



Marianne Frost, M.S.
Vice President, Regulatory Affairs
Biohaven Pharmaceuticals, Inc.
215 Church Street
New Haven, CT 06510

RE: NDA 212728
NURTEC ODT (rimegepant) orally disintegrating tablets, for sublingual or oral use
MA 71

Dear Ms. Frost:

The Food and Drug Administration has completed evaluation of your firm's response to our Untitled Letter dated March 8, 2021. Based on our evaluation, it appears that you have addressed the violations contained in this Untitled Letter.

This letter does not relieve you or your firm from the responsibility of taking all necessary steps to assure sustained compliance with the Federal Food, Drug, and Cosmetic Act and its implementing regulations or with other relevant legal authority. The Agency expects you and your firm to maintain compliance and will continue to monitor your state of compliance. This letter will not preclude any future regulatory action should violations be observed during subsequent surveillance or through other means.

If you have any questions or comments, please direct your response to the undersigned at the **Food and Drug Administration, Center for Drug Evaluation and Research, Office of Prescription Drug Promotion, 5901-B Ammendale Road, Beltsville, Maryland 20705-1266**. A courtesy copy can be sent by facsimile to (301) 847-8444. To ensure timely delivery of your submissions, please use the full address above and include a prominent directional notation (e.g., a sticker) to indicate that the submission is intended for OPDP. Please refer to MA 71 in addition to the NDA number in all future correspondence relating to this particular matter.

Sincerely,

{See appended electronic signature page}

Aline M. Moukhtara, RN, MPH
Team Leader
Division of Advertising & Promotion Review 1
Office of Prescription Drug Promotion

This is a representation of an electronic record that was signed electronically. Following this are manifestations of any and all electronic signatures for this electronic record.

/s/

ALINE M MOUKHTARA
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