

PERIACTIVE- sambucus nigra flower, echinacea purpurea, and centella asiatica mouthwash
Izun Pharmaceuticals Corp

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

PeriActive®

Drug Facts:

Active Ingredients (in each 15ml)	Purposes
▪ Sambucus nigra 3x	Irritation/gingivitis reducer gum repair stimulator
▪ Echinacea purpurea 4x	Irritation/gingivitis reducer
▪ Hydrocotyle asiatica 4x	Irritation/gingivitis reducer

Uses

- Helps protect against gingivitis
- Helps reduce gum bleeding and soreness
- Helps reduce plaque formation
- Helps reduce gum inflammation
- Promotes gingival healing

Warnings

Allergy alert

Do not use if allergic to any of the ingredients. Stop use and ask a dentist if:

- gingivitis, bleeding, or redness persists for more than 2 weeks
- you have painful or swollen gums, pus from the gum line, loose teeth, or increasing spacing between the teeth. These may be the signs of periodontitis, a serious form of gum disease.

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

In case more than is used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

Directions

adults and children 12 years and older	rinse for 60 seconds with 15ml (3 teaspoonful) twice a day
children 6-12 years	rinse for 60 seconds with 10ml (2 teaspoonful) twice a day
children under 6 years of age	do not use

- do not eat, drink or rinse your mouth for 30 minutes after use

- do not swallow
- children 6 years to under 12 years of age: supervise use

Other information

This rinse is not intended to replace brushing or flossing

- store at room temperature 15-30°C (59-86°F)
- keep out of direct sunlight

Inactive ingredients

cetylpyridinium chloride, cool mint flavor¹, disodium EDTA, FD&C Blue No. 1, lactic acid, maltodextrin, menthol, nat. cinnamon flavor¹, PEG-40 hydrogenated castor oil, propylene glycol, sodium benzoate, sorbitol, sucralose, water, wintergreen flavor¹

¹ contains one or more of these ingredients

Questions?

Call 1-800-956-2595

Distributed By: straumann

Institut Straumann AG

Peter Merian-Weg 12 | 4002 Basel | Switzerland

PRINCIPAL DISPLAY PANEL - 500 mL Bottle Label

PeriACTIVE®

DISTRIBUTED BY STRAUMANN

ADVANCED GUM CARE

ORAL RINSE

Powered by Nature

CLINICALLY PROVEN
to Reduce Inflammation

KILLS BACTERIA

REDUCES
Gum Bleeding & Soreness

PROTECTS
Against Plaque & Gingivitis

Fresh Mint

HOMEOPATHIC

500 mL e (16.9 fl. oz.)

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Institut Straumann AG
Peter Merian-Weg 12 | 4002 Basel | Switzerland
Tel +41 61 965 11 11 | Fax +41 61 965 11 01
E-mail: info@straumann.com | www.straumann.com

See bottom of bottle for expiry date

Safety-sealed for your protection.
Do not use if plastic sleeve is broken or missing.

(01)0085752008019 (17)21115(10)0001880

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gum repair stimulator
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Izun Pharmaceuticals Corp
One Rocket Plaza
New York, N.Y. 10020

GMDN CODE: 42345

PERIACTIVE

sambucus nigra flower, echinacea purpurea, and centella asiatica mouthwash

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49589-199
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Sambucus nigra flower (UNII: 07V4DX094T) (Sambucus nigra flower - UNII:07V4DX094T)	Sambucus nigra flower	3 [hp_X] in 1 mL
Echinacea purpurea (UNII: Q17G114Y98) (Echinacea purpurea - UNII:Q17G114Y98)	Echinacea purpurea	4 [hp_X] in 1 mL

Centella asiatica (UNII: 7M867G6T1U) (Centella asiatica - UNII:7M867G6T1U)	Centella asiatica	4 [hp_X] in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
cetylpyridinium chloride (UNII: D9OM4SK49P)	
edetate disodium (UNII: 7FLD91C86K)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
maltodextrin (UNII: 7CVR7L4A2D)	
polyoxyl 40 hydrogenated castor oil (UNII: 7YC686GQ8F)	
propylene glycol (UNII: 6DC9Q167V3)	
sodium benzoate (UNII: OJ245FE5EU)	
sorbitol (UNII: 506T60A25R)	
sucralose (UNII: 96K6UQ3ZD4)	
water (UNII: 059QF0KO0R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49589-199-50	12 in 1 CARTON	02/01/2019	
1		500 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49589-199-10	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		02/01/2019	

Labeler - Izun Pharmaceuticals Corp (624194523)

Revised: 2/2019

Izun Pharmaceuticals Corp