

Registration Number : DRP-1001

**SPECIAL CONDITION:**

Provided that nothing in the registration of the product herein granted shall be interpreted or construed as an endorsement or representation by FDA, that Registrant has the right or privilege to the use of the name or brand so registered; Registrant hereby agrees and affirms to indemnify and/or hold FDA free and harmless against any and all third-party claims on infringement of patent, trademark or intellectual property right arising from the registration of the product.

- |                                     |          |  |
|-------------------------------------|----------|--|
| <input type="checkbox"/>            | <b>A</b> | This is subject to batch notification.   |
| <input type="checkbox"/>            | <b>B</b> | This is subject to lot release certification.  |
| <input type="checkbox"/>            | <b>C</b> | This is subject to compliance with the requirements under FDA Circular No. 2013-004 for Monitored Release (MR) drug products.  |
| <input checked="" type="checkbox"/> | <b>D</b> | Subject to post-marketing surveillance of the marketing authorization holder's strict compliance to the Generic Labeling Requirements following the applicable provisions of A.O. No. 2016-0008 for drug products for human use and A.O. No. 105 s. 1991 for veterinary drug products. |
| <input type="checkbox"/>            | <b>E</b> | Submit a satisfactory Bioequivalence Study Report or Biowaiver (whichever is applicable) within the validity of this CPR in accordance with FDA Circular No. 2016-019.   |
| <input type="checkbox"/>            | <b>F</b> | Dangerous Drug - To be prescribed by a PDEA S-2 licensed practitioner in a DOH (yellow) prescription form. It is a habit-forming drug.   |
| <input type="checkbox"/>            | <b>G</b> | Dangerous Drug - To be prescribed by a PDEA S-2 licensed practitioner in a personalized ordinary prescription form. It is a habit-forming drug.  |
| <input type="checkbox"/>            | <b>H</b> | Patient Information Leaflet - Appropriate information for the consumers shall be written in Filipino and/or local dialects, as appropriate.  |
| <input type="checkbox"/>            | <b>I</b> | Submit a Certificate of Good Manufacturing Practice (GMP) Compliance of Foreign Drug Manufacturer(s) within the validity of this CPR in accordance with A. O. No. 2013-0022 and FDA Circular 2014-016.   |
| <input type="checkbox"/>            | <b>J</b> | Review of the submitted Bioequivalence Study Report or Biowaiver, whichever is applicable, shall be completed by the FDA within the validity of this CPR; correspondingly, this CPR shall be revoked if product interchangeability has not been established.                           |
| <input type="checkbox"/>            | <b>K</b> | Subject to satisfactory compliance to the post-approval commitments detailed in the letter accompanying this CPR.  |

**REMARKS:**

- The product has fulfilled the requirement of product interchangeability/equivalence evidence with the submission of a satisfactory Biowaiver based on the Biopharmaceutics Classification System (BCS).
- This Certificate of Product Registration (CPR) is revalidated to reflect its full validity until 30 January 2023. This supersedes CPR No. FDA-0295525 issued on 3 April 2018 valid until 30 January 2021.



FDA-20208066W4C652EACAVWYW9R

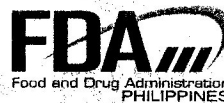


Republic of the Philippines

Department of Health

**FOOD AND DRUG ADMINISTRATION**

Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

**CERTIFICATE OF PRODUCT REGISTRATION**

Pursuant to the provisions of Republic Act (R.A.) No. 3720 as amended, known as the Foods, Drugs, Devices and Cosmetics Act, and consistent with R.A. No. 6675, known as the Generics Act of 1988, and R.A. No. 9711, otherwise known as the Food and Drug Administration Act of 2009, the product described hereunder has been found to conform with the requirements and standards for marketing authorization of pharmaceutical products per existing regulations in force as of date hereof.

**Registration Number** : **DRP-1001**

**Generic Name** : Metronidazole  
**Brand Name** : Flagex  
**Dosage Strength & Form** : 500 mg Tablet  
**Pharmacologic Category** : Antiprotozoal  
**Classification** : Prescription Drug (Rx)  
**Approved Shelf-life** : 24 months  
**Storage Condition** : Store at temperatures not exceeding 30°C.  
**Packaging** : Blister Pack x 10's (Box of 100's)

**Manufacturer** : New Myrex Laboratories, Inc.  
Bo. Catmon, Sta. Maria, Bulacan

The marketing authorization shall be valid until **30 January 2023** subject to the conditions listed on the reverse side. No change in the formulation, labelling and commercial presentation of this product shall be made at any time during the effectivity of this registration without prior written approval of this Office.

This marketing authorization is subject to suspension, cancellation or recall should any violation of R.A. No. 3720, R.A. No. 6675 and R.A. No. 9711 and/or regulations issued thereunder involving the product be committed.

Witness My Hand and Seal of this Office, this **19 November 2019**.

**By Authority of the Director General**  
**Per FDA Order No. 2016-005**

  
**ATTY. KATHERINE M. AUSTRIA-LOCK**

Officer-in-Charge  
Center for Drug Regulation and Research

REG. STATUS : Automatic Renewal (Revalidation)  
AMOUNT : Php 10,100.00  
OR NUMBER : 0947387  
DATE : 23 January 2018  
CODE : 433-220

BAR CODE :  
DOC TRACK :



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