

PARTICULARS TO APPEAR ON THE OUTER PACKAGING

CARTON BOX OF 1 BLISTER

1. NAME OF THE MEDICINAL PRODUCT

Mifepristone Linepharma 200 mg tablet
Mifepristone

2. STATEMENT OF ACTIVE SUBSTANCE(S)

1 tablet contains 200 mg mifepristone.

3. LIST OF EXCIPIENTS

4. PHARMACEUTICAL FORM AND CONTENTS

Tablet

1 tablet
30 tablets (as hospital pack).

5. METHOD AND ROUTE(S) OF ADMINISTRATION

Read the package leaflet before use.
Oral 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

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

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

<To be completed nationally>

12. MARKETING AUTHORISATION NUMBER(S)

13. BATCH NUMBER

Lot

14. GENERAL CLASSIFICATION FOR SUPPLY

Medicinal product subject to medical prescription.

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

Mifepristone Linepharma

MINIMUM PARTICULARS TO APPEAR ON BLISTERS OR STRIPS

ALU BLISTER

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Mifepristone

2. NAME OF THE MARKETING AUTHORISATION HOLDER

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3. EXPIRY DATE

EXP:

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Lot:

5. OTHER