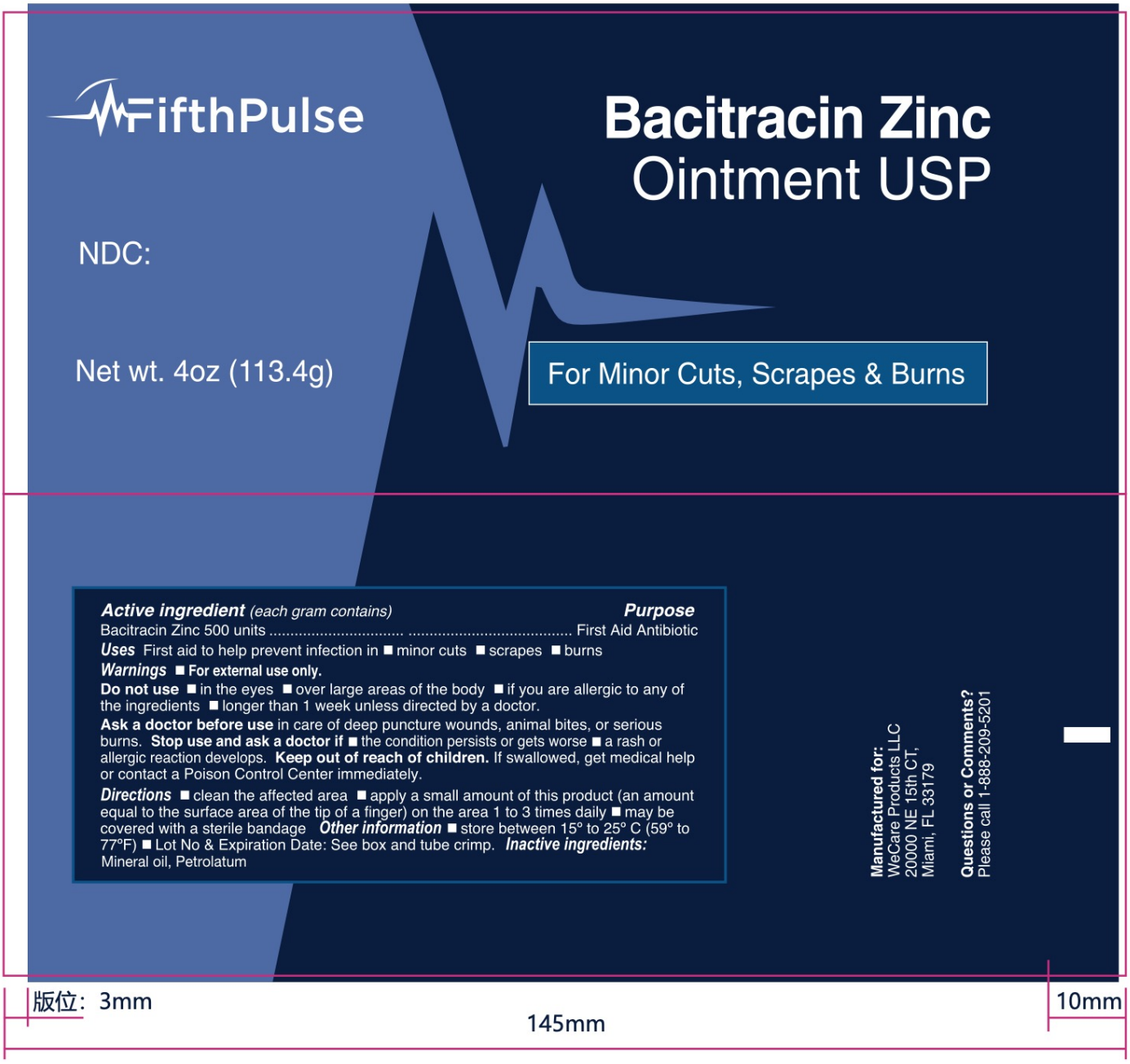
FIFTHPULSE BACITRACIN ZINC 4OZ- bacitracin zinc ointment
Front Pharmaceutical PLC



FifthPulse

Bacitracin Zinc Ointment USP

NDC:

Net wt. 4oz (113.4g)

For Minor Cuts, Scrapes & Burns

Active ingredient *(each gram contains)* ***Purpose***
Bacitracin Zinc 500 units .... First Aid Antibiotic

Uses First aid to help prevent infection in ■ minor cuts ■ scrapes ■ burns

Warnings ■ **For external use only.**

Do not use ■ in the eyes ■ over large areas of the body ■ if you are allergic to any of the ingredients ■ longer than 1 week unless directed by a doctor.

Ask a doctor before use in care of deep puncture wounds, animal bites, or serious burns. **Stop use and ask a doctor if** ■ the condition persists or gets worse ■ a rash or allergic reaction develops. **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center immediately.

Directions ■ clean the affected area ■ 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 ■ may be covered with a sterile bandage ***Other information*** ■ 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,
Miami, FL 33179

Questions or Comments?
Please call 1-888-209-5201

版位：3mm

145mm

10mm

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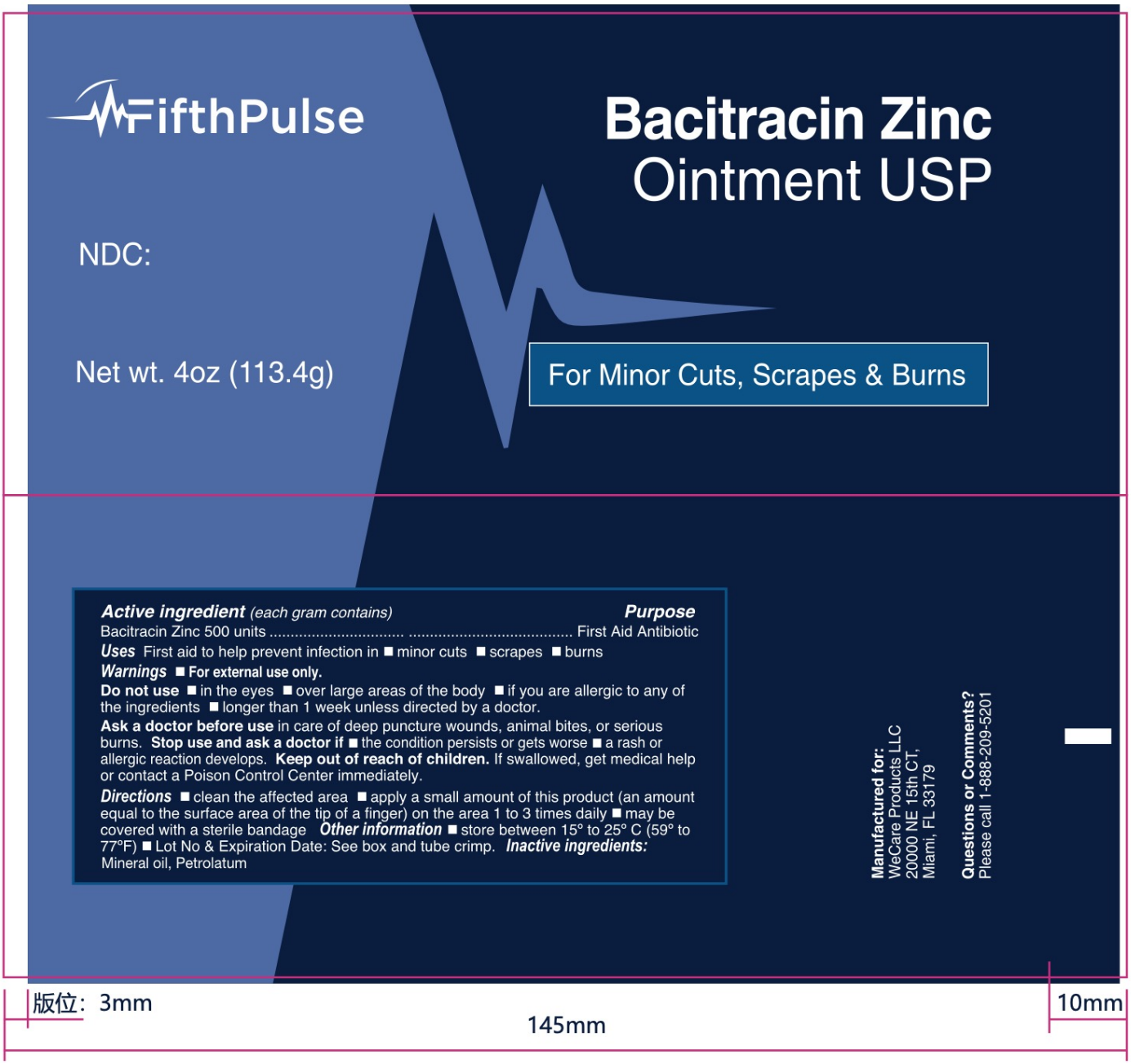

FIFTHPULSE BACITRACIN ZINC 4OZ

bacitracin zinc ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69571-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [iU] in 1 g

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics

Color		**Score**	
Shape		**Size**	145mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69571-012-02	113.4 g in 1 TUBE; Type 0: Not a Combination Product	07/15/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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(69571-012)

Revised: 7/2025 Front Pharmaceutical PLC