

HYDROXYZINE HYDROCHLORIDE- hydroxyzine hydrochloride solution
ATLANTIC BIOLOGICALS CORP.

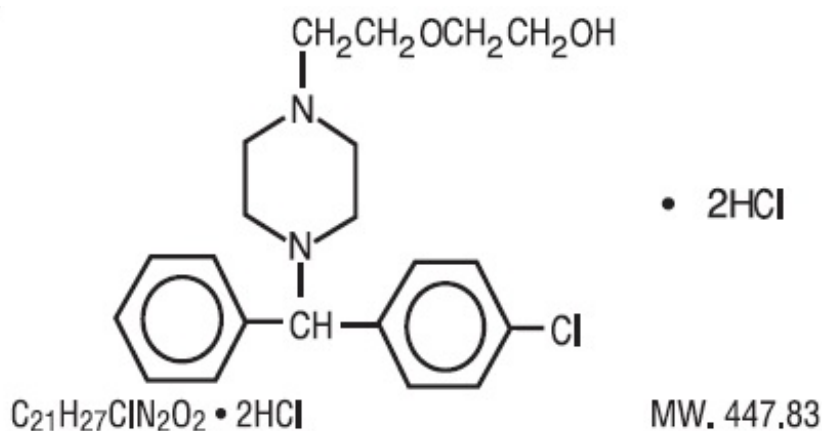
HydroXYzine HYDROCHLORIDE SYRUP
(HydroXYzine Hydrochloride Oral Solution, USP)

10 mg per 5 mL

Rx only

DESCRIPTION

Hydroxyzine hydrochloride is designated chemically as (\pm)-2-[2-[4-(*p*-Chloro- α -phenylbenzyl)-1-piperaziny]ethoxy]ethanol dihydrochloride



Inert ingredients: citric acid, peppermint flavor, sodium benzoate, sucrose and water. Citric acid and/or sodium citrate may be used to adjust the pH when necessary.

CLINICAL PHARMACOLOGY

Hydroxyzine hydrochloride is unrelated chemically to the phenothiazines, reserpine, meprobamate, or the benzodiazepines.

Hydroxyzine hydrochloride is not a cortical depressant, but its action may be due to suppression of activity in certain key regions of the subcortical area of the central nervous system. Primary skeletal muscle relaxation has been demonstrated experimentally. Bronchodilator activity, and antihistaminic and analgesic effects have been demonstrated experimentally and confirmed clinically. An antiemetic effect, both by the apomorphine test and the veriloid test, has been demonstrated. Pharmacological and clinical studies indicate that hydroxyzine in therapeutic dosage does not increase gastric secretion or acidity and in most cases has mild antisecretory activity. Hydroxyzine is rapidly absorbed from the gastrointestinal tract and its clinical effects are usually noted within 15 to 30 minutes after oral administration.

INDICATIONS AND USAGE

For symptomatic relief of anxiety and tension associated with psychoneurosis and as an adjunct in organic disease states in which anxiety is manifested.

Useful in the management of pruritus due to allergic conditions such as chronic urticaria and atopic and contact dermatoses, and in histamine-mediated pruritus.

As a sedative when used in premedication and following general anesthesia, **Hydroxyzine may potentiate meperidine**(Demerol[®]) and barbiturates, so their use in pre-anesthetic adjunctive therapy should be modified on an individual basis.

Atropine and other belladonna alkaloids are not affected by the drug. Hydroxyzine is not known to interfere with the action of digitalis in any way and it may be used concurrently with this agent.

The effectiveness of hydroxyzine as an antianxiety agent for long term use, that is more than 4 months, has not been assessed by systematic clinical studies. The physician should reassess periodically the usefulness of the drug for the individual patient.

CONTRAINDICATIONS

Oral hydroxyzine hydrochloride is contraindicated in patients with known hypersensitivity to hydroxyzine hydrochloride products, and in patients with known hypersensitivity to cetirizine hydrochloride or levocetirizine hydrochloride.

Hydroxyzine is contraindicated in patients with a prolonged QT interval.

Hydroxyzine, when administered to the pregnant mouse, rat and rabbit, induced fetal abnormalities in the rat and mouse at doses substantially above the human therapeutic range. Clinical data in human beings are inadequate to establish safety in early pregnancy. Until such data are available, hydroxyzine is contraindicated in early pregnancy.

Hydroxyzine is contraindicated for patients who have shown a previous hypersensitivity to any component of this medication.

WARNINGS

Nursing Mothers: It is not known whether this drug is excreted in human milk. Since many drugs are so excreted, hydroxyzine should not be given to nursing mothers.

PRECAUTIONS

THE POTENTIATING ACTION OF HYDROXYZINE MUST BE CONSIDERED WHEN THE DRUG IS USED IN CONJUNCTION WITH CENTRAL NERVOUS SYSTEM DEPRESSANTS SUCH AS NARCOTICS, NON-NARCOTIC ANALGESICS AND BARBITURATES. Therefore, when central nervous system depressants are administered concomitantly with hydroxyzine their dosage should be reduced. **QT Prolongation/Torsade de Pointes (TdP):** Cases of QT prolongation and Torsade de Pointes have been reported during post-marketing use of hydroxyzine. The majority of reports occurred in patients with other risk factors for QT prolongation/TdP (pre-existing heart disease, electrolyte imbalances or concomitant arrhythmogenic drug use). Therefore, hydroxyzine should be used with caution in patients with risk factors for QT prolongation, congenital long QT syndrome, a family history of long QT syndrome, other conditions that predispose to QT prolongation and ventricular arrhythmia, as well as recent myocardial infarction, uncompensated heart failure, and bradyarrhythmias.

Caution is recommended during the concomitant use of drugs known to prolong the QT interval. These include Class 1A (e.g., quinidine, procainamide) or Class III (e.g., amiodarone, sotalol) antiarrhythmics, certain antipsychotics (e.g., ziprasidone, iloperidone, clozapine, quetiapine, chlorpromazine), certain antidepressants (e.g., citalopram, fluoxetine), certain antibiotics (e.g., azithromycin, erythromycin, clarithromycin, gatifloxacin, moxifloxacin); and others (e.g., pentamidine, methadone, ondansetron, droperidol).

Since drowsiness may occur with use of this drug, patients should be warned of this possibility and cautioned against driving a car or operating dangerous machinery while taking hydroxyzine. Patients should be advised against the simultaneous use of other CNS depressant drugs, and cautioned that the effect of alcohol may be increased.

Geriatric Use: A determination has not been made whether controlled clinical studies of hydroxyzine hydrochloride included sufficient numbers of subjects aged 65 and over to define a difference in response from younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal or cardiac function, and of concomitant disease or other drug therapy.

The extent of renal excretion of hydroxyzine hydrochloride has not been determined. Because elderly patients are more likely to have decreased renal function, care should be taken in dose selections.

Sedating drugs may cause confusion and over sedation in the elderly; elderly patients generally should be started on low doses of hydroxyzine hydrochloride and observed closely.

Acute Generalized Exanthematous Pustulosis (AGEP): Hydroxyzine may rarely cause acute generalized exanthematous pustulosis (AGEP), a serious skin reaction characterized by fever and numerous small, superficial, non-follicular, sterile pustules, arising within large areas of edematous erythema. Inform patients about the signs of AGEP, and discontinue hydroxyzine at the first appearance of a skin rash, worsening of pre-existing skin reactions which hydroxyzine may be used to treat, or any other sign of hypersensitivity. If signs or symptoms suggest AGEP, use of hydroxyzine should not be resumed and alternative therapy should be considered. Avoid cetirizine or levocetirizine in patients who have experienced AGEP or other hypersensitivity reactions with hydroxyzine, due to the risk of cross-sensitivity.

ADVERSE REACTIONS

Side effects reported with the administration of hydroxyzine hydrochloride are usually mild and transitory in nature.

Skin and appendages: Oral hydroxyzine hydrochloride is associated with Acute Generalized Exanthematous Pustulosis (AGEP) and fixed drug eruptions in post marketing reports.

Anticholinergic: Dry mouth.

Central Nervous System: Drowsiness is usually transitory and may disappear in a few

days of continued therapy or upon reduction of the dose. Involuntary motor activity including rare instances of tremor and convulsions have been reported, usually with doses considerably higher than those recommended. Clinically significant respiratory depression has not been reported at recommended doses.

Cardiac System:QT prolongation, Torsade de Pointes.

Skin and appendages:Pruritis, rash, urticaria

OVERDOSAGE

The most common manifestation of overdose of hydroxyzine hydrochloride is hypersedation. Other reported signs and symptoms were convulsions, stupor, nausea and vomiting. As in the management of overdose with any drug, it should be borne in mind that multiple agents may have been taken. If vomiting has not occurred spontaneously, it should be induced. Immediate gastric lavage is also recommended. General supportive care, including frequent monitoring of the vital signs and close observation of the patient, is indicated. Hypotension, though unlikely, may be controlled with intravenous fluids and vasopressors (**do not use epinephrine as hydroxyzine counteracts its pressor action**). Caffeine and Sodium Benzoate Injection, USP, may be used to counteract central nervous system depressant effects.

There is no specific antidote. It is doubtful that hemodialysis would be of any value in the treatment of overdose with hydroxyzine. However, if other agents such as barbiturates have been ingested concomitantly, hemodialysis may be indicated. There is no practical method to quantitate hydroxyzine in body fluids or tissue after its ingestion or administration.

Hydroxyzine overdose may cause QT prolongation and Torsade de Pointes. ECG monitoring is recommended in cases of hydroxyzine overdose.

DOSAGE AND ADMINISTRATION

For symptomatic relief of anxiety and tension associated with psychoneurosis and as an adjunct in organic disease states in which anxiety is manifested: in adults, 50 to 100 mg four times daily.; children under 6 years, 50 mg daily in divided doses: children over 6 years: 50 to 100 mg daily in divided doses.

For use in the management of pruritus due to allergic conditions such as chronic urticaria and atopic and contact dermatoses, and in histamine-mediated pruritus: in adults, 25 mg three times to four times daily.; children under 6 years, 50 mg daily in divided doses and children over 6 years, 50 to 100 mg daily in divided doses.

As a sedative when used as a premedication and following general anesthesia: 50 to 100 mg in adults, and 0.6 mg/kg in children.

When treatment is initiated by the intramuscular route of administration, subsequent doses may be administered orally.

As with all medications, the dosage should be adjusted according to the patient's response to therapy.

HOW SUPPLIED

Hydroxyzine Hydrochloride Syrup (Hydroxyzine Hydrochloride Oral Solution, USP) 10 mg per 5 mL (teaspoonful) is a slightly yellow peppermint flavored liquid, supplied in 473 mL (16 fl. oz.) amber plastic bottles.

NDC 17856-0507-01 HYDROXYZINE HCL 10MG/5ML - 5 ML CUP 72 ct UD

Store at controlled room temperature 15° to 30°C (59° to 86°F). [See USP controlled room temperature].

Dispense in a tight, light-resistant container with child-resistant closure as defined in the USP.

Distributed by:

ATLANTIC BIOLOGICALS CORP.

MIAMI, FL 33179

17856-0507-01
HYDROXYZINE
HYDROCHLORIDE
ORAL SYRUP
10 MG / 5 ML
DELIVERS 5 ML



See package insert for indications and dosage schedule

Store at controlled room temperature 15°-30°C
(59°-86°F) [See USP controlled room
Temperature].
***Keep this and all Medication out of the reach
of children***



17856050701

17856-0507-01

Dosage: 10 MG/ 5ML

HYDROXYZINE HCL

Qty: 72 CUPS



GTIN: 00117856050711

S/N: 01371101

Exp: 10/20/21

Lot: 013711

RX

Packaged by: Unit Dose Solutions
Morrisville, NC 27590

Distributed by: Atlantic Biologicals Corp.
Miami FL 33179

Rev. 09/19

Call to Reorder: 800.509.7592

HYDROXYZINE HYDROCHLORIDE

hydroxyzine hydrochloride solution

Product Information

Product Type

HUMAN PRESCRIPTION
DRUG

**Item Code
(Source)**

NDC:17856-0507(NDC:54838-
502)

Route of Administration	ORAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	HYDROXYZINE DIHYDROCHLORIDE (UNII: 76755771U3) (HYDROXYZINE - UNII:30S50YM8OG)	HYDROXYZINE DIHYDROCHLORIDE	10 mg in 5 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)			
	SODIUM BENZOATE (UNII: OJ245FE5EU)			
	SUCROSE (UNII: C151H8M554)			
	WATER (UNII: 059QF0KO0R)			
	SODIUM CITRATE (UNII: 1Q73Q2JULR)			
Product Characteristics				
Color	yellow (slightly yellow)	Score		
Shape		Size		
Flavor	PEPPERMINT (peppermint flavor)	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17856-0507-1	72 in 1 BOX, UNIT-DOSE	05/08/2024	
1	NDC:17856-0507-2	5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA201674	08/21/2013		

Labeler - ATLANTIC BIOLOGICALS CORP. (047437707)

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