

**TUMS ULTRA- calcium carbonate tablet, chewable**  
**Haleon US Holdings LLC**

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***Drug Facts***

***Active ingredient (per tablet)***

Calcium Carbonate USP 1000 mg

***Purpose***

Antacid

***Uses***

relieves

- heartburn
- acid indigestion
- sour stomach
- upset stomach associated with these symptoms

***Warnings***

**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 7 tablets in 24 hours
- if pregnant do not take more than 5 tablets in 24 hours
- do not use the maximum dosage for more than 2 weeks except under the advice and supervision of a doctor

**Keep out of reach of children.**

***Directions***

- **adults and children 12 years of age and over:** chew 2-3 tablets as symptoms occur, or as directed by a doctor
- do not take for symptoms that persist for more than 2 weeks unless advised by a doctor

***Other information***

- **each tablet contains:** elemental calcium 400mg, sodium 2mg
- store below 30°C (86°F)

***Inactive ingredients (Assorted Fruit)***

adipic acid, corn starch, FD&C blue #1 lake, FD&C red #40 lake, FD&C yellow #5 (tartrazine) lake, FD&C yellow #6 lake, flavors, mineral oil, sodium polyphosphate, sucrose, talc

***Inactive ingredients (Assorted Tropical Fruit)***

corn starch, FD&C red #40 lake, FD&C yellow #5 (tartrazine) lake, FD&C yellow #6 lake, flavors, mineral oil, sodium polyphosphate, sucrose, talc

***Inactive ingredients (Assorted Berry)***

adipic acid, corn starch, FD&C blue #1 lake, FD&C red #40 lake, flavors, mineral oil, sodium polyphosphate, sucrose, talc

***Inactive ingredient (Peppermint)***

corn starch, flavor, mineral oil, sodium polyphosphate, sucrose, talc

***Inactive ingredients (Peppermint Tri-Color)***

corn starch, FD&C blue #1 lake, FD&C red #40 lake, FD&C yellow #5 (tartrazine) lake, flavor, mineral oil, sodium polyphosphate, sucrose, talc

***Questions?***

**1-800-897-7535 weekdays**

**Safety sealed- Do not use if printed inner seal beneath cap is missing or broken.**

[www.tums.com](http://www.tums.com)

**GlaxoSmithKline**

Moon Twp, PA 15108

**Gluten-Free**

**Principal Display Panel**

**NDC 0135-0118-83**

**TUMS®**

CALCIUM CARBONATE

ANTACID

**ASSORTED FRUIT**

**ULTRA STRENGTH 1000**

**72 CHEWABLE TABLETS**

**GOES TO WORK IN SECONDS!**

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PAREVE

103662XA (Front Label)

103593XA (Back Label)

**GOES TO WORK IN SECONDS!**

NDC 0135-0118-83

CALCIUM CARBONATE

**TUMS<sup>®</sup>**

ANTACID

**Assorted  
Fruit**

**ULTRA 1000  
STRENGTH**

**72 CHEWABLE  
TABLETS**



**103662XA**

**Principal Display Panel**

**NDC 0135-0180-02**

**TUMS®**

CALCIUM CARBONATE

ANTACID

**TROPICAL FRUIT**

**ULTRA STRENGTH 1000**

**72 CHEWABLE TABLETS**

**GOES TO WORK IN SECONDS!**

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PAREVE

103634XA (Front Label)

103591XA (Back Label)

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NDC 0135-0180-02



CALCIUM CARBONATE

**TUMS**<sup>®</sup>

ANTACID

***Tropical  
Fruit***

**ULTRA 1000**  
STRENGTH

**72 CHEWABLE  
TABLETS**



**103634XA**

Principal Display Panel  
NDC 0135-0181-02

**TUMS®**

CALCIUM CARBONATE

ANTACID

**ASSORTED BERRIES**

**ULTRA STRENGTH 1000**

**72 CHEWABLE TABLETS**

**GOES TO WORK IN SECONDS!**

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PAREVE

103665XA (Front Label)

103588XA (Back Label)

GOES TO WORK IN SECONDS!

NDC 0135-0181-02

CALCIUM CARBONATE

**TUMS**<sup>®</sup>

ANTACID

*Assorted  
Berries*

**ULTRA 1000**  
STRENGTH

**72** CHEWABLE  
TABLETS



PAREVE

**103665XA**

Principal Display Panel



**NDC 0135-0228-06**

**TUMS®**

CALCIUM CARBONATE

ANTACID

**PEPPERMINT**

**ULTRA STRENGTH 1000**

**72 CHEWABLE TABLETS**

**GOES TO WORK IN SECONDS!**

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103664XA (Front Label)

103592XA (Back Label)

GOES TO WORK IN SECONDS!

NDC 0135-0228-06



CALCIUM CARBONATE

**TUMS**<sup>®</sup>

ANTACID

*Peppermint*

**ULTRA 1000**  
STRENGTH

**72** CHEWABLE  
TABLETS



PAREVE

**103664XA**

Principal Display Panel

NDC 0135-0540-01

**TUMS®**

ANTACID

CALCIUM CARBONATE

**PEPPERMINT**

**ULTRA STRENGTH 1000**

**86 CHEWABLE TABLETS**

**20% MORE FREE**

**PAREVE**

©2014 GSK

104475XA (Front Label)

104476XA (Back Label)

**20%  
MORE FREE**

NDC 0135-0540-01

CALCIUM CARBONATE

**TUMS**<sup>®</sup>

ANTACID

**Peppermint**

**ULTRA 1000**  
STRENGTH

**86**

~~72~~

CHEWABLE  
TABLETS



104475XA

# TUMS ULTRA

calcium carbonate tablet, chewable

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0135-0118
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ50PE7D, CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1000 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>ADIPIIC ACID</b> (UNII: 76A0JE0FKJ)	
<b>SODIUM POLYMETAPHOSPHATE</b> (UNII: P1BM4ZH95L)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	

## Product Characteristics

<b>Color</b>	PINK (orange, yellow, green)	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	19mm
<b>Flavor</b>	CHERRY (assorted fruit, orange, lemon, lime)	<b>Imprint Code</b>	TUMS
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0118-01	86 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
2	NDC:0135-0118-04	265 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
3	NDC:0135-0118-14	160 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
4	NDC:0135-0118-	72 in 1 BOTTLE; Type 0: Not a Combination	02/12/2010	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/12/2010	

## TUMS ULTRA

calcium carbonate tablet, chewable

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0135-0180
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ5OPE7D, CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1000 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>SODIUM POLYMETAPHOSPHATE</b> (UNII: P1BM4ZH95L)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	

### Product Characteristics

<b>Color</b>	ORANGE (red-orange, cream to off-white, light yellow)	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	19mm
<b>Flavor</b>	TROPICAL FRUIT PUNCH (assorted tropical fruit, mandarin orange, orange-pineapple, strawberry-banana)	<b>Imprint Code</b>	TUMS
<b>Contains</b>			

### Packaging

Marketing Start	Marketing End
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0180-01	86 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
2	NDC:0135-0180-02	72 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
3	NDC:0135-0180-14	160 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/12/2010	

## TUMS ULTRA

calcium carbonate tablet, chewable

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0135-0181
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ50PE7D, CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1000 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>ADIPIC ACID</b> (UNII: 76A0JE0FKJ)	
<b>SODIUM POLYMETAPHOSPHATE</b> (UNII: P1BM4ZH95L)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	

### Product Characteristics

Color	PINK (mauve, bluish)	Score	no score
Shape	ROUND	Size	19mm
Flavor	BERRY (assorted berry, strawberry, raspberry, mixed berry)	Imprint Code	TUMS
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0181-01	3 in 1 CARTON	02/12/2010	
1		12 in 1 PACKAGE; Type 0: Not a Combination Product		
2	NDC:0135-0181-02	72 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
3	NDC:0135-0181-03	12 in 1 PACKAGE; Type 0: Not a Combination Product	02/12/2010	
4	NDC:0135-0181-05	265 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
5	NDC:0135-0181-06	86 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
6	NDC:0135-0181-07	1 in 1 CARTON	02/12/2010	
6		12 in 1 PACKAGE; Type 0: Not a Combination Product		
7	NDC:0135-0181-14	160 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/12/2010	

## TUMS ULTRA

calcium carbonate tablet, chewable

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0135-0228
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ50PE7D, CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1000 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	



MINERAL OIL (UNII: T5L8T28FGP)

SODIUM POLYMETAPHOSPHATE (UNII: P1BM4ZH95L)

### Product Characteristics

Color	WHITE	Score	no score
Shape	ROUND	Size	19mm
Flavor	PEPPERMINT	Imprint Code	TUMS
Contains			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0228-01	12 in 1 PACKAGE; Type 0: Not a Combination Product	02/12/2010	
2	NDC:0135-0228-04	10 in 1 PACKAGE; Type 0: Not a Combination Product	02/12/2010	
3	NDC:0135-0228-05	160 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
4	NDC:0135-0228-06	72 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	
5	NDC:0135-0228-07	86 in 1 BOTTLE; Type 0: Not a Combination Product	02/10/2015	04/30/2018

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/12/2010	

## TUMS ULTRA

calcium carbonate tablet, chewable

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0135-0540
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ5OPE7D, CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1000 mg

### Inactive Ingredients

Ingredient Name	Strength
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<b>SUCROSE</b> (UNII: C151H8M554)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>SODIUM POLYMETAPHOSPHATE</b> (UNII: P1BM4ZH95L)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	

### Product Characteristics

<b>Color</b>	WHITE, RED, GREEN	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	19mm
<b>Flavor</b>	PEPPERMINT	<b>Imprint Code</b>	TUMS
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0540-01	86 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2010	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/12/2010	

**Labeler** - Haleon US Holdings LLC (079944263)