PARTICULARS TO APPEAR ON THE OUTER PACKAGING

CARTON

1. NAME OF THE MEDICINAL PRODUCT

Donepezil SymPhar, 5 mg, film-coated tablets

(Donepezil hydrochloride)

2. STATEMENT OF ACTIVE SUBSTANCE(S)

One film-coated tablet contains 5 mg of Donepezil hydrochloride.

3. LIST OF EXCIPIENTS

Contains lactose monohydrate.
See leaftlet for further information.

4. PHARMACEUTICAL FORM AND CONTENTS

28 film-coated tablets
56 film-coated tablets
84 film-coated tablets

5. METHOD AND ROUTE(S) OF ADMINISTRATION

For oral administration only. 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

Expiry date:

9. SPECIAL STORAGE CONDITIONS

Do not store above 30ºC.

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

SymPhar Sp. z o.o., ul.Włoska 1, 00-777 Warszawa

12. MARKETING AUTHORISATION NUMBER(S)

13. BATCH NUMBER

Batch number:

14. GENERAL CLASSIFICATION FOR SUPPLY

Medicinal product subject to medical prescription.

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

Donepezil SymPhar 5 mg