

UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
Washington, D.C. 20549

**FORM 8-K**

CURRENT REPORT  
Pursuant to Section 13 OR 15(d) of  
The Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): August 6, 2020

**BRISTOL-MYERS SQUIBB COMPANY**

(Exact name of registrant as specified in its charter)

Delaware  
(State or other jurisdiction of  
incorporation or organization)

001-01136  
(Commission File Number)

22-0790350  
(I.R.S Employer  
Identification No.)

430 E. 29th Street, 14th Floor  
New York, NY, 10016  
(Address of principal executive offices) (zip code)

Registrant's telephone number, including area code: (212) 546-4000

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

**Securities registered pursuant to Section 12(b) of the Act:**

<u>Title of each class</u>	<u>Trading Symbol(s)</u>	<u>Name of each exchange on which registered</u>
Common Stock, \$0.10 Par Value	BMV	New York Stock Exchange
1.000% Notes due 2025	BMV25	New York Stock Exchange
1.750% Notes due 2035	BMV35	New York Stock Exchange
Bristol-Myers Squibb Contingent Value Rights	BMV RT	New York Stock Exchange
Celgene Contingent Value Rights	CELG RT	New York Stock Exchange

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.





## Bristol Myers Squibb Reports Second Quarter 2020 Financial Results

- Reports Second Quarter Revenues of \$10.1 Billion
- Posts GAAP Loss Per Share of \$0.04 and Non-GAAP EPS of \$1.63
- Advances Pipeline with Positive Topline Results for *Zeposia* in Ulcerative Colitis; Approvals for *Opdivo* plus *Yervoy* in Lung Cancer
- Updates 2020 GAAP EPS Guidance Range from \$0.37-\$0.57 to (\$0.06)-\$0.09 and Non-GAAP EPS Guidance Range from \$6.00-\$6.20 to \$6.10-\$6.25

(NEW YORK, August 6, 2020) – [Bristol Myers Squibb](#) (NYSE:BMJ) today reports results for the second quarter of 2020, which reflect strong product sales, continued advancement of the pipeline and robust operating performance.

“Our second quarter results reflect the passion and focus of our employees, who continue to introduce new medicines, support patients with serious diseases and deliver strong results during the COVID-19 pandemic,” said [Giovanni Caforio, M.D.](#), chairman and chief executive officer, Bristol Myers Squibb. “Our teams drove strong commercial execution while continuing to progress our integration initiatives. With several new product launches and the achievement of multiple milestones from our late-stage pipeline, I am confident that we are building a leading biopharma with a renewed portfolio of transformational medicines. Our financial flexibility and continued opportunities to invest in innovation position us well to deliver for the long-term.”

	<b>Second Quarter</b>		
<b>\$ amounts in millions, except per share amounts</b>	<b>2020</b>	<b>2019</b>	<b>Change</b>
Total Revenues	\$10,129	\$6,273	61 %
GAAP Diluted (Loss)/EPS	(0.04)	0.87	N/A
Non-GAAP Diluted EPS	1.63	1.18	38 %
Total Pro Forma Revenues*	10,129	10,160	0 %

\* The pro forma revenues assume the company's acquisition of Celgene (Celgene Acquisition) and its divestiture of Otezla<sup>®</sup> to Amgen Inc. (Otezla<sup>®</sup> Divestiture) occurred on January 1, 2019 and exclude foreign currency hedge gains and losses. Management believes that measuring revenue rates on a comparable pro forma basis is an appropriate way for investors to best understand the underlying performance of the business. See “Worldwide Pro Forma Revenue” in Quarterly Package of Financial Information for this quarter, which is available on [bms.com/investors/financial-reporting/quarterly-results](https://www.bms.com/investors/financial-reporting/quarterly-results), for information on the revenue of the company and Celgene on a stand-alone basis for the prior-year period. Otezla<sup>®</sup> is a trademark of Amgen Inc.

## **SECOND QUARTER FINANCIAL RESULTS**

**All comparisons are made versus the same period in 2019 unless otherwise stated.**

- Bristol Myers Squibb posted second quarter revenues of \$10.1 billion, an increase of 61% on a reported basis, or 63% when adjusted for foreign exchange. The increase was driven primarily by the impact of the Celgene Acquisition, which was completed on November 20, 2019. Revenues remained consistent on a pro forma basis, as sales were estimated to be negatively impacted by approximately \$600 million due mainly to COVID-19 related channel inventory work downs from the first quarter, as well as lower demand resulting from reduced new patient starts and fewer patient visits to physicians in the pandemic.
- U.S. revenues increased 77% to \$6.5 billion in the quarter. International revenues increased 40% to \$3.6 billion in the quarter. When adjusted for foreign exchange impact, international revenues increased 43%.
- Gross margin as a percentage of revenue increased from 68.6% to 73.4% in the quarter primarily due to product mix, partially offset by the unwinding of inventory purchase price accounting adjustments.
- Marketing, selling and administrative expenses increased 51% to \$1.6 billion in the quarter primarily due to \$600 million of costs associated with the broader portfolio resulting from the Celgene Acquisition.
- Research and development expenses increased 90% to \$2.5 billion in the quarter primarily due to \$1.1 billion of costs associated with the broader portfolio resulting from the Celgene Acquisition.
- Amortization of acquired intangible assets was \$2.4 billion in the quarter primarily due to the Celgene Acquisition.
- Income taxes were \$1.7 billion on pre-tax earnings of \$1.6 billion in the quarter primarily due to tax charges resulting from an internal transfer of certain intangible assets and the Otezla<sup>®</sup> Divestiture and purchase price adjustments. The effective tax rate was 19.0% in the same period a year ago.

- The company reported net loss attributable to Bristol Myers Squibb of \$85 million, or \$0.04 per share, in the second quarter, compared to net earnings of \$1.4 billion, or \$0.87 per share, for the same period a year ago. The results in the current quarter include costs and expenses resulting from purchase price accounting, contingent value rights fair value adjustments and other acquisition and integration expenses.
- The company reported non-GAAP net earnings attributable to Bristol Myers Squibb of \$3.8 billion, or \$1.63 per share, in the second quarter, compared to net earnings of \$1.9 billion, or \$1.18 per share, for the same period a year ago. A discussion of the non-GAAP financial measures is included under the “Use of Non-GAAP Financial Information” section.
- Cash, cash equivalents and marketable debt securities were \$22.2 billion and debt was \$46.7 billion, as of June 30, 2020.

## **SECOND QUARTER PRODUCT REVENUE HIGHLIGHTS**

\$ amounts in millions

Product	Quarter Ended June 30, 2020 on Reported Basis	% Change from Quarter Ended June 30, 2019 on Reported Basis	% Change from Quarter Ended June 30, 2019 on Pro Forma Basis**
<i>Revlimid</i>	\$2,884	N/A*	6%
<i>Eliquis</i>	\$2,163	6%	6%
<i>Opdivo</i>	\$1,653	(9)%	(9)%
<i>Orencia</i>	\$750	(4)%	(4)%
<i>Pomalyst/Imnovid</i>	\$745	N/A*	21%
<i>Sprycel</i>	\$511	(6)%	(6)%
<i>Yervoy</i>	\$369	1%	1%
<i>Abraxane</i>	\$308	N/A*	(2)%
<i>Empliciti</i>	\$97	7%	7%
<i>Reblozyl</i>	\$55	N/A*	N/A
<i>Inrebic</i>	\$15	N/A*	N/A
<i>Zeposia</i>	\$1	N/A*	N/A

\* Products were acquired as part of the Celgene Acquisition.

\*\* Pro forma product revenues assume the Celgene Acquisition and the Otezla® Divestiture occurred on January 1, 2019 and exclude foreign currency hedge gains and losses. Management believes that measuring product revenue rates on a comparable pro forma basis is an appropriate way for investors to best understand the underlying performance of the business. See “Worldwide Pro Forma Revenues” in the Quarterly Package of Financial Information for this quarter, which is available on [bms.com/investors/financial-reporting/quarterly-results](https://bms.com/investors/financial-reporting/quarterly-results), for information on the product revenue of the company and Celgene for the prior-year period.

## **FIRST HALF PRODUCT REVENUE HIGHLIGHTS**

\$ amounts in millions

Product	Six Months Ended June 30, 2020 on Reported Basis	% Change from Six Months Ended June 30, 2019 on Reported Basis	% Change from Six Months Ended June 30, 2019 on Pro Forma Basis**
<i>Revlimid</i>	\$5,799	N/A*	10%
<i>Eliquis</i>	\$4,804	21%	21%
<i>Opdivo</i>	\$3,419	(6)%	(6)%
<i>Orencia</i>	\$1,464	3%	3%
<i>Pomalyst/Imnovid</i>	\$1,458	N/A*	25%
<i>Sprycel</i>	\$1,032	3%	3%
<i>Yervoy</i>	\$765	2%	2%
<i>Abraxane</i>	\$608	N/A*	2%
<i>Empliciti</i>	\$194	11%	11%
<i>Reblozyl</i>	\$63	N/A*	N/A
<i>Inrebic</i>	\$27	N/A*	N/A
<i>Zeposia</i>	\$1	N/A*	N/A

\* Products were acquired as part of the Celgene Acquisition.

\*\* Pro forma product revenues assume the Celgene Acquisition and the Otezla® Divestiture occurred on January 1, 2019 and exclude foreign currency hedge gains and losses. Management believes that measuring product revenue rates on a comparable pro forma basis is an appropriate way for investors to best understand the underlying performance of the business. See "Worldwide Pro Forma Revenues" in the Quarterly Package of Financial Information for this quarter, which is available on [bms.com/investors/financial-reporting/quarterly-results](https://bms.com/investors/financial-reporting/quarterly-results), for information on the product revenue of the company and Celgene for the prior-year period.

## **SECOND QUARTER PRODUCT AND PIPELINE UPDATE**

### **Cardiovascular**

#### ***Eliquis***

#### ***Patent Update***

- In August, the Bristol-Myers Squibb-Pfizer Alliance announced the U.S. District Court decision to uphold both the composition of matter patent (US 6,967,208) and formulation patent (US 9,326,945) covering Eliquis® (apixaban). ([link](#))

## Oncology and Hematology

### ***Opdivo***

#### *Regulatory*

- In June, the company announced the U.S. Food and Drug Administration (U.S. FDA) approval of *Opdivo* (nivolumab) for the treatment of patients with unresectable advanced, recurrent or metastatic esophageal squamous cell carcinoma (ESCC) after prior fluoropyrimidine-and platinum-based chemotherapy. ([link](#))
- In May, the company announced *Opdivo* plus *Yervoy* (ipilimumab) given with two cycles of platinum-doublet chemotherapy was approved by the U.S. FDA for the first-line treatment of adult patients with metastatic or recurrent non-small cell lung cancer (NSCLC) with no EGFR or ALK genomic tumor aberrations. This approval was based on the Phase 3 CheckMate -9LA study. ([link](#))
- In May, the company announced the U.S. FDA approved *Opdivo* plus *Yervoy* for the first-line treatment of adult patients with metastatic NSCLC whose tumors express PD-LI $\geq$ 1% as determined by an FDA-approved test, with no EGFR or ALK genomic tumor aberrations. This approval was based on data from Part 1a of the Phase 3 Checkmate -227 study. ([link](#))

### ***Reblozyl***

#### *Regulatory*

- In June, the company and Acceleron Pharma Inc. announced the European Commission (EC) approved *Reblozyl* (luspatercept) for the treatment of transfusion-dependent anemia in adult patients with myelodysplastic syndromes (MDS) or beta thalassemia. ([link](#))

### **ide-cel**

#### *Regulatory*

- In July, the company and bluebird bio, Inc. announced that the companies submitted the Biologics License Application (BLA) to the U.S. FDA for idecabtagene vicleucel (ide-cel; bb2121) for patients with heavily pre-treated relapsed and refractory multiple myeloma. This submission follows the company's receipt of a Refusal to File letter from the U.S. FDA in May 2020 following the original BLA submission from March 2020. ([link](#))

- In May, the company announced that the European Medicines Agency (EMA) validated its Marketing Authorization Application (MAA) for ide-cel; bb2121, the company's investigational B-cell maturation antigen (BCMA)-directed chimeric antigen receptor (CAR) T cell immunotherapy co-developed with bluebird bio, Inc., for the treatment of adult patients with multiple myeloma who have received at least three prior therapies, including an immunomodulatory agent, a proteasome inhibitor and an anti-CD38 antibody. ([link](#))

## **CC-486**

### *Regulatory*

- In May, the company announced that the EMA validated its MAA for CC-486 for the maintenance treatment of adult patients with acute myeloid leukemia (AML), who achieved complete remission (CR) or CR with incomplete blood count recovery (CRi), following induction therapy with or without consolidation treatment, and who are not candidates for, or who choose not to proceed to, hematopoietic stem cell transplantation. ([link](#))

## **Pomalyst**

### *Regulatory*

- In May, the company announced that the U.S. FDA approved *Pomalyst* (pomalidomide) for patients with AIDS-related Kaposi sarcoma whose disease has become resistant to highly active antiretroviral therapy (HAART), or in patients with Kaposi sarcoma who are HIV-negative. ([link](#))

## **Liso-cel**

### *Regulatory*

- In July, the company announced that the EMA validated the MAA for liso-cel (lisocabtagene maraleucel), a CD19-directed chimeric antigen receptor (CAR) T cell therapy for the treatment of adults with relapsed or refractory large B-cell lymphoma after at least two prior therapies. ([link](#))



## **Medical Conferences**

- In May, at the American Society of Clinical Oncology (ASCO) 2020 Virtual Scientific Program, the company announced important new data and analysis across its cancer portfolio ([link](#)), including:
  - First disclosure of data from the Phase 3 CheckMate -9LA trial evaluating *Opdivo* plus *Yervoy* given concomitantly with two cycles of chemotherapy, for the first-line treatment of metastatic NSCLC. ([link](#))
  - Three-year follow-up results from the Phase 3 Checkmate -227 trial, demonstrating that *Opdivo* plus *Yervoy* provided sustained improvements in overall survival (OS) and additional efficacy measures as a first-line treatment for patients with metastatic NSCLC. ([link](#))
  - First presentation of data from the Phase 2 KarMMA study with bluebird bio, Inc. evaluating the efficacy and safety of the companies' investigational B-cell maturation antigen (BCMA)-directed chimeric antigen receptor (CAR) T cell immunotherapy, idecabtagene vicleucel (ide-cel; bb2121), in patients with relapsed and refractory multiple myeloma. ([link](#))
- In June, at the 25<sup>th</sup> European Hematology Association (EHA), the company announced important new data and analysis from 60 company-sponsored studies, highlighting the company's approaches to treating blood cancers and other diseases. ([link](#))

## **Immunology**

### ***Zeposia***

#### *Commercial*

- In June, the company announced the commercial launch and availability of *Zeposia* (ozanimod), a new oral treatment for relapsing forms of multiple sclerosis, in the U.S. *Zeposia* was approved by the U.S. FDA on March 25, 2020. ([link](#))

#### *Regulatory*

- In May, the company announced the EC approval *Zeposia* for the treatment of adult patients in the European Union with relapsing forms of multiple sclerosis. ([link](#))

### *Clinical*

- In June, the company announced results from True North, a pivotal Phase 3 trial evaluating oral *Zeposia* as an induction and maintenance therapy for adult patients with moderate to severe ulcerative colitis. True North met both primary endpoints of clinical remission in induction at Week 10 and in maintenance at Week 52. ([link](#))

### **Orencia**

#### *Clinical*

- In June, at the European E-Congress of Rheumatology (EULAR) 2020, the company announced results from the open-label switch period of Early AMPLE, a Phase IV exploratory biomarker study assessing the differences by which *Orencia* (abatacept) and adalimumab, interfere with disease progression in moderate-to-severe early rheumatoid arthritis (RA) patients who tested positive (seropositive) for certain autoantibodies. ([link](#))

### **COVID-19 Pandemic Response**

During the current world health crisis, the company continues to take all necessary actions to promote public health by carrying out its mission of providing life-saving medicines to the patients who depend on the company and supporting relief efforts across the globe. ([link](#))

### **Financial Guidance**

Bristol Myers Squibb is updating its 2020 GAAP EPS guidance range from \$0.37 - \$0.57 to (\$0.06) - \$0.09. In addition, the company is updating its 2020 non-GAAP EPS guidance range of \$6.00 - \$6.20 to \$6.10 - \$6.25. Adjusted 2020 GAAP and non-GAAP line items are:

	<b><u>GAAP</u></b>	<b><u>non-GAAP</u></b>
Revenue	\$40.5B - \$42.0B	\$40.5B - \$42.0B
Gross margin as a percentage of revenue	Approximately 74%	Approximately 80%
Marketing, selling, and administrative expense	\$6.5B - \$6.7B	\$6.5B - \$6.7B
Research and development expense	\$9.7B - \$9.9B	\$9.2B - \$9.4B
Other (income)/expense, net	\$0.9B - \$1.1B	(\$0.1B) - \$0.1B
Effective tax rate	Approximately 100%	16-17%
Weighted average diluted shares	Approximately 2.3 Billion	Approximately 2.3 Billion
EPS guidance	(\$0.06) - \$0.09	\$6.10 - \$6.25

The 2020 guidance assumes the peak impact of the current COVID-19 crisis on the business would occur in the second quarter of 2020, with a return to a more stable business environment in the third quarter and minimal impact from the fourth quarter of 2020 onwards. Additional key factors assumed in guidance now include:

- Mid-July foreign exchange and interest rates apply.
- Products that saw significant advanced buying at the end of the first quarter will see that inventory work-down during the rest of the year, mostly in the second quarter, which the company experienced, and to a lesser degree in the third and fourth quarters.
- A reduction in new-to-brand prescriptions, and on physician administered product demand during the second quarter, recovering during the third quarter and fully recovered in the fourth quarter.
- All clinical trial activities are planned to resume by the end of the year where local country restrictions have been lifted.

The financial guidance excludes the impact of any potential future strategic acquisitions and divestitures and any specified items that have not yet been identified and quantified. The 2020 non-GAAP EPS guidance further excludes other specified items as discussed under “Use of Non-GAAP Financial Information.” A reconciliation of non-GAAP financial measures to the most comparable GAAP measure and the reasons why management believes the use of these measures is important are provided in supplemental materials available on the company’s website. The financial guidance is subject to risks and uncertainties applicable to all forward-looking statements as described elsewhere in this press release.

### **Company and Conference Call Information**

Bristol Myers Squibb is a global biopharmaceutical company whose mission is to discover, develop and deliver innovative medicines that help patients prevail over serious diseases. For more information about Bristol Myers Squibb, visit us at [BMS.com](http://BMS.com) or follow us on [LinkedIn](#), [Twitter](#), [YouTube](#), [Facebook](#), and [Instagram](#).

There will be a conference call on August 6 at 8:30 a.m. ET during which company executives will review financial information and address inquiries from investors and analysts. Investors and the general public are invited to listen to a live webcast of the call at <http://investor.bms.com> or by dialing in the U.S. toll free 800-458-4121 or international 786-789-4772, confirmation code: 8970168, or using this [link](#), which becomes active 15 minutes prior to the

scheduled start time and entering your information to be connected. Materials related to the call will be available at the same website prior to the conference call.

A replay of the call will be available beginning at 12:00 p.m. ET on August 6 through 12:00 p.m. ET on August 20, 2020. The replay will also be available through <http://investor.bms.com> or by dialing in the U.S. toll free 888-203-1112 or international 719-457-0820, confirmation code: 8970168.

### **Use of Non-GAAP Financial Information**

This earnings release contains non-GAAP financial measures, including non-GAAP earnings and related EPS information that are adjusted to exclude certain costs, expenses, gains and losses and other specified items that are evaluated on an individual basis. Reconciliations of these non-GAAP financial measures to the most comparable GAAP measures are available on the company's website at [www.bms.com](http://www.bms.com).

These non-GAAP items are adjusted after considering their quantitative and qualitative aspects and typically have one or more of the following characteristics, such as being highly variable, difficult to project, unusual in nature, significant to the results of a particular period or not indicative of future operating results. Similar charges or gains were recognized in prior periods and will likely reoccur in future periods, including amortization of acquired intangible assets beginning in the fourth quarter of 2019, including product rights that generate a significant portion of our ongoing revenue, unwind of inventory fair value adjustments, acquisition and integration expenses, restructuring costs, accelerated depreciation and impairment of property, plant and equipment and intangible assets, R&D charges or other income resulting from upfront or contingent milestone payments in connection with the acquisition or licensing of third-party intellectual property rights, costs of acquiring a priority review voucher, divestiture gains or losses, stock compensation resulting from accelerated vesting of Celgene awards, certain retention-related employee compensation charges related to the Celgene Acquisition, pension, legal and other contractual settlement charges, interest expense on the notes issued in May 2019 incurred prior to the Celgene Acquisition and interest income earned on the net proceeds of those notes, equity investment and contingent value rights fair value adjustments and amortization of fair value adjustments of debt acquired from Celgene in our 2019 exchange offer, among other items. Deferred and current income taxes attributed to these items are also adjusted for considering their individual impact to the overall tax expense, deductibility and jurisdictional tax rates. Certain other significant tax items are also excluded such as the impact resulting from internal transfer of intangible assets and the Otezla<sup>®</sup> Divestiture. This earnings release also provides international revenues excluding the impact of foreign exchange.

Non-GAAP information is intended to portray the results of the company's baseline performance, supplement or enhance management, analysts and investors overall understanding of the company's underlying financial performance and facilitate comparisons among current, past and future periods. For example, non-GAAP earnings and EPS information are indications of the company's baseline performance before items that are considered by us to not be reflective of the company's ongoing results. In addition, this information is among the primary indicators that we use as a basis for evaluating performance, allocating resources, setting incentive compensation targets and planning and forecasting for future periods. This information is not intended to be

considered in isolation or as a substitute for net earnings or diluted EPS prepared in accordance with GAAP and may not be the same as or comparable to similarly titled measures presented by other companies due to possible differences in method and in the items being adjusted. We encourage investors to review our financial statements and publicly-filed reports in their entirety and not to rely on any single financial measure.

Amortization of acquired intangible assets were previously included in non-GAAP earnings and EPS information. These amounts have become significant to the financial results subsequent to the Celgene Acquisition and as a result, have been excluded in the non-GAAP results to better reflect our core operating performance. Comparable prior period non-GAAP results have not been revised to include this adjustment as the related amounts were insignificant (\$24 million and \$48 million for the three and six months ended June 30, 2019, respectively).

### **Website Information**

We routinely post important information for investors on our website, BMS.com, in the “Investors” section. We may use this website as a means of disclosing material, non-public information and for complying with our disclosure obligations under Regulation FD. Accordingly, investors should monitor the Investors section of our website, in addition to following our press releases, SEC filings, public conference calls, presentations and webcasts. We may also use social media channels to communicate with our investors and the public about our company, our products and other matters, and those communications could be deemed to be material information. The information contained on, or that may be accessed through, our website or social media channels are not incorporated by reference into, and are not a part of, this document.

### **Cautionary Statement Regarding Forward-Looking Statements**

This earnings release and the related attachments (as well as the oral statements made with respect to information contained in this release and the attachments) contain certain “forward-looking” statements within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended, regarding, among other things, statements relating to goals, plans and projections regarding the company’s financial position, results of operations, market position, product development and business strategy. These statements may be identified by the fact they use words such as “should,” “could,” “expect,” “anticipate,” “estimate,” “target,” “may,” “project,” “guidance,” “intend,” “plan,” “believe,” “will” and other words and terms of similar meaning and expression in connection with any discussion of future operating or financial performance, although not all forward-looking statements contain such terms. One can also identify forward-looking statements by the fact that they do not relate strictly to historical or current facts. These statements are likely to relate to, among other things, the company’s ability to execute successfully its strategic plans, including its business development strategy generally and in relation to its ability to realize the projected benefits of the Celgene Acquisition, the full extent of the impact of the COVID-19 pandemic on the company’s operations and the development and commercialization of its products, the expiration of patents or data protection on certain products, including assumptions about the company’s ability to retain patent exclusivity of certain products and the impact, and the result of governmental investigations. No forward-looking statement can be guaranteed, including that the company’s future clinical studies will support the data described in this release, product candidates will receive necessary clinical and manufacturing regulatory approvals, pipeline products will prove to be commercially successful, clinical and manufacturing regulatory approvals

will be sought or obtained within currently expected timeframes or contractual milestones will be achieved.

Such forward-looking statements are based on historical performance and current expectations and projections about the company's future financial results, goals, plans and objectives and involve inherent risks, assumptions and uncertainties, including internal or external factors that could delay, divert or change any of them in the next several years, that are difficult to predict, may be beyond the company's control and could cause the company's future financial results, goals, plans and objectives to differ materially from those expressed in, or implied by, the statements. Such risks, uncertainties and other matters include, but are not limited to, risks relating to integrating the company's and Celgene's business and operations, including with respect to human capital management, portfolio rationalization, finance and accounting systems, sales operations and product distribution, pricing systems and methodologies, data security systems, compliance programs and internal controls processes, on the company's ability to realize the anticipated benefits from the Celgene Acquisition; the impact of the company's significant additional indebtedness that it incurred and its issuance of additional shares in connection with the Celgene Acquisition on its ability to operate the combined company; various risks related to public health outbreaks, epidemics and pandemics, including the impact of the COVID-19 pandemic on the company's operations, the possibility of the COVID-19 pandemic delaying the timing of the FDA's approval decisions and that the company cannot reasonably assess or predict at this time the full extent of the adverse effect that the COVID-19 pandemic will have on its business, financial condition, results of operations and cash flows; challenges inherent in new product development, including obtaining and maintaining regulatory approval; increasing pricing pressures from market access, pharmaceutical pricing controls and discounting and other restrictions in the United States, the European Union and other regions around the world (including changes in rules and practices of managed care organizations and institutional and governmental purchasers and the proposals contained in the "American Patient First Blueprint" and the executive orders issued by the U.S. federal government in July designed to regulate prices and payment for pharmaceutical products); the possibility of difficulties and delays in product introduction and commercialization; the company's ability to obtain and protect market exclusivity rights and enforce patents and other intellectual property rights; the risk of certain novel approaches to disease treatment (such as CAR T therapy); industry competition from other manufacturers; the risk of an adverse patent litigation decision or settlement and exposure to other litigation and/or regulatory actions; the impact of any U.S. healthcare reform and legislation or regulatory action in the U.S. and markets outside the U.S. affecting pharmaceutical product pricing, reimbursement or access; changes in tax law and regulations; any decline in the company's future royalty streams; the failure of the company's suppliers, vendors, outsourcing partners, alliance partners and other third parties to meet their contractual, regulatory and other obligations; the company's ability to execute its financial, strategic and operational plans; the company's ability to identify potential strategic acquisitions, licensing opportunities or other beneficial transactions; the company's ability to attract and retain key personnel; the company's ability to effectively manage acquisitions, divestitures, alliances and other portfolio actions and to successfully realize the expected benefits of such actions; the company's dependency on several key products; potential difficulties, delays and disruptions in manufacturing, distribution or sale of products, including without limitation, interruptions caused by damage to the company's and the company's suppliers' manufacturing sites; regulatory decisions impacting labeling, manufacturing processes and/or other matters; the impact on the company's competitive position from counterfeit or unregistered versions of its products or stolen products; the adverse impact of cyber-attacks on the company's information systems or products,

including unauthorized disclosure of trade secrets or other confidential data stored in the company's information systems and networks; political and financial instability of international economies and sovereign risk; interest rate and currency exchange rate fluctuations, credit and foreign exchange risk management; and issuance of new or revised accounting standards. In addition, the 2020 financial guidance provided in this release relies on assumptions about the duration and severity of the COVID-19 pandemic, timing of the return to a more stable business environment, patient and physician behaviors, buying patterns and clinical trial activities (together, the "Recovery Process"), among other things. If the actual Recovery Process differs materially from our assumptions, the impact of COVID-19 on our business could be worse than expected and our results may be negatively impacted.

Forward-looking statements in this earnings release should be evaluated together with the many risks and uncertainties that affect the company's business and market, particularly those identified in the cautionary statement and risk factors discussion in the company's Annual Report on Form 10-K for the year ended December 31, 2019, as updated by the company's subsequent Quarterly Reports on Form 10-Q, Current Reports on Form 8-K and other filings with the Securities and Exchange Commission. The forward-looking statements included in this document are made only as of the date of this document and except as otherwise required by applicable law, the company undertakes no obligation to publicly update or revise any forward-looking statement, whether as a result of new information, future events, changed circumstances or otherwise.

BRISTOL-MYERS SQUIBB COMPANY  
 PRODUCT REVENUES  
 FOR THE THREE MONTHS ENDED JUNE 30, 2020 AND 2019  
 (Unaudited, dollars in millions)

	Worldwide Revenues			U.S. Revenues <sup>(b)</sup>		
	2020	2019	% Change	2020	2019	% Change
<b>Prioritized Brands</b>						
Revlimid	\$ 2,884	\$ —	N/A	\$ 2,048	\$ —	N/A
Eliquis	2,163	2,042	6 %	1,363	1,269	7 %
Opdivo	1,653	1,823	(9)%	956	1,112	(14)%
Orencia	750	778	(4)%	554	566	(2)%
Pomalyst/Imnovid	745	—	N/A	522	—	N/A
Sprycel	511	544	(6)%	308	307	—
Yervoy	369	367	1 %	254	253	—
Abraxane	308	—	N/A	218	—	N/A
Empliciti	97	91	7 %	59	63	(6)%
Reblozyl	55	—	N/A	55	—	N/A
Inrebic	15	—	N/A	15	—	N/A
Zeposia	1	—	N/A	1	—	N/A
<b>Established Brands</b>						
Baraclude	121	147	(18)%	3	7	(57)%
Vidaza	126	—	N/A	—	—	N/A
Other Brands <sup>(a)</sup>	331	481	(31)%	131	90	46 %
<b>Total</b>	<b>\$ 10,129</b>	<b>\$ 6,273</b>	<b>61 %</b>	<b>\$ 6,487</b>	<b>\$ 3,667</b>	<b>77 %</b>

(a) Includes Sustiva, Reyataz, Daklinza and all other BMS and Celgene products acquired as part of the Celgene acquisition that have lost exclusivity in major markets, over-the-counter brands and royalty revenue. Other Brands includes \$83 million worldwide and \$66 million U.S. revenues relating to Celgene products for the three months ended June 30, 2020.

(b) Includes Puerto Rico.



BRISTOL-MYERS SQUIBB COMPANY  
 PRODUCT REVENUES  
 FOR THE SIX MONTHS ENDED JUNE 30, 2020 AND 2019  
 (Unaudited, dollars in millions)

	Worldwide Revenues			U.S. Revenues <sup>(b)</sup>		
	2020	2019	% Change	2020	2019	% Change
<b>Prioritized Brands</b>						
Revlimid	\$ 5,799	\$ —	N/A	\$ 4,014	\$ —	N/A
Eliquis	4,804	3,967	21 %	3,140	2,475	27 %
Opdivo	3,419	3,624	(6)%	1,964	2,236	(12)%
Orencia	1,464	1,418	3 %	1,054	1,015	4 %
Pomalyst/Imnovid	1,458	—	N/A	1,011	—	N/A
Sprycel	1,032	1,003	3 %	608	547	11 %
Yervoy	765	751	2 %	511	528	(3)%
Abraxane	608	—	N/A	423	—	N/A
Empliciti	194	174	11 %	118	121	(2)%
Reblozyl	63	—	N/A	63	—	N/A
Inrebic	27	—	N/A	27	—	N/A
Zeposia	1	—	N/A	1	—	N/A
<b>Established Brands</b>						
Baraclude	243	288	(16)%	6	14	(57)%
Vidaza	284	—	N/A	2	—	N/A
Other Brands <sup>(a)</sup>	749	968	(23)%	311	180	73 %
<b>Total</b>	<u>\$ 20,910</u>	<u>\$ 12,193</u>	71 %	<u>\$ 13,253</u>	<u>\$ 7,116</u>	86 %

(a) Includes Sustiva, Reyataz, Daklinza and all other BMS and Celgene products acquired as part of the Celgene acquisition that have lost exclusivity in major markets, over-the-counter brands and royalty revenue. Other Brands includes \$205 million worldwide and \$169 million U.S. revenues relating to Celgene products for the six months ended June 30, 2020.

(b) Includes Puerto Rico.

BRISTOL-MYERS SQUIBB COMPANY  
CONSOLIDATED STATEMENTS OF EARNINGS  
FOR THE THREE AND SIX MONTHS ENDED JUNE 30, 2020 AND 2019  
(Unaudited, dollars and shares in millions except per share data)

	Three Months Ended June 30,		Six Months Ended June 30,	
	2020 <sup>(c)</sup>	2019	2020 <sup>(c)</sup>	2019
Net product sales	\$ 9,817	\$ 6,031	\$ 20,358	\$ 11,744
Alliance and other revenues	312	242	552	449
<b>Total Revenues</b>	<b>10,129</b>	<b>6,273</b>	<b>20,910</b>	<b>12,193</b>
Cost of products sold <sup>(a)</sup>	2,699	1,972	6,361	3,796
Marketing, selling and administrative	1,628	1,076	3,234	2,082
Research and development	2,522	1,325	4,894	2,673
Amortization of acquired intangible assets	2,389	24	4,671	48
Other (income)/expense, net	(736)	100	427	(161)
<b>Total Expenses</b>	<b>8,502</b>	<b>4,497</b>	<b>19,587</b>	<b>8,438</b>
Earnings Before Income Taxes	1,627	1,776	1,323	3,755
Provision for Income Taxes	1,707	337	2,169	601
<b>Net (Loss)/Earnings</b>	<b>(80)</b>	<b>1,439</b>	<b>(846)</b>	<b>3,154</b>
Noncontrolling Interest	5	7	14	12
<b>Net (Loss)/Earnings Attributable to BMS</b>	<b>\$ (85)</b>	<b>\$ 1,432</b>	<b>\$ (860)</b>	<b>\$ 3,142</b>
<b>Weighted-Average Common Shares Outstanding:</b>				
Basic	2,263	1,636	2,261	1,635
Diluted	2,263	1,637	2,261	1,637
<b>(Loss)/Earnings per Common Share:</b>				
Basic	\$ (0.04)	\$ 0.88	\$ (0.38)	\$ 1.92
Diluted	(0.04)	0.87	(0.38)	1.92
<b>Other (income)/expense, net</b>				
Interest expense <sup>(b)</sup>	\$ 357	\$ 123	\$ 719	\$ 168
Pension and postretirement	(2)	26	(6)	70
Royalties and licensing income	(311)	(303)	(721)	(611)
Divestiture losses/(gains)	9	8	(7)	8
Acquisition expenses	—	303	—	468
Contingent consideration	(165)	—	391	—
Investment income	(25)	(119)	(86)	(175)
Integration expenses	166	106	340	128
Provision for restructuring	115	10	275	22
Equity investment gains	(818)	(71)	(479)	(246)
Litigation and other settlements	(1)	—	31	1
Transition and other service fees	(50)	(2)	(111)	(4)
Intangible asset impairment	21	15	21	15
Reversion excise tax	—	—	76	—
Other	(32)	4	(16)	(5)
<b>Other (income)/expense, net</b>	<b>\$ (736)</b>	<b>\$ 100</b>	<b>\$ 427</b>	<b>\$ (161)</b>

(a) Excludes amortization of acquired intangible assets.

(b) Includes amortization of purchase price adjustments to Celgene debt.

(c) Includes Celgene results of operations for the entire period.



BRISTOL-MYERS SQUIBB COMPANY  
 SPECIFIED ITEMS  
 FOR THE THREE AND SIX MONTHS ENDED JUNE 30, 2020 AND 2019  
 (Unaudited, dollars in millions)

	Three Months Ended June 30,		Six Months Ended June 30,	
	2020 <sup>(b)</sup>	2019	2020 <sup>(b)</sup>	2019
Inventory purchase price accounting adjustments	\$ 714	\$ —	\$ 2,134	\$ —
Employee compensation charges	1	—	3	—
Site exit and other costs	13	139	29	151
<b>Cost of products sold</b>	<b>728</b>	<b>139</b>	<b>2,166</b>	<b>151</b>
Employee compensation charges	12	—	27	—
Site exit and other costs	(1)	—	5	1
<b>Marketing, selling and administrative</b>	<b>11</b>	<b>—</b>	<b>32</b>	<b>1</b>
License and asset acquisition charges	300	25	325	25
IPRD impairments	—	—	—	32
Inventory purchase price accounting adjustments	—	—	17	—
Employee compensation charges	15	—	33	—
Site exit and other costs	39	19	95	38
<b>Research and development</b>	<b>354</b>	<b>44</b>	<b>470</b>	<b>95</b>
<b>Amortization of acquired intangible assets</b>	<b>2,389</b>	<b>—</b>	<b>4,671</b>	<b>—</b>
Interest expense <sup>(a)</sup>	(41)	83	(82)	83
Pension and postretirement	—	44	—	93
Royalties and licensing income	(18)	—	(101)	—
Divestiture losses/(gains)	9	8	(7)	8
Acquisition expenses	—	303	—	468
Contingent consideration	(165)	—	391	—
Investment income	—	(54)	—	(54)
Integration expenses	166	106	340	128
Provision for restructuring	115	10	275	22
Equity investment gains	(818)	(71)	(479)	(246)
Reversion excise tax	—	—	76	—
<b>Other (income)/expense, net</b>	<b>(752)</b>	<b>429</b>	<b>413</b>	<b>502</b>
<b>Increase to pretax income</b>	<b>2,730</b>	<b>612</b>	<b>7,752</b>	<b>749</b>
Income taxes on items above	(3)	(105)	(294)	(148)
Income taxes attributed to Otezla <sup>®</sup> divestiture	255	—	255	—
Income taxes attributed to internal transfer of intangible assets	853	—	853	—
<b>Income taxes</b>	<b>1,105</b>	<b>(105)</b>	<b>814</b>	<b>(148)</b>
<b>Increase to net earnings</b>	<b>\$ 3,835</b>	<b>\$ 507</b>	<b>\$ 8,566</b>	<b>\$ 601</b>

(a) Includes amortization of purchase price adjustments to Celgene debt.

(b) Includes Celgene results of operations for the entire period.

BRISTOL-MYERS SQUIBB COMPANY  
RECONCILIATION OF CERTAIN GAAP LINE ITEMS TO CERTAIN NON-GAAP LINE ITEMS  
FOR THE THREE AND SIX MONTHS ENDED JUNE 30, 2020 AND 2019  
(Unaudited, dollars and shares in millions except per share data)

	Three Months Ended June 30, 2020			Six Months Ended June 30, 2020		
	GAAP <sup>(a)</sup>	Specified Items <sup>(a)(b)</sup>	Non-GAAP <sup>(a)</sup>	GAAP	Specified Items <sup>(a)(b)</sup>	Non-GAAP <sup>(a)</sup>
Gross Profit	\$ 7,430	\$ 728	\$ 8,158	\$ 14,549	\$ 2,166	\$ 16,715
Marketing, selling and administrative	1,628	(11)	1,617	3,234	(32)	3,202
Research and development	2,522	(354)	2,168	4,894	(470)	4,424
Amortization of acquired intangible assets	2,389	(2,389)	—	4,671	(4,671)	—
Other (income)/expense, net	(736)	752	16	427	(413)	14
Earnings Before Income Taxes	1,627	2,730	4,357	1,323	7,752	9,075
Provision for Income Taxes	1,707	(1,105)	602	2,169	(814)	1,355
Noncontrolling interest	5	—	5	14	—	14
Net (Loss)/Earnings Attributable to BMS used for Diluted EPS Calculation	\$ (85)	\$ 3,835	\$ 3,750	\$ (860)	\$ 8,566	\$ 7,706
Weighted-Average Common Shares Outstanding - Diluted	2,263	2,297	2,297	2,261	2,298	2,298
Diluted (Loss)/Earnings Per Share	\$ (0.04)	\$ 1.67	\$ 1.63	\$ (0.38)	\$ 3.73	\$ 3.35
Effective Tax Rate	104.9 %	(91.1)%	13.8 %	163.9 %	(149.0)%	14.9 %
	Three Months Ended June 30, 2019			Six Months Ended June 30, 2019		
	GAAP	Specified Items <sup>(b)</sup>	Non-GAAP	GAAP	Specified Items <sup>(b)</sup>	Non-GAAP
Gross Profit	\$ 4,301	\$ 139	\$ 4,440	\$ 8,397	\$ 151	\$ 8,548
Marketing, selling and administrative	1,076	—	1,076	2,082	(1)	2,081
Research and development	1,325	(44)	1,281	2,673	(95)	2,578
Amortization of acquired intangible assets	24	—	24	48	—	48
Other (income)/expense, net	100	(429)	(329)	(161)	(502)	(663)
Earnings Before Income Taxes	1,776	612	2,388	3,755	749	4,504
Provision for Income Taxes	337	105	442	601	148	749
Noncontrolling interest	7	—	7	12	—	12
Net Earnings Attributable to BMS used for Diluted EPS Calculation	\$ 1,432	\$ 507	\$ 1,939	\$ 3,142	\$ 601	\$ 3,743
Weighted-Average Common Shares Outstanding - Diluted	1,637	1,637	1,637	1,637	1,637	1,637
Diluted Earnings Per Share	\$ 0.87	\$ 0.31	\$ 1.18	\$ 1.92	\$ 0.37	\$ 2.29
Effective Tax Rate	19.0 %	(0.5)%	18.5 %	16.0 %	0.6 %	16.6 %

(a) Includes Celgene results of operations for the entire period.

(b) Refer to the Specified Items schedule for further details. Effective tax rate on the Specified Items represents the difference between the GAAP and Non-GAAP effective tax rate.

BRISTOL-MYERS SQUIBB COMPANY  
NET DEBT CALCULATION  
AS OF JUNE 30, 2020 AND DECEMBER 31, 2019  
(Unaudited, dollars in millions)

	June 30, 2020	December 31, 2019
Cash and cash equivalents	\$ 19,934	\$ 12,346
Marketable debt securities - current	1,724	3,047
Marketable debt securities - non-current	523	767
<b>Cash, cash equivalents and marketable debt securities</b>	<b>22,181</b>	<b>16,160</b>
Short-term debt obligations	(4,819)	(3,346)
Long-term debt	(41,853)	(43,387)
<b>Net debt position</b>	<b>\$ (24,491)</b>	<b>\$ (30,573)</b>

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**BRISTOL-MYERS SQUIBB COMPANY**  
**QUARTERLY TREND ANALYSIS OF REVENUES**  
(Unaudited, dollars in millions)

Revenues	2019							2020							% Change		FX Impact <sup>(c)</sup>	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(a)</sup>	Year <sup>(a)</sup>	1st Qtr <sup>(b)</sup>	2nd Qtr <sup>(b)</sup>	6 Months <sup>(b)</sup>	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD	Qtr vs. Qtr	YTD vs. YTD
United States	\$ 3,449	\$ 3,667	\$ 7,116	\$ 3,472	\$ 10,588	\$ 4,754	\$ 15,342	\$ 6,766	\$ 6,487	\$ 13,253					77%	86%	—	—
Europe	1,480	1,491	2,971	1,445	4,416	1,850	6,266	2,567	2,136	4,703				43%	58%	(3)%	(3)%	
Rest of the World	874	988	1,862	976	2,838	1,175	4,013	1,335	1,334	2,669				35%	43%	(4)%	(4)%	
Other	117	127	244	114	358	166	524	113	172	285				35%	17%	—	—	
<b>Total</b>	<b>\$ 5,920</b>	<b>\$ 6,273</b>	<b>\$ 12,193</b>	<b>\$ 6,007</b>	<b>\$ 18,200</b>	<b>\$ 7,945</b>	<b>\$ 26,145</b>	<b>\$ 10,781</b>	<b>\$ 10,129</b>	<b>\$ 20,910</b>					<b>61%</b>	<b>71%</b>	<b>(2)%</b>	<b>(2)%</b>

% of Revenues	2019							2020							% Change		FX Impact <sup>(c)</sup>	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(a)</sup>	Year <sup>(a)</sup>	1st Qtr <sup>(b)</sup>	2nd Qtr <sup>(b)</sup>	6 Months <sup>(b)</sup>	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD	Qtr vs. Qtr	YTD vs. YTD
United States	58.2 %	58.4 %	58.3 %	57.8 %	58.2 %	59.8 %	58.7 %	62.8 %	64.0 %	63.4 %								
Europe	25.0 %	23.8 %	24.4 %	24.1 %	24.3 %	23.3 %	24.0 %	23.8 %	21.1 %	22.5 %								
Rest of the World	14.8 %	15.8 %	15.3 %	16.2 %	15.6 %	14.8 %	15.3 %	12.4 %	13.2 %	12.8 %								
Other	2.0 %	2.0 %	2.0 %	1.9 %	1.9 %	2.1 %	2.0 %	1.0 %	1.7 %	1.3 %								
<b>Total</b>	<b>100.0 %</b>	<b>100.0 %</b>	<b>100.0 %</b>	<b>100.0 %</b>	<b>100.0 %</b>	<b>100.0 %</b>	<b>100.0 %</b>	<b>100.0 %</b>	<b>100.0 %</b>	<b>100.0 %</b>								

(a) Includes Celgene product revenues from November 20, 2019 through December 31, 2019.

(b) Includes Celgene product revenues for the entire period.

(c) Foreign exchange impacts were derived by applying the prior period average currency rates to the current period sales.



**BRISTOL-MYERS SQUIBB COMPANY**  
**EARNINGS FROM OPERATIONS**  
(Unaudited, dollars and shares in millions except per share data)

	2019							2020							% Change	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(c)</sup>	Year <sup>(c)</sup>	1st Qtr <sup>(d)</sup>	2nd Qtr <sup>(d)</sup>	6 Months <sup>(d)</sup>	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD
Net product sales	\$ 5,713	\$ 6,031	\$ 11,744	\$ 5,768	\$ 17,512	\$ 7,662	\$ 25,174	\$ 10,541	\$ 9,817	\$ 20,358					63%	73%
Alliance and other revenues	207	242	449	239	688	283	971	240	312	552					29%	23%
<b>Total Revenues</b>	<b>5,920</b>	<b>6,273</b>	<b>12,193</b>	<b>6,007</b>	<b>18,200</b>	<b>7,945</b>	<b>26,145</b>	<b>10,781</b>	<b>10,129</b>	<b>20,910</b>					<b>61%</b>	<b>71%</b>
Cost of products sold <sup>(a)</sup>	1,824	1,972	3,796	1,790	5,586	2,492	8,078	3,662	2,699	6,361					37%	68%
Marketing, selling and administrative	1,006	1,076	2,082	1,055	3,137	1,734	4,871	1,606	1,628	3,234					51%	55%
Research and development	1,348	1,325	2,673	1,378	4,051	2,097	6,148	2,372	2,522	4,894					90%	83%
Amortization of acquired intangible assets	24	24	48	25	73	1,062	1,135	2,282	2,389	4,671					**	**
Other (income)/expense, net	(261)	100	(161)	410	249	689	938	1,163	(736)	427					**	**
<b>Total Expenses</b>	<b>3,941</b>	<b>4,497</b>	<b>8,438</b>	<b>4,658</b>	<b>13,096</b>	<b>8,074</b>	<b>21,170</b>	<b>11,085</b>	<b>8,502</b>	<b>19,587</b>					<b>89%</b>	<b>**</b>
<b>Earnings/(Loss) Before Income Taxes</b>	<b>1,979</b>	<b>1,776</b>	<b>3,755</b>	<b>1,349</b>	<b>5,104</b>	<b>(129)</b>	<b>4,975</b>	<b>(304)</b>	<b>1,627</b>	<b>1,323</b>					<b>(8)%</b>	<b>(65)%</b>
Provision for Income Taxes	264	337	601	(17)	584	931	1,515	462	1,707	2,169					**	**
<b>Net (Loss)/Earnings</b>	<b>1,715</b>	<b>1,439</b>	<b>3,154</b>	<b>1,366</b>	<b>4,520</b>	<b>(1,060)</b>	<b>3,460</b>	<b>(766)</b>	<b>(80)</b>	<b>(846)</b>					<b>**</b>	<b>**</b>
Noncontrolling Interest	5	7	12	13	25	(4)	21	9	5	14					(29)%	17%
<b>Net (Loss)/Earnings Attributable to BMS</b>	<b>\$ 1,710</b>	<b>\$ 1,432</b>	<b>\$ 3,142</b>	<b>\$ 1,353</b>	<b>\$ 4,495</b>	<b>\$ (1,056)</b>	<b>\$ 3,439</b>	<b>\$ (775)</b>	<b>\$ (85)</b>	<b>\$ (860)</b>					<b>**</b>	<b>**</b>
<b>Diluted (Loss)/Earnings per Common Share*</b>	<b>\$ 1.04</b>	<b>\$ 0.87</b>	<b>\$ 1.92</b>	<b>\$ 0.83</b>	<b>\$ 2.75</b>	<b>\$ (0.55)</b>	<b>\$ 2.01</b>	<b>\$ (0.34)</b>	<b>\$ (0.04)</b>	<b>\$ (0.38)</b>					<b>**</b>	<b>**</b>
Weighted-Average Common Shares Outstanding - Diluted	1,637	1,637	1,637	1,634	1,636	1,918	1,712	2,258	2,263	2,261					—	—
Dividends declared per common share	\$ 0.41	\$ 0.41	\$ 0.82	\$ 0.41	\$ 1.23	\$ 0.45	\$ 1.68	\$ 0.45	\$ 0.45	\$ 0.90					10%	10%

  

% of Total Revenues	2019							2020						
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(c)</sup>	Year <sup>(c)</sup>	1st Qtr <sup>(d)</sup>	2nd Qtr <sup>(d)</sup>	6 Months <sup>(d)</sup>	3rd Qtr	9 Months	4th Qtr	Year
Gross Margin	69.2 %	68.6 %	68.9 %	70.2 %	69.3 %	68.6 %	69.1 %	66.0 %	73.4 %	69.6 %				

  

Other Ratios	
Effective tax rate	13.3 %    19.0 %    16.0 %    (1.3) %    11.4 %    (721.7) %    30.5 %    (152.0) %    104.9 %    163.9 %

  

Other (income)/expense, net	2019							2020							% Change	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(c)</sup>	Year <sup>(c)</sup>	1st Qtr <sup>(d)</sup>	2nd Qtr <sup>(d)</sup>	6 Months <sup>(d)</sup>	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD
Interest expense <sup>(b)</sup>	\$ 45	\$ 123	\$ 168	\$ 209	\$ 377	\$ 279	\$ 656	\$ 362	\$ 357	\$ 719					**	**
Pension and postretirement	44	26	70	1,537	1,607	(8)	1,599	(4)	(2)	(6)					**	**
Royalties and licensing income	(308)	(303)	(611)	(356)	(967)	(393)	(1,360)	(410)	(311)	(721)					3%	18%
Divestiture losses/(gains)	—	8	8	(1,179)	(1,171)	3	(1,168)	(16)	9	(7)					13%	**
Acquisition expenses	165	303	468	7	475	182	657	—	—	—					(100)%	(100)%
Contingent consideration	—	—	—	—	—	523	523	556	(165)	391					N/A	N/A
Investment income	(56)	(119)	(175)	(173)	(348)	(116)	(464)	(61)	(25)	(86)					(79)%	(51)%
Integration expenses	22	106	128	96	224	191	415	174	166	340					57%	**
Provision for restructuring	12	10	22	10	32	269	301	160	115	275					**	**
Equity investment (gains)/losses	(175)	(71)	(246)	261	15	(294)	(279)	339	(818)	(479)					**	95%
Litigation and other settlements	1	—	1	(1)	—	77	77	32	(1)	31					N/A	**
Transition and other service fees	(2)	(2)	(4)	(7)	(11)	(26)	(37)	(61)	(50)	(111)					**	**
Intangible asset impairment	—	15	15	—	15	—	15	—	21	21					40%	40%
Reversion excise tax	—	—	—	—	—	—	—	76	—	76					N/A	N/A
Other	(9)	4	(5)	6	1	2	3	16	(32)	(16)					**	**
<b>Other (income)/expense, net</b>	<b>\$ (261)</b>	<b>\$ 100</b>	<b>\$ (161)</b>	<b>\$ 410</b>	<b>\$ 249</b>	<b>\$ 689</b>	<b>\$ 938</b>	<b>\$ 1,163</b>	<b>\$ (736)</b>	<b>\$ 427</b>					<b>**</b>	<b>**</b>

\* Quarterly amounts may not add to the year-to-date amounts, as each period is computed on a discrete basis.

\*\* In excess of +/- 100%.

(a) Excludes amortization of acquired intangible assets.

(b) Includes amortization of purchase price adjustments to Celgene debt.

(c) Includes Celgene results of operations from November 20, 2019 through December 31, 2019.

(d) Includes Celgene results of operations for the entire period.

**BRISTOL-MYERS SQUIBB COMPANY**  
**RECONCILIATION OF GAAP AND NON-GAAP GROWTH DOLLARS AND PERCENTAGES EXCLUDING FOREIGN EXCHANGE IMPACT**  
**FOR THE PERIOD ENDED JUNE 30, 2020**  
**(Unaudited, dollars in millions)**

QUARTER-TO-DATE	2020 <sup>(b)</sup>	2019	\$ Change	% Change	Favorable / (Unfavorable) FX Impact \$*	2020 Excluding FX	Favorable / (Unfavorable) FX Impact %*	% Change Excluding FX
Revenues	\$ 10,129	\$ 6,273	\$ 3,856	61 %	\$ (85)	\$ 10,214	(2) %	63 %
Gross profit	7,430	4,301	3,129	73 %	N/A	N/A	N/A	N/A
Gross profit excluding specified items <sup>(a)</sup>	8,158	4,440	3,718	84 %	N/A	N/A	N/A	N/A
<b>Gross profit excluding specified items as a % of revenues</b>	<b>80.5 %</b>	<b>70.8 %</b>						
Marketing, selling and administrative	1,628	1,076	552	51 %	17	1,645	2 %	53 %
Marketing, selling and administrative excluding specified items <sup>(a)</sup>	1,617	1,076	541	50 %	17	1,634	2 %	52 %
<b>Marketing, selling and administrative excluding specified items as a % of revenues</b>	<b>16.0 %</b>	<b>17.2 %</b>						
Research and development	2,522	1,325	1,197	90 %	6	2,528	1 %	91 %
Research and development excluding specified items <sup>(a)</sup>	2,168	1,281	887	69 %	6	2,174	1 %	70 %
<b>Research and development excluding specified items as a % of revenues</b>	<b>21.4 %</b>	<b>20.4 %</b>						

YEAR-TO-DATE	2020 <sup>(b)</sup>	2019	\$ Change	% Change	Favorable / (Unfavorable) FX Impact \$*	2020 Excluding FX	Favorable / (Unfavorable) FX Impact %*	% Change Excluding FX
Revenues	\$ 20,910	\$ 12,193	\$ 8,717	71 %	\$ (167)	\$ 21,077	(2) %	73 %
Gross profit	14,549	8,397	6,152	73 %	N/A	N/A	N/A	N/A
Gross profit excluding specified items <sup>(a)</sup>	16,715	8,548	8,167	96 %	N/A	N/A	N/A	N/A
<b>Gross profit excluding specified items as a % of revenues</b>	<b>79.9 %</b>	<b>70.1 %</b>						
Marketing, selling and administrative	3,234	2,082	1,152	55 %	29	3,263	2 %	57 %
Marketing, selling and administrative excluding specified items <sup>(a)</sup>	3,202	2,081	1,121	54 %	29	3,231	1 %	55 %
<b>Marketing, selling and administrative excluding specified items as a % of revenues</b>	<b>15.3 %</b>	<b>17.1 %</b>						
Research and development	4,894	2,673	2,221	83 %	10	4,904	—	83 %
Research and development excluding specified items <sup>(a)</sup>	4,424	2,578	1,846	72 %	10	4,434	—	72 %
<b>Research and development excluding specified items as a % of revenues</b>	<b>21.2 %</b>	<b>21.1 %</b>						

\* Foreign exchange impacts were derived by applying the prior period average currency rates to the current period sales and expenses.

(a) Refer to the Specified Items schedule for further details.

(b) Includes Celgene results of operations for the entire period.

**BRISTOL-MYERS SQUIBB COMPANY**  
**WORLDWIDE REVENUES**  
**QUARTERLY REVENUES TREND ANALYSIS**  
(Unaudited, dollars in millions)

	2019							2020							\$ Change		% Change	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(b)</sup>	Year <sup>(b)</sup>	1st Qtr <sup>(c)</sup>	2nd Qtr <sup>(c)</sup>	6 Months <sup>(c)</sup>	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD	Qtr vs. Qtr	YTD vs. YTD
<b>Prioritized Brands</b>																		
Revlimid	\$ —	\$ —	\$ —	\$ —	\$ —	\$ 1,299	\$ 1,299	\$ 2,915	\$ 2,884	\$ 5,799					\$ 2,884	\$ 5,799	N/A	N/A
Eliquis	1,925	2,042	3,967	1,928	5,895	2,034	7,929	2,641	2,163	4,804					121	837	6%	21%
Opdivo	1,801	1,823	3,624	1,817	5,441	1,763	7,204	1,766	1,653	3,419					(170)	(205)	(9)%	(6)%
Orencia	640	778	1,418	767	2,185	792	2,977	714	750	1,464					(28)	46	(4)%	3%
Pomalyst/Imnovid	—	—	—	—	—	322	322	713	745	1,458					745	1,458	N/A	N/A
Sprycel	459	544	1,003	558	1,561	549	2,110	521	511	1,032					(33)	29	(6)%	3%
Yervoy	384	367	751	353	1,104	385	1,489	396	369	765					2	14	1%	2%
Abraxane	—	—	—	—	—	166	166	300	308	608					308	608	N/A	N/A
Empliciti	83	91	174	89	263	94	357	97	97	194					6	20	7%	11%
Reblozyl	—	—	—	—	—	—	—	8	55	63					55	63	N/A	N/A
Inrebic	—	—	—	—	—	5	5	12	15	27					15	27	N/A	N/A
Zeposia	—	—	—	—	—	—	—	—	1	1					1	1	N/A	N/A
<b>Established Brands</b>																		
Baraclude	141	147	288	145	433	122	555	122	121	243					(26)	(45)	(18)%	(16)%
Vidaza	—	—	—	—	—	58	58	158	126	284					126	284	N/A	N/A
Other Brands <sup>(a)</sup>	487	481	968	350	1,318	356	1,674	418	331	749					(150)	(219)	(31)%	(23)%
<b>Total</b>	<u>\$ 5,920</u>	<u>\$ 6,273</u>	<u>\$ 12,193</u>	<u>\$ 6,007</u>	<u>\$ 18,200</u>	<u>\$ 7,945</u>	<u>\$ 26,145</u>	<u>\$ 10,781</u>	<u>\$ 10,129</u>	<u>\$ 20,910</u>					<u>\$ 3,856</u>	<u>\$ 8,717</u>	<u>61%</u>	<u>71%</u>

(a) Includes Sustiva, Reyataz, Daklinza and all other BMS and Celgene products acquired as part of the Celgene acquisition that have lost exclusivity in major markets, over-the-counter (OTC) brands and royalty revenue. Other Brands includes \$83 million and \$205 million relating to Celgene products in the three and six months ended June 30, 2020, respectively.

(b) Includes Celgene product revenues from November 20, 2019 through December 31, 2019.

(c) Includes Celgene product revenues for the entire period.

**BRISTOL-MYERS SQUIBB COMPANY**  
**WORLDWIDE PRO FORMA REVENUES**  
**QUARTERLY REVENUES TREND ANALYSIS**  
(Unaudited, dollars in millions)

	2019							2020							S Change		% Change	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(e)</sup>	Year	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD	Qtr vs. Qtr	YTD vs. YTD
<b>Prioritized Brands</b>																		
Revlimid <sup>(a)</sup>	\$ 2,562	\$ 2,718	\$ 5,280	\$ 2,758	\$ 8,038	\$ 2,785	\$ 10,823	\$ 2,915	\$ 2,884	\$ 5,799					\$ 166	\$ 519	6%	10%
Eliquis	1,925	2,042	3,967	1,928	5,895	2,034	7,929	2,641	2,163	4,804					121	837	6%	21%
Opdivo	1,801	1,823	3,624	1,817	5,441	1,763	7,204	1,766	1,653	3,419					(170)	(205)	(9)%	(6)%
Orencia	640	778	1,418	767	2,185	792	2,977	714	750	1,464					(28)	46	(4)%	3%
Pomalyst/Imnovid <sup>(a)</sup>	554	617	1,171	662	1,833	692	2,525	713	745	1,458					128	287	21%	25%
Sprycel	459	544	1,003	558	1,561	549	2,110	521	511	1,032					(33)	29	(6)%	3%
Yervoy	384	367	751	353	1,104	385	1,489	396	369	765					2	14	1%	2%
Abraxane <sup>(a)</sup>	285	314	599	317	916	336	1,252	300	308	608					(6)	9	(2)%	2%
Empliciti	83	91	174	89	263	94	357	97	97	194					6	20	7%	11%
Rebzozyl <sup>(b)</sup>	—	—	—	—	—	—	—	8	55	63					55	63	N/A	N/A
Inrebic <sup>(b)</sup>	—	—	—	2	2	9	11	12	15	27					15	27	N/A	N/A
Zeposia <sup>(b)</sup>	—	—	—	—	—	—	—	—	1	1					1	1	N/A	N/A
<b>Established Brands</b>																		
Baraclude	141	147	288	145	433	122	555	122	121	243					(26)	(45)	(18)%	(16)%
Vidaza <sup>(a)</sup>	148	162	310	146	456	149	605	158	126	284					(36)	(26)	(22)%	(8)%
Other Brands <sup>(c)</sup>	552	557	1,109	420	1,529	393	1,922	418	331	749					(226)	(360)	(41)%	(32)%
<b>Total<sup>(d)</sup></b>	<b>\$ 9,534</b>	<b>\$ 10,160</b>	<b>\$ 19,694</b>	<b>\$ 9,962</b>	<b>\$ 29,656</b>	<b>\$ 10,103</b>	<b>\$ 39,759</b>	<b>\$ 10,781</b>	<b>\$ 10,129</b>	<b>\$ 20,910</b>					<b>\$ (31)</b>	<b>\$ 1,216</b>	<b>—</b>	<b>6%</b>

(a) Products were acquired as part of the Celgene acquisition. Reflects product revenues for the period prior to November 20, 2019, which was the date of the acquisition. All product revenues prior to November 20, 2019 have been recast to exclude foreign currency hedge gains and losses.

(b) Products were acquired as part of the Celgene acquisition. Reflects product revenues for the period prior to November 20, 2019, which was the date of the acquisition.

(c) Includes Sustiva, Reyataz, Daklinza and all other BMS and Celgene products acquired as part of the Celgene acquisition that have lost exclusivity in major markets, over-the-counter (OTC) brands and royalty revenue. Reflects Celgene product revenues for the period prior to November 20, 2019, which was the date of the acquisition, for such Celgene products.

(d) All historically reported Celgene revenues have been recast to exclude Otezla<sup>®</sup> product revenues.

(e) Celgene product revenues for the period of October 1, 2019 through November 19, 2019 are included below:

Revlimid	\$	1,486
Pomalyst/Imnovid		370
Abraxane		170
Inrebic		4
Vidaza		91
Other Brands		37
<b>Total</b>	<b>\$</b>	<b>2,158</b>

**BRISTOL-MYERS SQUIBB COMPANY**  
**U.S. REVENUES**  
**QUARTERLY REVENUES TREND ANALYSIS**  
(Unaudited, dollars in millions)

	2019							2020							% Change	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(c)</sup>	Year <sup>(c)</sup>	1st Qtr <sup>(d)</sup>	2nd Qtr <sup>(d)</sup>	6 Months <sup>(d)</sup>	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD
<b>Prioritized Brands</b>																
Revlimid	\$ —	\$ —	\$ —	\$ —	\$ —	\$ 899	\$ 899	\$ 1,966	\$ 2,048	\$ 4,014					N/A	N/A
Eliquis	1,206	1,269	2,475	1,124	3,599	1,156	4,755	1,777	1,363	3,140					7%	27%
Opdivo	1,124	1,112	2,236	1,088	3,324	1,020	4,344	1,008	956	1,964					(14)%	(12)%
Orencia	449	566	1,015	554	1,569	577	2,146	500	554	1,054					(2)%	4%
Pomalyst/Imnovid	—	—	—	—	—	226	226	489	522	1,011					N/A	N/A
Sprycel	240	307	547	325	872	319	1,191	300	308	608					—	11%
Yervoy	275	253	528	222	750	254	1,004	257	254	511					—	(3)%
Abraxane	—	—	—	—	—	122	122	205	218	423					N/A	N/A
Empliciti	58	63	121	62	183	63	246	59	59	118					(6)%	(2)%
Reblozyl	—	—	—	—	—	—	—	8	55	63					N/A	N/A
Inrebic	—	—	—	—	—	5	5	12	15	27					N/A	N/A
Zeposia	—	—	—	—	—	—	—	—	1	1					N/A	N/A
<b>Established Brands</b>																
Baraclude	7	7	14	2	16	4	20	3	3	6					(57)%	(57)%
Vidaza	—	—	—	—	—	1	1	2	—	2					N/A	N/A
Other Brands <sup>(a)</sup>	90	90	180	95	275	108	383	180	131	311					46%	73%
<b>Total<sup>(b)</sup></b>	<b>\$ 3,449</b>	<b>\$ 3,667</b>	<b>\$ 7,116</b>	<b>\$ 3,472</b>	<b>\$ 10,588</b>	<b>\$ 4,754</b>	<b>\$ 15,342</b>	<b>\$ 6,766</b>	<b>\$ 6,487</b>	<b>\$ 13,253</b>					<b>77%</b>	<b>86%</b>

(a) Includes Sustiva, Reyataz, Daklinza and all other BMS and Celgene products acquired as part of the Celgene acquisition that have lost exclusivity in major markets, OTC brands and royalty revenue. Other Brands includes \$66 million and \$169 million relating to Celgene products in the three and six months ended June 30, 2020, respectively.

(b) Includes Puerto Rico.

(c) Includes Celgene product revenues from November 20, 2019 through December 31, 2019.

(d) Includes Celgene product revenues for the entire period.

**BRISTOL-MYERS SQUIBB COMPANY**  
**U.S. PRO FORMA REVENUES**  
**QUARTERLY REVENUES TREND ANALYSIS**  
(Unaudited, dollars in millions)

	2019							2020							% Change	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(d)</sup>	Year	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD
<b>Prioritized Brands</b>																
Revlimid <sup>(a)</sup>	\$ 1,686	\$ 1,810	\$ 3,496	\$ 1,902	\$ 5,398	\$ 1,914	\$ 7,312	\$ 1,966	\$ 2,048	\$ 4,014					13%	15%
Eliquis	1,206	1,269	2,475	1,124	3,599	1,156	4,755	1,777	1,363	3,140				7%	27%	
Opdivo	1,124	1,112	2,236	1,088	3,324	1,020	4,344	1,008	956	1,964				(14)%	(12)%	
Orencia	449	566	1,015	554	1,569	577	2,146	500	554	1,054				(2)%	4%	
Pomalyst/Imnovid <sup>(a)</sup>	390	447	837	469	1,306	489	1,795	489	522	1,011				17%	21%	
Sprycel	240	307	547	325	872	319	1,191	300	308	608				—	11%	
Yervoy	275	253	528	222	750	254	1,004	257	254	511				—	(3)%	
Abraxane <sup>(a)</sup>	196	207	403	206	609	237	846	205	218	423				5%	5%	
Empliciti	58	63	121	62	183	63	246	59	59	118				(6)%	(2)%	
Reblozyl <sup>(a)</sup>	—	—	—	—	—	—	—	8	55	63				N/A	N/A	
Inrebic <sup>(a)</sup>	—	—	—	2	2	9	11	12	15	27				N/A	N/A	
Zeposia <sup>(a)</sup>	—	—	—	—	—	—	—	—	1	1				N/A	N/A	
<b>Established Brands</b>																
Baraclude	7	7	14	2	16	4	20	3	3	6				(57)%	(57)%	
Vidaza <sup>(a)</sup>	3	3	6	2	8	2	10	2	—	2				(100)%	(67)%	
Other Brands <sup>(b)</sup>	135	145	280	147	427	136	563	180	131	311				(10)%	11%	
<b>Total<sup>(c)</sup></b>	<b>\$ 5,769</b>	<b>\$ 6,189</b>	<b>\$ 11,958</b>	<b>\$ 6,105</b>	<b>\$ 18,063</b>	<b>\$ 6,180</b>	<b>\$ 24,243</b>	<b>\$ 6,766</b>	<b>\$ 6,487</b>	<b>\$ 13,253</b>				<b>5%</b>	<b>11%</b>	

(a) Products were acquired as part of the Celgene acquisition. Reflects product revenues for the period prior to November 20, 2019, which was the date of the acquisition.

(b) Includes Sustiva, Reyataz, Daklinza and all other BMS and Celgene products acquired as part of the Celgene acquisition that have lost exclusivity in major markets, OTC brands and royalty revenue. Reflects Celgene product revenues for the period prior to November 20, 2019, which was the date of the acquisition, for such Celgene products.

(c) Includes Puerto Rico. All historically reported Celgene revenues have been recast to exclude Otezla<sup>®</sup> product revenues.

(d) Celgene product revenues for the period of October 1, 2019 through November 19, 2019 are included below:

Revlimid	\$	1,015
Pomalyst/Imnovid		263
Abraxane		115
Inrebic		4
Vidaza		1
Other Brands		28
<b>Total</b>	<b>\$</b>	<b>1,426</b>

**BRISTOL-MYERS SQUIBB COMPANY**  
**INTERNATIONAL REVENUES**  
**QUARTERLY REVENUES TREND ANALYSIS**  
(Unaudited, dollars in millions)

	2019							2020							% Change <sup>(c)</sup>	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(b)</sup>	Year <sup>(b)</sup>	1st Qtr <sup>(d)</sup>	2nd Qtr <sup>(d)</sup>	6 Months <sup>(d)</sup>	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD
<b>Prioritized Brands</b>																
Revlimid	\$ —	\$ —	\$ —	\$ —	\$ —	\$ 400	\$ 400	\$ 949	\$ 836	\$ 1,785					N/A	N/A
Eliquis	719	773	1,492	804	2,296	878	3,174	864	800	1,664					3%	12%
Opdivo	677	711	1,388	729	2,117	743	2,860	758	697	1,455					(2)%	5%
Orencia	191	212	403	213	616	215	831	214	196	410					(8)%	2%
Pomalyst/Imnovid	—	—	—	—	—	96	96	224	223	447					N/A	N/A
Sprycel	219	237	456	233	689	230	919	221	203	424					(14)%	(7)%
Yervoy	109	114	223	131	354	131	485	139	115	254					1%	14%
Abraxane	—	—	—	—	—	44	44	95	90	185					N/A	N/A
Empliciti	25	28	53	27	80	31	111	38	38	76					36%	43%
<b>Established Brands</b>																
Baraclude	134	140	274	143	417	118	535	119	118	237					(16)%	(14)%
Vidaza	—	—	—	—	—	57	57	156	126	282					N/A	N/A
Other Brands <sup>(a)</sup>	397	391	788	255	1,043	248	1,291	238	200	438					(49)%	(44)%
<b>Total</b>	<b>\$ 2,471</b>	<b>\$ 2,606</b>	<b>\$ 5,077</b>	<b>\$ 2,535</b>	<b>\$ 7,612</b>	<b>\$ 3,191</b>	<b>\$ 10,803</b>	<b>\$ 4,015</b>	<b>\$ 3,642</b>	<b>\$ 7,657</b>					<b>40%</b>	<b>51%</b>

(a) Includes Sustiva, Reyataz, Daklinza and all other BMS and Celgene products acquired as part of the Celgene acquisition that have lost exclusivity in major markets, OTC brands and royalty revenue. Other Brands includes \$17 million and \$36 million relating to Celgene products in the three and six months ended June 30, 2020, respectively.

(b) Includes Celgene product revenues from November 20, 2019 through December 31, 2019.

(c) The foreign exchange impact on international revenues was unfavorable 3% for both the second quarter and year-to-date. The foreign exchange impact on Prioritized Brands is included below.

(d) Includes Celgene product revenues for the entire period.

	Quarter-to-Date			Year-to-Date		
	Revenue Change %	Favorable/ (Unfavorable) FX Impact %	Revenue Change % Excluding FX	Revenue Change %	Favorable/ (Unfavorable) FX Impact %	Revenue Change % Excluding FX
Eliquis	3%	(2)%	5%	12%	(2)%	14%
Opdivo	(2)%	(5)%	3%	5%	(5)%	10%
Orencia	(8)%	(3)%	(5)%	2%	(3)%	5%
Sprycel	(14)%	(3)%	(11)%	(7)%	(3)%	(4)%
Yervoy	1%	(5)%	6%	14%	(5)%	19%
Empliciti	36%	—	36%	43%	(1)%	44%

**BRISTOL-MYERS SQUIBB COMPANY**  
**INTERNATIONAL PRO FORMA REVENUES**  
**QUARTERLY REVENUES TREND ANALYSIS**  
(Unaudited, dollars in millions)

	2019							2020							% Change	
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(d)</sup>	Year	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr	Year	Qtr vs. Qtr	YTD vs. YTD
<b>Prioritized Brands</b>																
Revlimid <sup>(a)</sup>	\$ 876	\$ 908	\$ 1,784	\$ 856	\$ 2,640	\$ 871	\$ 3,511	\$ 949	\$ 836	\$ 1,785					(8)%	—
Eliquis	719	773	1,492	804	2,296	878	3,174	864	800	1,664					3%	12%
Opdivo	677	711	1,388	729	2,117	743	2,860	758	697	1,455					(2)%	5%
Orencia	191	212	403	213	616	215	831	214	196	410					(8)%	2%
Pomalyst/Imnovid <sup>(a)</sup>	164	170	334	193	527	203	730	224	223	447					31%	34%
Sprycel	219	237	456	233	689	230	919	221	203	424					(14)%	(7)%
Yervoy	109	114	223	131	354	131	485	139	115	254					1%	14%
Abraxane <sup>(a)</sup>	89	107	196	111	307	99	406	95	90	185					(16)%	(6)%
Empliciti	25	28	53	27	80	31	111	38	38	76					36%	43%
<b>Established Brands</b>																
Baraclude	134	140	274	143	417	118	535	119	118	237					(16)%	(14)%
Vidaza <sup>(a)</sup>	145	159	304	144	448	147	595	156	126	282					(21)%	(7)%
Other Brands <sup>(b)</sup>	417	412	829	273	1,102	257	1,359	238	200	438					(51)%	(47)%
<b>Total<sup>(c)</sup></b>	<b>\$ 3,765</b>	<b>\$ 3,971</b>	<b>\$ 7,736</b>	<b>\$ 3,857</b>	<b>\$ 11,593</b>	<b>\$ 3,923</b>	<b>\$ 15,516</b>	<b>\$ 4,015</b>	<b>\$ 3,642</b>	<b>\$ 7,657</b>					<b>(8)%</b>	<b>(1)%</b>

(a) Products were acquired as part of the Celgene acquisition. Reflects product revenues for the period prior to November 20, 2019, which was the date of the acquisition. All product revenues prior to November 20, 2019 have been recast to exclude foreign currency hedge gains and losses.

(b) Includes Sustiva, Reyataz, Daklinza and all other BMS and Celgene products acquired as part of the Celgene acquisition that have lost exclusivity in major markets, OTC brands and royalty revenue. Reflects Celgene product revenues for the period prior to November 20, 2019, which was the date of the acquisition, for such Celgene products.

(c) All historically reported Celgene revenues have been recast to exclude Otezla<sup>®</sup> product revenues.

(d) Celgene product revenues for the period of October 1, 2019 through November 19, 2019 are included below:

Revlimid	\$	471
Pomalyst/Imnovid		107
Abraxane		55
Vidaza		90
Other Brands		9
<b>Total</b>	<b>\$</b>	<b>732</b>



**BRISTOL-MYERS SQUIBB COMPANY**  
**SPECIFIED ITEMS**  
(Unaudited, dollars in millions)

	2019							2020						
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(b)</sup>	Year <sup>(b)</sup>	1st Qtr <sup>(c)</sup>	2nd Qtr <sup>(c)</sup>	6 Months <sup>(c)</sup>	3rd Qtr	9 Months	4th Qtr	Year
Inventory purchase price accounting adjustments	\$ —	\$ —	\$ —	\$ —	\$ —	\$ 660	\$ 660	\$ 1,420	\$ 714	\$ 2,134				
Employee compensation charges	—	—	—	—	—	1	1	2	1	3				
Site exit and other costs	12	139	151	22	173	24	197	16	13	29				
<b>Cost of products sold</b>	<b>12</b>	<b>139</b>	<b>151</b>	<b>22</b>	<b>173</b>	<b>685</b>	<b>858</b>	<b>1,438</b>	<b>728</b>	<b>2,166</b>				
Employee compensation charges	—	—	—	—	—	27	27	15	12	27				
Site exit and other costs	1	—	1	—	1	8	9	6	(1)	5				
<b>Marketing, selling and administrative</b>	<b>1</b>	<b>—</b>	<b>1</b>	<b>—</b>	<b>1</b>	<b>35</b>	<b>36</b>	<b>21</b>	<b>11</b>	<b>32</b>				
License and asset acquisition charges	—	25	25	—	25	—	25	25	300	325				
IPRD impairments	32	—	32	—	32	—	32	—	—	—				
Inventory purchase price accounting adjustments	—	—	—	—	—	—	—	17	—	17				
Employee compensation charges	—	—	—	—	—	33	33	18	15	33				
Site exit and other costs	19	19	38	20	58	109	167	56	39	95				
<b>Research and development</b>	<b>51</b>	<b>44</b>	<b>95</b>	<b>20</b>	<b>115</b>	<b>142</b>	<b>257</b>	<b>116</b>	<b>354</b>	<b>470</b>				
<b>Amortization of acquired intangible assets</b>	<b>—</b>	<b>—</b>	<b>—</b>	<b>—</b>	<b>—</b>	<b>1,062</b>	<b>1,062</b>	<b>2,282</b>	<b>2,389</b>	<b>4,671</b>				
Interest expense <sup>(a)</sup>	—	83	83	166	249	73	322	(41)	(41)	(82)				
Pension and postretirement	49	44	93	1,545	1,638	(3)	1,635	—	—	—				
Royalties and licensing income	—	—	—	(9)	(9)	(15)	(24)	(83)	(18)	(101)				
Divestiture losses/(gains)	—	8	8	(1,179)	(1,171)	3	(1,168)	(16)	9	(7)				
Acquisition expenses	165	303	468	7	475	182	657	—	—	—				
Contingent consideration	—	—	—	—	—	523	523	556	(165)	391				
Investment income	—	(54)	(54)	(99)	(153)	(44)	(197)	—	—	—				
Integration expenses	22	106	128	96	224	191	415	174	166	340				
Provision for restructuring	12	10	22	10	32	269	301	160	115	275				
Equity investment (gains)/losses	(175)	(71)	(246)	261	15	(294)	(279)	339	(818)	(479)				
Litigation and other settlements	—	—	—	—	—	75	75	—	—	—				
Reversion excise tax	—	—	—	—	—	—	—	76	—	76				
Other	—	—	—	—	—	2	2	—	—	—				
<b>Other (income)/expense, net</b>	<b>73</b>	<b>429</b>	<b>502</b>	<b>798</b>	<b>1,300</b>	<b>962</b>	<b>2,262</b>	<b>1,165</b>	<b>(752)</b>	<b>413</b>				
<b>Increase to pretax income</b>	<b>137</b>	<b>612</b>	<b>749</b>	<b>840</b>	<b>1,589</b>	<b>2,886</b>	<b>4,475</b>	<b>5,022</b>	<b>2,730</b>	<b>7,752</b>				
Income taxes on items above	(43)	(105)	(148)	(275)	(423)	(264)	(687)	(291)	(3)	(294)				
Income taxes attributed to Otezla® divestiture	—	—	—	—	—	808	808	—	255	255				
Income taxes attributed to internal transfer of intangible assets	—	—	—	—	—	—	—	—	853	853				
<b>Income taxes</b>	<b>(43)</b>	<b>(105)</b>	<b>(148)</b>	<b>(275)</b>	<b>(423)</b>	<b>544</b>	<b>121</b>	<b>(291)</b>	<b>1,105</b>	<b>814</b>				
<b>Increase to net earnings</b>	<b>\$ 94</b>	<b>\$ 507</b>	<b>\$ 601</b>	<b>\$ 565</b>	<b>\$ 1,166</b>	<b>\$ 3,430</b>	<b>\$ 4,596</b>	<b>\$ 4,731</b>	<b>\$ 3,835</b>	<b>\$ 8,566</b>				

(a) Includes amortization of purchase price adjustments to Celgene debt.

(b) Includes Celgene results of operations from November 20, 2019 through December 31, 2019.

(c) Includes Celgene results of operations for the entire period.

**BRISTOL-MYERS SQUIBB COMPANY**  
**RECONCILIATION OF CERTAIN GAAP LINE ITEMS TO CERTAIN NON-GAAP LINE ITEMS**  
(Unaudited, dollars in millions)

	2019							2020						
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(b)</sup>	Year <sup>(b)</sup>	1st Qtr <sup>(c)</sup>	2nd Qtr <sup>(c)</sup>	6 Months <sup>(c)</sup>	3rd Qtr	9 Months	4th Qtr	Year
<b>Gross Profit</b>	\$ 4,096	\$ 4,301	\$ 8,397	\$ 4,217	\$ 12,614	\$ 5,453	\$ 18,067	\$ 7,119	\$ 7,430	\$ 14,549				
Specified items <sup>(a)</sup>	12	139	151	22	173	685	858	1,438	728	2,166				
<b>Gross profit excluding specified items</b>	4,108	4,440	8,548	4,239	12,787	6,138	18,925	8,557	8,158	16,715				
<b>Marketing, selling and administrative</b>	1,006	1,076	2,082	1,055	3,137	1,734	4,871	1,606	1,628	3,234				
Specified items <sup>(a)</sup>	(1)	—	(1)	—	(1)	(35)	(36)	(21)	(11)	(32)				
<b>Marketing, selling and administrative excluding specified items</b>	1,005	1,076	2,081	1,055	3,136	1,699	4,835	1,585	1,617	3,202				
<b>Research and development</b>	1,348	1,325	2,673	1,378	4,051	2,097	6,148	2,372	2,522	4,894				
Specified items <sup>(a)</sup>	(51)	(44)	(95)	(20)	(115)	(142)	(257)	(116)	(354)	(470)				
<b>Research and development excluding specified items</b>	1,297	1,281	2,578	1,358	3,936	1,955	5,891	2,256	2,168	4,424				
<b>Amortization of acquired intangible assets</b>	24	24	48	25	73	1,062	1,135	2,282	2,389	4,671				
Specified items <sup>(a)</sup>	—	—	—	—	—	(1,062)	(1,062)	(2,282)	(2,389)	(4,671)				
<b>Amortization of acquired intangible assets excluding specified items</b>	24	24	48	25	73	—	73	—	—	—				
<b>Other (income)/expense, net</b>	(261)	100	(161)	410	249	689	938	1,163	(736)	427				
Specified items <sup>(a)</sup>	(73)	(429)	(502)	(798)	(1,300)	(962)	(2,262)	(1,165)	752	(413)				
<b>Other expense/(income), net excluding specified items</b>	(334)	(329)	(663)	(388)	(1,051)	(273)	(1,324)	(2)	16	14				

(a) Refer to the Specified Items schedule for further details.

(b) Includes Celgene results of operations from November 20, 2019 through December 31, 2019.

(c) Includes Celgene results of operations for the entire period.

**BRISTOL-MYERS SQUIBB COMPANY**  
**RECONCILIATION OF GAAP TO NON-GAAP EPS**  
(Unaudited, dollars and shares in millions except per share data)

	2019							2020						
	1st Qtr	2nd Qtr	6 Months	3rd Qtr	9 Months	4th Qtr <sup>(b)</sup>	Year <sup>(b)</sup>	1st Qtr <sup>(c)</sup>	2nd Qtr <sup>(c)</sup>	6 Months <sup>(c)</sup>	3rd Qtr	9 Months	4th Qtr	Year
<b>Earnings/(Loss) before income taxes</b>	\$ 1,979	\$ 1,776	\$ 3,755	\$ 1,349	\$ 5,104	\$ (129)	\$ 4,975	\$ (304)	\$ 1,627	\$ 1,323				
<b>Specified items<sup>(a)</sup></b>	137	612	749	840	1,589	2,886	4,475	5,022	2,730	7,752				
<b>Earnings before income taxes excluding specified items</b>	2,116	2,388	4,504	2,189	6,693	2,757	9,450	4,718	4,357	9,075				
<b>Provision for income taxes</b>	264	337	601	(17)	584	931	1,515	462	1,707	2,169				
<b>Income taxes on specified items<sup>(a)</sup></b>	43	105	148	275	423	264	687	291	3	294				
<b>Income taxes attributed to Otezla<sup>®</sup> divestiture<sup>(a)</sup></b>	—	—	—	—	—	(808)	(808)	—	(255)	(255)				
<b>Income taxes attributed to internal transfer of intangible assets<sup>(a)</sup></b>	—	—	—	—	—	—	—	—	(853)	(853)				
<b>Provision for income taxes excluding tax on specified items and income taxes attributed to Otezla<sup>®</sup> divestiture and internal transfer of intangible assets</b>	307	442	749	258	1,007	387	1,394	753	602	1,355				
<b>Noncontrolling Interest</b>	5	7	12	13	25	(4)	21	9	5	14				
<b>Specified items<sup>(a)</sup></b>	—	—	—	—	—	—	—	—	—	—				
<b>Noncontrolling Interest excluding specified items</b>	5	7	12	13	25	(4)	21	9	5	14				
<b>Net (loss)/earnings attributable to BMS used for Diluted EPS Calculation - GAAP</b>	1,710	1,432	3,142	1,353	4,495	(1,056)	3,439	(775)	(85)	(860)				
<b>Specified items<sup>(a)</sup></b>	94	507	601	565	1,166	3,430	4,596	4,731	3,835	8,566				
<b>Net earnings attributable to BMS used for Diluted EPS Calculation excluding specified items - Non-GAAP</b>	1,804	1,939	3,743	1,918	5,661	2,374	8,035	3,956	3,750	7,706				
<b>Weighted-average Common Shares Outstanding - Diluted-GAAP</b>	1,637	1,637	1,637	1,634	1,636	1,918	1,712	2,258	2,263	2,261				
<b>Weighted-average Common Shares Outstanding - Diluted-Non-GAAP</b>	1,637	1,637	1,637	1,634	1,636	1,941	1,712	2,298	2,297	2,298				
<b>Diluted (Loss)/Earnings Per Share - GAAP*</b>	\$ 1.04	\$ 0.87	\$ 1.92	\$ 0.83	\$ 2.75	\$ (0.55)	\$ 2.01	\$ (0.34)	\$ (0.04)	\$ (0.38)				
<b>Diluted Earnings Per Share attributable to specified items<sup>(a)</sup></b>	0.06	0.31	0.37	0.34	0.71	1.77	2.68	2.06	1.67	3.73				
<b>Diluted Earnings Per Share - Non-GAAP*</b>	\$ 1.10	\$ 1.18	\$ 2.29	\$ 1.17	\$ 3.46	\$ 1.22	\$ 4.69	\$ 1.72	\$ 1.63	\$ 3.35				
<b>Effective Tax Rate</b>	13.3 %	19.0 %	16.0 %	(1.3) %	11.4 %	(721.7) %	30.5 %	(152.0) %	104.9 %	163.9 %				
<b>Specified items<sup>(a)</sup></b>	1.2 %	(0.5) %	0.6 %	13.1 %	3.6 %	735.7 %	(15.7) %	168.0 %	(91.1) %	(149.0) %				
<b>Effective Tax Rate excluding specified items</b>	14.5 %	18.5 %	16.6 %	11.8 %	15.0 %	14.0 %	14.8 %	16.0 %	13.8 %	14.9 %				

\* Quarterly amounts may not add to the year-to-date amounts, as each period is computed on a discrete basis.

(a) Refer to the Specified Items schedule for further details. Effective tax rate on the Specified Items represents the difference between the GAAP and Non-GAAP effective tax rate.

(b) Includes Celgene results from operations from November 20, 2019 through December 31, 2019.

(c) Includes Celgene results of operations for the entire period.

**BRISTOL-MYERS SQUIBB COMPANY**  
**SELECTED BALANCE SHEET INFORMATION**  
(Unaudited, dollars in millions)

	March 31, 2019	June 30, 2019	September 30, 2019	December 31, 2019 <sup>(a)</sup>	March 31, 2020	June 30, 2020	September 30, 2020	December 31, 2020
Cash and cash equivalents	\$ 7,335	\$ 28,404	\$ 30,489	\$ 12,346	\$ 15,817	\$ 19,934		
Marketable debt securities - current	1,429	953	2,053	3,047	2,505	1,724		
Marketable debt securities - non-current	1,233	994	925	767	651	523		
<b>Cash, cash equivalents and marketable debt securities</b>	<b>9,997</b>	<b>30,351</b>	<b>33,467</b>	<b>16,160</b>	<b>18,973</b>	<b>22,181</b>		
Short-term debt obligations	(381)	(545)	(569)	(3,346)	(3,862)	(4,819)		
Long-term debt	(5,635)	(24,433)	(24,390)	(43,387)	(42,844)	(41,853)		
<b>Net (debt)/cash position</b>	<b>\$ 3,981</b>	<b>\$ 5,373</b>	<b>\$ 8,508</b>	<b>\$ (30,573)</b>	<b>\$ (27,733)</b>	<b>\$ (24,491)</b>		

(a) Includes Celgene balances as of December 31, 2019.

**BRISTOL-MYERS SQUIBB COMPANY**  
**2020 FULL YEAR PROJECTED DILUTED EPS FROM OPERATIONS**  
**EXCLUDING PROJECTED SPECIFIED ITEMS**

	Full Year 2020		
	Pre-tax	Tax	After-tax
Projected Diluted (Loss)/Earnings Attributable to Shareholders per Common Share - GAAP			(\$0.06) to \$0.09
<b>Projected Specified Items:</b>			
Purchase price accounting adjustments <sup>(a)</sup>	5.31	0.29	5.02
Acquisition, restructuring and integration expenses <sup>(b)</sup>	0.61	0.13	0.48
Equity investment losses and contingent consideration	(0.04)	(0.06)	0.02
Research and development license and asset acquisition charges	0.15	—	0.15
Employee compensation charges <sup>(c)</sup>	0.04	0.01	0.03
Divestiture gains and licensing income	(0.03)	—	(0.03)
Other	0.01	—	0.01
Income taxes attributed to Otezla® divestiture and internal transfer of intangible assets	—	(0.48)	0.48
<b>Total</b>	<b>6.05</b>	<b>(0.11)</b>	<b>6.16</b>
Projected Diluted Earnings Attributable to Shareholders per Common Share - Non-GAAP			<u>\$6.10 to \$6.25</u>

(a) Includes amortization of acquired intangible assets, unwind of inventory fair value adjustments and amortization of debt assumed from Celgene.

(b) Includes acquisition-related financing, transaction, restructuring and integration expenses recognized in Cost of products sold, Research and development and Other (income)/expense, net.

(c) Includes items recognized in Cost of products sold, Marketing, selling and administrative and Research and development.

The following table summarizes the company's 2020 financial guidance:

Line item	GAAP	Non-GAAP
Revenues	\$40.5 billion - \$42.0 billion	\$40.5 billion - \$42.0 billion
Gross margin as a percent of revenue	Approximately 74%	Approximately 80%
Marketing, selling and administrative expense	\$6.5 billion - \$6.7 billion	\$6.5 billion - \$6.7 billion
Research and development expense	\$9.7 billion - \$9.9 billion	\$9.2 billion - \$9.4 billion
Other (income)/expense, net	\$0.9 billion - \$1.1 billion	(\$0.1 billion) - \$0.1 billion
Effective tax rate	Approximately 100%	16% - 17%
Weighted average diluted shares	Approximately 2.3 billion	Approximately 2.3 billion

The GAAP financial results for the full year of 2020 will include specified items, including purchase price accounting adjustments, acquisition and integration expenses, charges associated with restructuring, downsizing and streamlining worldwide operations, research and development license and asset acquisition charges, divestiture gains or losses, stock compensation resulting from accelerated vesting of Celgene awards, certain retention-related employee compensation charges related to the Celgene transaction, equity investment and contingent value rights fair value adjustments and tax items resulting from internal transfer of intangible assets and the Otezla® divestiture, among other items. The financial guidance for 2020 excludes the impact of any potential future strategic acquisitions and divestitures and other specified items that have not yet been identified and quantified. For a fuller discussion of items that could impact full year GAAP results, as well as the use of non-GAAP financial information, see Bristol Myers Squibb Reports Second Quarter 2020 Financial Results on August 6, 2020, including "2020 Financial Guidance" and "Use of non-GAAP Financial Information" therein.