

**ANTISEPTIC SOLUTION- antiseptic solution solution**  
**Grupo Salypro de Mexico, S.A. de C.V.**

*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.*

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**These highlights do not include all the information needed to use VIKUT safely and effectively. See full prescribing information for [www.vikutsolution.com](http://www.vikutsolution.com)**

**WARNING**

For external use on the skin only.  
Severe injury may result from use internally or as a douche. Avoid contact with mucous membranes."

**INDICATIONS AND USAGE**

Astringent, Alkaline, for superficial and deep festering wounds. for 1st, 2nd and 3rd ulcers and burns. grade. Ideal for people with diabetes.Úpply directly without diluting, on the affected area or use an impregnated gauze.

**DOSAGE AND ADMINISTRATION**

It Is Recommended to apply 3 to 5 times a day.

**DOSAGE FORMS AND STRENGTHS**

Antiseptic Solution 40ml

**CONTRAINDICATIONS**

Allergy or hypersensitivity to the active ingredient.

**WARNINGS AND PRECAUTIONS**

There may be moderate burning or pain after the first applications of the solution, which usually disappear with the continuation of the treatment.

**DRUG INTERACTIONS**

Free drug interactions. No patient presented adverse events related to treatment with Vikút®.

## **USE IN SPECIFIC POPULATIONS**

VIKÚT solution is indicated for pre and post-operative antisepsis and the delimitation of the surgical field. Antisepsis of minor and deep wounds, burns (1, 2, and 3rd degree), lacerations, pyoderma, acne and bacterial and fungal infections of the skin, scalp. In wounds where it is desired to accelerate the healing process such as traumatic surgical wounds, episiotomies, varicose ulcers, decubitus ulcers, where in addition to accelerating the healing process it is desired to eliminate or prevent the presence of pathogenic germs that could delay the process healing.

## **LABEL**

81760-310-01 40 mL Solution



One solution, different problems  
**Antiseptic Solution**  
**Astringent and Alkaline**

Health Register: No. 2179C2018 SSA

Disinfects and regenerates:  
wounds, scrapes, uicers and burns

On container of  
40 ml

Formula:  
Each 100 ml contains:  
Potassium Permanganate.....5.12 g  
Alcohol.....60.75 ml  
Benzoic Alcohol.....4.05 g  
Salicylic Acid.....2.02 g  
Vehicle q.s.....100 ml

**"Rx only"**

The formula is patented. It is owned by Grupo Salypro de Mexico, S.A. de C.V.

Route of administration: Topical

Indications: Astringent, Alkaline, for superficial and deep festering wounds.  
For 1st, 2nd and 3rd ulcers and burns, grade. Ideal for people with diabetes.  
Apply directly without diluting, on the affected area or use an impregnated gauze.

IT IS RECOMMENDED TO APPLY 3 TO 5 TIMES A DAY.

Caution: Do not use in people sensitive to potassium permanganate.  
Keep out of reach of children. Store at room temperature in a dry place.  
In case of local irritation or discomfort, discontinue use of the product.  
If symptoms are persistent, consult your doctor.

"Warning - For external use on the skin only.

Severe injury may result from use internally or as a douche.  
Avoid contact with mucous membranes."

Made in México : Aqua Medica, S.A. de C.V. Carretera Federal México-Cuautla  
Km. 65.8 No 8 Tetelcingo, C.P. 62757, Cuautla morelos  
Manufacturer's Licence number (s): 183300507A0125

**ANTISEPTIC SOLUTION**

antiseptic solution solution

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:81760-310
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>POTASSIUM PERMANGANATE</b> (UNII: 00OT1QX5U4) (PERMANGANATE ION - UNII:2BL953CCZ2)	POTASSIUM PERMANGANATE	5.12 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	28.06 g in 100 g
<b>METHYL ALCOHOL</b> (UNII: Y4S76JW15)	60.75 g in 100 g
<b>ACETYLSALICYLSALICYLIC ACID</b> (UNII: VBE72MCP5L)	2.02 g in 100 g
<b>.ALPHA.-(.ALPHA.-AMINOPROPYL)BENZYL ALCOHOL</b> (UNII: S8TT5K3C8Y)	4.05 g in 100 g

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81760-310-01	40 g in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	04/01/2021	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/01/2021	

**Labeler** - Grupo Salypro de Mexico, S.A. de C.V. (951596655)

### Establishment

Name	Address	ID/FEI	Business Operations
Grupo Salypro de Mexico, S.A. de C.V.		951596655	manufacture(81760-310)

Revised: 1/2022

Grupo Salypro de Mexico, S.A. de C.V.