

NDA 022195 NDA 022207

## NOTIFICATION OF NON-COMPLIANCE WITH PREA

Hikma Pharmaceuticals USA, Inc. 1809 Wilson Road Columbus OH, 43228

Attention: James Connell

Associate Director, Regulatory Affairs

Dear Mr. Connell:

Please refer to your new drug application (NDA) submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (the Act) for morphine sulfate oral solution and tablets, which was approved on March 17, 2008.

The Agency has determined that you have failed to meet the postmarketing requirement (PMR) of the Pediatric Research Equity Act (PREA) for this application because you have not yet submitted your pediatric assessment for PMR 204-4, which was deferred until January 31, 2023.

Under the provisions of section 505B(d)(1) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) [21 U.S.C. 355c(d)(1)], you must respond in writing within 45 calendar days of the date of this letter. Your response should include the reason(s) for the delayed pediatric assessment and a date by which you expect to submit the assessment. You may also include a request for a deferral extension, if applicable, which should be identified as a "**DEFERRAL EXTENSION REQUESTED**" in your response.

In accordance with the FD&C Act, FDA will post this letter and your response to the website at <a href="https://www.fda.gov/drugs/development-resources/non-compliance-letters-under-505bd1-federal-food-drug-and-cosmetic-act">https://www.fda.gov/drugs/development-resources/non-compliance-letters-under-505bd1-federal-food-drug-and-cosmetic-act</a> with redactions for any trade secrets and confidential commercial information 60 calendar days from the date of this letter.

Please identify your response to this letter as a "RESPONSE TO PREA NON-COMPLIANCE LETTER." To facilitate our review, submit this information to your NDA with a cross-reference letter to the investigational new drug application (IND) to which your protocol has been submitted.

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If you have any questions, email the Regulatory Project Manager, Cooma Asonye, at Ajoa.Asonye@fda.hhs.gov.

Sincerely,

{See appended electronic signature page}

Rigoberto Roca, MD Director Division of Anesthesiology, Addiction Medicine and Pain Medicine Office of Neuroscience Center for Drug Evaluation and Research \_\_\_\_\_

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

RIGOBERTO A ROCA 06/14/2023 07:59:14 AM

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