ISOXSUPRINE HYDROCHLORIDE - isoxsuprine hydrochloride tablet
Vista Pharmaceuticals, Inc.

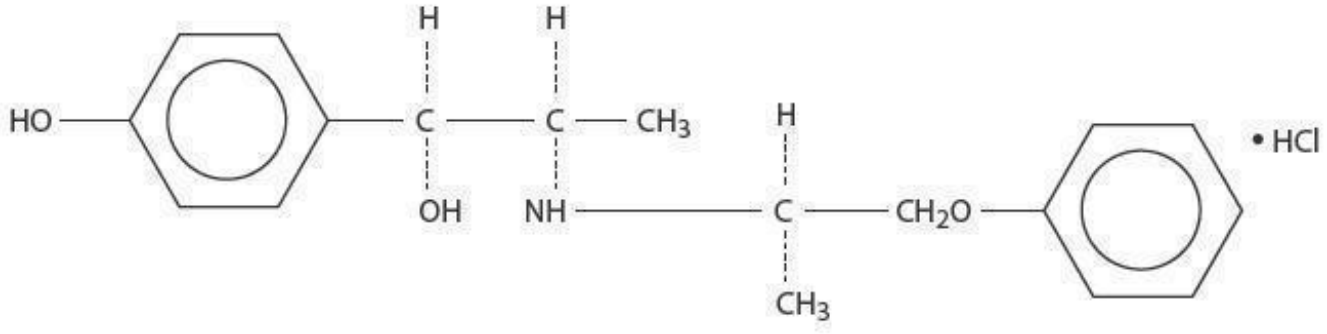
Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

ISOXSUPRINE HYDROCHLORIDE TABLETS, USP

CAUTION: Federal Law prohibits dispensing without prescription

DESCRIPTION

lsoxsuprine HCI occurs as a white odorless, crystalline powder, having a bitter taste, It has a following structural formula



INDICATIONS

Based on a review of this drug by the National Academy of Sciences -National Research Council and / or other information, the FDA has classified the medications as follows :

Possibly Effective :
1. For the relief of symptoms associated with cerebral vascular insufficiency
2. In peripheral vascular disease of arteriosclerosis obliterans, thromboangitis obliterans (Buerger's Disease) and Raynaud's disease.

Final classification of the less than - effective indications requires further investigation.

COMPOSITION

Each tablet contains lsoxsuprine HCI 20 mg.

CONTRAINDICATIONS AND CAUTIONS

There are no known contraindications to oral use when administered in recommended doses. Should not be given immediately postpartum or in the presence of arterial bleeding.

ADVERSE REACTIONS

On rare occasions oral administration of the drug has been associated in time with the occurrences of hypotension, tachycardia, nausea, vomiting, dizziness, abdominal distress, and severe rash. If rash appears the drug should be discontinued.
Although available evidence suggests a temporal association of these reactions with Isoxsuprine, a casual relationship can be neither confirmed nor refused.

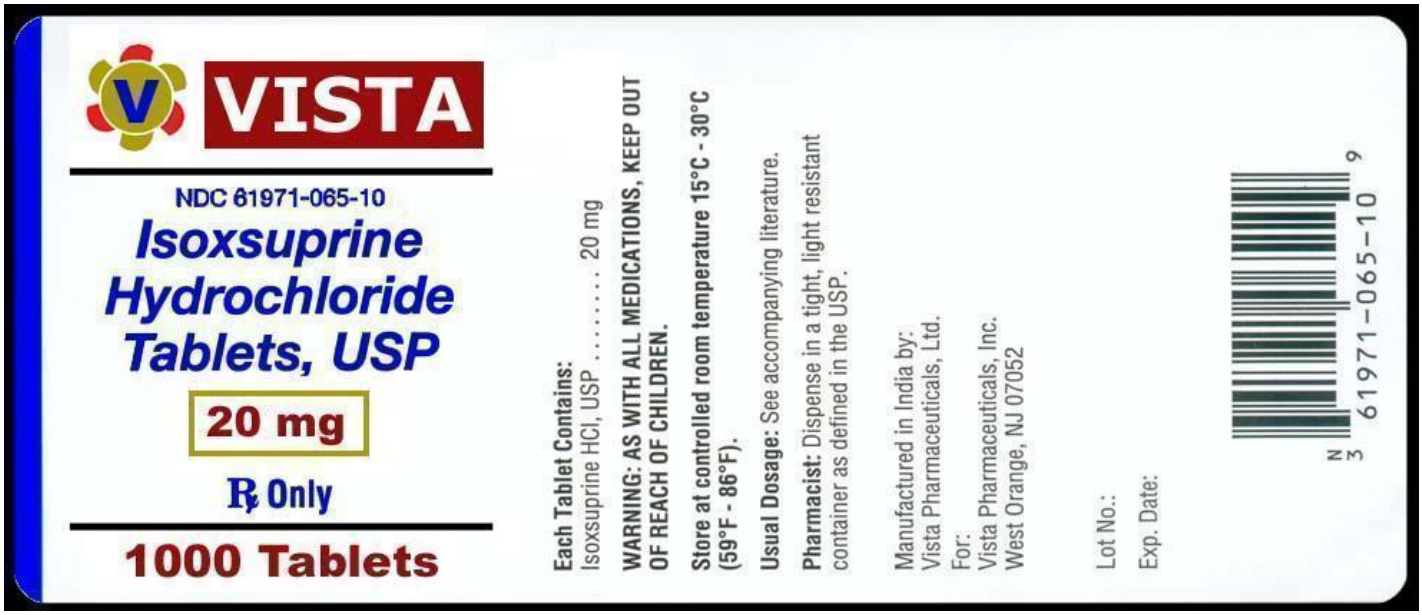
DOSAGE AND ADMINISTRATION

10 to 20mg three or four times daily

HOW SUPPLIED

lsoxsuprine HCI 20mg tablets are supplied in HDPE containers of 1,000's

Manufactured in India by
Vista Pharmaceuticals, Limited.



ISOXSUPRINE HYDROCHLORIDE

isoxsuprine hydrochloride tablet

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG LABEL	**Item Code (Source)**	NDC:61971-065
Route of Administration	ORAL	**DEA Schedule**	

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Isoxsuprine hydrochloride (Isoxsuprine)	Isoxsuprine hydrochloride	20 mg

Inactive Ingredients	
Ingredient Name	**Strength**
LACTOSE	
CELLULOSE, MICROCRYSTALLINE	

CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS	
POVIDONE K30	
SODIUM STARCH GLYCOLATE TYPE A POTATO	
MAGNESIUM STEARATE	

Product Characteristics

Color	white	**Score**	2 pieces
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	20;VISTA065
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61971-065-10	1000 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/19/1997	

Labeler - Vista Pharmaceuticals, Inc. (943932806)

Establishment

Name	Address	ID/FEI	Business Operations
Vista Pharmaceuticals, Limited.		916648541	manufacture(61971-065), analysis(61971-065)

Revised: 12/2012 Vista Pharmaceuticals, Inc.