CENTER FOR DRUG EVALUATION AND RESEARCH

APPLICATION NUMBER:

209176Orig1s000

PROPRIETARY NAME REVIEW(S)

PROPRIETARY NAME MEMORANDUM

Division of Medication Error Prevention and Analysis (DMEPA)
Office of Medication Error Prevention and Risk Management (OMEPRM)
Office of Surveillance and Epidemiology (OSE)
Center for Drug Evaluation and Research (CDER)

*** This document contains proprietary information that cannot be released to the public***

Date of This Review: September 1, 2016

Application Type and Number: NDA 209176

Product Name and Strength: Radicava (edaravone) Injection

0.3 mg/mL

Total Product Strength: 30 mg/100 mL

Product Type: Single-ingredient

Rx or OTC: Rx

Applicant/Sponsor Name: Mitsubishi Tanabe

Panorama #: 2016-8945213

DMEPA Primary Reviewer: Ebony Whaley, PharmD, BCPPS

DMEPA Team Leader: Lolita White, PharmD

Reference ID: 3980833

1 INTRODUCTION

This memorandum is to reassess the proposed proprietary name, Radicava, which was found conditionally acceptable under PIND 126396 on May 9, 2016.^a We note that all product characteristics remain the same.

2 METHODS AND DISCUSSION

2.1 MISBRANDING ASSESSMENT

The Office of Prescription Drug Promotion (OPDP) determined that the proposed name would not misbrand the proposed product. DMEPA and the Division of Neurology Products (DNP) concurred with the findings of OPDP's assessment of the proposed name.

2.2 SAFETY ASSESSMENT

For re-assessment of the proposed proprietary name, DMEPA evaluated the previously identified names of concern considering any lessons learned from recent post-marketing experience, which may have altered our previous conclusion regarding the acceptability of the proposed proprietary name. Additionally, DMEPA searched the USAN stem list to determine if the name contains any USAN stems as of the last USAN updates. The July 29, 2016 search of USAN stems did not find any USAN stems in the proposed proprietary name.

3 CONCLUSIONS

Our re-assessment did not identify any names that represent a potential source of drug name confusion. Therefore, we maintain that the proposed proprietary name is acceptable from a promotional and safety perspective.

If you have any questions or need clarifications, please contact Corwin Howard, OSE project manager, at 240-402-8654.

3.1 COMMENTS TO THE APPLICANT

We have completed our review of the proposed proprietary name, Radicava, and have concluded that this name is acceptable.

If any of the proposed product characteristics as stated in your July 5, 2016 submission are altered prior to approval of the marketing application, the name must be resubmitted for review.

Reference ID: 3980833

^a Harris, J. Proprietary Name Review for Radicava (PIND 126396) Silver Spring (MD): Food and Drug Administration, Center for Drug Evaluation and Research, Office of Surveillance and Epidemiology, Division of Medication Error Prevention and Analysis (US); 2016 MAY 9. Panorama No. 2016-2818367.

REFERENCES

1. USAN Stems (http://www.ama-assn.org/ama/pub/physician-resources/medical-science/united-states-adopted-names-council/naming-guidelines/approved-stems.page)

USAN Stems List contains all the recognized USAN stems.