



FOLI-REC[®] LÍQUIDO



Product information

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These pages are general information pages. No guarantee is given as to the completeness of the information contained or its compliance with national regulatory requirements. Users should consult the local site of their countries to obtain information that complies with applicable national regulations.

1. NAME OF VETERINARY MEDICINAL PRODUCT

FOLI-REC® Líquido

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each 100 ml contains:

Recombinant equine chorionic gonadotropin (reCG) 7,000 I.U.
Excipients q.s.p. 100,00 mL

3. PHARMACEUTICAL FORM

Ready-to-use injectable solution.

4. INDICATIONS

FOLI-REC® is indicated to promote a bigger ovulatory capacity, it can be used in superovulation protocols and as complement in heat induction and ovulation synchronization protocols, when used along with other reproductive hormones such as: progestogens, oestradiol benzoate, prostaglandins F2 α or its synthetic analogues and gonadotropin releasing hormones (GnRH).

5. IMMUNOLOGICAL PROPERTIES

Ready-to-use product.

FOLI-REC® should be administered intramuscularly in a single dose, with the following dosages:

- Cows Oestrus Synchronization – 49 to 140 I.U. (0.7 to 2.0 mL);
- Superovulation – 2,000 I.U. (28.5 mL).

6. CONTRAINDICATIONS AND LIMITATIONS OF USE

The use of FOLI-REC® is contraindicated in animals with hypersensitivity to any of the formulation components. The product has not been tested in pregnant cows. Do not use the product in young, sexually immature females. High doses produce superovulation and are not advisable for animals to be inseminated or mated (natural breeding) due to the possibility of multiple pregnancies, except for embryo transfer purposes.

The success of FOLI-REC® use is dependent on adequate hormonal reproductive protocols.

7. PRECAUTIONS

In animals:

Exclusive use in female cattle.

Do not use the product after its expiration date and pay attention to product's expiration date after opening the packaging. Keep the product in its original packaging and use sterile syringes and needles, observing good aseptic practices. Follow the recommended dosages when using the product. Only the veterinarian can make changes to the product dosages.

In humans:

When using the product, protect yourself with rubber gloves (nitrile gloves). Do not handle the product with bare hands. Do not eat, drink, or smoke during application. Do not mix with other products. Do not reuse packaging. Remains of products and packaging must be discarded as recommended by current legislation, avoiding contamination of the environment. It is recommended that the product not be administered by pregnant women, due to the risk of accidental injection. In this case, consult a doctor immediately and show the product leaflet. There is no antidote. Read the leaflet carefully before administering the product.

8. ADVERSE REACTIONS

When used according to the labeled dose, no adverse events are expected. However, individual hypersensitivity reactions may occur. In those cases, it is recommended to stop the treatment immediately and establish symptomatic therapy based on epinephrine, anti-histaminic and glucocorticoids.

9. WITHDRAWAL PERIODS

Cattle:

Meat and offal: 0 days

Milk: 0 days

10. STORAGE CONDITIONS

Store in refrigerated conditions between 2° - 8°C, away from sunlight and out of reach of children and pets.

11. PRESENTATIONS

Glass vials containing 30 mL or 100 mL of product. Not all pack sizes may be marketed.

12. SHELF LIFE

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

Shelf-life after first opening the immediate packaging: 30 days.

Sale under prescription and administration under the guidance of the veterinarian.