DOPAMINE- dopamine hydrochloride liquid
Deseret Biologicals, Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

DRUG FACTS:

ACTIVE INGREDIENTS:

Dopamine Hydrochloride 8X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

PURPOSE:

Dopamine Hydrochloride - Headache, Reading Difficulty, Poor Memory

USES:

• For the temporary relief of symptoms including:

• headache • reading difficulty • poor memory

These statements are based upon homeopathic principles. They have not been reviewed by the Food and Drug Administration.

WARNINGS:

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center right away.

Tamper Evident: Sealed for Your Protection. Do not use if seal is broken or missing.

KEEP OUT OF REACH OF CHILDREN:

In case of overdose, contact a physician or Poison Control Center right away.

DIRECTIONS:

1-10 drops under the tongue, 3 times a day or as directed by a health professional. Consult a physician for use in children under 12 years of age.

INACTIVE INGREDIENTS:

Demineralized water, 25% ethanol

QUESTIONS:

Dist. By: Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070

www.desbio.com

800-827-8204

PACKAGE LABEL DISPLAY:

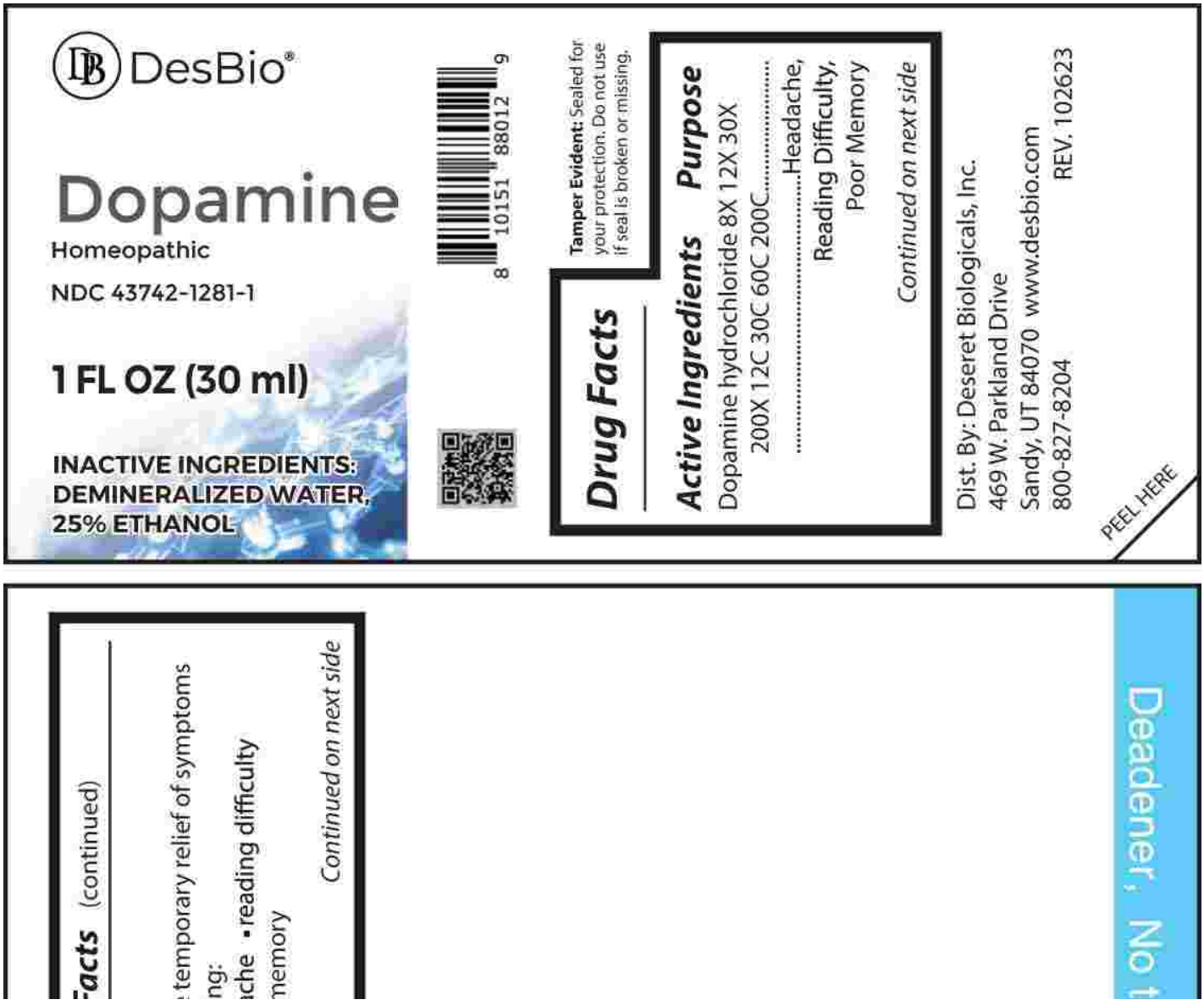
DesBio

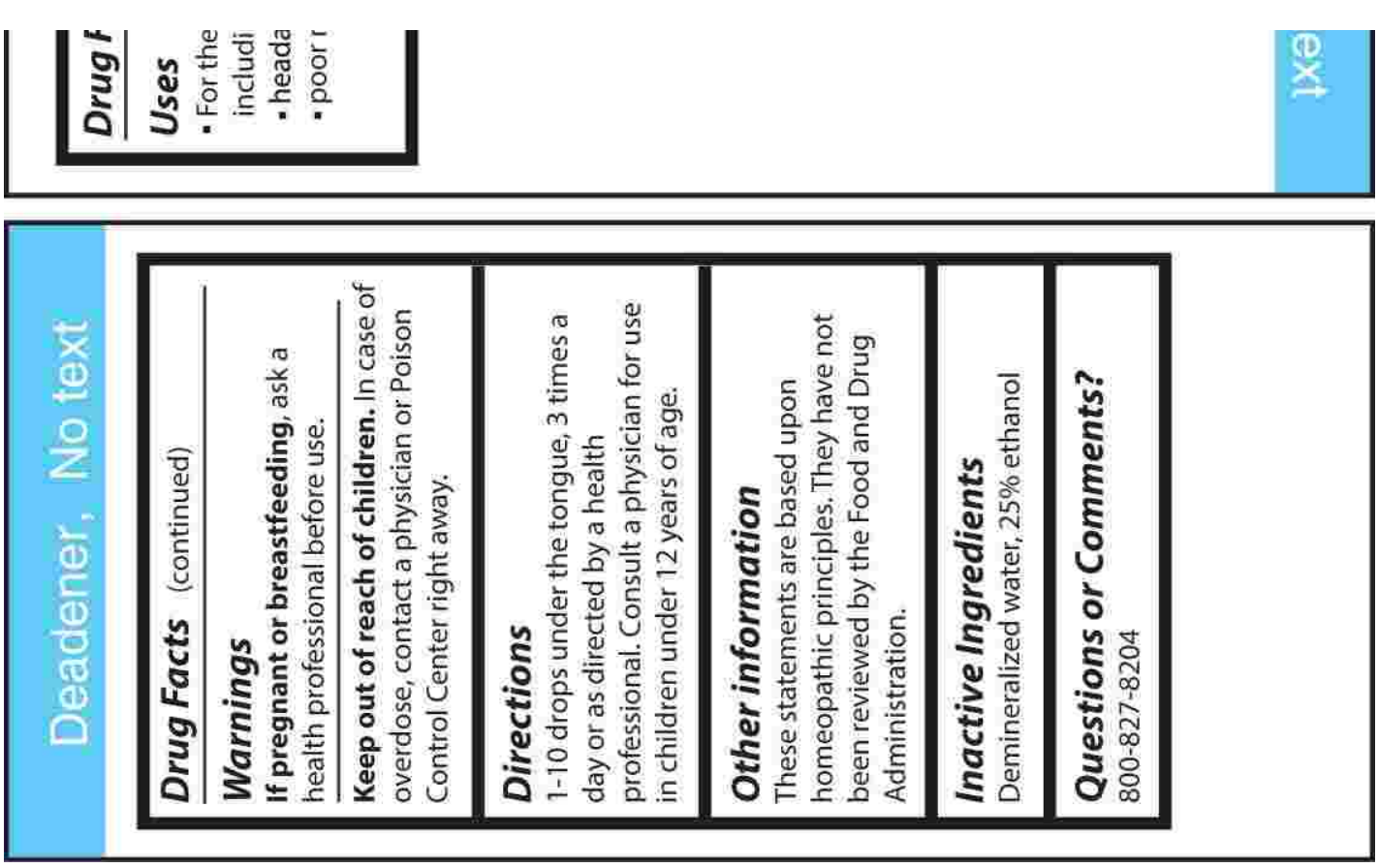
Dopamine

Homeopathic

NDC 43742-1281-1

1 FL OZ (30 ml)





DOPAMINE

dopamine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43742-1281
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DOPAMINE HYDROCHLORIDE (UNII: 7L3E358N9L) (DOPAMINE - UNII:VTD58H1Z2X)	DOPAMINE HYDROCHLORIDE	8 [hp_X] in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:43742-1281-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/27/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/27/2018	

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company, Inc. (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company, Inc.		844330915	manufacture(43742-1281) , api manufacture(43742-1281) , label(43742-1281) , pack(43742-1281)

Revised: 2/2024 Deseret Biologicals, Inc.