XYLOCAINE® 2% VISCOUS

NAME OF THE DRUG

Xylocaine 2% Viscous contains lignocaine hydrochloride as the active ingredient. Australian Approved Name: Lignocaine hydrochloride

DESCRIPTION

Xylocaine 2% Viscous is an aqueous topical anaesthetic for use on mucous membranes.

Each mL of Xylocaine Viscous contains lignocaine hydrochloride 21.2 mg (equivalent to lignocaine hydrochloride anhydrous 20 mg), methyl hydroxybenzoate, propyl hydroxybenzoate, sodium hydroxide, saccharin sodium, cherry extract, carmellose sodium and water for injection. The cherry extract contains water, flavour, citric acid and amaranth.

PHARMACOLOGY

Lignocaine, the active ingredient of Xylocaine Viscous, stabilises the neuronal membrane and prevents the initiation and conduction of nerve impulses, thereby effecting local anaesthetic action.

The onset of action of Xylocaine Viscous occurs within 3-5 minutes on mucous membranes. Its low surface tension ensures an even film over the surface of the mucous membrane so that the Xylocaine comes into intimate contact with the total surface. High viscosity ensures sufficiently prolonged contact with the mucous membrane. It is ineffective when applied to intact skin.

Lignocaine may be absorbed following topical administration to mucous membranes, its rate of absorption and amount of dose absorbed depending upon concentration and total dose administered, the specific site of application and duration of exposure. In general, the rate of absorption occurs most rapidly after intratracheal administration.

Lignocaine is well absorbed from the gastrointestinal tract, but little intact drug appears in the circulation because of bio transformation in the liver. Lignocaine is metabolised rapidly by the liver, and metabolites and unchanged drug are excreted by the kidney.

Excessive blood levels may cause changes in cardiac output, total peripheral resistance and mean arterial pressure. These changes may be attributable to a direct depressant effect of the anaesthetic agent on various components of the cardiovascular system.
Biotransformation includes oxidative N-dealkylation, ring hydroxylation, cleavage of the amide linkage and conjugation. The pharmacological / toxicological actions of the metabolites are similar to, but not less potent than, those of lignocaine. Approximately 90% of lignocaine is excreted in the form of various metabolites, and less than 10% is excreted unchanged. The primary metabolite in urine is a conjugate of 4-hydroxy-2,6-dimethylaniline.

The plasma binding of lignocaine is dependent on drug concentration, and the fraction bound decreases with increasing concentration. At concentrations of 1 to 4 μg of free base/mL, 60 to 80% of lignocaine is protein bound. Binding is also dependent on the plasma concentrations of the alpha-1-acid glycoprotein.

Lignocaine crosses the blood-brain and placental barriers, presumably by passive diffusion.

Studies of lignocaine metabolism following iv bolus injection have shown that the elimination half-life is usually 1.5 to 2 hours. The half-life may be prolonged 2-fold or more in patients with liver dysfunction. Renal dysfunction does not affect lignocaine kinetics, but may increase the accumulation of metabolites.

Factors such as acidosis and the use of CNS stimulants and depressants affect the CNS levels of lignocaine required to produce overt systemic effects. Objective adverse manifestations become increasingly apparent with increasing venous plasma levels above 6.0 μg free base/mL. In the rhesus monkey arterial blood levels of 18 to 21 μg/mL have been shown to be the threshold for convulsive activity.

INDICATIONS

Xylocaine 2% Viscous Solution is indicated for the relief of pain and discomfort associated with:

- irritated or inflamed mucous membranes of the mouth, pharynx and upper gastrointestinal tract, e.g. post-tonsillectomy sore throat, dumping syndrome;
- introduction of instruments and catheters into the respiratory and gastrointestinal tract.

CONTRAINDICATIONS

Known history of hypersensitivity to lignocaine or other local anaesthetics of the amide type or to other components of the viscous solution.

Hypersensitivity to methyl and/or propyl hydroxybenzoate or to their metabolite para aminobenzoic acid.
PRECAUTIONS

Warning:
Excessive dosage, or short intervals between doses, can result in high levels of lignocaine or its metabolites and serious adverse effects. Patients should be instructed to strictly adhere to the recommended dosage and administration guidelines. The management of serious adverse reactions may require the use of resuscitative equipment, oxygen and other resuscitative drugs. Patients should not exceed the recommended dose or use Xylocaine 2% Viscous for prolonged periods except on the advice of their physician.

The lowest dose that results in effective anaesthesia should be used to avoid high plasma levels and serious adverse effects. Tolerance to elevated blood levels varies with the status of the patient.

Excessive dosage or short intervals between doses may result in high plasma levels and serious adverse effects. Following too high or repeated doses of viscous lignocaine in children under the age of three, serious side effects have been reported. Patients should be instructed to adhere strictly to the recommended dosage. This is especially important in children where the doses vary with weight.

Dosage reduction
Debilitated, elderly patients or patients with partial or complete heart block and/or acutely ill patients, and children should be given reduced doses commensurate with their age and physical status.

Excessive absorption
Absorption from wound surfaces and mucous membranes is relatively high, especially in the bronchial tree. Because of the possibility of significant systemic absorption, Xylocaine Viscous should be used with caution in patients with traumatised mucosa and/or sepsis in the region of the proposed application.

If the dose or site of administration is likely to result in high blood levels, lignocaine, in common with other local anaesthetics, should be used with caution in patients with epilepsy, impaired cardiac conduction, bradycardia, impaired hepatic function, in severe shock, the elderly, patients in poor general health and patients with severe renal dysfunction.

Patients treated with anti-arrhythmic drugs class III (eg, amiodarone) should be kept under close surveillance and ECG monitoring considered, since cardiac effects may be additive.

Eating and drinking
The use of topical anaesthetic agents in the oral cavity may interfere with swallowing and thus enhance the danger of aspiration of food or drink. For this reason, food or drink should not be ingested within 60 minutes of using local anaesthetics in the mouth or throat area. Numbness of the tongue or buccal
mucosa may increase the danger of biting or heat trauma. Food, chewing gum or hot drinks should not be taken while the mouth or throat area is anaesthetised.

**Malignant hyperthermia**

Many drugs used during the conduct of anaesthesia are considered potential triggering agents for familial malignant hyperthermia. It has been shown that the use of amide local anaesthetics in malignant hypothermia patients is generally safe, but cases of malignant hyperthermia have occasionally documented after use.

**Endotracheal tube lubrication**

When used for endotracheal tube lubrication care should be taken to avoid introduction of the viscous solution into the lumen of the tube. The solution may dry on the inner surface leaving residue which tend to clump with flexion, narrowing the lumen. There have been rare reports in which this residue has caused the lumen to occlude.

**Porphyric patients**

Xylocaine 2% Viscous is probably porphyrinogenic and should only be used on patients with acute porphyria where there are strong or urgent indications. Appropriate precautions should be taken for all porphyric patients.

* **Carcinogenic and Mutagenic Potential**

  Genotoxicity tests with lignocaine are inconclusive. In genotoxicity studies, a metabolite of lignocaine, 2,6-xylidine, showed evidence of activity in some tests but not in other tests. This metabolite has been shown to have carcinogenic potential (nasal and subcutaneous tumours) in preclinical toxicological studies evaluating chronic exposure.

**Use in pregnancy - Category A**

Lignocaine crosses the placental barrier and may be taken up by foetal tissues. When used for surface anaesthesia, lignocaine blood levels after normal doses are low so little drug is available for placental transfer.

There are, however, no adequate and well-controlled studies in pregnant women. Reproduction studies have been performed in rats at doses of 500 mg/kg/day and have revealed no evidence of harm to the foetus caused by lignocaine.

It is reasonable to assume that a large number of pregnant women and women of child-bearing age have used lignocaine. No specific disturbances to the reproduction process have so far been reported.

**Labour and delivery**

Lignocaine is not contraindicated in labour and delivery.

**Using in lactation**

Lignocaine enters the breast milk, but in such small quantities that there is generally no risk of affecting the child at therapeutic dose levels.
Effects on ability to drive and operate machines

Depending on the dose, local anaesthetics may have a very mild effect on mental function and may temporarily impair locomotion and co-ordination.

Interactions

Antiarrhythmic drugs

Lignocaine should be used with caution in patients receiving other local anaesthetics or agents structurally related to amide-type local anaesthetics, e.g. antiarrhythmic drugs such as mexiletine, since the toxic effects are additive.

Specific interaction studies with lidocaine and anti-arrhythmic drugs class III (e.g. amiodarone) have not been performed, but caution is advised.

Enzyme inducing drugs

Cimetidine or betablockers have been shown to cause potentially toxic plasma concentrations when lignocaine is given in repeated high doses over a long period of time. Therefore, caution should be taken if lignocaine was administered at higher than recommended doses over extended period of time.

Phenytoin and other antiepileptic drugs such as phenobarbitone, primidone and carbamazepine appear to enhance the metabolism of lignocaine but the significance of this effect is not known. Phenytoin and lignocaine have additive cardiac depressant effects.

ADVERSE REACTIONS

Systemic adverse reactions are rare and may result from high plasma levels caused by excessive dosage or rapid absorption, or may result from a hypersensitivity, idiosyncrasy or reduced tolerance on the part of the patient. Such reactions are systemic in nature and involve the central nervous system and/or the cardiovascular system.

Central Nervous System

CNS reactions are excitatory and/or depressant and may be characterised by lightheadedness, nervousness, apprehension, euphoria, confusion, dizziness, drowsiness, tinnitus, blurred vision, vomiting, sensations of heat, cold or numbness, twitching, tremors, convulsions, unconsciousness and possibly respiratory arrest. The excitatory reactions may be brief or may not occur at all, in which case the first manifestations of toxicity may be drowsiness, progressing to unconsciousness and respiratory arrest.

Drowsiness following administration of lignocaine is usually an early sign of a high blood level of the drug and may occur as a result of rapid absorption.

Cardiovascular

Cardiovascular reactions are usually depressant and may be characterised by hypotension, myocardial depression, bradycardia and possibly cardiac arrest.
Allergic reactions

Allergic reactions may occur as a result of sensitivity either to the local anaesthetic agent or to other ingredients in the formulation. Allergic reactions as a result of sensitivity to lignocaine are rare (<0.1%). The detection of sensitivity by skin testing is of doubtful value.

The extremely rare cases of allergy to local anaesthetic preparations have included bronchospasm, chest pain, dyspnoea, pruritus, rash, rhinitis, increased sweating, urticaria, sleepiness, dizziness, paraesthesia, oedema, and in the most severe instances anaphylactic shock. Several cases of contact dermatitis have been reported with the use of lignocaine.

DOSAGE AND ADMINISTRATION

As with any local anaesthetic, reactions and complications are best averted by employing the minimal effective dosage. Debilitated, acutely ill or elderly patients and children should be given doses commensurate with their age and physical condition.

Shake the bottle well before use.

For symptomatic treatment of irritated or inflamed mucous membranes of the mouth and pharynx, the usual adult dose is 15 mL undiluted. For use in the mouth, the solution should be swished around the mouth for approximately 30 seconds and spat out. For use on the pharynx, the solution should be gargled and may be swallowed. This dose should not be administered at intervals of less than three hours.

Maximum dosage

Although the incidence of adverse effects with lignocaine is quite low, caution should be exercised when using large amounts.

Adults: No more than 15 mL (300 mg lignocaine HCl) every 3 hours or 120 mL in a 24 hour period.

Children over 3 years of age: No more than 4.5 mg/kg of bodyweight or 5 mL (100 mg lignocaine HCl) every 3 hours or 40 mL in a 24 hour period.

Children under 3 years of age: No more than 1.25 mL applied with a cotton swab every 3 hours or 10 mL in a 24 hour period.

OVERDOSAGE

In the event of an overdose, contact the Poisons Information Centre on 13 11 26.
Management of Local Anaesthetic Emergencies

The first consideration is prevention, best accomplished by careful and constant monitoring of cardiovascular and respiratory vital signs and the patient's state of consciousness after each local anaesthetic administration. At the first sign of change, oxygen should be administered.

Treatment

If convulsions occur then immediate attention is required for the maintenance of a patent airway and assisted or controlled ventilation with oxygen, via a positive airway pressure delivery system mask. Adequacy of the circulation should then be evaluated, bearing in mind that drugs used to treat convulsions depress the circulation when administered iv.

Should convulsions persist despite adequate respiratory support, and if the status of the circulation permits, appropriate anticonvulsant medication such as an ultra-short acting barbiturate (eg. thiopental) or a benzodiazepine (eg. diazepam) may be administered iv. The clinician should be familiar with these anticonvulsant drugs prior to use of local anaesthetics.

Dialysis is of negligible value in the treatment of acute overdosage with lignocaine.

PRESENTATION

Xylocaine 2% Viscous Solution, 200 mL bottle.

STORAGE

Store below 25°C.

NAME AND ADDRESS OF MANUFACTURER

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* Please note changes in Product Information.

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