PARTICULARS TO APPEAR ON THE OUTER PACKAGE CARDBOARD BOX

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

PG 600, Powder and solvent for solution for injection

2. STATEMENT OF ACTIVE SUBSTANCES

Each reconstituted 5ml dose contains:

Active substances:

Gonadotrophin, chorionic Ph. Eur.	200 IU
Serum gonadotrophin	400 IU

Solvent: Sterile phosphate buffered water for injection.

3. PACKAGE SIZE

5 x 1 dose with 5 x 5 ml solvent
1 x 5 doses with 1 x 25 ml solvent

4. TARGET SPECIES

Pigs.

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Intramuscular or subcutaneous use.

7. WITHDRAWAL PERIODS

Withdrawal period:
Meat and offal: Zero days.

8. EXPIRY DATE

Exp. {mm/yyyy}
Once broached use by…

9. SPECIAL STORAGE PRECAUTIONS

Store in a refrigerator (2 °C – 8 °C).
Protect from light.
Reconstituted product should be stored no longer than 24 hours at 2 °C – 8 °C.
Discard unused solution.
This product does not contain a preservative.
Keep the container in the outer carton

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

Intervet International B.V.

14. MARKETING AUTHORISATION NUMBER

Vm 06376/4095

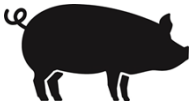
15. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS – VIAL LABEL (ACTIVE SUBSTANCE)

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

PG 600, Powder for solution for injection



2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Each dose contains:

Gonadotrophin, chorionic Ph. Eur.	200 IU
Serum gonadotrophin	400 IU

1 Dose
5 Doses

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

Once broached use by…
Reconstituted product should be stored no longer than 24 hours at 2 °C - 8 °C.

PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS - IMMEDIATE LABEL SOLVENT - 5 ml and 25 ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Solvent for PG 600

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Sterile phosphate buffered water for injection
5 ml
25 ml

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}
Reconstituted product should be stored no longer than 24 hours at 2 °C – 8 °C.

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

PG 600, Powder and solvent for solution for injection

2. Composition

Each reconstituted 5 ml dose contains:

Active substances:

Gonadotrophin, chorionic Ph. Eur.	200 IU
Serum gonadotrophin	400 IU

Powder for injection: White to off-white powder.
Solvent: Clear, colourless.

3. Target species

Pigs.

4. Indications for use

The action of serum gonadotrophin is similar to that of the follicle stimulating hormone (FSH) secreted by the anterior pituitary gland. In the same way chorionic gonadotrophin may be compared with luteinising hormone (LH).
The combination of gonadotrophins in the veterinary medicinal product is indicated for the promotion of a fertile oestrous cycle in gilts and in sows post-weaning and the treatment of suboestrus or anoestrus in barren sows due to hormonal imbalance.

5. Contraindications

Do not inject into the subcutaneous fat.

6. Special warnings

Special warnings:
None.

Special precautions for safe use in the target species:
When administered by the subcutaneous route, care must be taken to avoid injection into subcutaneous fat. In the unlikely event of an individual anaphylactoid reaction, 1 – 3 ml adrenaline 1:1000 solution should be given by intramuscular injection.
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 and show the package leaflet or the label to the physician.

Wash hands after use.

Pregnancy:
Not indicated for use during pregnancy. The injection of the veterinary medicinal product into a pregnant sow or gilt will fail to produce heat and will not cause abortion.

Interaction with other medicinal products and other forms of interaction:
None known.

Overdose:
No specific treatment or antidote recommended.

Major incompatibilities:
Do not mix with any other veterinary medicinal product, except the solvent supplied for use with the veterinary medicinal product.

7. Adverse events

Pigs:
None known.

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 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

Intramuscular or subcutaneous use.

Sows and gilts: One dose (5 ml) of reconstituted veterinary medicinal product should be aseptically injected subcutaneously or preferably, intramuscularly, at the base of the ear using a 1.5” needle, which must be directed horizontally.

Gilts: Administration of a single dose of the veterinary medicinal product to non-cycling gilts over the age of five months will normally result in a fertile oestrus within five days.

Sows post-weaning: To promote early post-partum oestrus (particularly where early weaning is practised) it is recommended that a single injection of the veterinary medicinal product is given within 48 hours of weaning.

Anoestrus or acyclic sows: Cases of suboestrus or anoestrus due to hormonal imbalance may respond to a single dose of the veterinary medicinal product, exhibiting normal heat within five days of injection.

If corpora lutea are present on the ovary when the veterinary medicinal product is administered, the sow or gilt will fail to respond. For this reason, if an anoestrus sow

or gilt fails to respond to the veterinary medicinal product, wait 10 days before repeating the injection.

9. Advice on correct administration

Reconstitution: Reconstitute by injecting approximately 2 ml of the solvent provided into the hormone vial, ensure the freeze-dried plug is fully dissolved, and re-inject the resulting solution into the solvent vial. Shake well before use.
Reconstituted solution is colourless.

10. Withdrawal periods

Withdrawal period: Meat and offal: Zero days.

11. Special storage precautions

Keep out of the sight and reach of children.

Store in a refrigerator (2 °C – 8 °C).
Keep the container in the outer carton.
Protect from light.

Reconstituted solution should be stored for no longer than 24 hours at 2 °C – 8 °C.
Discard unused solution.
This product does not contain a preservative.
Do not use this veterinary medicinal product after the expiry date which is stated on the carton after Exp. The expiry date refers to the last day of that month.
Keep the container in the outer carton

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 06376/4095

Pack sizes:
Cardboard box with 5 x single dose powder vials and 5 x 5 ml solvent vials.
Cardboard box with 1 x 5 doses powder vial and 1 x 25 ml solvent vial.

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:
Intervet International B.V.
Wim de Körverstraat 35
5831 AN Boxmeer
Netherlands

Manufacturer responsible for batch release:
Intervet International B.V.
Wim de Körverstraat 35
5831 AN Boxmeer
Netherlands

Or

Intervet International GmbH
Feldstrasse 1a
85716 Unterschleissheim
Germany

Local representative:
MSD Animal Health UK Limited
Walton Manor, Walton
Milton Keynes, MK7 7AJ
United Kingdom

Contact details to report suspected adverse reactions:
UK(GB)
MSD Animal Health UK Ltd.
Tel.: +44 (0)1908 685685

UK(NI)
Intervet Ireland Ltd.
Tel.: +353 (0)1 2970220

17. Other information

POM-V

Gavin Hall
Approved: 13 October 2025