ONO PHARMACEUTICAL CO.,LTD.

February 17, 2020

Opdivo (nivolumab) Plus Yervoy (ipilimumab) Demonstrates Continued Survival Benefit at 42-Month Follow-up in Patients with Previously Untreated Advanced or Metastatic Renal Cell Carcinoma

This information is intended to notify the press release issued on February 15 by Bristol-Myers Squibb. Please click https://www.bms.com/media/press-releases.html for the original press release.

First paragraph extracted from the original press release:

(PRINCETON, NJ, February 15, 2020) – Bristol-Myers Squibb Company (NYSE: BMY) today announced updated results from the Phase 3 CheckMate -214 study evaluating the combination of Opdivo (nivolumab) plus Yervoy (ipilimumab) versus sunitinib in patients with previously untreated advanced or metastatic renal cell carcinoma (RCC). With a minimum follow-up of 42 months, the combination of Opdivo plus Yervoy continues to show superior overall survival (OS), objective response rates (ORR), duration of response (DOR) and complete response (CR) rates. The safety profile for Opdivo plus Yervoy was consistent with prior findings and no new safety signals or drug-related deaths occurred with extended follow-up. The data will be featured in an oral presentation (Abstract #609) on Saturday, February 15, 2020 at the American Society of Clinical Oncology 2020 Genitourinary Cancers Symposium in San Francisco.

About Opdivo

Opdivo is a programmed death-1 (PD-1) immune checkpoint inhibitor that is designed to uniquely harness the body's own immune system to help restore anti-tumor immune response by blocking the interaction between PD-1 and its ligands. By harnessing the body's own immune system to fight cancer, Opdivo has become an important treatment option across multiple cancers since the approval for the treatment of melanoma in Japan in July 2014. Opdivo is currently approved in more than 65 countries, including Japan, South Korea, Taiwan, China, the US and European Union.

In Japan, Ono Pharmaceutical Co., Ltd. ("ONO") launched Opdivo for the treatment of unresectable melanoma in September 2014. Thereafter, Opdivo received an approval for additional indications of unresectable, advanced or recurrent non-small cell lung cancer in December 2015, unresectable or metastatic renal cell cancer in August 2016, relapsed or refractory classical Hodgkin lymphoma in December 2016, recurrent or metastatic head and neck cancer in March 2017, unresectable advanced or recurrent gastric cancer which has progressed after chemotherapy in September 2017, unresectable advanced or recurrent malignant pleural mesothelioma which has progressed after chemotherapy and adjuvant treatment of melanoma, etc. in August 2018.

In addition, ONO has submitted supplemental applications for the treatment of MSI-H colorectal cancer and esophageal cancer, and is conducting clinical development program including esophageal cancer, esophago-gastric junction cancer, small cell lung cancer, hepatocellular carcinoma, glioblastoma, urothelial cancer, ovarian cancer, bladder cancer, colorectal cancer, pancreatic cancer, biliary tract cancer, etc.

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