ORCHID ANTIBACTERIAL VANILLA AND APPLE BLOSSOM- ethyl alcohol liquid H E B

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

ETHYL ALCOHOL 62%

PURPOSE

ANTISEPTIC

USES

TO HELP REDUCE BACTERIA ON THE SKIN

WARNINGS

FOR EXTERNAL USE ONLY

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE WITH WATER

STOP USING THIS PRODUCT AND ASK DOCTOR IF

IRRITATION OR REDNESS DEVELOPS AND LASTS

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

DIRECTIONS

- APPLY SMALL AMOUNT TO YOUR PALM AND RUB HANDS TOGETHER BRISKLY UNTIL DRY
- CHILDREN UNDER 6 SHOULD BE SUPERVISED WHEN USING THIS PRODUCT

OTHER INFORMATION

STORE AT A TEMPERATURE BELOW 110°F (43°C)

QUESTIONS/COMMENTS?

1-866-695-3030

INACTIVE INGREDIENTS

WATER (AQUA), PROPYLENE GLYCOL, GLYCERIN, ISOPROPYL MYRISTATE,

AMINOMETHYL PROPANOL, CARBOMER, TOCOPHERYL ACETATE, FRAGRANCE (PARFUM), VANILLA PLANIFOLIA FRUIT EXTRACT, PYRUS MALUS (APPLE) FRUIT EXTRACT, ALOE BARBADENSIS LEAF JUICE, MANNITOL, CELLULOSE, HYDROXYPROPYL METHYLCELLULOSE, RETINYL PALMITATE, IRON OXIDES (CI 77491), TITANIUM DIOXIDE (CI 77891), RED 33 (CI 17200)

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ORCHID ANTIBACTERIAL VANILLA AND APPLE BLOSSOM ethyl alcohol liquid Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-165 Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
PROPYLENE GLYCOL (UNII: 6 DC9 Q167V3)			
GLYCERIN (UNII: PDC6A3C0OX)			
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)			
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)			
CARBOMER 934 (UNII: Z135WT9208)			
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)			
VANILLA (UNII: Q74T35078H)			
APPLE (UNII: B423VGH5S9)			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			
MANNITOL (UNII: 3OWL53L36A)			
POWDERED CELLULOSE (UNII: SMD1X3XO9M)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
VITAMIN A PALMITATE (UNII: 1D1K0 N0 VVC)			
FERRIC OXIDE RED (UNII: 1K09F3G675)			
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)			
D&C RED NO. 33 (UNII: 9DBA0SBB0L)			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:37808-165-08	236 mL in 1 BOTTLE, PLASTIC		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	0 1/15/20 15			

Labeler - HEB (007924756)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment						
Name	Address	ID/FEI	Business Operations			
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(37808-165)			

Revised: 1/2015 HE B