LIQUIDGUARD- phosphates liquid Nano Care Deutschland AG

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.

The liquid guard is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Phosphates 2.5 % v/v. Purpose: Liquid Guard

Liquid Guard

Liquid Guard to help reduce bacteria that potentially can cause disease.

Warnings

For external use only. Keep out of reach of eyes, ears, and mouth

Do not use

Keep out od reach of childrens

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Place enough product to clean the infected area. Do not use of wet surfaces / skin.

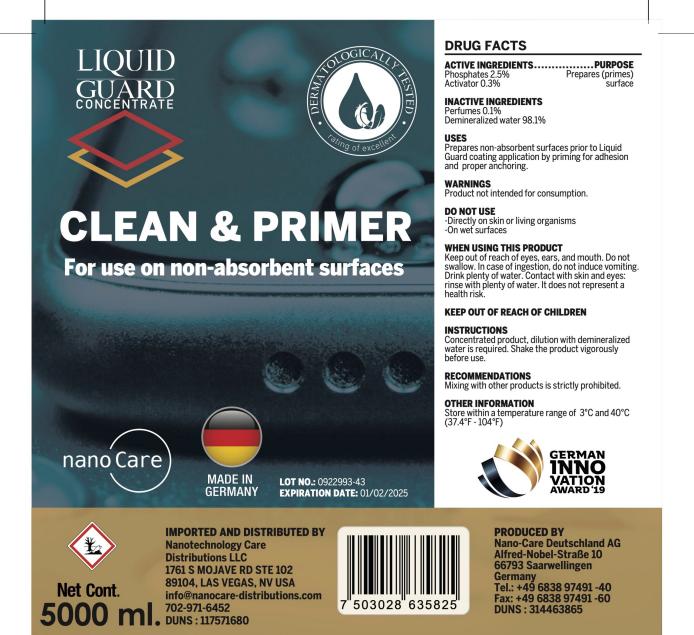
Store between 15-30C (59-86F)

Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

purified water USP

Maximum 1000000000 mL in 1 CONTAINER (79836-500-01)



5000 mL NDC: 79836-500-01

LIQUIDGUARD

phosphates liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79836-500
Route of Administration	TOPICAL		
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Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

PHO SPHATE IO N (UNII: NK08 V8 K8 HR) (PHO SPHATE IO N - UNII: NK08 V8 K8 HR)	PHOSPHATE ION	2.5 g in 100 mL
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Inactive Ingredients

Ingredient Name	Strength
ingredient Name	Strength

WATER (UNII: 059QF0KO0R) 98.1 mL in 100 mL

Packaging

	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:79836-500-	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/30/2020	

Labeler - Nano Care Deutschland AG (314463865)

Registrant - Nano Care Deutschland AG (314463865)

Establishment				
Name	Address	ID/FEI	Business Operations	
Nano Care Deutschland AG		314463865	manufacture(79836-500)	

Revised: 8/2020 Nano Care Deutschland AG