PARTICULARS TO APPEAR ON THE OUTER PACKAGING

Carton for blister

1. NAME OF THE MEDICINAL PRODUCT

<ARIPIPRAZOLE> 5 mg tablets
<ARIPIPRAZOLE> 10 mg tablets
<ARIPIPRAZOLE> 15 mg tablets
<ARIPIPRAZOLE> 30 mg tablets

Aripiprazole

2. STATEMENT OF ACTIVE SUBSTANCE(S)

Each tablet contains 5 mg aripiprazole
Each tablet contains 10 mg aripiprazole
Each tablet contains 15 mg aripiprazole
Each tablet contains 30 mg aripiprazole

3. LIST OF EXCIPIENTS

Contains lactose. See package leaflet for further information.

4. PHARMACEUTICAL FORM AND CONTENTS

Tablet

7 tablets
10 tablets
14 tablets
20 tablets
28 tablets
30 tablets
50 tablets
56 tablets
60 tablets
72 tablets
90 tablets
98 tablets
100 tablets

5. METHOD AND ROUTE(S) OF ADMINISTRATION

Oral 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

8. EXPIRY DATE

EXP

9. SPECIAL STORAGE CONDITIONS

Store in the original package in order to protect from moisture.

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)

< [To be completed nationally]>

13. BATCH NUMBER

Lot

14. GENERAL CLASSIFICATION FOR SUPPLY

< [To be completed nationally]>

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

<ARIPIPRAZOLE> 5 mg
<ARIPIPRAZOLE> 10 mg
<ARIPIPRAZOLE> 15 mg
<ARIPIPRAZOLE> 30 mg

17. UNIQUE IDENTIFIER – 2D BARCODE

2D barcode carrying the unique identifier included.

18. UNIQUE IDENTIFIER – HUMAN READABLE DATA

PC:
SN:
NN: