SANITIZING WIPES- alcohol cloth
Condor Co., Ltd.

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.

Sanitizing Wipes

Benzalkonium chloride 0.1%

Antiseptic, Hand Sanitizer

Antiseptic, Hand Sanitizer

Hand sanitizer to help reduce bacteria on the skin.

For use when soap and water are not available.

For external use only. Flammable. Keep away from heat or flame

in children less than 2 months of age
on open skin wounds

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.

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.

Open packet, remove and unfold pre-moistened tissue. Wipe hands or affected area thoroughly, then discard properly after use. Reseal back after use to keep wipes fresh.

Store between 15-30C (59-86F)
Avoid freezing and excessive heat above 40C (104F)

Phenoxy Ethanol, Polyhexamethylene Biguanide Hydrochloride, Purified Water, Propylene Glycol, Ethyl Alcohol, Vitamin E, Lemon Terpenes, Aloe Vera Extract.

10 wipes NDC: 80144-001-01

Directions
- Open packet, remove and unfold pre-moistened tissue. Wipe hands or affected area thoroughly, then discard properly after use. Reseal back after use to keep wipes fresh.

Inactive Ingredients
- Phenoxy Ethanol, Polyhexamethylene Biguanide Hydrochloride, Purified Water, Propylene Glycol, Ethyl Alcohol, Vitamin E, Lemon Terpenes, Aloe Vera Extract.

3 79508 00005 9

Made in China

Distributed in the USA by DOC HYGIENE
www.DocHygiene.com
Questions or Comments?
info@DocHygiene.com

SCAN TO SHOP

DOC HYGIENE™

EXTRA STRENGTH

SANITIZING WIPES

KILLS 99.9% OF GERMS

10 SANITIZING WIPES

5.9″ X 7″
(15cm X 18cm)
Lemon Scent

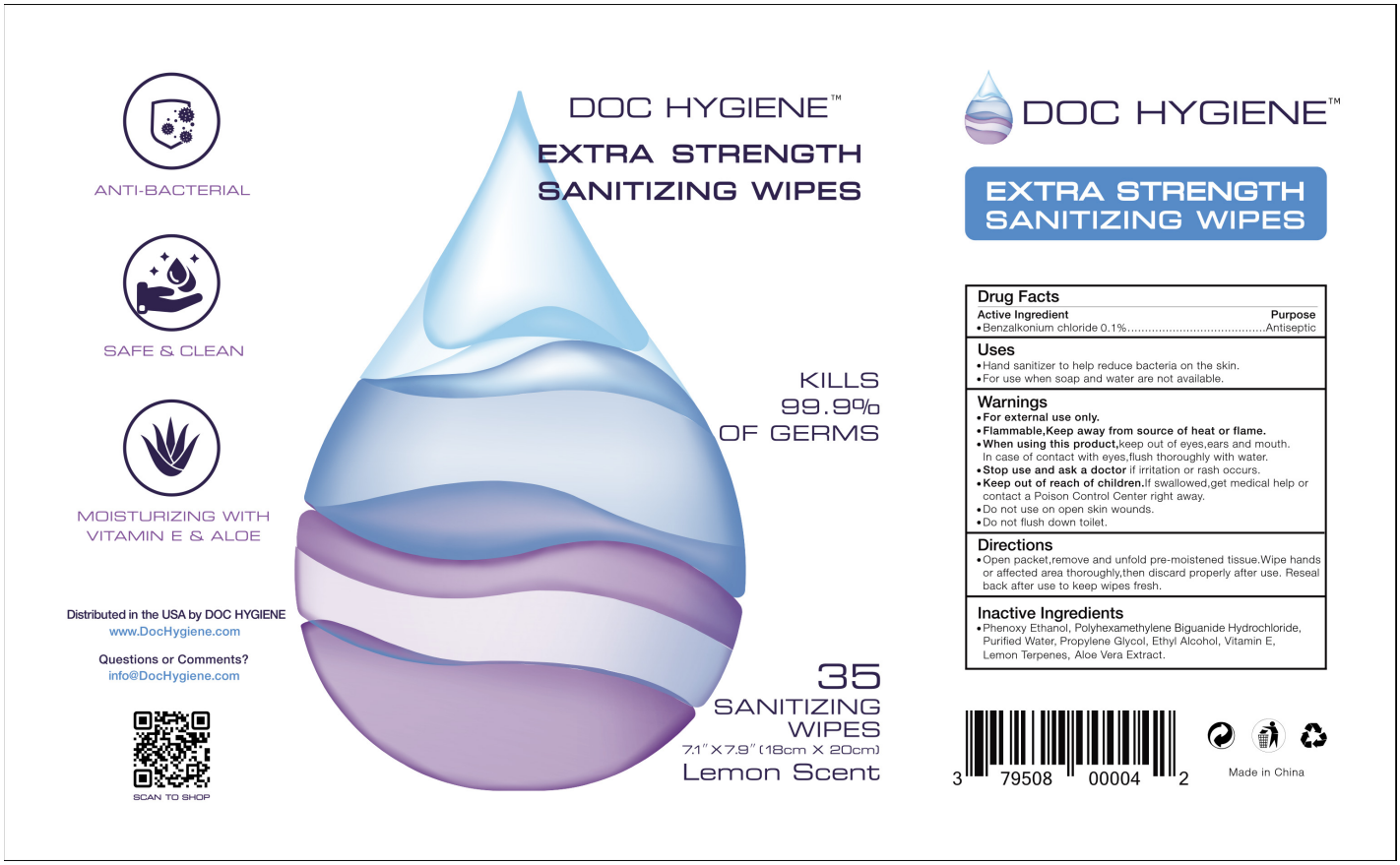
Drug Facts

Active Ingredient — **Purpose**
- Benzalkonium chloride 0.1% ..... Antiseptic

Uses
- Hand sanitizer to help reduce bacteria on the skin.
- For use when soap and water are not available.

Warnings
- **For external use only.**
- **Flammable, Keep away from source of heat or flame.**
- **When using this product,** keep out of eyes, ears and mouth. In case of contact with eyes, flush thoroughly with water.
- **Stop use and ask a doctor,** if irritation or rash occurs.
- **Keep out of reach of children,** If swallowed, get medical help per contact a Poison Control Center right away.
- Do not use on open skin wounds. Do not flush down toilet.

35 wipes NDC: 80144-001-02



SANITIZING WIPES

alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80144-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 mg in 100 mg

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	95.61 mg in 100 mg
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.05 mg in 100 mg
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	0.05 mg in 100 mg
POLIHEXANIDE HYDROCHLORIDE (UNII: 4XI6112496)	0.02 mg in 100 mg
ALCOHOL (UNII: 3K9958V90M)	4 mg in 100 mg
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.05 mg in 100 mg
LEMON TERPENES (UNII: 5DHA4TVW63)	0.02 mg in 100 mg
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.1 mg in 100 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80144-001-01	10 mg in 1 POUCH; Type 0: Not a Combination Product	08/19/2020	
2	NDC:80144-001-02	35 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/19/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/19/2020	

Labeler - Condor Co., Ltd. (560075779)

Registrant - Condor Co., Ltd. (560075779)

Establishment

Name	Address	ID/FEI	Business Operations
Condor Co., Ltd.		560075779	manufacture(80144-001)

Revised: 8/2020 Condor Co., Ltd.