WHITE PETROLATUM- petrolatum ointment
Dynarex Corporation

1140 White Petrolatum NDC 67777-005-00

1141 White Petrolatum NDC 67777-005-10

1145 White Petrolatum NDC 67777-005-50

1147 White Petrolatum NDC 67777-005-70

Active Ingredient

White Petrolatum 100%

Purpose

Skin Protectant

Uses

- Temporarily protects minor cuts, scrapes, burns
- Temporarily protects and helps relieve chafed, chapped, or cracked skin and lips
- Helps prevent and protect from the drying effects of wind and cold weather
- Protects chafed skin due to diaper rash

Warnings

For external use only

Do not use on

- Deep or puncture wounds
- Animal bites
- Serious burns

When using this product

- Avoid contact with eyes

Stop use and ask a doctor if

- Condition worsens
- Symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-12222) right away.

Directions

• For diaper rash: Change wet and soiled diapers promptly, cleanse the diaper area, and allow to dry. Apply ointment liberally as often as necessary with each diaper change, especially at bedtime or anytime when exposure to wet diapers may be prolonged.

• For other uses: Apply as needed.

Other Information

• Store at room temperature 15º-30ºC (59º-86ºF)

• Avoid excessive heat

• Tamper evident. Do not use if seal is torn, cut, or opened

Questions?

1-888-396-2739 Monday - Friday, 9AM - 5PM EST.

Inactive Ingredient

Label

1140 White Petrolatum

Label

1140 White Petrolatum

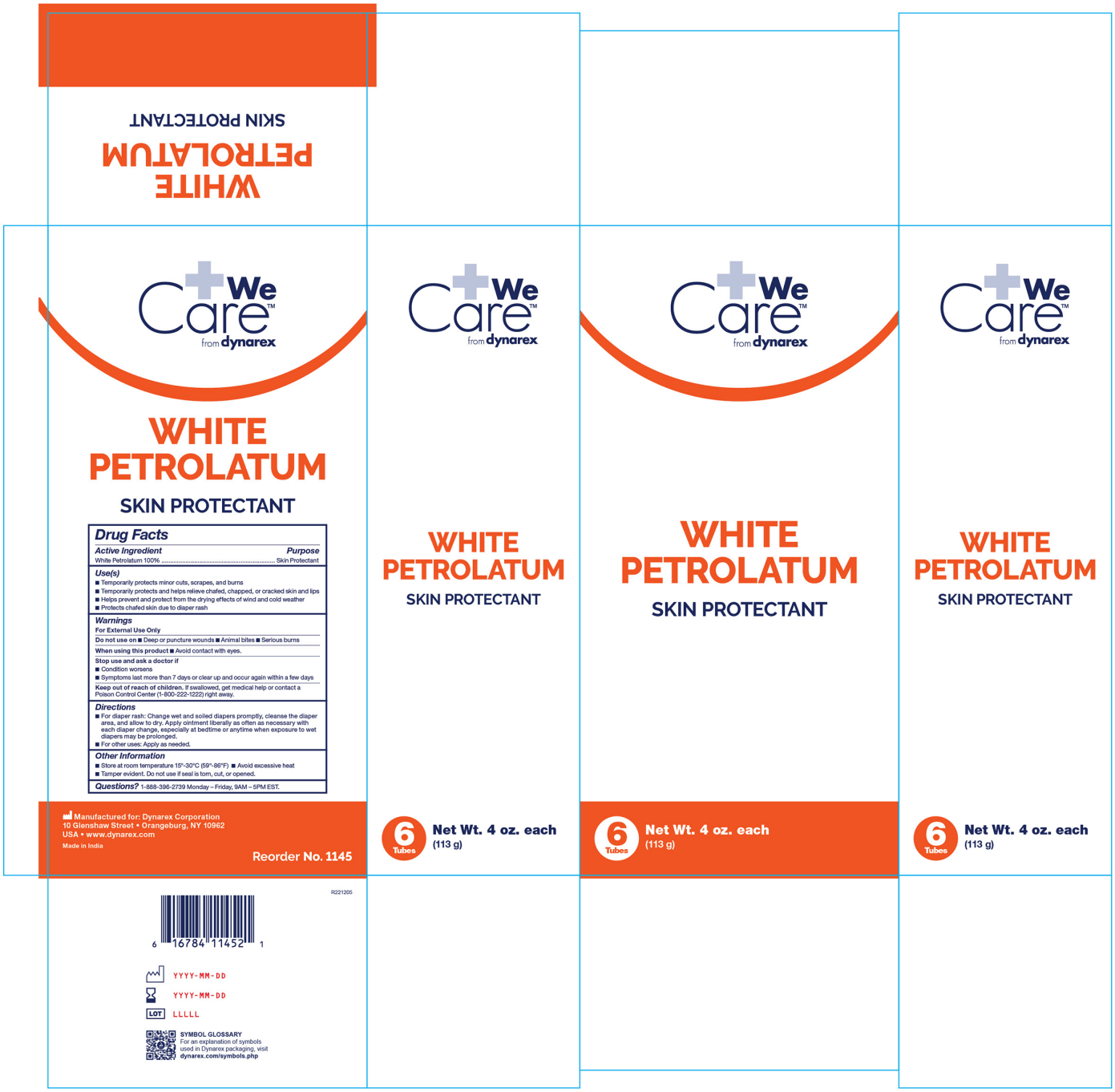
Label

1141 White Petrolatum

Label

1147 White Petrolatum

Label



1145 White Petrolatum

WHITE PETROLATUM

petrolatum ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

WHITE PETROLATUM (UNII: B6E5W8RQJ4) (WHITE PETROLATUM - UNII:B6E5W8RQJ4)	WHITE PETROLATUM	1 g in 1 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-005-00	864 in 1 CASE	10/26/2018	
1	NDC:67777-005-01	144 in 1 BOX		
1		5 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:67777-005-10	72 in 1 CASE	10/26/2018	
2	NDC:67777-005-11	1 in 1 BOX		
2		28.4 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:67777-005-50	72 in 1 CASE	10/26/2018	
3	NDC:67777-005-51	6 in 1 BOX		
3		113 g in 1 TUBE; Type 0: Not a Combination Product		
4	NDC:67777-005-70	12 in 1 CASE	10/26/2018	
4	NDC:67777-005-71	425 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	10/26/2018	

Labeler - Dynarex Corporation (008124539)

Revised: 1/2024 Dynarex Corporation