PARTICULARS TO APPEAR ON THE OUTER PACKAGING AND IMMEDIATE PACKAGING

CONTAINER LABEL/CARTON LABEL

1. NAME OF THE MEDICINAL PRODUCT

<Invented Name> 750 mg Film-coated Tablets
Glucosamine sulfate

2. STATEMENT OF ACTIVE SUBSTANCE(S)

Each tablet contains 942 mg glucosamine sulfate sodium chloride equivalent to 750 mg glucosamine sulfate or 589 mg glucosamine.

3. LIST OF EXCIPIENTS

Excipients: also contains sodium, soya lecithin and lactose monohydrate. See leaflet for further information

4. PHARMACEUTICAL FORM AND CONTENTS

8 Film-Coated Tablets
10 Film-Coated Tablets
12 Film-Coated Tablets
14 Film-Coated Tablets
20 Film-Coated Tablets
28 Film-Coated Tablets
30 Film-Coated Tablets
56 Film-Coated Tablets
60 Film-Coated Tablets
112 Film-Coated Tablets
120 Film-Coated Tablets
168 Film-Coated Tablets
180 Film-Coated Tablets
336 Film-Coated Tablets
360 Film-Coated 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 REACH AND SIGHT OF CHILDREN

Keep out of the ~~reach and sight~~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.
Once opened the tablets should be used within 6 months.
Use by

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

BN or Lot

14. GENERAL CLASSIFICATION FOR SUPPLY

<To be completed nationally>~~POM~~
~~Medicinal product not subject to medical prescription~~
~~Medicinal product subject to medical prescription~~
~~POM~~

15. INSTRUCTIONS ON USE

Take one tablet twice a day or 2 tablets once a day.
The tablets should be swallowed whole with water.
Relief of symptoms in mild to moderate osteoarthritis of the knee

16. INFORMATION IN BRAILLE

<Invented Name> 750 mg tablets

17. OTHER

PARTICULARS TO APPEAR ON THE OUTER PACKAGING

CARTON LABELLING FOR BLISTERS

1. NAME OF THE MEDICINAL PRODUCT

<Invented Name> 750 mg Film-coated Tablets
Glucosamine sulfate

2. STATEMENT OF ACTIVE SUBSTANCE(S)

Each tablet contains 942 mg glucosamine sulfate sodium chloride equivalent to 750 mg glucosamine sulfate or 589 mg glucosamine.

3. LIST OF EXCIPIENTS

Excipients: also contains sodium, soya lecithin and lactose monohydrate. See leaflet for further information

4. PHARMACEUTICAL FORM AND CONTENTS

8 Film-Coated Tablets
10 Film-Coated Tablets
12 Film-Coated Tablets
14 Film-Coated Tablets
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56 Film-Coated Tablets
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112 Film-Coated Tablets
120 Film-Coated Tablets
168 Film-Coated Tablets
180 Film-Coated Tablets
336 Film-Coated Tablets
360 Film-Coated 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 REACH AND SIGHT OF CHILDREN

Keep out of the ~~reach and sight~~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

BN or Lot

14. GENERAL CLASSIFICATION FOR SUPPLY

<To be completed nationally>
~~Medicinal product not subject to medical prescription~~
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~~POM~~

15. INSTRUCTIONS ON USE

Take one tablet twice a day or 2 tablets once a day.
The tablets should be swallowed whole with water.
Relief of symptoms in mild to moderate osteoarthritis of the knee

16. INFORMATION IN BRAILLE

<Invented Name> 750mg tablets

17. OTHER