

**DENTEMP CANKER COVER- canker cover tablet, extended release
DOC Brands**

Dentemp Canker Cover

ACTIVE INGREDIENT

Active Ingredient (per each tablet)

Menthol 2.5 mg

PURPOSE

Purpose

Oral Pain Reliever

Indications for Use

Indications for the temporary relief of pain associated with Canker Sores

PREGNANCY OR BREASTFEEDING

If pregnant, or breast-feeding or taking prescription drugs, ask a health care professional

before use

WARNINGS

Warnings

Do not use this product for more than 7 days unless directed by a dentist or doctor.

Stop use and ask a dentist or doctor if .sore mouth symptoms do not improve in 7 days

.irritation, pain or redness worsens .swelling, rash or fever develops

Do not exceed recommended dosage.

Warnings

Keep out of reach of children. In case of accidental overdose, get medical help or contact a

Poison Control Center (1-800-222-1222) right away.

DIRECTIONS

Directions .adults and children 5 years of age and older, apply up to 3 tablets per day,

as needed. Place the tablet on a clean, dry finger with the white side up. Place the white side on

the sore and hold in place for 10 seconds. If sore is difficult to reach (in the fold between the

cheek and gum or near the teeth or lip) break the tablet along the score and use half. Some

discomfort may occur during the first few minutes, but will quickly subside, followed by hours of

soothing relief. Within 30 minutes the tablet forms a clear, gel-like bandage that seals and

protects the sore for hours before dissolving. Do not remove the tablet before it dissolves. In

case of discomfort, the tablet may be removed by gently peeling the tablet from the sides while

washing with warm water. Do not use any instrument to remove the tablet.

.children under 5 years, ask a doctor

STORAGE

Other information store in a cool, dry place.

INACTIVE INGREDIENTS

Inactive ingredients annato, carbomer homopolymer type a, citrus oil, hydroxypropyl cellulose, magnesium chloride, povidone K30, povidone K90, silicon dioxide, xylitol

QUESTIONS

Questions or comments? 833-363-2763 or consumeraffairs@doc-brands.com

DENTEMP CANKER COVER PRINCIPAL DISPLAY PANEL



Forms Protective Gel Barrier
Protects from Irritants
& Promotes Healing

Relieves Pain From:

- ✓ Canker Sores
- ✓ Mouth Sores
- ✓ Mouth Irritation



HELPS HEAL AS FAST AS 1 DAY*

NO sucrose, glucose, lactose or preservatives. Less than 1 calorie.

Dentemp® Canker Cover®

Oral Pain Reliever

Clinically Proven*

To Relieve Pain & Heal Sores
Helps Heal as Fast as 1 Day
Soothing Relief Up to 12 Hours
Stays On While Eating



6 COUNT

Healing Process:



11.437

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*Clinical on File
U.S. Pat. No. 7,943,169
www.dentemp.com
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Wayne, PA 19087 USA
Made in USA | D-211-007



Drug Facts

Active ingredient (per each tablet) Menthol 2 mg, Oral pain reliever

Indications For the temporary relief of pain associated with canker sores.

Warnings
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Stop use and ask a dentist or doctor if sore mouth symptoms do not improve in 7 days.
Irritation, pain or redness worsens *swelling, rash or fever develops.
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Dentemp® Canker Cover®
Oral Pain Reliever

DENTEMP CANKER COVER

canker cover tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73653-211
Route of Administration	ORAL, TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.5 mg in 150 mg

Inactive Ingredients

Ingredient Name	Strength
POVIDONE K30 (UNII: U725QWY32X)	
POVIDONE K90 (UNII: RDH86HJV5Z)	
XYLITOL (UNII: VCQ006KQ1E)	
ANNATTO (UNII: 6PQP1V1B6O)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
CITRUS LIMON FRUIT OIL (UNII: 0HNC1J1YED)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
CARBOMER HOMOPOLYMER TYPE A (UNII: F68VH75CJC)	

Product Characteristics

Color	pink (one side pink, one side white)	Score	no score
Shape	ROUND	Size	10mm
Flavor	CITRUS	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73653-211-07	6 in 1 CARTON	02/20/2020	
1	NDC:73653-211-06	150 mg in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	02/20/2020	

Labeler - DOC Brands (081254601)

