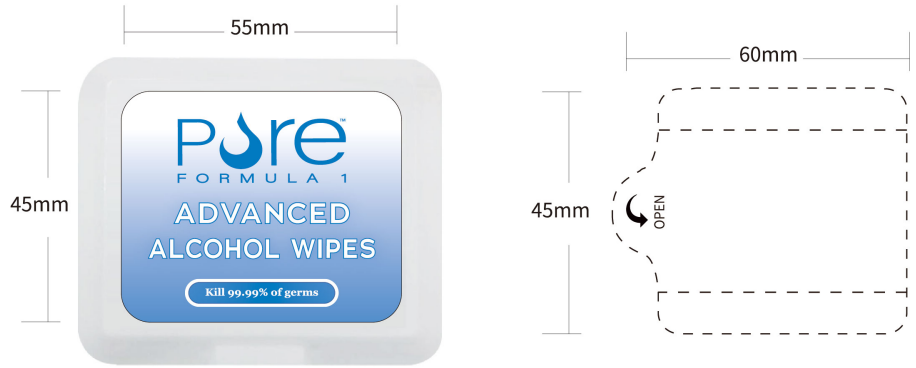
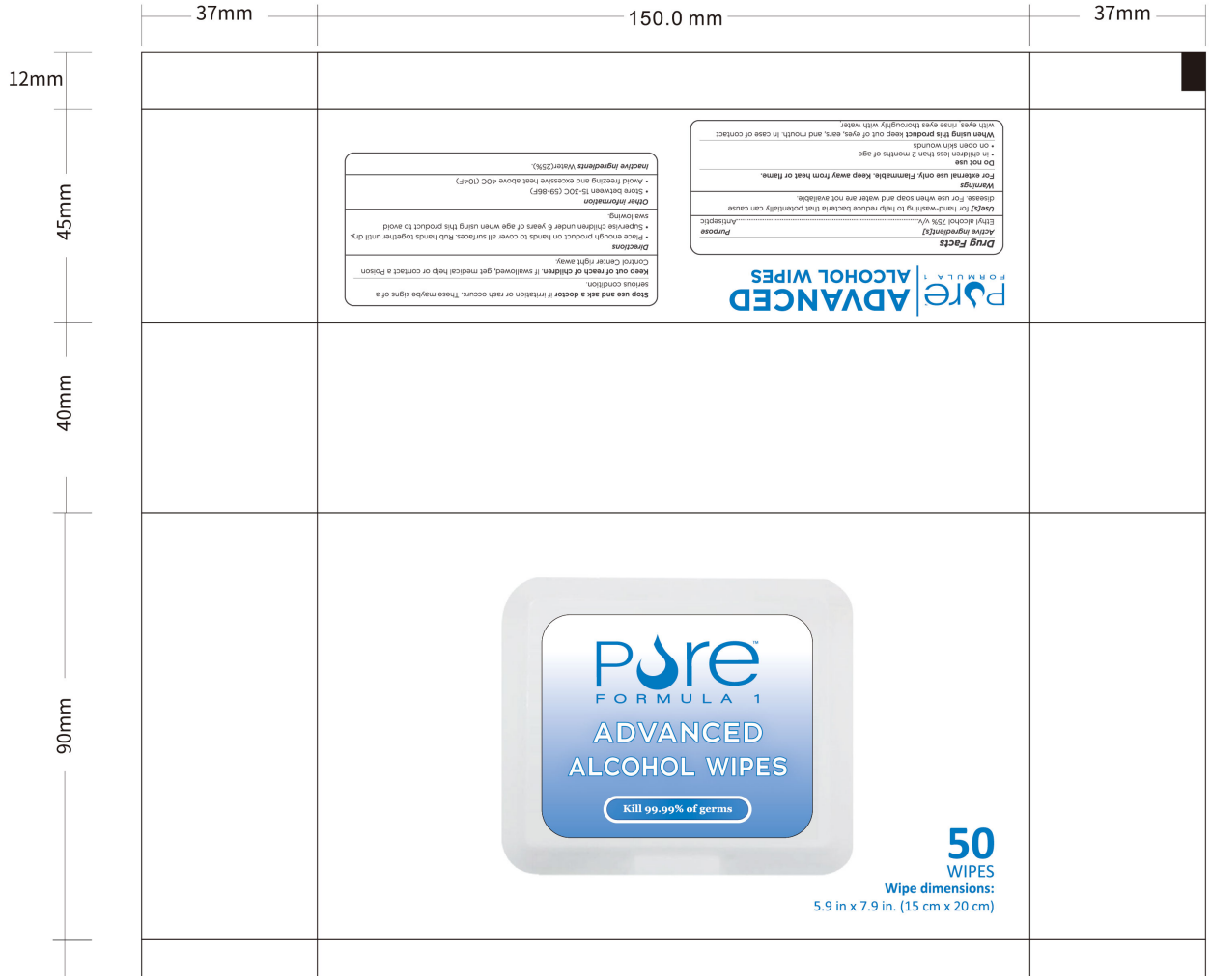
PURE FORMULA 1 ADVANCED ALCOHOL WIPES 01- alcohol cloth
Shenzhen Lantern Science Co.,Ltd.

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.

Pure Formula 1 advanced alcohol wipes

drug facts





Active ingredient

Active ingredient purpose

Ethyl alcohol 75%v/v Antiseptic

Use

for hand-washing to help reduce bacteria that potentially can cause disesase.For use when soap and waterare not available.

warnings

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

Do not use

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 maybe signs of a serious condition.

Keep out of reach of children

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

Directions

Place enough product on hands to cover all surfaces,Rub hands together until dry.

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

Other information

Store between15-30C(59-86F)

Avoid freezing and excessive heat above 40C(104F)

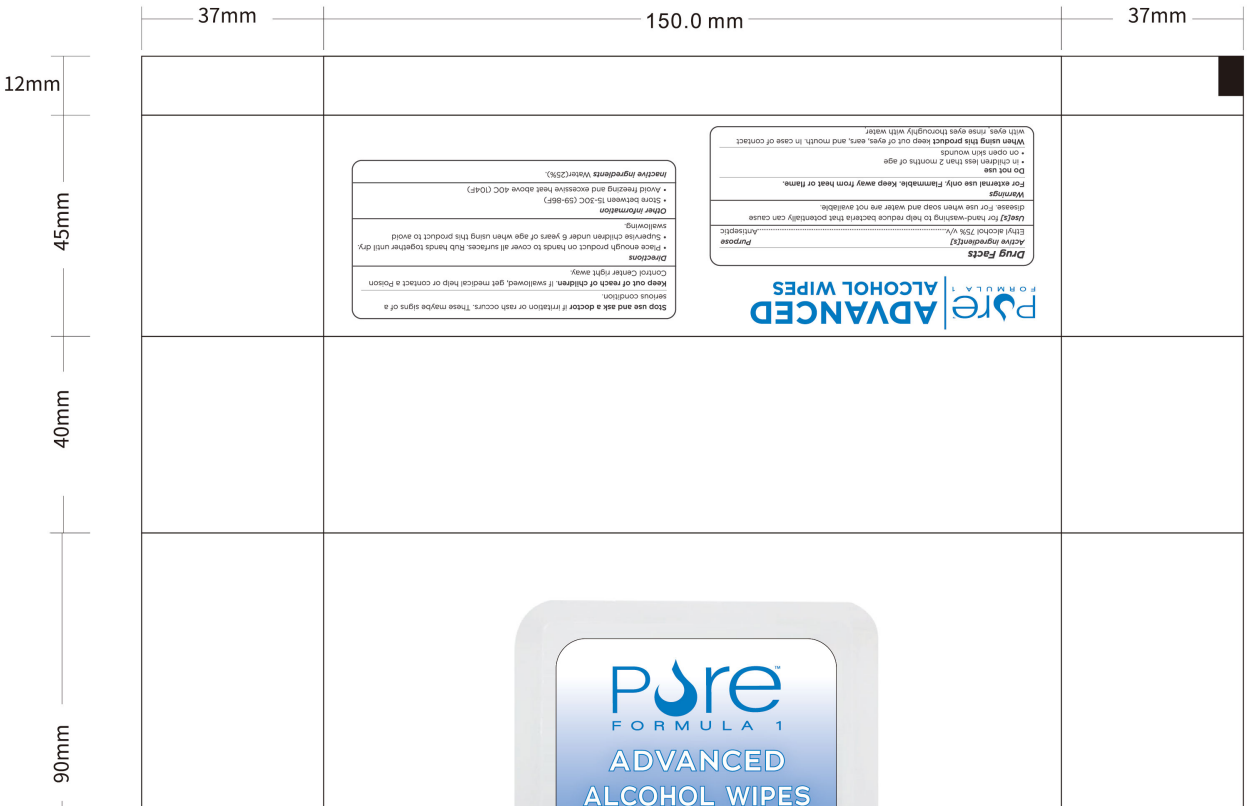
Inactive ingredient

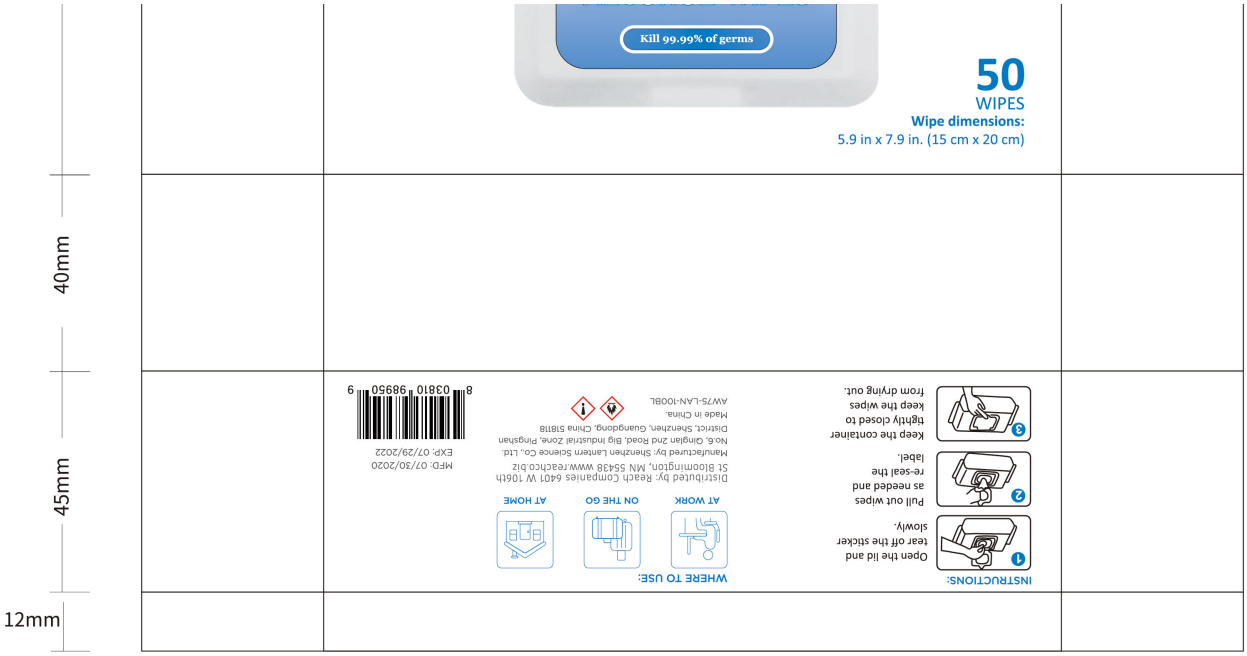
Water(25%)

usage

Place enough product on hands to cover all surfaces,Rub hands together until dry.

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





PURE FORMULA 1 ADVANCED ALCOHOL WIPES 01

alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54860-298
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 in 100

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	25

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-298-01	50 in 1 PACKAGE; Type 0: Not a Combination Product	08/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/03/2020	

Labeler - Shenzhen Lantern Science Co.,Ltd. (421222423)

Registrant - Lantern Beauty America,Inc. (117371139)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-298)

Revised: 8/2020 Shenzhen Lantern Science Co.,Ltd.