

RAZZMATAZZ ANTACID- calcium carbonate tablet, chewable
Provision Medical Products

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.

UseS

For the relief of-Sour Stomach, Upset Stomach, Heartburn, Acid Indigestion
If pregnant or breast-feeding, ask a health professional before use.

KEEP OUT OF REACH OF CHILDREN. In case of overdose, get medical help or contact a Poison Control Center right away.

Prompt medical attention is critical for adults as well as for children, even if you do not notice any signs or symptoms.

Warnings:

Ask a doctor or health professional before use if you are: currently taking a prescription drug. Antacids may react with certain prescription drugs.

Directions:Do not use more thsn directed. Adults and children 12 years or older. Take 2 tablets every 4 to 6 hours or as needed, do not exceed 18 tablets in 24 hours, or as directed by a doctor.

Children under 12 years-Do not give to children under 12 years unless directed by a doctor.

Inactive Ingredients:

corn startch, FD&C Red 40, flavor, magnesium stearate, silicon dioxide, startch, sucrose

ACTIVE INGREDIENT-calcium carbonate 420 MG
ANTACID



RAZZMATAZZ ANTACID

calcium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69103-2516
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6Z B)	CALCIUM CARBONATE	420 mg

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white (WHITE WTH GREEN FLECKS)	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	FR14
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69103-2516-5	250 in 1 CARTON	03/31/2015	07/01/2024
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:69103-2516-6	100 in 1 CARTON	03/31/2015	07/01/2024
2		2 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	03/31/2015	07/01/2024

Labeler - Provision Medical Products (036936831)

Registrant - Provision Medical Products (036936831)

Establishment

Name	Address	ID/FEI	Business Operations
ULTRAtab Laboratories, Inc.		151051757	manufacture(69103-2516)