TUCKS TRIPLE RELIEF HEMORRHOIDAL- white petrolatum, glycerin, pramoxine hydrochloride, and phenylephrine hydrochloride cream
Blistex Inc.

Tucks® Triple Relief Hemorrhoidal

Drug Facts

Active ingredients	***Purpose***
Glycerin 15% (w/w)	Protectant
Phenylephrine HCl 0.25% (w/w)	Vasoconstrictor
Pramoxine HCl 1.0% (w/w)	Local anesthetic
White Petrolatum 16% (w/w)	Protectant

Uses

- for the temporary relief of pain, soreness, or burning
- helps relieve the local itching and discomfort associated with hemorrhoids
- temporarily shrinks hemorrhoidal tissue
- temporarily provides a coating for relief of anorectal discomforts
- temporarily protects the inflamed, irritated anorectal surface to help make bowel movements less painful
- temporarily forms a protective coating over inflamed tissues to help prevent drying of tissues
- may provide a cooling sensation

Warnings

For external use only

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- difficulty in urination due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are

- presently taking a prescription drug for high blood pressure or depression.

When using this product

- do not exceed the recommended daily dosage unless directed to do so by a doctor
- do not put directly in the rectum by using fingers or any mechanical device or applicator

Stop use and ask a doctor if

- bleeding occurs
- condition worsens or does not improve within 7 days
- allergic reactions develop due to ingredients in this product
- the symptom being treated does not subside or if redness, irritation, swelling, pain, or other symptoms develop or increase

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults:

- when practical, clean the affected area with mild soap and warm water and rinse thoroughly
- gently dry by patting or blotting with toilet tissue or a soft cloth before application of this product
- when first opening the tube, remove foil seal. Do not use if foil seal is broken.
- apply externally or in the lower portion of the anal canal only
- apply externally to the affected area up to 4 times daily
- for application in the lower anal canal, remove cover from dispensing cap. Attach dispensing cap to tube. Lubricate dispensing cap well, then gently insert dispensing cap partially into the anus.
- thoroughly clean dispensing cap after each use and replace cover
- children under 12 years of age: ask a doctor

Other information

www.tucks.com

Inactive ingredients

arachidyl alcohol, arachidyl glucoside, beeswax, behenyl alcohol, disodium EDTA, ethylhexylglycerin, hamamelis virginiana (witch hazel) water, microcrystalline wax, phenoxyethanol, polyglyceryl-3 diisostearate, polysorbate 60, purified water, sorbitan isostearate, stearyl behenate

PRINCIPAL DISPLAY PANEL - 26 g Tube Carton

NEW

TUCKS®

TRIPLE RELIEF
HEMORRHOIDAL CREAM

Hemorrhoidal Cream

MAXIMUM STRENGTH
PAIN RELIEF

helps relieve pain
from burning, itching
& discomfort

shrinks swollen
hemorrhoidal tissue

cools & soothes
sensitive areas

BRAND USED
BY HOSPITALS

with
WITCH
HAZEL

Net Wt. 0.9 oz. (26 g) - 1 tube

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10157-2126
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
WHITE PETROLATUM (UNII: B6E5W8RQJ4) (WHITE PETROLATUM - UNII:B6E5W8RQJ4)	WHITE PETROLATUM	16 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	15 g in 100 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	1 g in 100 g
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	0.25 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
STEARYL BEHENATE (UNII: A68S9UI39D)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
HAMAMELIS VIRGINIANA TOP WATER (UNII: NT00Y05A2V)	
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46P231IQV8)	
ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU)	
DOCOSANOL (UNII: 9G1OE216XY)	
ARACHIDYL GLUCOSIDE (UNII: 6JVW35JOOJ)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:10157-2126-1	1 in 1 CARTON	01/06/2025	
1		26 g in 1 TUBE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC monograph drug	M015	01/06/2025	

Labeler - Blistex Inc. (005126354)

Establishment

Name	Address	ID/FEI	Business Operations
Blistex Inc.		005126354	MANUFACTURE(10157-2126)

Revised: 12/2024 Blistex Inc.