Polymyxin B Sulfate
Bacitracin Zinc
Neomycin Sulfate

Terramycin® PLUS

710 mcg/ 10 mg/ 5mg per g Topical Ointment

Topical Antimicrobial Combinations

1.0 NAME OF THE PRODUCT

Terramycin® Plus

2.0 DESCRIPTION OF THE PRODUCT

Polymyxin B Sulfate, Bacitracin Zinc, Neomycin Sulfate (Terramycin ® PLUS) is available in tube as 3.5 mg and 5 mg.

3.0 WHAT IS IN THE MEDICINE?

Polymyxin B Sulfate, Bacitracin Zinc, Neomycin Sulfate. Terramycin ® PLUS (Polymyxin B Sulfate, Bacitracin Zinc, Neomycin Sulfate)

4.0 STRENGTH OF THE MEDICINE

Each gram contains Polymyxin B Sulfate 710 mcg, Bacitracin Zinc 10 mg, Neomycin Sulfate 5 mg.

5.0 WHAT IS THIS MEDICINE USED FOR?

First aid for the prevention and treatment of infection in minor cuts, scrapes, burns, abrasions, insect bites, pimples, pustules and other skin breaks.

6.0 HOW MUCH AND HOW OFTEN SHOULD YOU USE THIS MEDICINE?

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.

7.0 WHEN SHOULD YOU NOT TAKE THIS MEDICINE?

Not intended for use on existing scars.

8.0 CARE THAT SHOULD BE TAKEN WHEN TAKING THIS MEDICINE?

- For external use only.
- Do not use in the eyes or apply over large areas of the body.
- In case of deep puncture wounds, animal bites or serious burns, consult a physician.
- Stop use and consult a physician if the condition persists or gets worse, or if a rash or other allergic reaction develop.
- Do not use longer than 1 week unless directed by a physician.

9.0 UNDESIRABLE EFFECTS OF THIS MEDICINE
Not Applicable

10.0 WHAT OTHER MEDICINE OR FOOD SHOULD BE AVOIDED WHILE TAKING THIS MEDICINE?
Not applicable

11.0 WHAT SHOULD YOU DO IF YOU MISS A DOSE?
Not Applicable

12.0 SIGNS AND SYMPTOMS OF OVERDOSE
Not Applicable

13.0 WHAT TO DO WHEN YOU HAVE TAKEN MORE THAN THE RECOMMENDED DOSAGE?
Not Applicable

14.0 HOW SHOULD YOU KEEP THIS MEDICINE?
Store at temperatures not exceeding 30°C

15.0 WHEN SHOULD YOU CONSULT YOUR DOCTOR?
If you experience any of the side effects, consult your doctor.

16.0 FDA REGISTRATION NUMBER
710 mcg/ 10 mg/ 5 mg per g Topical Ointment - DRP-3570

17.0 DATE OF FIRST AUTHORIZATION/RENEWAL OF AUTHORIZATION
710 mcg/ 10 mg/ 5 mg per g Topical Ointment: 20 November 2018

Keep out of reach of children.

For suspected adverse drug reaction, report to the FDA: www.fda.gov.ph

Seek medical attention immediately at the first sign of any adverse drug reaction.

Manufactured by:

Interphil Laboratories, Inc.
Canlubang Industrial Estate,
Bo. Pittland, Cabuyao, Laguna

Marketing Authorization Holder:

Pfizer, Inc.
19F-20F, 8 Rockwell Building, Hidalgo Drive,
Rockwell Center, Poblacion, Makati City 1210
Metro Manila, Philippines

Under Authority of Pfizer Inc., N.Y., U.S.A.

Revision No.: PIL 0.1
Revision date: 03 June 2022
Reference: NA/MAH office address update
Reference Date: NA