



BLA 761381/S-001

**SUPPLEMENT APPROVAL/  
FULFILLMENT OF POSTMARKETING  
REQUIREMENT**

Bristol-Myers Squibb Company  
Attention: Jateh Major, PharmD  
Associate Director  
P.O. Box 5326  
Princeton, NJ 08543-5326

Dear Dr. Major:

Please refer to your December 27, 2024, supplemental biologics license application (sBLA) and your amendments, submitted under section 351(a) of the Public Health Service Act for Opdivo Qvantig (nivolumab and hyaluronidase-nvhy); subcutaneous injection.

This Prior Approval supplemental biologics license application provides for the following:

1. Updates the Opdivo Qvantig prescribing information with the following new indication:

OPDIVO QVANTIG, as monotherapy, is indicated for the first-line treatment of adult patients with unresectable or metastatic hepatocellular carcinoma (HCC) following treatment with intravenous nivolumab and ipilimumab combination therapy.

2. Fulfills post marketing requirement (PMR) 4762-1 and the conversion from accelerated approval to regular approval of the following indication:

OPDIVO QVANTIG, as monotherapy, is indicated for the treatment of adult patients with unresectable or metastatic hepatocellular carcinoma (HCC) who have been previously treated with sorafenib following treatment with intravenous nivolumab and ipilimumab combination therapy.

There is a limitation of use that OPDIVO QVANTIG is not indicated in combination with ipilimumab for the treatment of patients with unresectable or metastatic HCC.

**APPROVAL & LABELING**

We have completed our review of this application, as amended. It is approved, effective on the date of this letter, for use as recommended in the enclosed agreed-upon labeling.

## **CONTENT OF LABELING**

As soon as possible, but no later than 14 days from the date of this letter, submit, via the FDA automated drug registration and listing system (eLIST), the content of labeling [21 CFR 601.14(b)] in structured product labeling (SPL) format, as described at FDA.gov,<sup>1</sup> that is identical to the enclosed labeling (text for the Prescribing Information and Medication Guide) and include the labeling changes proposed in any pending “Changes Being Effected” (CBE) supplements.

Information on submitting SPL files using eLIST may be found in the guidance for industry *SPL Standard for Content of Labeling Technical Qs and As*.<sup>2</sup>

The SPL will be accessible via publicly available labeling repositories.

Also within 14 days, amend all pending supplemental applications that include labeling changes for this BLA, including pending “Changes Being Effected” (CBE) supplements, for which FDA has not yet issued an action letter, with the content of labeling [21 CFR 601.12(f)] in Microsoft Word format that includes the changes approved in this supplemental application, as well as annual reportable changes. To facilitate review of your submission(s), provide a highlighted or marked-up copy that shows all changes, as well as a clean Microsoft Word version. The marked-up copy should provide appropriate annotations, including supplement number(s) and annual report date(s).

## **REQUIRED PEDIATRIC ASSESSMENTS**

Under the Pediatric Research Equity Act (PREA) (21 U.S.C. 355c), all applications for new active ingredients (which includes new salts and new fixed combinations), new indications, new dosage forms, new dosing regimens, or new routes of administration are required to contain an assessment of the safety and effectiveness of the product for the claimed indication(s) in pediatric patients unless this requirement is waived, deferred, or inapplicable.

We are waiving the pediatric assessment requirements for this application because the condition rarely or never occurs in pediatric patients

## **SUBPART E FULFILLED**

We approved this BLA, on December 27, 2024, under the regulations at 21 CFR 601 Subpart E for Accelerated Approval of Biological Products for Serious or Life-Threatening Illnesses. Approval of this supplement fulfills your commitments made under 21 CFR 601.

---

<sup>1</sup> <http://www.fda.gov/ForIndustry/DataStandards/StructuredProductLabeling/default.htm>

<sup>2</sup> We update guidances periodically. For the most recent version of a guidance, check the FDA Guidance Documents Database <https://www.fda.gov/RegulatoryInformation/Guidances/default.htm>.

## **FULFILLMENT OF POSTMARKETING REQUIREMENT**

We have received your submission dated June 21, 2024, to BLA 125554, containing the final report for the following postmarketing requirement listed in the December 27, 2024, approval letter.

- 4762-1 Submit the final report demonstrating an improvement in overall survival from a multicenter, randomized trial to confirm the clinical benefit of nivolumab in combination with ipilimumab over standard therapy in patients with advanced hepatocellular carcinoma, that may inform product labeling. Submit the datasets with the final report.

We have reviewed your submission and conclude that the above requirement was fulfilled. You are not required to report on the status of closed (released or fulfilled) PMRs/PMCs in your annual report required under 21.CFR 601.70 of the FD&CA.

We remind you that there is a postmarketing requirement and a postmarketing commitment listed in the December 27, 2024, approval letter that are still open.

## **PROMOTIONAL MATERIALS**

You may request advisory comments on proposed introductory advertising and promotional labeling. For information about submitting promotional materials, see the final guidance for industry *Providing Regulatory Submissions in Electronic and Non-Electronic Format-Promotional Labeling and Advertising Materials for Human Prescription Drugs*.<sup>3</sup>

You must submit final promotional materials and Prescribing Information, accompanied by a Form FDA 2253, at the time of initial dissemination or publication [21 CFR 601.12(f)(4)]. Form FDA 2253 is available at [FDA.gov](http://FDA.gov).<sup>4</sup> Information and Instructions for completing the form can be found at [FDA.gov](http://FDA.gov).<sup>5</sup>

All promotional materials that include representations about your drug product must be promptly revised to be consistent with the labeling changes approved in this supplement, including any new safety-related information [21 CFR 601.12(f)(4)].

The revisions in your promotional materials should include prominent disclosure of the important new safety-related information that appears in the revised labeling. Within 7 days of receipt of this letter, submit your statement of intent to comply with 21 CFR 601.12(f)(4).

---

<sup>3</sup> For the most recent version of a guidance, check the FDA guidance web page at <https://www.fda.gov/media/128163/download>.

<sup>4</sup> <http://www.fda.gov/downloads/AboutFDA/ReportsManualsForms/Forms/UCM083570.pdf>

<sup>5</sup> <http://www.fda.gov/downloads/AboutFDA/ReportsManualsForms/Forms/UCM375154.pdf>

## **REPORTING REQUIREMENTS**

We remind you that you must comply with reporting requirements for an approved BLA (in 21 CFR 600.80 and in 21 CFR 600.81).

If you have any questions, contact Gina Davis, Senior Regulatory Health Project Manager, at [Gina.Davis@fda.hhs.gov](mailto:Gina.Davis@fda.hhs.gov).

Sincerely,

*{See appended electronic signature page}*

Steven Lemery, M.D., MHS  
Director  
Division of Oncology 3  
Office of Oncologic Diseases  
Center for Drug Evaluation and Research

### ENCLOSURES:

- Content of Labeling
  - Prescribing Information
  - Medication Guide

-----  
**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/  
-----

STEVEN J LEMERY  
10/27/2025 04:45:42 PM