MINT ANTACID- calcium carbonate tablet, chewable
ADVANCED FIRST AID, INC.

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.

ACTIVE INGREDIENT IN EACH TABLET- CALCIUM CARBONATE 420 MG

ANTACID

Uses:

for the relief of:

•heartburn • sour stomach • acid indigestion

Warnings:

Do not:

•Take more than 18 tablets in a 24 hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice of a physician.

Stop use and ask a doctor if:

•You are currently taking any prescription drug. Antacids may react with certain prescription drugs.

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

KEEP OUT OF REACH OF CHILDREN.

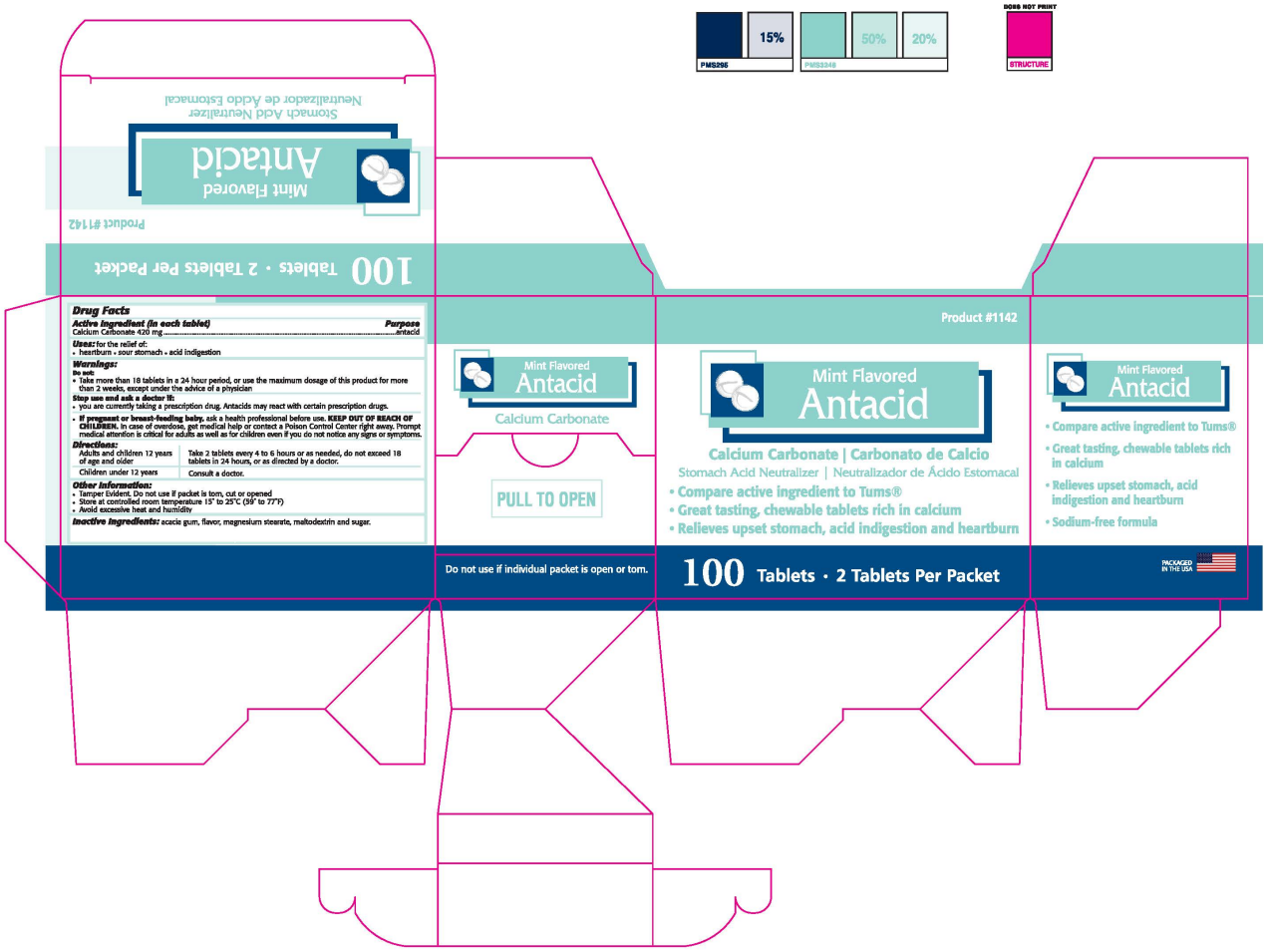
In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions:

Adults and children 12 years of age and 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: Consult a doctor.

Inactive Ingredients: acacia gum, flavor, magnesium stearate, maltodextrin, starch and sugar.



MINT ANTACID

calcium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67060-303
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ5OPE7D)	CALCIUM CARBONATE	420

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
ACACIA (UNII: 5C5403N26O)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	

Product Characteristics

Color	white (WHITE)	**Score**	no score
Shape	ROUND	**Size**	12mm
Flavor	MINT	**Imprint Code**	FR8
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67060-303-68	100 in 1 CARTON	04/07/2015	
1	NDC:67060-303-02	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	04/07/2015	

Labeler - ADVANCED FIRST AID, INC. (114477180)

Registrant - ADVANCED FIRST AID, INC. (114477180)

Establishment

Name	Address	ID/FEI	Business Operations
ULTRA SEAL CORPORATION		085752004	pack(67060-303)

Establishment

Name	Address	ID/FEI	Business Operations
ULTRA TAB LABORATORIES, INC.		151051757	manufacture(67060-303)

Revised: 3/2019 ADVANCED FIRST AID, INC.