SALLY HANSEN OUCH-RELIEF NUMBING WIPES- benzocaine cloth Coty US 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.

Drug Facts (Numbing Wipes)

Active ingredient

6% Benzocaine

Purpose

Topical Analgesic

Uses

for temporary relief of pain and itching associated with minor skin irritations due to hair removal

Warnings

Allergy Alert: do not use this product if you have a history of allergy to local anesthetics such as procaine, butacaine, benzocaine or other "caine" anesthetics

For external use only. Not intended for use by children under 13 years of age

When using this product

avoid contact with eyes

Stop use and ask a doctor if

- condition worsens
- symptoms last for more than 7 days or clear up and occur again within a few days

Do not use

over raws surfaces, or blistered areas

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- Rub well (see enclosed instructions)
- Use wipes no more than 3 to 4 times daily. Discard wipe after use

Adults and children 13 years of age and older apply to :he area to be treated 10 minutes before hair removal Children under 13 years of age consult a doctor

Other information

As with all products containing benzocaine, localized allergic reactions may occur after prolonged or repeated use

Inactive ingredients

ALCOHOL DENAT., AQUA/WATER/EAU, PROPYLENE GLYCOL, METHYLPARABEN

Questions or comments?

Call us at 1-800-953-5080 9:00AM - 5:00PM EST

Principal Display Panel - Carton Label

OUCH-RELIEF™ NUMBING WIPES

PRE-WAX

PAIN-RELIEF TECHNOLOGY

FOR FACE & BODY

Pre-Treatment Help Reduce Pain While Waxing

Soothe & Prepare Skin Before Waxing

Ready & Easy to Use

9 PRE-WAX WIPES

Total 9 Ouch-Relief™ Numbing Wipes, Instructions

Dermatologist & Salon Tested



SALLY HANSEN OUCH-RELIEF NUMBING WIPES

benzocaine cloth

Product Information							
Product Type	HUMAN OTC DRUG	Item Code (S	ource)	NDC:661	.84-156		
Route of Administration	TOPICAL						
Active Ingredient/Active	Majaty						
Active Ingredient/Active	моюту						
Ingre	dient Name		Basis of Stre	ength	Strength		

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
WATER (UNII: 059QF0KO0R)		
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
METHYLPARABEN (UNII: A2I8C7HI9T)		

l	P	Packaging						
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
	1	NDC:66184-156- 01	9 in 1 CARTON; Type 0: Not a Combination Product	01/01/2015				

Marketing In	arketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part333B	01/01/2015				

Labeler - Coty US LLC (039056361)

Revised: 10/2021 Coty US LLC