

PARTICULARS TO APPEAR ON THE OUTER PACKAGING

Outer carton

1. NAME OF THE MEDICINAL PRODUCT

Implanon NXT®
68 mg implant

etonogestrel

2. STATEMENT OF ACTIVE SUBSTANCE(S)

1 contraceptive implant contains: 68 mg etonogestrel

3. LIST OF EXCIPIENTS

Other ingredients:

Ethylene vinyl acetate copolymer, Barium sulphate, Magnesium stearate.

See leaflet for further information

Medicinal product subject to medical prescription

4. PHARMACEUTICAL FORM AND CONTENTS

Implant

1 applicator containing 1 implant

5. METHOD AND ROUTE(S) OF ADMINISTRATION

Read the package leaflet before use.

For subdermal use

6. SPECIAL WARNING THAT THE MEDICINAL PRODUCT MUST BE STORED OUT OF THE SIGHT AND REACH OF CHILDREN

Keep out of the sight and reach of children.

7. OTHER SPECIAL WARNING(S), IF NECESSARY

8. EXPIRY DATE

EXP.:

9. SPECIAL STORAGE CONDITIONS

Store in the original blister package.

Do not store above 30°C.

10. SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED MEDICINAL PRODUCTS OR WASTE MATERIALS DERIVED FROM SUCH MEDICINAL PRODUCTS, IF APPROPRIATE

11. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

N.V. Organon, Kloosterstraat 6,
5349 AB Oss, The Netherlands
Tel: 00800-66550123
Email: dpoc.benelux@organon.com

12. MARKETING AUTHORISATION NUMBER(S)

13. BATCH NUMBER

Lot:

14. GENERAL CLASSIFICATION FOR SUPPLY

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

17. UNIQUE IDENTIFIER – 2D BARCODE

2D barcode carrying the unique identifier included.

18. UNIQUE IDENTIFIER – HUMAN READABLE DATA

SN
MFG
GTIN: 00366582507994

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGING

Blister

1. NAME OF THE MEDICINAL PRODUCT

Implanon NXT®
68 mg implant

etonogestrel

2. STATEMENT OF ACTIVE SUBSTANCE(S)

1 contraceptive implant contains: 68 mg etonogestrel

3. LIST OF EXCIPIENTS

4. PHARMACEUTICAL FORM AND CONTENTS

1 applicator containing 1 implant

5. METHOD AND ROUTE(S) OF ADMINISTRATION

For subdermal use.
Read the package leaflet before use.

6. SPECIAL WARNING THAT THE MEDICINAL PRODUCT MUST BE STORED OUT OF THE SIGHT AND REACH OF CHILDREN

Keep out of the sight and reach of children.

7. OTHER SPECIAL WARNING(S), IF NECESSARY

For single use only.
Contents are sterile unless package is damaged or opened.

8. EXPIRY DATE

EXP.:

9. SPECIAL STORAGE CONDITIONS

Store in the original blister package.

Do not store above 30°C.

10. SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED MEDICINAL PRODUCTS OR WASTE MATERIALS DERIVED FROM SUCH MEDICINAL PRODUCTS, IF APPROPRIATE

11. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

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13. BATCH NUMBER

Lot:

14. GENERAL CLASSIFICATION FOR SUPPLY

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

PARTICULARS TO APPEAR ON THE PATIENT ALERT CARD

1. NAME OF THE MEDICINAL PRODUCT

Implanon NXT®
68 mg implant
etonogestrel

2. OTHER SPECIAL WARNING(S), IF NECESSARY

Important notice:

Occasionally, gently palpate the implant to be sure that you know its location. If you cannot feel the implant at any time, contact your doctor as soon as possible.

The holder of this card is using a progestogen-only subdermal contraceptive implant. The implant is located at the inner side of the upper arm.

In, for example, the case of a casualty do not make an attempt to remove the implant.

Implanon NXT is X-ray visible.

See reverse side for further information.

KEEP THIS CARD IN A SAFE PLACE!

3. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

For further information, please contact:

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5349 AB Oss, The Netherlands
Tel: 00800-66550123
Email: dpoc.benelux@organon.com

4. BATCH NUMBER

Lot:

5. OTHER

Clinical details:

Name

Date of insertion

Date for removal

Arm

Left Right

Implanon NXT® is a registered trademark in one or more countries.

PARTICULARS TO APPEAR ON THE PATIENT LABEL

1. NAME OF THE MEDICINAL PRODUCT

Implanon NXT®
68 mg implant

2. BATCH NUMBER

Lot:

3. OTHER

For patient file

PARTICULARS TO APPEAR ON THE PHYSICIAN LABEL

1. NAME OF THE MEDICINAL PRODUCT

Implanon NXT®
68 mg implant

2. BATCH NUMBER

Lot:

3. OTHER

For physician file