FIFTHPULSE BACITRACIN ZINC 10Z- bacitracin zinc ointment Front Pharmaceutical PLC

	MarifthPulse NDC: Net wt. 10z (28.4g) Bacitracin Z Ointment U	SP			
	Active ingredient (each gram contains) Purpose Bacitracin Zinc 500 units First Aid Antibiotic Uses First aid to help prevent infection in I minor cuts I scrapes I burns First Aid Antibiotic Warnings I For external use only. Do not use I in the eyes I over large areas of the body I if you are allergic to any of the ingredients I longer than 1 week unless directed by a doctor. Sta doctor before use in care of deep puncture wounds, animal bites, or serious burns. Stop use and ask a doctor if I the condition persists or gets worse I arash or allergic reaction develops. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately. Directions I clean the affected area I apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily I may be covered with a sterile bandage Other information I store between 15° to 25° C (59° to 77°F) Lot No & Expiration Date: See box and tube crimp. Inactive ingredients: Mineral oil, Petrolatum	Manufactured for: WeCare Products LLC 20000 NE 15th CT, Questions or Comments? Miarni, FL 33179 Please call 1-888-209-5201			
版位: 3mm 105mm					
ווווכטו					













FIFTHPULSE BACITRACIN ZINC 10Z

bacitracin zinc ointment

Product Information									
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:69571-013					
Route of Administration	TOPICAL								
A stine la sue die st/A stine Maistre									
Active Ingredient/Active Moiety									
Ingredient Name Basis of			Basis of Stre	ngth	Strength				
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I) BAG			BACITRACIN		500 [iU] in 1 g				
Inactive Ingredients									
Ingredient Name				Strength					
PETROLATUM (UNII: 4T6H12BN9U)									

Product Characteristics Color Score 105mm Shape Size 105mm Flavor Imprint Code 105mm Contains Imprint Code 105mm Package Description Marketing Start Date 1 NDC:69571-013- 01 28.4 g in 1 TUBE; Type 0: Not a Combination Product 07/15/2025											
ColorSocreInterpretenderInterpretenderShapeSize105mmFlavorImprint CodeImprint CodeColorInterpretenderImprint CodeStateMarketing Start DateMarketing DateNDC:69571-013- 0128.4 g in 1 TUBE; Type 0: Not a Combination Product07/15/2025											
Shape Size 105mm Flavor Imprint Code Imprint Code Imprint Code Contains Imprint Code Imprint Code Imprint Code Imprint Code Imprint Code Imprin											
Flavor Imprint Code Contains Imprint Code Package Description Marketing Start Marketing Start Date Date 1 NDC:69571-013- 01 28.4 g in 1 TUBE; Type 0: Not a Combination 07/15/2025											
Item Code Package Description Marketing Start Date Marketing Date NDC:69571-013- 28.4 g in 1 TUBE; Type 0: Not a Combination Product 07/15/2025 07/15/2025											
Item Code Package Description Marketing Start Date Marketing Start Date 1 NDC:69571-013- 01 28.4 g in 1 TUBE; Type 0: Not a Combination Product 07/15/2025											
#Item CodePackage DescriptionMarketing Start DateMarketing Date1NDC:69571-013- 0128.4 g in 1 TUBE; Type 0: Not a Combination 											
# Item Code Package Description Marketing Start Date Marketing Date 1 NDC:69571-013- 01 28.4 g in 1 TUBE; Type 0: Not a Combination Product 07/15/2025 07/15/2025											
# Item Code Package Description Date Date 1 NDC:69571-013- 01 28.4 g in 1 TUBE; Type 0: Not a Combination Product 07/15/2025	Packaging										
01 Product	-										
Marketing Information											
Marketing Information											
Marketing Information											
MarketingApplication Number or MonographMarketing StartMarketCategoryCitationDateDate											
OTC Monograph Drug M004 07/15/2025											

Labeler - Front Pharmaceutical PLC (530897792)

Registrant - Front Pharmaceutical PLC (530897792)

Establishment								
Name	Address	ID/FEI	Business Operations					
Front Pharmaceutical PLC		530897792	manufacture(69571-013)					

Revised: 7/2025

Front Pharmaceutical PLC