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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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

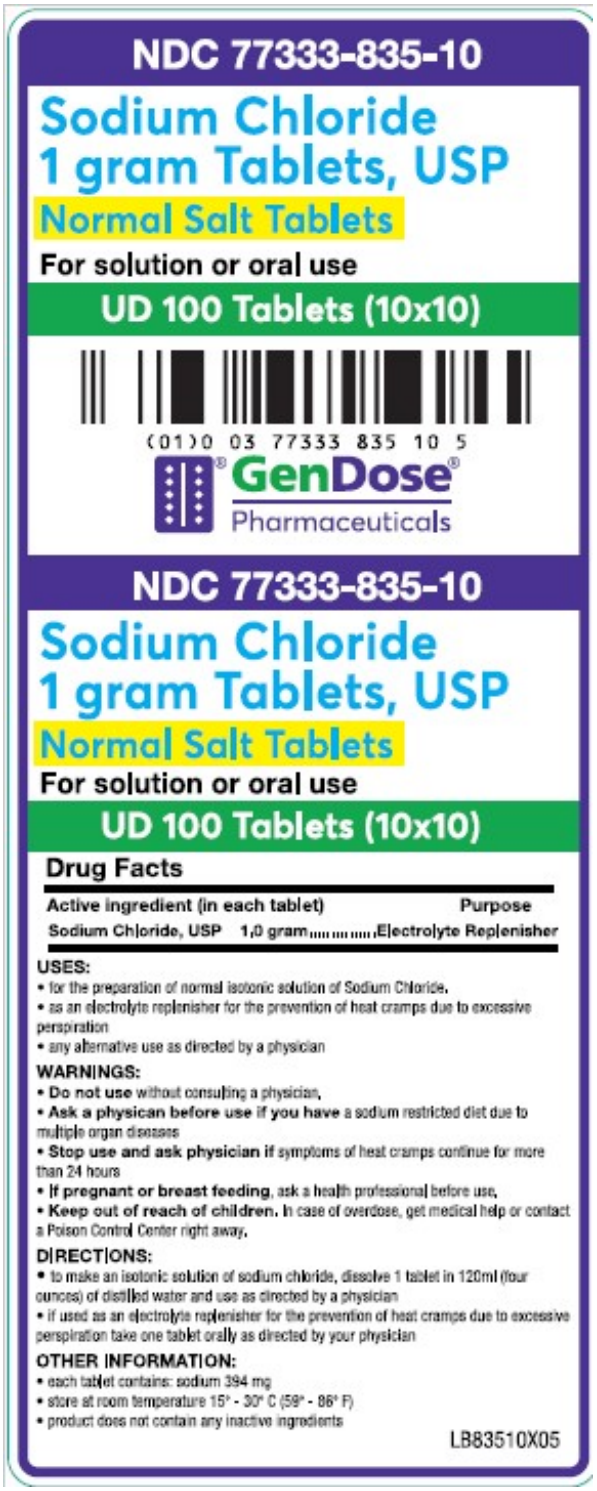
- TO MAKE AN ISOTONIC SOLUTION OF SODIUM CHLORIDE, DISSOLVE 1 TABLET IN 120 ML (FOUR OUNCES) OF DISTILLED WATER AND USE AS DIRECTED BY A PHYSICIAN.
- IF USED AS AN ELECTROLYTE REPLENISHER FOR THE PREVENTION OF HEAT CRAMPS DUE TO EXCESSIVE PERSPIRATION TAKE ONE TABLET ORALLY AS DIRECTED BY YOUR PHYSICIAN.

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
Route of Administration	ORAL		

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	NDC:77333-835-25	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 final	part349	05/15/2020	

Labeler - Gendose Pharmaceuticals, LLC (080257510)

Registrant - Gendose Pharmaceuticals, LLC (080257510)

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