FLAWLESS HAND SANITIZER- propolis extract spray
NBIO CO., LTD.

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.

ACTIVE INGREDIENT

Propolis Extract 0.5%

INACTIVE INGREDIENTS

Water, Sodium Silicate

PURPOSE

Sanitizer

WARNINGS

■ For external use only.

When using this product
■ Do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
■ Avoid contact with broken skin.
■ Do not inhale or ingest.

Stop use and ask a doctor
■ If irritation or rash appears and lasts.

KEEP OUT OF REACH OF CHILDREN

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

Uses

■ Hand sanitizer to help reduce bacteria that potentially can cause disease.
■ For use when soap and water are not available.

Directions

■ Please spray it at a distance of 10~20cm so that your hands are enough covered. Rub hands together until dry.
■ Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

■ Store between 41~86□(Between 5~30□)
■ Avoid freezing and excessive heat above 104□(40□) and direct sunlight.

QUESTIONS

■ www.nbio.kr / 82-31-432-5670

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

FLAWLESS HAND SANITIZER

propolis extract spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77560-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PROPOLIS WAX (UNII: 6Y8XYV2NOF) (PROPOLIS WAX - UNII:6Y8XYV2NOF)	PROPOLIS WAX	1.25 g in 250 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Sodium Silicate (UNII: IJF18F77L3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77560-010-01	250 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2020	

Labeler - NBIO CO., LTD. (687218375)

Registrant - NBIO CO., LTD. (687218375)

Establishment

Name	Address	ID/FEI	Business Operations
NBIO CO., LTD.		687218375	manufacture(77560-010)

Revised: 5/2020 NBIO CO., LTD.