SODIUM CHLORIDE 1 GRAM- sodium chloride tablet Gendose Pharmaceuticals, LLC

GENDOSE - SODIUM CHLORIDE 1 GRAM TABLETS, USP (77333-835)

ACTIVE INGREDIENT (IN EACH TABLET)

SODIUM CHLORIDE, USP 1 GRAM

PURPOSE

ELECTROLYTE REPLENISHER

USES

- FOR TTHE PREPARATION OF NORMAL ISOTONIC SOLUTION OF SODIUM CHLORIDE.
- AS AN ELECTROLYTE REPLENISHER FOR THE PREVENTION OF HEAT CRAMPS DUE TO EXCESSIVE PERSPIRATION.
- ANY ALTERNATIVE USE AS DIRECTED BY A PHYSICIAN.

WARNINGS

- DO NOT USE WITHOUT CONSULTING A PHYSICIAN.
- ASK A PHYSICIAN BEFORE USE IF YOU HAVE A SODIUM RESTRICTED DIET DUE TO MULTIPLE ORGAN DISEASES.
- STOP USE AND ASK A PHYSICIAN IF SYMPTOMS OF HEAT CRAMPS CONTIUE FOR MORE THAN 24 HOURS.
- IF PREGNANT OR BREAST FEEDING, ASK A PHYSICIAN BEFORE USE.
- KEEP OUT OF REACH OF CHILDREN. IN CASE OF OVERDOSE, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

DIRECTIONS

- TO MAKE AN ISOTONIC SOLUTION OF SODIUM CHLORIDE, DISSOLVE 1 TABLET IN 120 ML (FOUR OUNCES) OF DISTILLED WATER AND USE AS DIRECTED BY A PHYSICIAN.
- IF USED AS AN ELECTROLYTE REPLENISHER FOR THE PREVENTION OF HEAT CRAMPS DUE TO EXCESSIVE PERSPIRATION TAKE ONE TABLET ORALLY AS DIRECTED BY YOUR PHYSICIAN.

INACTIVE INGREDIENTS

NONE

OTHER INFORMATION

- EACH TABLET CONTAINS: SODIUM 394 MG
- STORE AT ROOM TEMPERATURE 15° 30°C (59° 86°F)
- PRODUCT DOES NOT CONTAINS ANY INACTIVE INGREDIENTS.



SODIUM CHLORIDE 1 GRAM

sodium chloride tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77333-835

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
-----------------	--------------------------	----------

SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)

SODIUM CHLORIDE

1 g

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	13mm
Flavor		Imprint Code	G13
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77333-835- 10	100 in 1 BOX	05/15/2020	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M018	05/15/2020			

Labeler - Gendose Pharmaceuticals, LLC (080257510)

Revised: 1/2024

Gendose Pharmaceuticals, LLC