MAGIC XTREME XTRA SOFT ALCOHOL WIPE 1- ethyl alcohol cloth
ENSOBRETADOS Y DERIVADOS S.A. DE C.V.

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.

MAGIC XTREME XTRA SOFT ALCOHOL WIPE 1

DRUG FACTS

ACTIVE INGREDIENT

Ethyl Alcohol 70%

PURPOSE

Antibacterial

USES

Hand sanitizer to decrease bacteria on the skin.

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 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.

Keep out of reach of children If swallowed get medical help or contact a Poison Control Center right away.

DIRECTIONS

■ Wet hands thoroughly. Allow to dry without wiping.

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

OTHER INFORMATION

■ Store between 15-30C (59-86F)

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

INACTIVE INGREDIENTS

Water (Aqua), Glycerin.

QUESTIONS OR COMMENTS?

1-855-349-7735 or visit **www.eydmx.com**

MAGIC XTREME XTRA SOFT ALCOHOL WIPE 1 (NDC 78885-001-11)

MAGIC XTREME XTRA SOFT ALCOHOL WIPE 1

ethyl alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78885-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78885-012-00	1 mL in 1 PACKET; Type 0: Not a Combination Product	01/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/15/2021	

Labeler - ENSOBRETADOS Y DERIVADOS S.A. DE C.V. (813382660)

Registrant - ENSOBRETADOS Y DERIVADOS S.A. DE C.V. (813382660)

Establishment

Name	Address	ID/FEI	Business Operations
ENSOBRETADOS Y DERIVADOS S.A. DE C.V.		813382660	manufacture(78885-012)

Revised: 1/2021 ENSOBRETADOS Y DERIVADOS S.A. DE C.V.