LABELLING

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGING
Bottle Label

1. Name of the medicinal product

[HA718 trade name]* Dolutegravir (sodium) Tablets

2. Statement of active substance

Each film-coated tablet contains 50 mg dolutegravir (as sodium).

3. List of excipients

Each film-coated tablet contains mannitol and sunset yellow FCF (E 110).
See patient information leaflet for further information.

4. Pharmaceutical form and contents

Film Coated Tablets

30 tablets
90 tablets

5. Method and route of administration

Oral 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 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 container.

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

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 supplier

Laurus Labs Limited
2nd Floor, Serene Chambers, Road No.7
Banjara Hills, Hyderabad, Telangana-500034
India

12. WHO Reference Number (Prequalification Programme)

HA718

13. Manufacturer's batch number

<Batch> <Lot> {number}

14. (Advice on) General classification for supply

Medicinal product subject to medical prescription.

15. Instructions on use