CHUNGWIDAN-F- cinnamon, nutmeg liquid
Lydia Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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cinnamon, nutmeg

digestion

Keep out of reach of children

Children under 7 years of age:consult a doctor

Take 1-2 envelop as symptoms accur,as directed by a doctor

Ask a doctor before use if you have taking a prescription drug.

Antacids may interact with certain prescription drugs.

Mint,Sodium Citrate Hydrate,Purified Water

for oral use

CHUNGWIDAN-F

Drug Facts

Active Ingredients	***Purpose***
Cinnamon 500mg	Topical analgesic
Nutmeg 300mg	Topical analgesic

Uses
• Digestion

Warnings
Ask a doctor before use if you have taking a prescription drug.
Antacids may interact with certain prescription drugs.

Directions
Children under 7 years of age:consult a doctor
Take 1-2 envelop as symptoms accur,as directed by a doctor

Other Information
Store at room temperature

Inactive Ingredient
Mint,Sodium Citrate Hydrate,Purified Water

Questions or comments ?
Call weekdays from 9 a.m to 5 p.m EST at (213) 266-2776
Distributed By: Joyful Worldwide,Inc
8345 Garden Grove Blvd #106 Garden Grove,CA92844, USA

Made in South Korea

CHUNGWIDAN-F

cinnamon, nutmeg liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72988-0029
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CINNAMON (UNII: 5S29HWU6QB) (CINNAMON - UNII:5S29HWU6QB)	CINNAMON	500 mg in 30 mL
NUTMEG (UNII: AEE24M3MQ9) (NUTMEG - UNII:AEE24M3MQ9)	NUTMEG	300 mg in 30 mL

Inactive Ingredients	
Ingredient Name	**Strength**
MINT (UNII: FV98Z8GITP)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72988-0029-1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/06/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/06/2022	

Labeler - Lydia Co., Ltd. (695735569)

Registrant - Lydia Co., Ltd. (695735569)

Establishment

Name	Address	ID/FEI	Business Operations
Lydia Co., Ltd.		695735569	manufacture(72988-0029)

Revised: 6/2022 Lydia Co., Ltd.