a veterinary surgeon.

Use of corticosteroids in horses has been reported to induce laminitis. Therefore horses treated with such preparations should be monitored frequently during the treatment period. During therapy effective doses suppress the hypothalamo-pituitary-adrenal axis. Following cessation of treatment, symptoms of adrenal insufficiency extending to adrenocortical atrophy can arise and this may render the animal unable to deal adequately with stressful situations. Consideration should therefore be given to means of minimising problems of adrenal insufficiency following the withdrawal of treatment (for further discussion see standard texts).

Because of the pharmacological properties of the active ingredient, special care should be taken when the veterinary medicinal product is used in animals with a weakened immune system. Corticosteroids may delay wound healing and the immunosuppressant actions may weaken resistance to or exacerbate existing infections. In the presence of bacterial infection, antibacterial drug cover is usually required when steroids are used. In the presence of viral infections, steroids may worsen or hasten the progress of the disease.

Except in cases of acetoanaemia and induction of parturition, corticosteroid administration is to induce an improvement in clinical signs rather than a cure.

The underlying disease should be further investigated. When treating groups of animals, use a draw-off needle to avoid excessive broaching of the stopper.

Following intra-articular administration, use of the joint should be minimised for one month and surgery on the joint should not be performed within eight weeks of use of this route of administration.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

Care should be taken to avoid accidental self-injection.

In case of accidental self-injection, seek medical advice immediately and show the package leaflet or the label to the doctor.

The veterinary medicinal product should not be administered by pregnant women.

People with known hypersensitivity to dexamethasone or to any of the excipients should avoid contact with the veterinary medicinal product.

Avoid contact with skin and eyes. In the event of accidental eye or skin contact, wash the area thoroughly with clean running water.

Wash hands after use.

Pregnancy:

Apart from the use of the veterinary medicinal product to induce parturition in cattle, corticosteroids are not recommended for use in pregnant animals. Administration in early pregnancy is known to have caused foetal abnormalities in laboratory animals. Administration in late pregnancy may cause early parturition or abortion.

If the veterinary medicinal product is used for induction of parturition in cattle, then a high incidence of retained placentae may be experienced and possible subsequent metritis and/or subfertility. Such use of dexamethasone, particularly at early time points, may be associated with reduced viability of the calf.

Lactation:

Use of the veterinary medicinal product in lactating cows may cause a reduction in milk yield.

Interaction with other medicinal products and other forms of interaction:

Concurrent use with non-steroidal anti-inflammatory drugs may exacerbate gastrointestinal tract ulceration.

Because corticosteroids can reduce the immune response to vaccination, dexamethasone should not be used in combination with vaccines or within two weeks after vaccination.

Administration of dexamethasone may induce hypokalaemia and hence increase the risk of toxicity from cardiac glycosides. The risk of hypokalaemia may be increased if dexamethasone is administered together with potassium depleting diuretics.

Concurrent use with anticholinesterase may lead to increased muscle weakness in patients with myasthenia gravis.

Glucocorticoids antagonise the effects of insulin.

Concurrent use with phenobarbital, phenytoin and rifampicin can reduce the effects of dexamethasone.

Overdose:

An overdose can induce drowsiness and lethargy in horses.

Major incompatibilities:

In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

7. Adverse events

Cattle, horses, pigs, dogs and cats:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):
Hypersensitivity reaction; Cushingoid symptoms ¹ ; Polyuria (increased urination) ² ; Polydipsia (increased thirst) ² , Polyphagia (increased appetite) ² ; Sodium retention ³ , Water retention ³ , Hypokalaemia (low blood potassium) ³ , Hyperglycaemia (high blood sugar) ⁴ , Changes in blood biochemical and haematological parameters, Elevated liver enzymes; Cutaneous calcinosis (calcium deposit in the skin); Gastro-intestinal ulceration ⁵ , Pancreatitis (inflammation of the pancreas) ⁶ ; Hepatomegaly (enlarged liver) ⁷ ; Laminitis ⁸ ; Milk production decrease ⁹ .

¹ Cushingoid symptoms involving significant alteration of fat, carbohydrate, protein and mineral metabolism, e.g. redistribution of body fat, muscle weakness and wastage and osteoporosis may result.

² After systemic administration and particularly during the early stages of therapy.

³ In long term use.

⁴ Transient.

⁵ Reported in animals treated with corticosteroids and in animals with spinal cord trauma.

⁶ Increased risk of acute pancreatitis.

⁷ With increased serum hepatic enzymes.

⁸ In horses.

⁹ In cattle.

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed in this package leaflet, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder <or the local representative of the marketing authorisation holder> using the contact details at the end of this leaflet, or via your national reporting system: {national system details}.

8. Dosage for each species, routes and method of administration

Routes of administration:

Horses: intravenous (**i.v.**), intramuscular (**i.m.**) or intra-articular use.

Cattle, pigs, dogs and cats: intramuscular use (**i.m.**).

For the treatment of inflammatory or allergic conditions: The following average doses are advised. However the advised dose used should be determined by the severity of the signs and the length of time for which they have been present.

<i>Species</i>	<i>Dosage</i>
Horses, cattle, pigs	0.06 mg/kg body weight corresponding to 1.5 ml/50 kg
Dogs, cats	0.1 mg/kg body weight corresponding to 0.5 ml/10 kg

For the treatment of primary ketosis in cattle (acetoaemia) 0.02 to 0.04 mg/kg body weight corresponding to a dose of 5-10 ml per cow given by intramuscular injection is advocated dependent on the size of the cow and the duration of the signs. Care should be taken not to overdose Channel Island breeds. Larger doses will be required if the signs have been present for some time or if relapsed animals are being treated.

For the induction of parturition - to avoid foetal oversize and mammary oedema in cattle.

A single intramuscular injection of 0.04 mg/kg body weight corresponding to 10 ml per cow after day 260 of pregnancy.

Parturition will normally occur within 48-72 hours.

For the treatment of arthritis, bursitis or tenosynovitis by intra-articular injection in the horse.

Dose 1- 5 ml.

These quantities are not specific and are quoted purely as a guide. Injections into joint spaces or bursae should be preceded by the removal of an equivalent volume of synovial fluid. Strict asepsis is essential.

9. Advice on correct administration

Normal aseptic technique should be observed. To measure small volumes of less than 1 ml, a suitably graduated syringe should be used to ensure accurate administration of the correct dose.

10. Withdrawal periods

Cattle:

Meat and offal: 8 days.

Milk: 72 hours.

Pigs:

Meat and offal: 2 days.

Horses:

Meat and offal: 8 days.

Milk: Not authorised for use in horses producing milk for human consumption.

11. Special storage precautions

Keep out of the sight and reach of children.

Keep the vial in the outer carton in order to protect from light.

Do not use this veterinary medicinal product after the expiry date which is stated on the carton and vial after Exp . The expiry date refers to the last day of that month.

Shelf life after first opening the immediate packaging: 28 days.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater or household waste.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required.

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. Marketing authorisation numbers and pack sizes

MA number:

Pack sizes:

Box with one 50 ml vial.

Box with one 100 ml vial.

Not all pack sizes may be marketed.

15. Date on which the package leaflet was last revised

<{MM/YYYY}>

<{DD/MM/YYYY}>

<{DD month YYYY}>

Detailed information on this veterinary medicinal product is available in the [Union Product Database \(https://medicines.health.europa.eu/veterinary\)](https://medicines.health.europa.eu/veterinary).

16. Contact details

Marketing authorisation holder :

Laboratorios Syva, S.A.
Calle Marqués de la Ensenada, 16
28004 MADRID
ESPAÑA

Manufacturer responsible for batch release:

Laboratorios Syva, S.A.
Avenida del Párroco Pablo Díez, 49-57
San Andrés del Rabanedo
24010 LEÓN
ESPAÑA

Local representatives and contact details to report suspected adverse reactions:

For any information about this veterinary medicinal product, please contact the local representative of the marketing authorisation holder.

17. Other information