

March 20, 2024

Office of the Vermont Attorney General 109 State Street Montpelier, VT 05609

RE: Notice of a New Prescription Drug Pursuant to 18 V.S.A. § 4637

Dear Office of the Vermont Attorney General,

Amneal Pharmaceuticals, LLC ("Amneal") is issuing this notice pursuant to 18 V.S.A. § 4637(c), which requires prescription drug manufacturers to provide the Office of the Attorney General (the "Office") with certain information following the release of a drug in the commercial market whose Wholesale Acquisition Cost ("WAC") exceeds the threshold set for a specialty drug under the Medicare Part D Program.

Amneal released Ropivacaine Hydrochloride Inj ("the Product") into the market on March 20, 2024, at a WAC that exceeds the threshold set for a specialty drug under the Medicare Part D program. This notice provides the additional required information by 18 V.S.A. § 4637(c) regarding the Product. Amneal notes that the Office of the Attorney General has not yet set forth and released a format for submissions under this section. Furthermore, as authorized by 18 V.S.A. § 4637(d), Amneal has limited the information reported to that which is in the public domain or publicly available.

| NDC | Product | Package Size | WAC |
|---------------|-------------------------------|--------------|------------|
| 70121-1734-03 | Ropivacaine Hydrochloride Inj | 12 | \$1,420.20 |

| Statutory Requirement | Reported Information | |
|---|---|--|
| A description of the marketing and | Pursuant to 18 V.S.A. § 4637(d), Amneal has | |
| pricing plans used in the launch of the | elected to limit its response to this item due to the | |
| new drug in the United States and | requested information not being publicly available | |
| internationally 18 V.S.A. § 4637(c)(1) | and in the public domain. | |
| The estimated volume of patients that | NDC 70121-1734-03: 58,000 | |
| may be prescribed the drug 18 V.S.A. § 4637(c)(2) | | |
| Was the drug granted breakthrough | Breakthrough Therapy Designation – No | |
| therapy designation or priority review by the | Priority Review – No | |
| federal Food and Drug Administration prior to | | |
| final approval? 18 V.S.A. § 4637(c)(3) | | |
| The date and price of acquisition if the | Not Applicable | |
| drug was not developed by the | | |
| manufacturer 18 V.S.A. § 4637(c)(4) | | |

Sincerely,

William E. Riker

William & Priber

Senior Director, Pricing & Contracts