

Terlipressin EVER Pharma is indicated for the treatment of bleeding oesophageal varices and Hepato-renal syndrome type I

Terlipressin EVER Pharma was developed to offer a more convenient and safer to handle product than the originator Glypressin®. Available as a ready to use solution in a vial, there is no need to either reconstitute the product with solvent or break open ampoules.

Both initial and maintenance dose ranges are conveniently and economically covered by providing a 10ml vial in addition to the current 5ml vial.

Ready to use product - for immediate use without reconstitution

Available in vials - safer and more convenient than ampoules

5ml and new 10ml presentation covering initial and maintenance dose ranges

Additional indication - for Hepato-renal syndrome type I



Terlipressin acetate



Active Ingredient	Terlipressin acetate
Excipients	Sodium chloride , Acetic acid, Sodium hydroxide (for pH-adjustment), Hydrochloric acid (for pH-adjustment), Water for injections
Primary Packaging	Colourless glass vial
Closure	Bromobutyl rubber stopper and sealed with aluminium flip-off cap
Strength	5ml of injection solution contains 1mg terlipressin acetate corresponding to 0.85mg terlipressin
	10ml of injection solution contains 2mg terlipressin acetate corresponding to 1.7mg terlipressin
Active Strengths	Each ml contains 0.2mg terlipressin acetate corresponding to 0.17mg terlipressin
Presentations	5ml and 10ml
Pack sizes	1 x 5ml, 5 x 5ml, 1 x 10ml, 5 x 10ml
Indications	Treatment of bleeding oesophageal varices, emergency treatment of type 1 hepatorenal syndrome, as defined by IAC (International Ascites Club) criteria
Stability Unopened	24 months, Store in a refrigerator (2°C to 8°C). Do not freeze
After 1st Opening	Use Immediately

Terlipressin acetate EVER Pharma 0.2 mg/ml solution for injection Composition: Each 1 ml contains 0.2 mg/ml solution for injection pH-adjustment), Water for injections: Therapeutic indications: Treatment of bleeding oesophageal varices. Emergency treatment of type 1 hepatorenal syndrome, as defined by IAC (International Ascites Club) criteria. Contraindications: Hypersensitivity to the active substance or to any of the excipients listed in section 6.1., Pregnancy. Side effects: common: headache, ventricular and supra-ven-tricular arrhythmia, bradycardia, signs of lschaemia in the ECG, hypertension, hypotension, peripheral ischaemia, peripheral vasoconstriction, facial palor, transient adominal cramps, transient diarrhea, paleness, abdominal cramps (in women), uncommon: hyponatraemia, triggering of a convulsive disorder, angina pectorias, acute hypertension rise, in particular in patients already suffering from hypertension (generally, it decreases spontaneously), atrial fibrillation, ventricular stratysciles, tachycardia, chest pain, myocardial infarction, fluid overload with pulmonary oedema, intestinal ischaemia, peripheral cyanosis, hot flushes, pain in the chest, bronchospasm, respiratory distress, respiratory failure, transient mounting, hymphangitis, rare: hypergylocardial ischaemia, styspoea, coal cutaneous necrosis. More information about pharmaceutical form, posology and method of administration, special warnings and precautions for use, interaction with other medicinal products and other forms of interaction, fertility, pregnancy and licatation, effects on ability to drive and use machines, undestinable effects, overdose, pharmacodynamics properties, pharmacokinetic properties, precilical safety data, incompatibilities, shelf life, special precautions for storage, nature and contents of the container and special products and disposal is available in the summary of product characteristics. Only available on prescription. Last update: June 2016, Marketing Authorisation Holder: EVER Valin

