HAND2MIND SANITIZING WET WIPES- benzalkonium chloride solution
Taizhou Duorui Biotechnology 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.

80800-002 wipes

Active Ingredient(s)

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

For use when soap and water are not available

Warnings

For external use only.

When using this product, keep out of eyes, ears or 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 immediate medical help or contact a Poison Control Center right away.

Directions

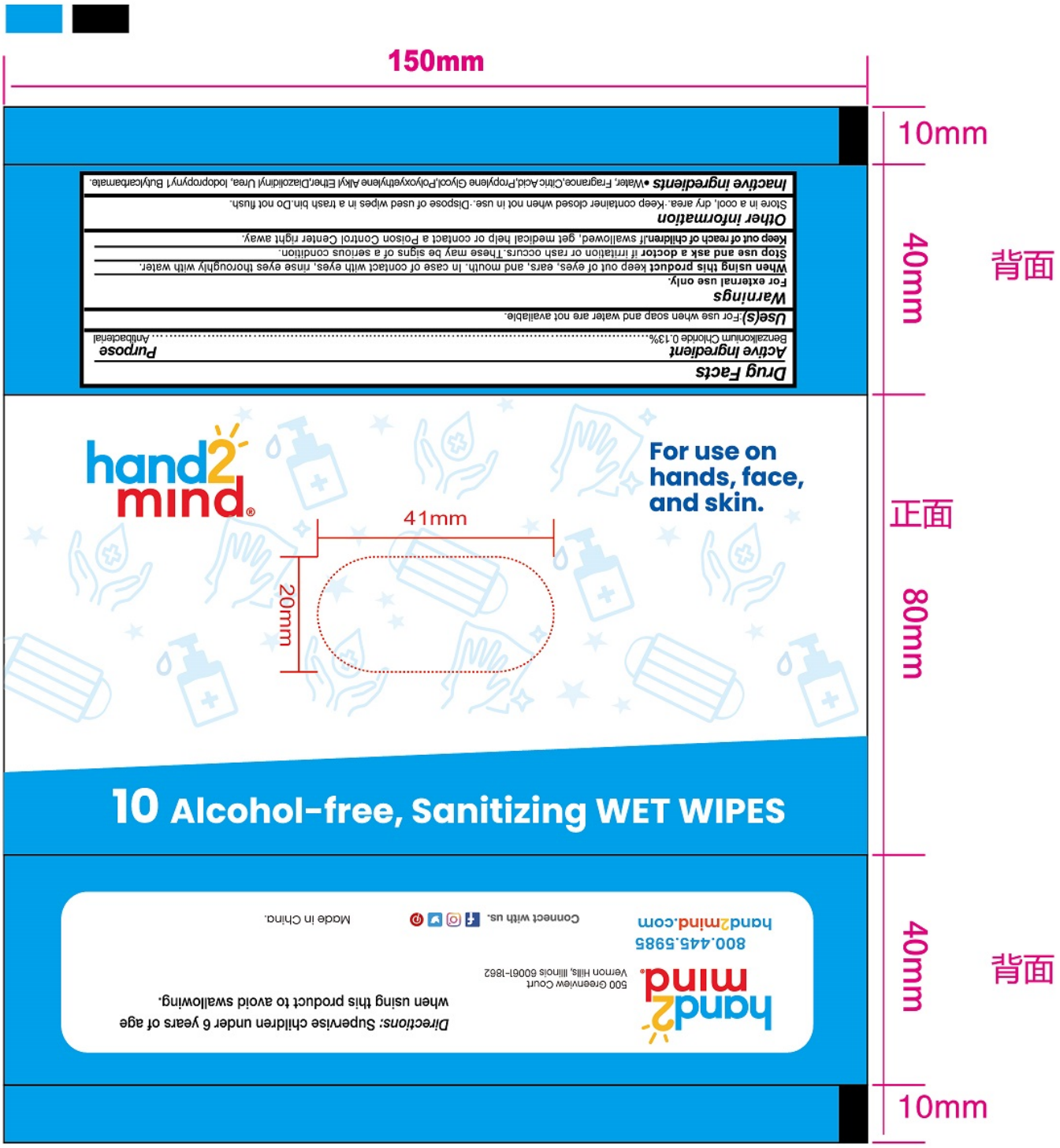
- Supervise children under 6 years of age when using this product to avoid swallowing

Other Information

Store in a cool, dry area. Keep container closed when not in use. Dispose of used wipes in trash bin. Do not flush.

Inactive ingredients

Water, Fragrance, Citric Acid, Propylene Glycol, Polyoxyethylene Alkyl Ether, Diazolidinyl Urea, Iodopropynyl Butylcarbamate



HAND2MIND SANITIZING WET WIPES

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80800-002

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80800-002-01	1 in 1 PACKAGE	03/12/2021	
1		3.46 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
2	NDC:80800-002-02	10 in 1 PACKAGE	03/12/2021	
2		3.46 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
3	NDC:80800-002-03	20 in 1 PACKAGE	03/12/2021	
3		3.46 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
4	NDC:80800-002-04	30 in 1 PACKAGE	03/12/2021	
4		3.46 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
5	NDC:80800-002-05	60 in 1 PACKAGE	03/12/2021	
5		3.46 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
6	NDC:80800-002-06	110 in 1 PACKAGE	03/12/2021	
6		3.46 mL in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/12/2021	

Labeler - Taizhou Duorui Biotechnology Co., Ltd. (713362289)

Establishment

Name	Address	ID/FEI	Business Operations
Taizhou Duorui Biotechnology Co., Ltd.		713362289	manufacture(80800-002)

Revised: 3/2021

Taizhou Duorui Biotechnology Co., Ltd.