STOKO REFRESH FOAMING INSTANT HAND SANITIZER- ethyl alcohol liquid
Deb USA, 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.

Drug Facts

Active Ingredient

Ethanol (68%)

Purpose

Hand Antiseptic

Uses

Hand antiseptic to decrease bacteria on the skin.

Warnings

Warnings

Flammable, For external use only. Avoid contact with eyes. In case of accidental eye contact, flush eyes thoroughly with water. Discontinue use if irritation and redness develops.

Stop use and ask doctor if

Skin or eye irritation persists for more than 72 hours

Keep out of reach of children.

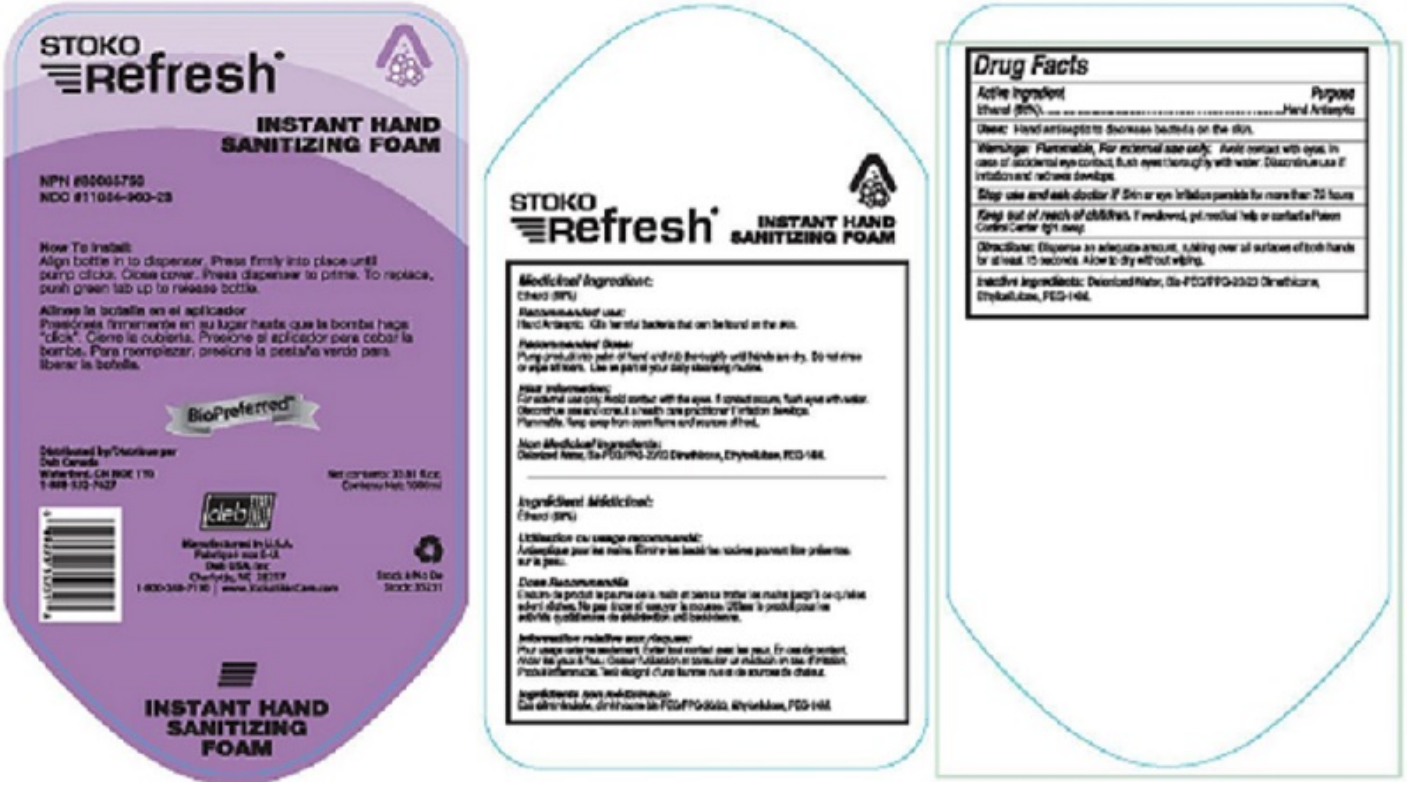
If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Dispense an adequate amount, rubbing over all surfaces of both hands for at least 15 seconds. Allow to dry without wiping.

Inactive Ingredients

Deionized Water, Bis-PEG/PPG-20/20 Dimethicone, Ethylcellulose, PEG-14M.



STOKO Refresh®
INSTANT HAND SANITIZING FOAM
NPN 80005750
NDC 11084-960-28
How To Install:
Align bottle in to dispenser. Press firmly into place until
pump clicks. Close cover. Press dispenser to prime.
To replace, push green tab up to release bottle.
Alinee la botella en el aplicador

Presiónela firmemente en su lugar hasta que la bomba haga “click”. Cierre la cubierta. Presione el aplicador para cebar la bomba. Para reemplazar, presione la pestaña verde para liberar la botella.
Bio Preferred.
Distributed by/Distribué par
Deb Canada
Waterford,ON N0E1Y0
1-888-332-7627
Net Contents 33.82 fl oz
CONTENU NET 1000 ml
Manufactured in U.S.A.
Fabriquée aux É.-U.
Deb USA, Inc.
Charlotte, NC, 28217
1-800-248-7190 / stoko.debgroup.com/us
Stock # / No De
Stock: 35231
INSTANT HAND SANITIZING FOAM
Medical Ingredient: Ethanol 68% v/v
Recommended use:
Hand Antiseptic. Kills harmful bacteria that can be found on the skin.
Recommended Dose:
Pump product into palm of hand and rub thoroughly until hands are dry. Do not rinse or wipe off foam. Use as part of your daily cleansing routine.
Risk Information:
For external use only. Avoid contact with eyes. If contact occurs, flush eyes with water. Discontinue use and consult a health care practitioner if irritation develops.
Flammable. Keep away from open flame and sources of heat.
Non Medicinal Ingredients: Deionized Water, Bis-PEG/PPG-20/20 Dimethicone, Ethylcellulose, PEG-14M.
Ingrédient médicinal:
Éthanol 68 % v/v
Utilisation ou usage recommandé:
Antiseptique pour les mains. Élimine les bactéries nocives pouvant être présentes sur la peau.
Dose Recommandée:
Enduire de produit la paume des mains et bien se frotter les mains jusqu’à ce qu’elles soient sèches. Ne pas rincer ni essuyer la mousse. Utiliser le produit pour les activités quotidiennes de désinfection anti-bactérienne
Information relative aux risques:
Pour usage externe seulement.
Éviter tout contact avec les yeux. En cas de contact, rincer les yeux a l’eau. Cesser l’utilisation et consulter un médecin en cas d’irritation.
Produit inflammable. Tenir éloigné d’une flamme nue et de sources de chaleur.
Ingrédients non médicinaux: Eau déminéralisée, diméthicone bis-PEG/PPG-20/20, éthylcellulose, PEG-14M.

STOKO REFRESH FOAMING INSTANT HAND SANITIZER

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-960
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	68 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BIS-PEG/PPG-20/20 DIMETHICONE (UNII: 05W209DSBS)	
ETHYLCELLULOSE (50 MPA.S) (UNII: 6I475159RA)	
POLYETHYLENE OXIDE 600000 (UNII: 2126FD486L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-960-28	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2014	12/31/2022

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2014	12/31/2022

Labeler - Deb USA, Inc. (607378015)

Registrant - Deb-STOKO USA, LLC (055861874)

Establishment

Name	Address	ID/FEI	Business Operations
Aphena Pharma Solutions-Maryland LLC		829739833	manufacture(11084-960)

Revised: 12/2018 Deb USA, Inc.