PARTICULARS TO APPEAR ON THE OUTER PACKAGING
(CARTON FOR BLISTER and HDPE container)

1. NAME OF THE MEDICINAL PRODUCT

[Invented name] 1 mg tablets
Rasagiline

2. STATEMENT OF ACTIVE SUBSTANCE(S)

Each tablet contains 1 mg of rasagiline (as rasagiline tartrate).

3. LIST OF EXCIPIENTS

4. PHARMACEUTICAL FORM AND CONTENTS

Tablet

DE/H/4364/001/DC
Only blister
10 tablets
28 tablets
30 tablets
98 tablets
100 tablets
112 tablets

Only tablet container
30 tablets
100 tablets

DE/H/4387/001/DC
Only blister
10 tablets
28 tablets
30 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

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

Medicinal product subject to medical prescription.

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

[Invented name] 1 mg tablets

17. UNIQUE IDENTIFIER – 2D BARCODE

2D barcode carrying the unique identifier included.

18. UNIQUE IDENTIFIER - HUMAN READABLE DATA

PC: {number}
SN: {number}
NN: {number}

1.3.1 spc-label-pl - common-outer - 4,353 (DE/H/4364-4387/001 Change ID 164785 CDS)	20180626~~20180611~~
RASAGILINE TARTRATE 1 MG TABLET	722-2444.00