LABELING

PARTICULARS TO APPEAR ON THE OUTER PACKAGING

Label/ white HDPE bottle

1. NAME OF THE MEDICINAL PRODUCT

Lamivudine/Zidovudine 150 mg/300 mg Tablets*

2. STATEMENT OF ACTIVE SUBSTANCE

Each film-coated tablet contains 150 mg lamivudine and 300 mg zidovudine

3. LIST OF EXCIPIENTS

Contains less than 1 mmol sodium per tablet, that is to say essentially 'sodium-free'
See patient information leaflet for further information.

4. PHARMACEUTICAL FORM AND CONTENTS

60 film-coated 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 and sight of children.

7. OTHER SPECIAL WARNING(S), IF NECESSARY

8. EXPIRY DATE

EXP {MM/YYYY}

* Trade names are not prequalified by WHO. This is the national medicines regulatory authority's (NMRA) responsibility. Throughout this WHOPAR the Proprietary Name is given as an example only.

9. SPECIAL STORAGE CONDITIONS

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

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

Anhui Biochem Bio-Pharmaceutical Co., Ltd
No. 30 Hongfeng Road, Hi-Tech Development Zone, Hefei City,
Anhui Province, 230088,
People's Republic of China

12. WHO REFERENCE NUMBER (PREQUALIFICATION PROGRAMME)

HA656

13. MANUFACTURER'S BATCH NUMBER

<Batch> {number}

14. GENERAL CLASSIFICATION FOR SUPPLY

Medicinal product subject to medical prescription.

15. INSTRUCTIONS ON USE