For the use of Registered Medical Practitioner, Pharmacist, Hospital or a Laboratory only.

UNIZAR FORTE

(Losartan Potassium, Amlodipine & Hydrochlorothiazide Tablets)

COMPOSITION:

Each film coated tablet contains:

Losartan Potassium USP

Amlodipine Besylate USP

Equivalent to Amlodipine Hydrochlorothiazide USP

Colour: Ferric oxide red

Excipients:

50 mg

5 mg

12.5 mg

q.s.

INDICATIONS: UNIZAR FORTE it is indicated for the treatment of cardiovascular disease.

CONTRAINDICATIONS: Pregnancy, lactation; children with CrCl <30 ml/min/1.73m2.

PRECAUTIONS: Volume-depleted patients including patients on diuretics and salt restriction; renal artery stenosis; elderly; renal or hepatic impairment. Monitor serum-potassium concentration.

PREGNANCY AND LACTATION:

Category C: Either studies in animals have revealed adverse effects on the foetus (teratogenic or embryocidal or other) and there are no controlled studies in women or studies in women and animals are not available. Drugs should be given only if the potential benefit justifies the potential risk to the foetus.

Category D: There is positive evidence of human foetal risk, but the benefits from use in pregnant women may be acceptable despite the risk (e.g., if the drug is needed in a life-threatening situation or for a serious disease for which safer drugs cannot be used or are ineffective).

SIDE EFFECTS: Headache, dizziness, back pain, myalgia, respiratory tract disorders, asthenia/fatigue, first dose hypotension, rash, angioedema, neutropenia, GI disturbances, transient elevation of liver enzymes, impaired renal function, taste disturbances and hyperkalaemia.

DRUG INTERACTIONS: Hypotensive effect of losartan potentiated by diuretics and other antihypertensives. Risk of hyperkalaemia increases with concomitant ACE inhibitors, ciclosporin, potassium-sparing diuretics and K supplements. Hypotensive effect may be antagonised and increased risk of renal impairment when used with NSAIDs.

Potentially Fatal: Risk of lithium toxicity with losartan use.

DOSAGE AND ADMINISTRATION: As directed by the physician.

STORAGE: Store below 30°C in a dark place.

PRESENTATION: Blister Pack of 10 Tablets.

Marketed by:
MEDISEL (KENYA) LIMITED
Factory Street,
P.O.BOX 540, Thika,
Kenya.

Manufactured in India by:



AUROCHEM PHARMACEUTICALS (I) PVT LTD.
58, PalgharTaluka Industrial Co-op Estate Limited,
Palghar-401 404. Dist- Thane, INDIA.
www.aurochemlaboratories.com