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Medicinal product

- valsartan® 40 mg Film-Coated tablet
- valsartan® 80 mg Scored Film-Coated tablet
- valsartan® 160 mg Scored Film-Coated tablet

Pharmacologic Category

Angiotensin II Receptor Blocker Antihypertensive

Therapeutic indications

Treatment of heart failure

Management of hypertension

Reduction of cardiovascular mortality in patients with left ventricular dysfunction or failure post-myocardial infarction

Use: Off-Label

Acute coronary syndrome (secondary prevention of cardiovascular events); Improve kidney outcomes in hypertensive patients with chronic kidney disease (CKD) (diabetic and nondiabetic population)

Dosing

Hypertension: initial: 80 mg or 160 mg once daily, dose may be increased to achieve the desired effect; usual dosage range: 80 to 320 mg once daily; target dose:160 to 320 mg once daily; maximum dose: 320 mg/day Heart failure: Initial: 40 mg twice daily; titrate dose to 80 mg, and then to 160 mg twice daily, as tolerated; maximum dose: 320 mg/day

Left ventricular dysfunction or failure after MI: Initial: 20 mg twice daily; may increase within 7 days to 40 mg twice daily; titrate dose to target of 160 mg twice daily as tolerated; may initiate ≥12 hours following MI

Valsartan produces direct antagonism of the angiotensin II (AT2) receptors, unlike the ACE inhibitors. It displaces angiotensin If from the AT1 receptor and produces its blood pressure-lowering effects by antagonizing AT1-induced vasoconstriction, aldosterone release, catecholamine release, arginine vasopressin release, water intake, and hypertrophic responses. This action results in a more efficient blockade of the cardiovascular effects of angiotensin II and fewer side effects than the ACE inhibitors

Contraindications

Pregnancy

Breast-feeding

Hypersensitivity to the active substance or to any of the excipients

Severe hepatic impairment, biliary cirrhosis, and cholestasis

Concomitant use with Aliskiren-containing products in patients with diabetes mellitus or renal impairment

Interactions

Taking the following drugs or supplements while taking valsartan may cause adverse reactions: Potassium-sparing diuretics, Potassium supplements, Salt substitutes, NSAIDs, Lithium, ACE Inhibitors, Aliskiren, and Other

Significant adverse reaction

Allergic reaction, hyperkalemia, hypotension, Increased blood urea nitrogen, Dizziness

Special warnings and precautions

Fetal Toxicity: Valsartan poses serious risks to an unborn baby. Women should not take valsartan during pregnancy Hypotension: Valsartan can cause low blood pressure, leading to dizziness or lightheadedness Kidney Problems: Valsartan increases the risk of acute kidney failure. The kidney function should be monitored High Potassium Levels: Some patients may see an increase in their potassium levels while taking valsartan

Administration

Administer with or without food

Monitoring Parameters

Baseline and periodic blood pressure, electrolyte panels, renal function; in HF, serum potassium during dose escalation and periodically thereafter

Pregnancy and breastfeeding

Drugs that act on the renin-angiotensin system can cause injury and death to the developing fetus. Discontinue as soon as possible once pregnancy is detected

Valsartan is not recommended for use during breastfeeding and alternative treatments with better-established safety profiles during breastfeeding are preferable

- 1 https://www.uptodate.com/
- 2- https://www.drugs.com/ 3- https://www.medicines.org.uk/
- 4- https://www.medicalnewstoday.com/