

**SODIUM CHLORIDE NORMAL SALT- sodium chloride tablet**  
**Citragen Pharmaceuticals Inc**

*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.*

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**Sodium Chloride Tablets, USP**

***Drug Facts***

***Active ingredient (in each tablet)***

Sodium Chloride, USP 500 mg

***Purpose***

Electrolyte Replenisher

***Uses***

- for the 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 continue for more than 24 hours.

**If pregnant or breast-feeding,** ask a health professional 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 isotonic solution of sodium chloride, dissolve 2 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 or two tablet orally as directed by your physician.

**Other Information:**

- **each tablet contains:** sodium 197 mg
- store at room temperature 15°-30°C (59°-86°F)
- product does not contain any inactive ingredients

**Questions or comments?**

Phone: +1-510-249-9066 (9AM-5PM PST, Mon-Fri); e-mail: info@citragenpharma.com

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

**Manufactured by:**

CitraGen Pharmaceuticals, Inc.,  
Fremont, CA 94538.

www.citragenpharma.com

Rev. 07/20 R-00

**CitraGen Pharmaceuticals, Inc.**

NDC: 70369-010-09

**Sodium Chloride Tablets, USP 500 mg**

**Normal salt tablets**

**For solution or oral use**

200 tablets



NDC:70369-010-09

# Sodium Chloride Tablets, USP 500 mg

Normal salt tablets  
For solution or oral use

**200 Tablets**

<b>Drug Facts</b>		
<b>Active Ingredient (in each tablet)</b> Sodium Chloride, USP 500 mg.....	<b>Purpose</b> Electrolyte Replenisher	<b>Directions</b> •to make isotonic solution of sodium chloride, dissolve 2 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 or two tablet orally as directed by physician
<b>Uses:</b> •for the 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		<b>Other information:</b> •each tablet contains: sodium 197 mg •store at room temperature 15°-30°C (59° - 86°F) •Product does not contain any inactive ingredients
<b>Warnings</b> Do not use without consulting a physician Ask a physician before use if you have a sodium restricted diet due to multiple organ disease		
Stop use and ask a physician if symptoms of heat cramps continue for more than 24 hours		<b>Questions or comments?</b> Phone: +1-510-249-9066, (9AM-5PM PST, M-F); e-mail: info@citragenpharma.com
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Rev. 07/20



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Lot No. :  
Exp. Date:

<b>SODIUM CHLORIDE NORMAL SALT</b>			
sodium chloride tablet			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70369-010
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	500 mg

**Product Characteristics**

Color	white	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	CG010
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70369-010-09	200 in 1 BOTTLE; Type 0: Not a Combination Product	07/20/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	07/20/2020	

**Labeler** - Citragen Pharmaceuticals Inc (024949457)**Registrant** - Citragen Pharmaceuticals Inc (024949457)**Establishment**

Name	Address	ID/FEI	Business Operations
Citragen Pharmaceuticals Inc		024949457	manufacture(70369-010)

Revised: 12/2022

Citragen Pharmaceuticals Inc