Ramipril tablet

Each film coated tablet contains Ramipril BP 5 mg. Each film coated tablet contains Ramipril BP 10 mg.

COMPOSITION: Ramipril-1.25 Tablets Each tablet contains......Ramipril 1.25 mg

Ramipril-2.5 Tablets Each tablet contains.....Ramipril 2.5 mg

Ramipril-5 Tablets Each tablet contains......Ramipril 5 mg

INDICATIONS

* Reduction in risk of myocardial infarction, stroke and death from cardiovascular causes

* Hypertension

* Heart failure post myocardial infarction

DOSAGE AND ADMINISTRATION Reduction in risk of myocardial infarction, stroke and death from cardiovascular causes

Ramipril should be given at an initial dose of 2.5 mg, once a day for 1 week, 5 mg, once a day for the next 3 weeks and then increased as tolerated, to a maintenance dose of 10 mg, once daily. If the patient is hypertensive or recently post myocardial infarction, it can be given as a divided dose.

Hypertension

The recommended initial dose of Ramipril is 2.5 mg once daily in patients not receiving a diuretic. Dosage should be adjusted according to the blood pressure response. The usual maintenance dosage range is 2.5 to 20 mg/day administered as a single dose or in two equally divided doses. If blood pressure is not controlled with ramipril alone, a diuretic can be added.

Heart failure post myocardial infarction

The recommended starting dose of Ramipril is 2.5 mg twice daily. A patient who becomes hypotensive at this dose may be switched to 1.25 mg twice daily, and after one week at the starting dose, patients should be titrated (if tolerated) toward a target dose of 5 mg twice daily, with dosage increases being about 3 weeks apart. After the initial dose of ramipril, the patient should be observed under medical supervision for at least two hours and until blood pressure has stabilized for at least an additional hour. If possible, the dose of any concomitant diuretic should be reduced which may diminish the likelihood of hypotension.

In patients who are currently being treated with a diuretic, symptomatic hypotension occasionally can occur following the initial dose of ramipril. To reduce the likelihood of hypotension, the diuretic should, if possible, be discontinued two to three days prior to beginning therapy with ramipril. Then, if blood pressure is not controlled with ramipril alone, diuretic therapy should be resumed. If the diuretic cannot be discontinued, an initial dose of 1.25 mg ramipril should be used to avoid excess hypotension.

Dosage in renal impairment

In patients with creatinine clearance less than 40 mL/min/1.73m 2 (serum creatinine approximately more than 2.5 mg/dl) doses only 25% of those normally used should be expected to induce full therapeutic levels of ramiprilat.

Hypertension

For patients with hypertension and renal impairment, the recommended initial dose is 1.25 mg ramipril once daily. Dosage may be titrated upward until blood pressure is controlled or to a maximum total daily dose of 5 mg.

Heart failure post myocardial infarction

For patients with heart failure and renal impairment, the recommended initial dose is 1.25 mg ramipril once daily. The dose may be increased to 1.25 mg twice daily and up to a maximum dose of 2.5 mg twice daily depending upon clinical response and tolerability.

CONTRAINDICATIONS

Hypersensitivity to ramipril or any other ACE inhibitor (e.g. a patient who has experienced angioedema during therapy with any other ACE inhibitor).

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PACKAGING INFORMATION:

Ramipril-1.25Strip of 10 tablets Ramipril-2.5Strip of 10 tablets Ramipril-5Strip of 10 tablets



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About Taj Pharmaceutical Limited

Taj Pharmaceuticals Limited is a pharmaceutical company founded and based in India. The company manufacturers pharmaceutical formulations and API for India and other countries of world. The company was established in 1995 as an enterprise and in 2004 became a public limited company. As per Mumbai pharmaxil and Chemixil association the company manufacturers and exports to countries like Albania, Argentina, Austria, Chile and Iraq. In 1995 pharmaceuticals wing only has a schedule M certification for pharmaceuticals products manufacturing in India. Taj Pharmaceuticals established its manufacturing unit in Gujarat because of government policies in 1999 with WHO / GMP licence. The company in 2003 revived all the old manufacturing units and approached the FDA Gujarat for 4000 new pharmaceuticals drug permissions for the first time in India.

According to the Indian Trade Mark the company owns about 450 brands and 4600 generic manufacturing permissions in India. According to the export data analysis the company was the largest exporter of generic medicines to the Europe and Middle East countries.

www.tajpharma.com

The company medicines are present in France, Georgia, Egypt and CIF countries.



Taj Pharmaceuticals Limited

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