CAPSULE RIFABUTIN (150 mg.)

Description: Each Rifabutin Capsule shall contain Rifabutin USP 150 mg.
Rifabutin Capsules shall conform to the requirements of USP.

Labelling on Strips :
The label shall indicate the content of Rifabutin USP in each Capsule.
The label shall incorporate manufacturing license no., batch no., date of mfg., date of expiry of the individual drug and the storage requirements. The label shall also bear "Schedule H Drug".
The label shall conform to the requirements of Rule 96 of Drugs & Cosmetics Act.

Labelling on Millboard/Greyboard Box:
The label shall indicate the content of Rifabutin USP in each Capsule.
The label shall incorporate manufacturing license no., batch no. date of mfg., date of expiry, of the drug and storage requirements along with the number of the strips. The label shall also bear "Schedule H Drug".
The label shall conform to the requirements of Rule 96 of Drugs & Cosmetics Act.

Storage: Store protected from light and from excessive heat.

Shelf Life: Shelf life of Rifabutin would be 24 months.
At least $5/6^{th}$ of the total stipulated shelf life must remain at the time of purchase.
(In case of limited procurement, $5/6^{th}$ requirements may be relaxed depending upon the anticipated period of consumption).