PARTICULARS TO APPEAR ON THE OUTER PACKAGING

{Blister Pack Carton}

1. NAME OF THE MEDICINAL PRODUCT

..//.. 5 mg orodispersible tablets
..//.. 10 mg orodispersible tablets
..//.. 15 mg orodispersible tablets
..//.. 20 mg orodispersible tablets

escitalopram

2. STATEMENT OF ACTIVE SUBSTANCE(S)

..//.. 5 mg: Each orodispersible tablet contains 5 mg escitalopram, equivalent to 6.3875 mg escitalopram oxalate
..//.. 10 mg: Each orodispersible tablet contains 10 mg escitalopram, equivalent to 12.775 mg escitalopram oxalate
..//.. 15 mg: Each orodispersible tablet contains 15 mg escitalopram, equivalent to 19.1625 mg escitalopram oxalate
..//.. 20 mg: Each orodispersible tablet contains 20 mg escitalopram, equivalent to 25.55 mg escitalopram oxalate

3. LIST OF EXCIPIENTS

Also contains lactose monohydrate.
See enclosed leaflet for further information

4. PHARMACEUTICAL FORM AND CONTENTS

Orodispersible tablet

PT/H/0846/001, 003:
10 orodispersible tablets
12 orodispersible tablets
14 orodispersible tablets
15 orodispersible tablets
28 orodispersible tablets
30 orodispersible tablets
56 orodispersible tablets
60 orodispersible tablets
98 orodispersible tablets

100 orodispersible tablets
120 orodispersible tablets

PT/H/0846/002, 004: 10 orodispersible tablets
12 orodispersible tablets
14 orodispersible tablets
15 orodispersible tablets
28 orodispersible tablets
30 orodispersible tablets
56 orodispersible tablets
60 orodispersible tablets
98 orodispersible tablets
100 orodispersible tablets
120 orodispersible tablets

5. METHOD AND ROUTE(S) OF ADMINISTRATION

For 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: MM/YYYY

9. SPECIAL STORAGE CONDITIONS

Store in the original package in order to protect from moisture and 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)

[To be completed nationally]

13. BATCH NUMBER

Batch: XXXXXX

14. GENERAL CLASSIFICATION FOR SUPPLY

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

/../ 5 mg orodispersible tablet
/../ 10 mg orodispersible tablet
/../ 15 mg orodispersible tablet
/../ 20 mg orodispersible tablet