Updated: August 2025

# PARTICULARS TO APPEAR ON THE OUTER PACKAGING CARTON LABEL

## 1. Name of the medicinal product

[RH094 trade name]\* Misoprostol 200 µg tablets Misoprostol

#### 2. Statement of active substance

Each tablet contains 200 µg misoprostol

### 3. List of excipients

Each tablet contains hydrogenated castor oil. See patient information leaflet for further information.

#### 4. Pharmaceutical form and contents

1 x 4 tablets

2 x 4 tablets

3 x 4 tablets

15 x 4 tablets

30 x 4 tablets

#### 5. Method and route of administration

Oral, vaginal, sublingual or buccal use Read the patient information 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 sight and reach of children.

#### 7. Other special warning(s), if necessary

### 8. Expiry date

EXP {MM/YYYY}

### 9. Special storage conditions

Do not store above 30°C. Store in the original package.

# 10. Special precautions for disposal of unused medicinal products or waste materials derived from such medicinal products, if appropriate

\* Trade names are not prequalified by WHO. This is the national medicines regulatory authority's responsibility.

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## 11. Name and address of the supplier

Senador Laboratories Private Limited, Plot 2B & 2C, Biotech Park, Phase II, Lalgadi Malakpet, Medchal Malkajgiri, Hyderabad, Telangana, India, 500101

# 12. WHO Reference Number (Prequalification Programme)

RH094

#### 13. Manufacturer's batch number

<Batch> {number}

## 14. (Advice on) General classification for supply

Medicinal product subject to medical prescription.

### 15. Instructions on use

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# MINIMUM PARTICULARS TO APPEAR ON BLISTERS OR STRIP

## 1. Name of the medicinal product

[RH094 trade name]  $^{\dagger}$  Misoprostol 200  $\mu g$  tablets Misoprostol

# 2. Name of the supplier

Senador Laboratories Private Limited

# 3. Expiry date

EXP {MM/YYYY}

### 4. Manufacturer's batch number

<Batch> {number}

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<sup>&</sup>lt;sup>†</sup> Trade names are not prequalified by WHO. This is the national medicines regulatory authority's responsibility.