PARTICULARS TO APPEAR ON THE OUTER PACKAGE

Cardboard box containing 20 bottles of 100 ml,
Cardboard box containing 20 bottles of 250 ml,
Cardboard box containing 10 bottles of 500 ml,
Cardboard box containing 10 bottles of 1000 ml
Cardboard box containing 1 bottle of 100 ml
Cardboard box containing 1 bottle of 250 ml
Cardboard box containing 1 bottle of 500 ml
Cardboard box containing 1 bottle of 1000 ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Glucose B. Braun Vet Care 50 mg/ml solution for infusion

2. STATEMENT OF ACTIVE SUBSTANCES

Each ml contains:

Glucose 50.0 mg
(equivalent to glucose monohydrate 55.0 mg)

3. PACKAGE SIZE

20x100 ml
20x250 ml
10x500 ml
10x1000 ml
1x100 ml
1x250 ml
1x500 ml
1x1000 ml

4. TARGET SPECIES

Cattle, horses, sheep, goats, pigs, dogs and cats

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Intravenous use.

7. WITHDRAWAL PERIODS

Withdrawal periods:
Cattle, horses, sheep, goats, pigs:
Meat and offal: Zero days

Cattle, horses, sheep, goats:
Milk: Zero hours

8. EXPIRY DATE

EXP {mm/yyyy}
Once opened use immediately.

9. SPECIAL STORAGE CONDITIONS

Keep the bottle 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 REACHOF CHILDREN"

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

B. Braun Melsungen AG

14. MARKETING AUTHORISATION NUMBERS

Vm 03551/4002

15. BATCH NUMBER

Lot number:

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE

Bottles of 100, 250, 500 and 1000 ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Glucose B. Braun Vet Care 50 mg/ml solution for infusion

2. STATEMENT OF ACTIVE SUBSTANCES

Each ml contains:

Glucose 50.0 mg
(equivalent to glucose monohydrate 55.0 mg)

3. TARGET SPECIES

Cattle, horses, sheep, goats, pigs, dogs and cats

4. ROUTES OF ADMINISTRATION

Intravenous use.
Read the package leaflet before use.

5. WITHDRAWAL PERIODS

Withdrawal periods:
Cattle, horses, sheep, goats, pigs:
Meat and offal: Zero days

Cattle, horses, sheep, goats:
Milk: Zero hours

6. EXPIRY DATE

EXP {mm/yyyy}
Once opened use immediately.

7. SPECIAL STORAGE CONDITIONS

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

8. NAME OF THE MARKETING AUTHORISATION HOLDER

B. Braun Melsungen AG

9. BATCH NUMBER

Lot number:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Glucose B. Braun Vet Care 50 mg/ml solution for infusion for cattle, horses, sheep, goats, pigs, dogs and cats

2. Composition

Each ml contains:

Active substances:

Glucose 50.0 mg
(equivalent to glucose monohydrate 55.0 mg)

Caloric value	837 kJ/l = 200 kcal/l
Theoretical osmolarity	278 mOsm/l
pH value	3.5 - 5.5

Clear, colourless or almost colourless aqueous solution, free from visible particles.

3. Target species

Cattle, horses, sheep, goats, pigs, dogs and cats.

4. Indications for use

- Treatment of dehydration (in the absence of shock)
- Parenteral rehydration
- Correction of hypernatremia
- Correction of hyperkalaemia
- Transient supportive treatment of hypoglycaemia

5. Contraindications

Do not administer to hyperglycaemic animals.
Do not use for the correction of hypotonic dehydration. Do not use in animals with pre-existing peripheral oedema caused by a reduction in intravascular oncotic pressure.
This veterinary medicinal product is not suitable as a sole source of calorie requirements or as a substitute for oral or parenteral nutrition.

6. Special warnings

Special precautions for safe use in the target species:
This veterinary medicinal product does not contain electrolytes. Care should be taken to closely monitor electrolyte and phosphate balance in patients undergoing infusion of this product, and to adjust treatment accordingly.

This veterinary medicinal product should be used with particular caution in animals with the following conditions:

- Diabetes mellitus
- Intracranial or intraspinal bleeding
- Anuria
- Addisons disease

Severe or long standing hypernatraemia should be corrected gradually.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
None

Pregnancy and lactation:
The safety of the veterinary medicinal product has not been established during pregnancy and lactation. Use only according to the benefit-risk assessment by the responsible veterinarian.

Overdose:
Overperfusion can lead to overhydration, hypertension and extravascular fluid accumulation. Symptoms may include respiratory distress. In the case of overperfusion, reduce or cease fluid infusion and administer oxygen, diuretics and adjunctive treatment as necessary. Monitor respiration and heart rate, fluid output, electrolyte balance and blood glucose during administration.
The administration of excess glucose can lead to hyperglycaemia, glycosuria and polyuria.

Special restrictions for use and special conditions for use:
For administration only by a veterinarian.

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

The medicinal product is incompatible with calcium-disodium edetate, histamine diphosphate, sodium warfarin and sodium thiopental.

Glucose solutions should not be administered through the same infusion equipment simultaneously with, before, or after administration of blood, because of the possibility of pseudo-agglutination.

7. Adverse events

Target species: Cattle, horses, sheep, goats, pigs, dogs and cats

Very rare (<1 animal / 10,000 animals treated, including isolated reports):
Thrombosis
Undetermined frequency (cannot be estimated from the available data)
Glycosuria[1], osmotic diuresis[1]

[1] If the product is administered at a rate in excess of 10 ml/kg bodyweight/hour. Please refer to section 8 “Dosage for each species, routes and method of administration”.

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 its local representative using the contact details at the end of this leaflet, or via your national reporting system at:
Website: https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine
e-mail: adverse.events@vmd.gov.uk

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

Intravenous use. Administer slowly via intravenous infusion.
For single use only.
This product should not be administered at a rate in excess of 10 ml/kg bodyweight/hour, otherwise glycosuria and osmotic diuresis may result.
Infusion rates should be calculated according to the presenting condition, bodyweight and degree of dehydration of the animal being treated. The total fluid volume to be administered should consider existing deficits, maintenance requirements and ongoing losses.

9. Advice on correct administration

Do not use unless the solution is clear, free from visible particles, and if you notice signs of deterioration.
IV fluids should be warmed up to body temperature prior to administration.
Maintain aseptic precautions throughout administration.

10. Withdrawal periods

Cattle, horses, sheep, goats, pigs:
Meat and offal: Zero days

Cattle, horses, sheep, goats:
Milk: Zero hours

11. Special storage precautions

Keep out of the sight and reach of children.
Keep the bottle in the outer carton in order to protect from light.
Do not use this veterinary medicinal product after the expiry date stated on the label after EXP. The expiry date refers to the last day of that month.
Shelf-life after first opening the container: use immediately.

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

Vm 03551/4002

Pack sizes:

Cardboard box containing one bottle of 100 ml
Cardboard box containing one bottle of 250 ml
Cardboard box containing one bottle of 500 ml
Cardboard box containing one bottle of 1000 ml
Cardboard box containing 20 bottles of 100 ml
Cardboard box containing 20 bottles of 250 ml
Cardboard box containing 10 bottles of 500 ml
Cardboard box containing 10 bottles of 1000 ml

Not all pack sizes may be marketed.

15. PID LINK (Do not print heading)

[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

16. Contact details

Marketing authorisation holder:
B. Braun Melsungen AG
Carl-Braun Strasse 1
34212 Melsungen
Germany

Manufacturer responsible for batch release:
B. Braun Medical SA
Ctra. de Terrassa, 121
08191 Rubí (Barcelona), Spain

Local representatives and contact details to report suspected adverse reactions:
B. Braun Medical Ltd
Brookdale Road
Thorncliffe Park Estate
Sheffield
S35 2PW
Email: medinfo.bbmuk@bbraun.com
Tel: 0800 2980299

17. Other information

POM-V

Approved 06 January 2026
Gavin Hall