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

Dentemp Canker Cover 2ct

Menthol 2.5 mg

Oral Pain Reliever

For temporary relief of pain associated with Canker Sores

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.

If pregnant or breastfeeding or taking prescription drugs, ask a health care professional before use.

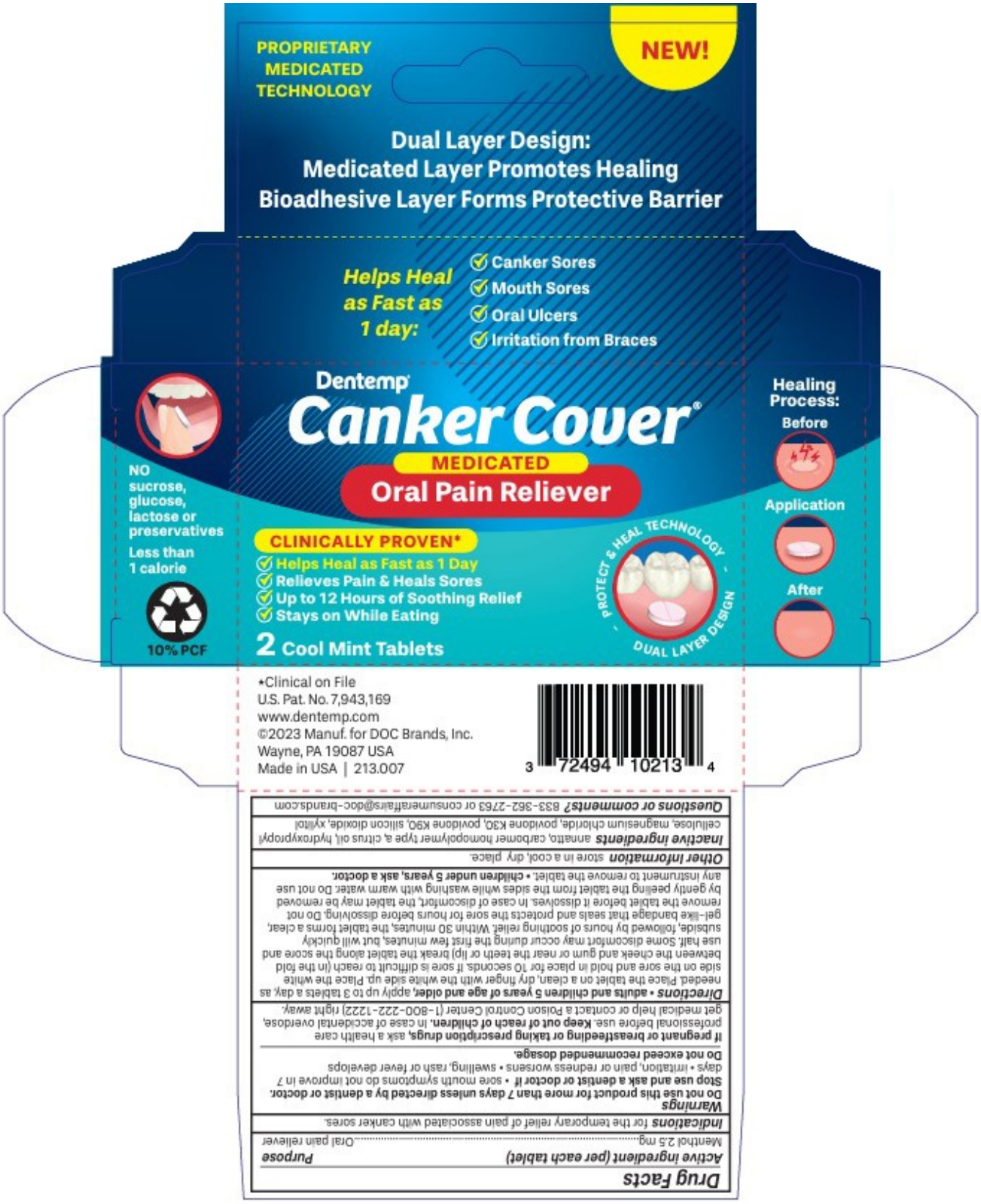
Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

adults and children 5 years of age and older, apply up to 3 tablets a 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.

store in a cool, dry place.

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

833-362-2763 or consumeraffairs@doc-brands.com



DENTEMP CANKER COVER

canker cover tablet, extended release

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73653-212

Route of Administration	TOPICAL, ORAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

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

Product Characteristics

Color	pink	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor	MINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73653-212-07	2 in 1 CARTON	02/01/2025	
1	NDC:73653-212-01	1 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	01/09/2025	

Labeler - DOC Brands (081254601)