PARTICULARS TO APPEAR ON THE OUTER PACKAGING

{NATURE/TYPE}

1. NAME OF THE MEDICINAL PRODUCT

<Product name> 5 mg tablets
<Product name> 10 mg tablets
<Product name> 15 mg tablets
<Product name> 20 mg tablets
<Product name> 30 mg tablets

Aripiprazole

2. STATEMENT OF ACTIVE SUBSTANCE(S)

Each tablet contains 5 mg of aripiprazole.
Each tablet contains 10 mg of aripiprazole.
Each tablet contains 15 mg of aripiprazole.
Each tablet contains 20 mg of aripiprazole.
Each tablet contains 30 mg of aripiprazole.

3. LIST OF EXCIPIENTS

Contains lactose. Read the package leaflet before use.

4. PHARMACEUTICAL FORM AND CONTENTS

Tablet

5 mg
7 tablets
7 x 1 tablet
14 tablets
14 x 1 tablet
15 tablets
28 tablets
28 x 1 tablet
30 tablets
49 tablets
56 tablets
56 x 1 tablet
60 tablets
98 tablets
98 x 1 tablet

100 tablets

10 mg
7 tablets
7 x 1 tablet
14 tablets
14 x 1 tablet
28 tablets
28 x 1 tablet
30 tablets
49 tablets
56 tablets
56 x 1 tablet
60 tablets
98 tablets
98 x 1 tablet
100 tablets

15 mg
7 tablets
7 x 1 tablet
14 tablets
14 x 1 tablet
28 tablets
28 x 1 tablet
30 tablets
49 tablets
56 tablets
56 x 1 tablet
60 tablets
98 tablets
98 x 1 tablet
100 tablets

20 mg
7 tablets
7 x 1 tablet
14 tablets
14 x 1 tablet
28 tablets
28 x 1 tablet
30 tablets
49 tablets
56 tablets
56 1 x tablet
98 tablets
98 x 1 tablet

30 mg
7 tablets
7 x 1 tablet

14 tablets
14 x 1 tablet
28 tablets
28 x 1 tablet
30 tablets
49 tablets
56 tablets
56 x 1 tablet
60 tablets
98 tablets
98 x 1 tablet
100 tablets

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

<To be completed nationally>

9. SPECIAL STORAGE CONDITIONS

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

<To be completed nationally>

14. GENERAL CLASSIFICATION FOR SUPPLY

<To be completed nationally>

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

<Product name> 5 mg tablets
<Product name> 10 mg tablets
<Product name> 15 mg tablets
<Product name> 20 mg tablets
<Product name> 30 mg tablets

Trilingual countries:
<Product name> 5 mg tabl.
<Product name> 10 mg tabl.
<Product name> 15 mg tabl.
<Product name> 20 mg tabl.
<Product name> 30 mg tabl.