

EMESTOP

Ampoule, Tablets & Syrup

Composition

Emestop Injection

Each ampoule of 2 ml contains Metoclopramide hydrochloride 10 mg.

Emestop Syrup

Each teaspoonful (5 ml) contains Metoclopramide hydrochloride 5 mg.

Emestop Tablets

Each tablet contains Metoclopramide hydrochloride 10 mg.

Action

Emestop, a dopamine receptor antagonist, acts both centrally and peripherally. Its antiemetic action is due to a central effect on the chemoreceptor trigger zone-vomiting centre). Peripherally, Emestop stimulates motility of the upper gastrointestinal tract without affecting gastric, biliary or pancreatic secretions. Experimental evidence has suggested that the effect on motility may be associated with enhanced cholinergic excitatory processes at the postganglionic neuromuscular junction, antagonism of non-adrenergic, non-cholinergic (i.e. dopaminergic) inhibitory nerves and/or a direct effect on smooth muscle.

The post-synaptic activity of Metoclopramide appears to result from its ability to enhance acetylcholine release from post-ganglionic neurons in the gastrointestinal tract, and from its ability to sensitize muscarinic receptors of gastrointestinal smooth muscle to the actions of acetylcholine.

Although the major effects of Metoclopramide appear to be cholinergically based, antagonism of gastrointestinal dopaminergic activity might enhance the cholinergic like activity of the drug.

The effect of Emestop is not dependent on intact vagal innervations, but anticholinergic drugs can abolish it.

Emestop increases the tone and amplitude of gastric and especially antral contractions. It relaxes the pyloric sphincter and the duodenal bulb and increases peristalsis of the duodenum and jejunum, resulting in accelerated gastric emptying and intestinal transit. Like phenothiazines and related drugs that are dopamine antagonists, Metoclopramide produces sedation and may produce extrapyramidal reactions. The drug inhibits the central and peripheral effects of apomorphine, induces release of prolactin and causes a transient increase in circulating aldosterone levels. Emestop also increases the resting tone of the lower esophageal sphincter but has little, if any, effect on the colon or the gall bladder.

Pharmacokinetics

The onset of the pharmacological action of Emestop is 1-3 minutes following an I.V. dose, 10-15 minutes following I.M. administration, and 30-60 minutes after an oral dose. The effect of Metoclopramide persists for 1-2 hours, and the plasma half-life of an oral dose is about 4 hours. This prolongs in patients with impaired renal function. About 85% of the radioactivity of an oral dose appears in the urine within 72 hours, with half appearing as unchanged drug or glucuronide and sulphate conjugates.

Indications

Digestive Disorders

Relief of symptoms such as gastrointestinal pain, heartburn, dyspepsia, flatulence, regurgitation in peptic ulcer, reflux esophagitis, gastritis, duodenitis, hiatus hernia, cholelithiasis post-cholecystectomy dyspepsia, post-anaesthetic vomiting, vomiting due to anti-migraine therapy and intolerance to drugs.

Nausea and Vomiting

- Nausea and vomiting associated with diabetic gastroparesis.

- Nausea and vomiting caused by emetogenic cancer chemotherapy.
- Nausea and vomiting in postoperative conditions, e.g. restoration of gastrointestinal motility in the post-vagotomy syndrome and/or in morphine-induced delay in gastric emptying.

Facilitation of Diagnostic Procedures

Enhancement of passage of barium meals and facilitation of duodenal intubation procedures.

Contraindications

- Metoclopramide is contraindicated in patients with known sensitivity or intolerance to its action. It should not be administered to patients with pheochromocytoma because it may induce a hypertensive crisis, probably due to the release of catecholamines from the tumour.
- Metoclopramide should not be used whenever stimulation of gastrointestinal motility might be dangerous, such as in the presence of gastrointestinal hemorrhage, mechanical obstruction or perforation.
- Metoclopramide is also contraindicated in epileptics or patients receiving other drugs likely to cause extrapyramidal reactions, since the frequency and severity of seizures or extrapyramidal symptoms may be increased.

Warnings

Pregnancy

Category B

Reproduction studies in small animals at 12-250 times the human dose have revealed no evidence of impaired fertility or significant harm to the fetus. However, there are no well controlled studies in pregnant women. Therefore, this drug should only be used when clearly needed, and when the potential benefits outweigh the unknown potential hazards to the fetus.

Nursing Mothers

Metoclopramide readily enters into breast milk and may concentrate to about twice the plasma level 2 hour's post administration. Therefore, it is recommended that nursing be discontinued if Emestop is indicated.

Adverse Reactions

The most frequent adverse reactions are restlessness, drowsiness, fatigue, lassitude, agitation, irritability, constipation, and diarrhea, and urticarial maculopapular rash, dryness of the mouth, methemoglobinemia, neck pain and rigidity. These reactions occur in approximately 10% of patients treated and are usually mild, transient and reversible upon drug withdrawal.

Extrapyramidal adverse reactions occur in approximately 1-9% of treated patients. More frequently, these reactions occur in children and young adults, and include dystonic reactions of various types which are not dose related (e.g. involuntary movement of limbs and facial grimacing, torticollis, oculogyric crisis, rhythmic protrusion of the tongue, bulbar type of speech, trismus) and which respond rapidly to treatment with antiparkinsonism drugs such as biperiden I.M. or diazepam.

Depression has been reported but this is rare.

Urinary incontinence has also been reported.

Edema (reversible) has also been reported as an adverse reaction with Metoclopramide. This may be due to the effect by Metoclopramide of increasing aldosterone levels with resultant retention of salt and water.

Precautions

The relief provided by Metoclopramide may delay recognition of serious diseases. Therefore, Metoclopramide should not be prescribed until a diagnosis has been made. Patients should be cautioned against engaging in potentially dangerous activities requiring mental alertness, such as driving a car or operating machinery, for a few hours after the drug has been administered. The same precaution applies to childhood activities such as riding a bicycle or playing near traffic.

Intravenous Metoclopramide should be administered slowly, as rapid administration may induce a transient but intense feeling of anxiety and restlessness, followed by drowsiness. Care should be exercised in diabetic patients, as drug-induced gastroparesis may lead to premature activity of administered insulins, resulting in hypoglycemia.

Metoclopramide elevates prolactin levels that persist during chronic administration. Although the clinical significance of this is not clear, extreme caution should be exercised in patients with elevated serum prolactin levels and/or in patients with known or suspected breast cancer.

Metoclopramide has been reported to induce tardive dyskinesia and dysarthria, especially in geriatric patients. This may develop during administration and/or following discontinuation of use. These effects, although possibly reversible, are extremely distressing and their resolution is often slow. Metoclopramide should be avoided in patients with a pre-existing dopaminergic deficiency such as idiopathic Parkinsonism. However, signs of Parkinsonism may even appear in neurologically normal patients if higher than recommended doses are administered or renal insufficiency is present. In patients with severe renal function impairment, the dose should be reduced by at least 60% of that normally prescribed. This is particularly important because Metoclopramide induced Parkinsonism is resistant to levodopa therapy.

Drug Interactions

Metoclopramide/Anticholinergics/ Narcotic Analgesics

Anticholinergic drugs and narcotic analgesics antagonize the effects of Metoclopramide on gastrointestinal motility.

Metoclopramide/ Alcohol/ Sedatives/ Hypnotics/ Narcotics/ Tranquilizers

Additive sedative effects may occur when Metoclopramide is given in conjunction with alcohol or drugs belonging to the classes in this list.

Metoclopramide/ Digoxin/ Cimetidine/ Penicillin

Metoclopramide may diminish the absorption of these agents from the intestine, thus decreasing their bioavailability.

Metoclopramide/ Paracetamol/ Acetylsalicylic Acid/ Levodopa/ Diazepam/ Ethanol/ Lithium/ Tetracycline/ Zopiclone

Metoclopramide may enhance the absorption of these agents from the small intestine.

Metoclopramide/ Insulin

Because Metoclopramide influences the delivery of food to the intestines and thus the rate of absorption, insulin dosage or timing thereof may require adjustment.

Dosage and Administration

The daily dosage of Emestop is based on 0.5 mg/kg body weight, given in equally divided doses. In normal circumstances, this should not be exceeded. It is particularly important to adhere to this dosage in children and young adults.

The parenteral route (I.M. or I.V.) is recommended if severe symptoms are present. Intravenous injections should be administered slowly over a 1-2 minute period. Intravenous infusions should be administered slowly over a period of not less than 15 minutes.

Emestop syrup should be given half an hour before, or one hour after meals. Emestop tablets should be taken with half a glass of water or other liquid, half an hour before, or one hour after meals.

Digestive Disorders

Emestop Ampoules

Adults

1 ampoule of 10 mg I.M. or I.V., 1-3 times daily, according to the severity of the condition.

Adolescents 15-20 years

1/2 - 1 ampoule of 10 mg I.M. or I.V. 1-3 times daily

Children 5-14 years

Up to 1/2 ampoule of 10 mg I.M. or I.V., 1-3 times daily

Emestop Tablets

Adults

One tablet, 3-4 times daily.

Adolescents 15-20 years

1/2 -One tablet, 2-3 times daily

Emestop Syrup

Adults

Two teaspoonsfuls, 3 or 4 times daily

Adolescents 15-20 years

1-2 teaspoonsfuls, 2-3 times daily

Children

5-14 years: 1/2 -1 teaspoonful, 2-3 times daily.

3-5 years: about 40 drop (2 mg), 2-3 times daily.

1-3 years: about 20 drop (1 mg), 2-3 times daily.

Nausea and Vomiting associated with Diabetic Gastroparesis

One tablet of Emestop, half an hour before each meal and at bedtime for 2-8 weeks, depending on the response and the likelihood of continued well-being on cessation of treatment

The initial route of administration depends on the severity of the observable symptoms. If only the earliest manifestations of gastric stasis are present, the oral route is indicated. However, if the symptoms are more severe, I.V. therapy should be instituted (for up to 10 days) until symptoms subside. After 10 days, oral administration should be used for maintenance.

Nausea and Vomiting caused by Emetogenic Cancer Chemotherapy

Emestop injection should be diluted in 50 ml of a parenteral solution .

Intravenous infusions should be administered slowly over a period of not less than 15 minutes, half an hour before beginning cancer chemotherapy, and repeated every 2 hours for 2 doses, then every 3 hours for 3 doses. The initial doses should be 2 mg/kg body weight if highly emetogenic drugs such as cisplatin or dacarbazine are used alone or in combination. For less emetogenic regimens, 1 mg/kg body weight per dose may be adequate.

Nausea and Vomiting in Postoperative Conditions

In postoperative conditions such as restoration of gastrointestinal motility in the vagotomy syndrome and/or morphine-induced delay in gastric emptying, the recommended dosages are those listed under Digestive Disorders.

Facilitation of Diagnostic Procedures

To Facilitate Small Bowel Intubation

If the tube has not passed the pylorus with conventional manoeuvres in 10 minutes, the intravenous route may administer a single dose of Emestop injection (undiluted) slowly over a 1-2 minute period. The recommended single dose of Emestop injection is 10 mg (adults), 2.5-5 mg (children 6-14 years of age) and 0.1 mg/kg body weight (children under 6 years of age).

To Aid in Radiological Examinations

In patients where delayed gastric emptying interferes with radiological examination of the stomach and/or small intestine, a single dose of Emestop injection (undiluted) may be administered slowly by the intravenous route over a 1-2 minute period. The recommended single dose of Emestop injection is 10 mg (adults), 2.5-5 mg (children 6-14 years of age) and 0.1 mg/kg body weight (children under 6 years of age).

Over Dosage

Manifestations

Symptoms of over dosage may include drowsiness, disorientation and extrapyramidal reactions.

Treatment

Anticholinergic drugs, antiparkinsonism drugs or antihistamine drugs with anticholinergic properties may be useful in controlling the extrapyramidal reactions.

Symptoms are generally self-limiting and usually disappear within 24 hours.

Pharmaceutical Precautions

Metoclopramide preparations are photosensitive and, therefore, should be stored protected from light. Metoclopramide injection is compatible for dilution with 5% Dextrose, normal saline, Ringer's Injection and Lactated Ringer's Injection. Diluted solutions of Metoclopramide can be maintained in a cool dark place, but not longer than 48 hours.

Presentation

Emestop Injection

Box of 5 ampoules.

Emestop Syrup

Bottle of 30 ml.

Emestop Tablets

Box of 20 tablets.