MAYINGLONG HEMORRHOIDS- zinc oxide, petrolatum, lanolin ointment
Mayinglong Pharmaceutical Group 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.

MAYINGLONG HEMORRHOIDS OINTMENT

Active Ingredients

Zinc Oxide 0.8g Astrigent

Petrolatum 7.85g Protectant

Purpose

Astrigent

Protectant

Protectant

Uses

helps relieve the local itching and discomfort associated with hemorrhoids

temporarily shrinks hemorrhoidal tissue and relieves burning

temporarily provides a coating for relief of anorectal discomforts

temporarily protects the inflamed, irritated anorectal surface to help make bowel movements less painful

Warning

For external and /or intrarectal use only. Not for oral use.

Stop use

bleeding occurs

condition worses or does not improve within 7 days

introduction of applicator into the rectum causes additional pain

Warning

Apply it carfully during pregnancy.

Warning

Keep this and all drugs out of reach of children.

Direction

Apply properly onto anus or apply on affected areas twice a day.

Other information

Store at 20-25 degree Centigrade (68-77 degree Fahrenheit)

Inactive ingredients

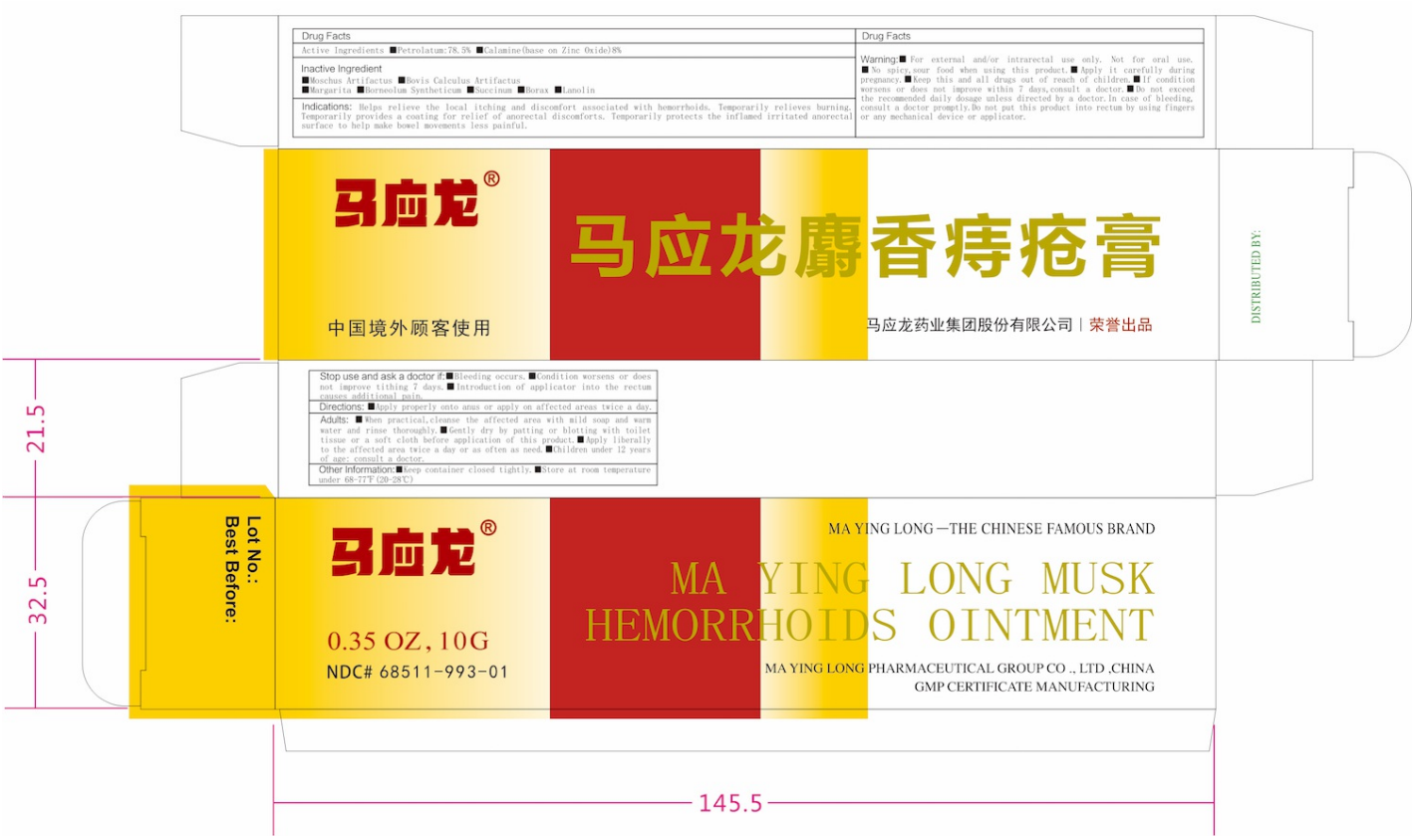
Musk Ketone

Pearl (Hyriopsis Cumming II)

Borneol

Sodium Borate

Drug Facts



MAYINGLONG HEMORRHOIDS

zinc oxide, petrolatum, lanolin ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68511-993
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	800 mg in 10000 mg
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	7850 mg in 10000 mg

Inactive Ingredients	
Ingredient Name	**Strength**
MUSK KETONE (UNII: 483V3E1L6J)	
PEARL (HYRIOPSIS CUMINGII) (UNII: A75L5FZ40U)	
BORNEOL (UNII: M89NIB437X)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
LANOLIN (UNII: 7EV65EAW6H)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:68511-993-01	10000 mg in 1 CARTON; Type 0: Not a Combination Product	10/25/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part346	12/15/2003	

Labeler - Mayinglong Pharmaceutical Group Co Ltd (526823828)

Registrant - Mayinglong Pharmaceutical Group Co Ltd (526823828)

Establishment

Name	Address	ID/FEI	Business Operations
Mayinglong Pharmaceutical Group Co Ltd		526823828	manufacture(68511-993)

Revised: 3/2018 Mayinglong Pharmaceutical Group Co Ltd