STOMACHIN ANTACID- magnesium carbonate and sodium bicarbonate powder Chang Kuo Chou Pharmaceutical 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.

Stomachin®

Drug Facts

Active ingredients (in each gram)	Purpose	
Magnesium carbonate 58mg	Antacid	
Sodium bicarbonate 809mg	Antacid	

Uses

relieves:

- acid indigestion
- heartburn
- sour stomach
- upset stomach from overindulgence in food or drink

Warnings

Allergy alert

Do not use if you are allergic to the antacid.

Do not use more than 3 packets in a 24-hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a physician.

Ask a doctor or pharmacist before use if you are

- presently taking a prescription drug. Antacid may interact with certain prescription drugs.
- On magnesium or sodium restricted diet.

Stop use and ask a doctor if

• symptoms persist or worsen.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Directions

Take 2 gram (one packet) with water, 3 times a day

Other information

- each gram contains : magnesium 16 mg and sodium 221 mg
- store at cool and dry place

Inactive ingredients

clove, licorice, menthol, rhubarb

Manufactured by : Chang Kuo Chou Pharmaceutical Co.,Ltd. Taoyuan, Taiwan

PRINCIPAL DISPLAY PANEL - 40 g Packet Box

NDC 43337-9329-9

Stomachin[®]

ANTACID POWDER

Fast Relief of:

Acid indigestion Upset stomach Heartburn

Net weight: 1.4oz.(40g), Total 20 packets

Product of Taiwan



NDC 43337-9329-9

提國周 选門故 Stomachin

提國周 选肖散 Stomachin

ANTACID POWDER

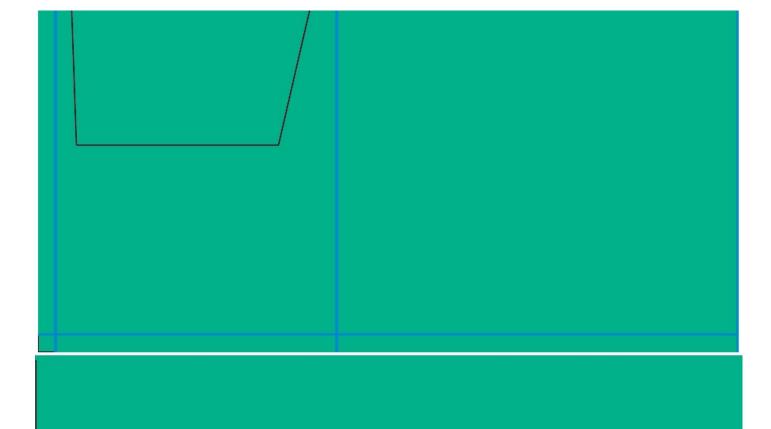
Fast Relief of:

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LOT: EXP:

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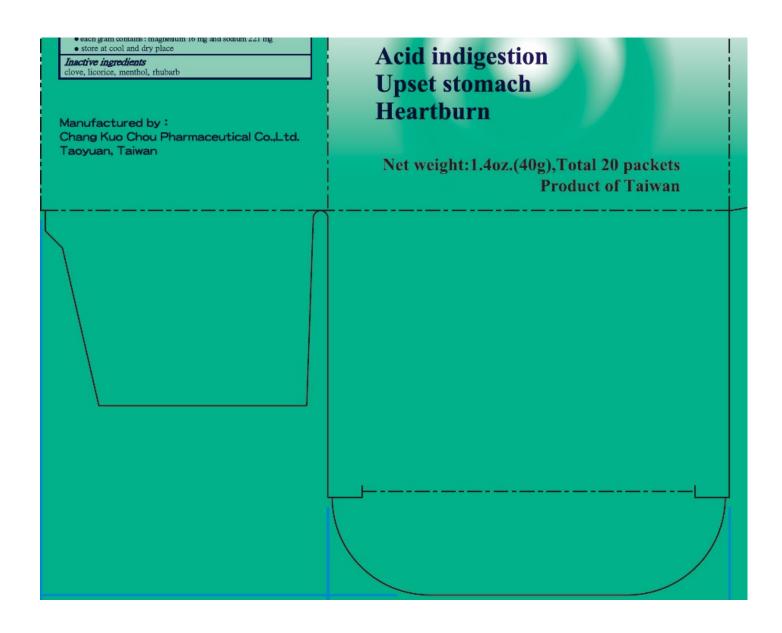
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NDC 43337-9329-9

Stomachin

ANTACID POWDER

Fast Relief of:



STOMACHIN ANTACID

magnesium carbonate and sodium bicarbonate powder

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43337-9329
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
MAGNESIUM CARBONATE (UNII: 0E53J927NA) (CARBONATE ION - UNII:7UJQ50PE7D)	MAGNESIUM CARBONATE	58 mg in 1 g		
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (BICARBONATE ION - UNII: HN1ZRA3Q20)	SODIUM BICARBONATE	809 mg in 1 g		

Inactive Ingredients	
Ingredient Name	Strength

CLOVE (UNII: K48IKT5321)	29 mg in 1 g
LICORICE (UNII: 61ZBX54883)	86 mg in 1 g
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	4 mg in 1 g
RHUBARB (UNII: G280W4MW6E)	14 mg in 1 g

Product Characteristics			
Color	WHITE (beige)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:43337- 9329-8	450 g in 1 CAN; Type 0: Not a Combination Product	04/30/2009		
2	NDC:43337- 9329-7	95 g in 1 CAN; Type 0: Not a Combination Product	04/30/2009		
3	NDC:43337- 9329-9	20 in 1 BOX	04/30/2009		
3		2 g in 1 PACKET; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	04/30/2009	

Labeler - Chang Kuo Chou Pharmaceutical Co., Ltd. (656241486)

Establishment				
Name	Address	ID/FEI	Business Operations	
Chang Kuo Chou Pharmaceutical Co., Ltd.		656241486	MANUFACTURE(43337-9329)	

Revised: 12/2022 Chang Kuo Chou Pharmaceutical Co., Ltd.