ALKA-SELTZER HEARTBURN RELIEFCHEWS- calcium carbonate tablet, chewable
Bayer HealthCare LLC.

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.

Alka-Seltzer ® Heartburn ReliefChews™

Drug Facts

Active ingredient (in each chewable tablet)

Calcium carbonate 750 mg

Purpose

Antacid

Uses

for the relief of:
- acid indigestion
- heartburn
- sour stomach
- upset stomach associated with these symptoms

Warnings

Do not use if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor or pharmacist before use if you are presently taking a prescription drug. Antacids may interact with certain prescription drugs.

When using this product
- do not take more than 10 chewable tablets in a 24-hour period
- do not use the maximum dosage of this product for more than 2 weeks except under the advice and supervision of a physician
- constipation may occur

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

Keep out of reach of children.

Directions
- adults and children 12 years and over: chew and swallow 1-2 chewable tablets every

2 to 4 hours as symptoms occur, or as directed by a doctor
- children under 12 years: consult a doctor
- do not take more than 10 chewable tablets in a 24-hour period

Other information
- **each chewable tablet contains:** calcium 300 mg
- contains FD&C Yellow No. 5 (tartrazine) as a color additive
- store at room temperature. Avoid humidity. Close cap tightly after use.

Inactive ingredients

acacia, beeswax, carmine, carnauba wax, citric acid, corn starch, corn syrup, DL-alpha tocopherol, FD&C blue #1 aluminum lake, FD&C red #40 aluminum lake, FD&C yellow #5 lake (tartrazine), FD&C yellow #6, FD&C yellow #6 aluminum lake, FD&C yellow #6 lake, flavors, hydrogenated coconut oil, medium chain triglycerides, methyl paraben, modified starch, phosphoric acid, pregelatinized modified starch, propyl paraben, propylene glycol, purified water, shellac, sodium benzoate, sorbic acid, sorbitol, soy lecithin, sucrose, titanium dioxide

Questions?

1-800-986-0369 (Mon – Fri 9AM – 5PM EST) or www.alkaseltzer.com

Dist. by:
Bayer HealthCare LLC
Whippany, NJ 07981

PRINCIPAL DISPLAY PANEL - 36 Tablet Bottle Label

assorted fruit

Alka-

Seltzer ®

Calcium Carbonate /

Antacid

EXTRA STRENGTH

HEARTBURN

RELIEFCHEWS™

Fast Powerful Relief

36 CHEWABLE TABLETS

ALKA-SELTZER HEARTBURN RELIEFCHEWS

calcium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0280-0221
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	750 mg

Inactive Ingredients

Ingredient Name	Strength
YELLOW WAX (UNII: 2ZA36H0S2V)	
ACACIA (UNII: 5C5403N26O)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
STARCH, CORN (UNII: O8232NY3SJ)	
CORN SYRUP (UNII: 9G5L16BK6N)	
.ALPHA.-TOCOPHEROL, DL- (UNII: 7QWA1RIO01)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYDROGENATED COCONUT OIL (UNII: JY81OXM1OM)	

MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SHELLAC (UNII: 46N107B71O)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBIC ACID (UNII: X045WJ989B)	
SORBITOL (UNII: 506T60A25R)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
SUCROSE (UNII: C151H8M554)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	yellow, orange, pink	**Score**	no score
Shape	ROUND	**Size**	18mm
Flavor	LEMON, ORANGE, STRAWBERRY	**Imprint Code**	AS
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0280-0221-36	36 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2014	
2	NDC:0280-0221-60	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2014	
3	NDC:0280-0221-12	120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2014	
4	NDC:0280-0221-90	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/12/2015	
5	NDC:0280-0221-08	8 in 1 POUCH; Type 0: Not a Combination Product	03/23/2016	
6	NDC:0280-0221-01	32 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2020	
7	NDC:0280-0221-02	66 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2020	
8	NDC:0280-0221-03	200 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/15/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	03/15/2014	

Labeler - Bayer HealthCare LLC. (112117283)