

REMBRANDT INTENSE STAIN- sodium fluoride paste, dentifrice

Ranir 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.

373 Rembrandt Intense Stain Mint Whitening Toothpaste

Active ingredient

Sodium Fluoride (0.243%)

Purpose

Anticavity

Use

Aids in the prevention of dental cavities.

Warnings

Keep out of reach of children under 6 years of age.

If more than amount used for brushing is accidentally swallowed, seek medical help or contact a Poison Control Center right away.

Directions

Adults and children 2 years of age and older: Brush teeth thoroughly (for at least 1 minute), preferably after each meal or at least 2 times a day, or as directed by a dentist or doctor.

Children under 6 years of age: Use a pea-sized amount and instruct in good brushing and rinsing habits (to minimize swallowing). Supervise children as necessary until capable of using without supervision.

Children under 2 years of age: Consult a dentist or doctor. Do not swallow.

Other information

- Store at 20°-25°C (68°-77°F).
- Do not use if inner foil is torn, cut or missing.

Inactive ingredients

Glycerin, Hydrated Silica, Water, Sorbitol, Tetrapotassium Pyrophosphate, Flavor, Titanium Dioxide, Sodium Lauroyl Sarcosinate, Cellulose Gum, Lauryl Glucoside, PVP, Sodium Sulfate, Cocamidopropyl Betaine, Xanthan Gum, Sodium Saccharin, Sucralose.

QUESTIONS OR COMMENTS?

Email consumercare@rembrandt.com, or call 800-548-3663.

Distributed by Ranir, LLC,

Grand Rapids, MI 49512

©2018 Ranir, LLC 80102AB

Directions: updated use instructions 7/2/2019

Principal Display Panel

I INTENSE STAIN®

REMBRANDT

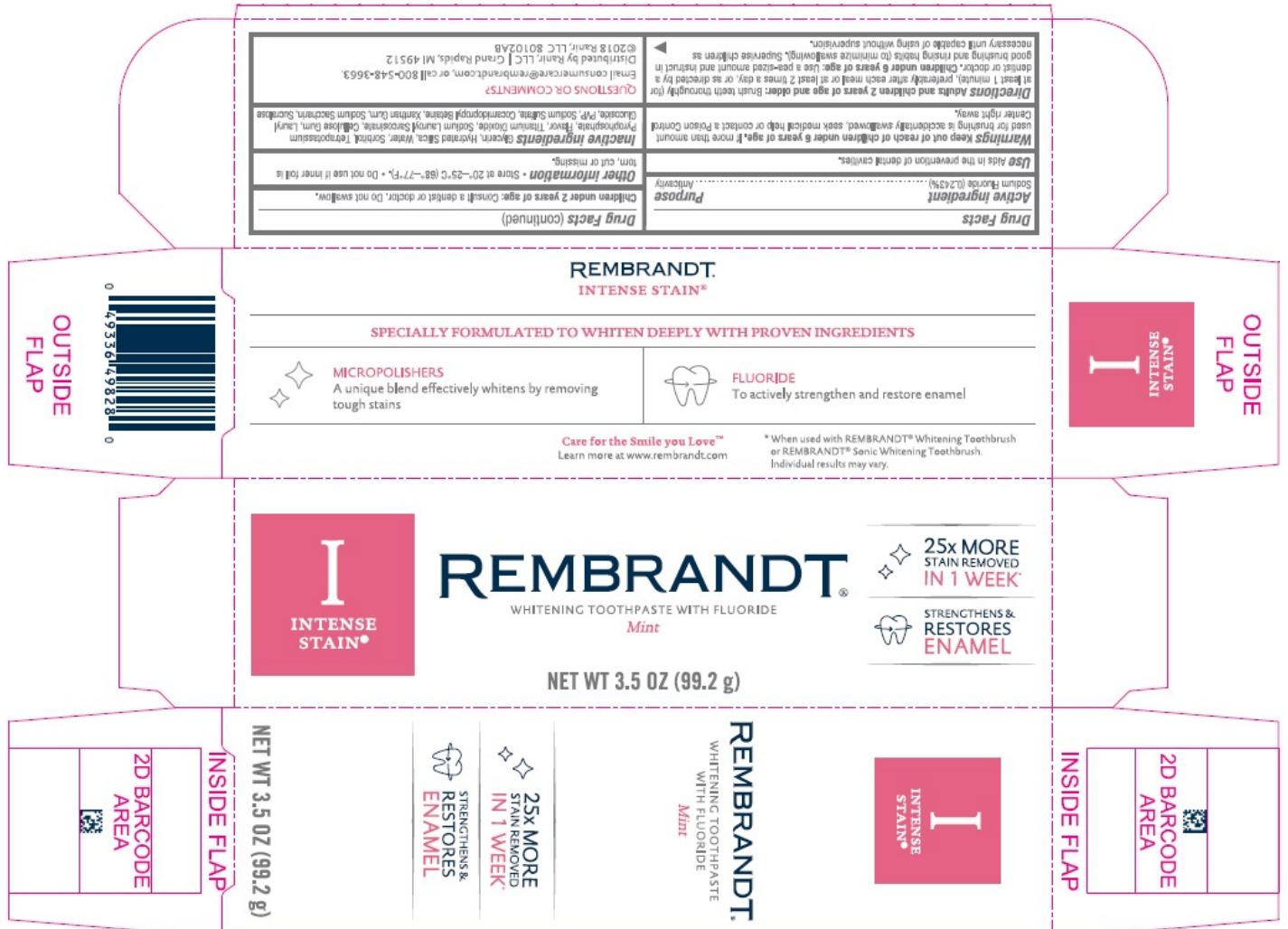
WHITENING TOOTHPASTE WITH FLUORIDE

Mint

25x MORE STAIN REMOVED IN 1 WEEK

STRENGTHENS & RESTORES ENAMEL

NET WT 3.5 OZ (99.2 g)



REMBRANDT INTENSE STAIN

sodium fluoride paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66923-373
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	24.3 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
POTASSIUM PYROPHOSPHATE (UNII: B9W4019H5G)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM LAUROYL SARCO SINATE (UNII: 632GS99618)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66923-373-02	1 in 1 CARTON	07/25/2016	
1	NDC:66923-373-01	99.2 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part355	07/25/2016	

