4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2025-N-3082]

Issuance of Priority Review Voucher; Rare Pediatric Disease Product; MODESYO (dordaviprone)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the issuance of a priority review voucher to the sponsor of a rare pediatric disease product application. The Federal Food, Drug, and Cosmetic Act (FD&C Act) authorizes FDA to award priority review vouchers to sponsors of approved rare pediatric disease product applications that meet certain criteria. FDA is required to publish notice of the award of the priority review voucher. FDA has determined that MODESYO (dordaviprone), approved August 6, 2025, manufactured by Chimerix, Inc., meets the criteria for a priority review voucher.

FOR FURTHER INFORMATION CONTACT: Quyen Tran, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993-0002, 301-796-2771.

SUPPLEMENTARY INFORMATION: FDA is announcing the issuance of a priority review voucher to the sponsor of an approved rare pediatric disease product application. Under section 529 of the FD&C Act (21 U.S.C. 360ff), FDA will award priority review vouchers to sponsors of approved rare pediatric disease product applications that meet certain criteria. FDA has determined MODESYO (dordaviprone) manufactured by Chimerix, Inc., meets the criteria for a priority review voucher. MODESYO (dordaviprone) capsules are indicated for the treatment of adult and pediatric patients 1 year of age and older with diffuse midline glioma harboring an H3 K27M mutation with progressive disease following prior therapy.

For further information about the Rare Pediatric Disease Priority Review Voucher

Program and for a link to the full text of section 529 of the FD&C Act, go to

https://www.fda.gov/ForIndustry/DevelopingProductsforRareDiseasesConditions/RarePediatric

DiseasePriorityVoucherProgram/default.htm. For further information about MODESYO

(dordaviprone), go to the "Drugs@FDA" website at

https://www.accessdata.fda.gov/scripts/cder/daf/.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

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