

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): **October 22, 2014**

Biogen Idec Inc.

(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction of incorporation)

0-19311
(Commission File Number)

33-0112644
(IRS Employer Identification No.)

225 Binney Street, Cambridge, Massachusetts 02142
(Address of principal executive offices; Zip Code)

Registrant's telephone number, including area code: **(617) 679-2000**

(Former name or former address, if changed since last report.)

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:

- 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))

Item 2.02 Results of Operations and Financial Condition.

On October 22, 2014, Biogen Idec Inc. issued a press release announcing its results of operations and financial condition for the third quarter ended September 30, 2014. A copy of the press release is furnished as Exhibit 99 and is incorporated herein by reference.

The press release is being furnished pursuant to Item 2.02 of this Current Report on Form 8-K and shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934 (the “Exchange Act”) or otherwise subject to the liabilities of that Section, nor shall such document be deemed incorporated by reference in any filing under the Securities Act of 1933 or the Exchange Act.

Item 9.01 Financial Statements and Exhibits.

The exhibit listed on the Exhibit Index immediately preceding such exhibit is furnished as part of this Current Report on Form 8-K.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

BIOGEN IDEC INC.

By: /s/ Robert A. Licht
Robert A. Licht
Senior Vice President

Date: October 22, 2014

EXHIBIT INDEX

Exhibit Number

Description

99

Biogen Idec's press release dated October 22, 2014.

Biogen Idec Media Contact: **Biogen Idec Investor Contacts:**

Jason Glashow Claudine Prowse, Ph.D.
Senior Director, Public Affairs Vice President, Investor Relations
Biogen Idec Biogen Idec
Tel: (781) 464-3260 Tel: (781) 464-2442

Biogen Idec

Ben Strain
Associate Director, Investor Relations

Tel: (781) 464-2442

**BIOGEN IDEC THIRD QUARTER 2014 REVENUES
INCREASE 37% TO \$2.5 BILLION**

PLEGRIDY™ Approved in the US and EU for Multiple Sclerosis

ELOCTATE™ Launched in the US for Hemophilia A

Rollout of TECFIDERA® Continues across the Globe

Cambridge, Mass., October 22, 2014 -- Biogen Idec Inc. (NASDAQ: BIIB) today reported third quarter 2014 results, including revenue of \$2.5 billion, a 37% increase compared to the third quarter of 2013. Third quarter 2014 non-GAAP diluted earnings per share (EPS) were \$3.80, an increase of 61% over the third quarter of 2013. Non-GAAP net income attributable to Biogen Idec for the third quarter was \$900 million, an increase of 60% over the third quarter of 2013.

On a reported basis, GAAP diluted EPS for the third quarter of 2014 were \$3.62, an increase of 77% over the third quarter of 2013. GAAP net income attributable to Biogen Idec for the third quarter of 2014 was \$857 million, an increase of 76% versus the same period in the prior year. A reconciliation of GAAP to Non-GAAP quarterly financial results and 2014 full year guidance can be found in Table 3 at the end of this release.

“The third quarter was a period of significant achievement as we continued to make progress against our corporate objectives,” said Chief Executive Officer George A. Scangos, Ph.D. “We introduced innovative therapies for MS and hemophilia and began to launch them in markets worldwide. We also launched TECFIDERA in several European countries, furthering its position as a leading oral MS therapy.

“As we continued to extend our commercial business we remained focused on building our future, bringing to our team leading talent in technology, ALS research, neurology and gene therapy. I am also very proud of being recognized for our focus on corporate citizenship as again we were named the leading biotechnology company on the Dow Jones Sustainability Index,” Dr. Scangos added.

Third Quarter 2014 Performance Highlights

- TECFIDERA revenues were \$787 million, consisting of \$638 million in U.S. sales and \$149 million in sales outside the U.S.
- Interferon revenues, including AVONEX[®] and PLEGRIDY, were \$745 million, consisting of \$482 million in U.S. sales and \$263 million in sales outside the U.S.
- TYSABRI[®] revenues were \$501 million, consisting of \$275 million in U.S. sales and \$226 million in sales outside the U.S.
- Net revenues relating to RITUXAN[®] and GAZYVA[®] from our unconsolidated joint business arrangement were \$291 million.
- ALPROLIX[®] revenues were \$25 million, and ELOCTATE[™] revenues were \$22 million.

Other Third Quarter 2014 Financial Results

- Revenues for FAMPYRA[®] and FUMADERM[™] were \$37 million.
- Royalty revenues were \$67 million.
- Corporate partner revenues were \$36 million.
- As of September 30, 2014, Biogen Idec had cash, cash equivalents and marketable securities totaling approximately \$3.2 billion.
- During the quarter, the Internal Revenue Service issued final regulations related to the Branded Pharmaceutical Drug (BPD) Fee, which had the effect of changing the recognition of the fee for accounting purposes, from the period in which the fee was paid, to the period when the sale occurs. Our products that are subject to the BPD fee include PLEGRIDY, TECFIDERA, TYSABRI and RITUXAN. As a result of these final regulations, we recognized an incremental BPD fee which reduced our share of revenues from unconsolidated joint business by \$21.0 million and increased selling, general and administrative expense by \$18.5 million for the periods 2013 through the end of this quarter.

2014 Financial Guidance

Biogen Idec updated its full year 2014 financial guidance.

This updated guidance consists of the following components:

- Revenue growth is expected to be approximately 38% to 41% compared to 2013, unchanged from prior guidance.
- R&D expense is expected to be approximately 20% to 21% of total revenue.
 - For the balance of the year, full year guidance for R&D expense includes approximately \$50 million intended for new early and mid-stage business development opportunities. This amount is reduced from prior guidance due to revised business development expectations through the end of the year.
- SG&A expense is expected to be approximately 22% to 23% of total revenue, unchanged from prior guidance.
- GAAP diluted EPS is expected to be between \$12.00 and \$12.10.

- Non-GAAP diluted EPS is expected to be between \$13.45 and \$13.55, an increase over prior guidance primarily due to updated business development expectations.

Biogen Idec may incur charges, realize gains or experience other events in 2014 that could cause actual results to vary from this guidance.

Multiple Sclerosis (MS) Events

- In July 2014, Biogen Idec received marketing authorization from the European Commission for PLEGRIDY, a pegylated interferon administered subcutaneously once every two weeks for adults with relapsing-remitting multiple sclerosis.
- In August 2014, the U.S. Food and Drug Administration approved PLEGRIDY as a new treatment for people with relapsing forms of multiple sclerosis.
- In September 2014, Biogen Idec presented extensive new data from its multiple sclerosis portfolio at the sixth Triennial Joint Meeting of the Americas Committee for Treatment and Research in Multiple Sclerosis and the European Committee for Research and Treatment in Multiple Sclerosis in Boston.
- TECFIDERA was recently launched in Denmark, the Netherlands, Switzerland, and Hungary.

Hemophilia Events

- In July 2014, Biogen Idec launched ELOCTATE in the U.S. for the treatment of adults and children with hemophilia A.
- In October 2014, Biogen Idec submitted a Marketing Authorisation Application (MAA) for ELOCTA to the European Medicines Agency (EMA). ELOCTA is the approved trade name in Europe for ELOCTATE. The MAA submission is subject to validation by the EMA.

Other Events

- During the third quarter of 2014, Biogen Idec announced the hiring of several key leaders including:
 - Adriana Karaboutis, Executive Vice President, Technology and Business Solutions, to lead Biogen Idec's information technology (IT) operations and advance the Company's use of technology and data to enhance overall engagement with patients and healthcare providers.
 - Donald R. Johns, M.D., Vice President, to lead Biogen Idec's amyotrophic lateral sclerosis (ALS) Innovation Hub (ALS iHub), a dedicated unit focused on accelerating the discovery and development of novel therapies for ALS by integrating research with clinical development.
 - Olivier Danos, Ph.D., Senior Vice President, Gene Therapy, to lead the Company's gene therapy research group, a team dedicated to identifying and developing new technologies for gene transfer and genome engineering.
- In September 2014, Biogen Idec announced that it was named the biotechnology industry leader on the Dow Jones Sustainability World Index. The Company was also named to the Dow Jones

Sustainability Index North America for the fifth consecutive year, one of only two biotech companies included.

Conference Call and Webcast

The Company's earnings conference call for the third quarter will be broadcast via the internet at 8:30 a.m. EDT on October 22, 2014, and will be accessible through the Investors section of Biogen Idec's homepage, www.biogenidec.com. Supplemental information in the form of a slide presentation will also be accessible at the same location on the internet at the time of the conference call and will be subsequently available on the website for at least one month.

About Biogen Idec

Through cutting-edge science and medicine, Biogen Idec discovers, develops and delivers to patients worldwide innovative therapies for the treatment of neurodegenerative diseases, hematologic conditions and autoimmune disorders. Founded in 1978, Biogen Idec is the world's oldest independent biotechnology company and patients worldwide benefit from its leading multiple sclerosis and innovative hemophilia therapies. For product labeling, press releases and additional information about the Company, please visit www.biogenidec.com.

Safe Harbor

This press release contains forward-looking statements, including statements about our business strategy and 2014 financial guidance. These forward-looking statements may be accompanied by such words as "anticipate," "believe," "could," "estimate," "expect," "forecast," "intend," "may," "plan," "potential," "project," "target," "will" and other words and terms of similar meaning. You should not place undue reliance on these statements.

These statements involve risks and uncertainties that could cause actual results to differ materially from those reflected in such statements, including: our dependence on sales from our principal products; uncertainty of success in execution of our commercialization of new products; failure to protect and enforce our data, intellectual property and other proprietary rights and the diminution of our ability to derive anticipated benefits from our products; difficulties in obtaining or changes in the availability of reimbursement for our products; uncertainty of success in developing other product candidates, including our ability to obtain product approvals in a timely manner or at all for new or current products; the occurrence of adverse safety events with our products; failure to compete effectively due to significant product competition in the markets for our products; dependence on collaborators and other third parties for the development and commercialization of products; problems with our manufacturing processes; failure to manage our growth and execute our growth initiatives; failure to comply with legal and regulatory requirements; the risks of doing business internationally; charges and other costs relating to our properties; risks and uncertainties relating to the timing, outcome and impact of legal, administrative and other proceedings and disputes; fluctuations in our effective tax rate; our ability to attract and retain qualified personnel; uncertainty and potential liabilities relating to product liability and intellectual property claims; the market, interest and credit risks associated with our portfolio of marketable securities; environmental risks; and the other risks and uncertainties that are described in the Risk Factors section of our most recent annual or quarterly report and in other reports we have filed with the SEC.

These statements are based on our current beliefs and expectations and speak only as of the date of this press release. We do not undertake any obligation to publicly update any forward-looking statements.

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TABLE 1
Biogen Idec Inc. and Subsidiaries
Condensed Consolidated Statements of Income
(unaudited, in thousands, except per share amounts)

| | For the Three Months Ended September 30, | | For the Nine Months Ended September 30, | |
|--|---|-------------------|--|---------------------|
| | 2014 | 2013 | 2014 | 2013 |
| Revenues: | | | | |
| Product, net | \$ 2,117,366 | \$ 1,453,554 | \$ 5,916,423 | \$ 3,935,251 |
| Unconsolidated joint business | 290,678 | 303,210 | 890,859 | 856,601 |
| Royalty | 67,148 | 54,144 | 145,348 | 125,076 |
| Corporate partner | 36,254 | 16,872 | 110,019 | 49,421 |
| Total revenues | <u>2,511,446</u> | <u>1,827,780</u> | <u>7,062,649</u> | <u>4,966,349</u> |
| Cost and expenses: | | | | |
| Cost of sales, excluding amortization of acquired intangible assets | 302,639 | 234,696 | 873,771 | 599,173 |
| Research and development | 417,174 | 410,017 | 1,393,331 | 1,021,820 |
| Selling, general and administrative | 570,436 | 405,584 | 1,658,732 | 1,189,194 |
| Amortization of acquired intangible assets | 122,431 | 99,998 | 382,515 | 233,524 |
| Collaboration profit sharing | — | — | — | 85,357 |
| (Gain) loss on fair value remeasurement of contingent consideration | (49,433) | (97) | (46,213) | (2,983) |
| Total cost and expenses | <u>1,363,247</u> | <u>1,150,198</u> | <u>4,262,136</u> | <u>3,126,085</u> |
| Gain on sale of rights | 4,379 | 6,949 | 12,138 | 17,319 |
| Income from operations | <u>1,152,578</u> | <u>684,531</u> | <u>2,812,651</u> | <u>1,857,583</u> |
| Other income (expense), net | (16,290) | (4,640) | (17,030) | (29,525) |
| Income before income tax expense and equity in loss of investee, net of tax | <u>1,136,288</u> | <u>679,891</u> | <u>2,795,621</u> | <u>1,828,058</u> |
| Income tax expense | 274,774 | 186,105 | 721,709 | 410,753 |
| Equity in loss of investee, net of tax | 5,394 | 6,170 | 14,932 | 12,270 |
| Net income | <u>856,120</u> | <u>487,616</u> | <u>2,058,980</u> | <u>1,405,035</u> |
| Net income (loss) attributable to noncontrolling interests, net of tax | (738) | — | 7,660 | — |
| Net income attributable to Biogen Idec Inc. | <u>\$ 856,858</u> | <u>\$ 487,616</u> | <u>\$ 2,051,320</u> | <u>\$ 1,405,035</u> |
| Net income per share: | | | | |
| Basic earnings per share attributable to Biogen Idec Inc. | <u>\$ 3.63</u> | <u>\$ 2.06</u> | <u>\$ 8.67</u> | <u>\$ 5.93</u> |
| Diluted earnings per share attributable to Biogen Idec Inc. | <u>\$ 3.62</u> | <u>\$ 2.05</u> | <u>\$ 8.64</u> | <u>\$ 5.89</u> |
| Weighted-average shares used in calculating: | | | | |
| Basic earnings per share attributable to Biogen Idec Inc. | <u>236,217</u> | <u>237,070</u> | <u>236,641</u> | <u>237,131</u> |
| Diluted earnings per share attributable to Biogen Idec Inc. | <u>236,972</u> | <u>238,349</u> | <u>237,449</u> | <u>238,508</u> |

TABLE 2
Biogen Idec Inc. and Subsidiaries
Condensed Consolidated Balance Sheets
(unaudited, in thousands)

| | As of September 30, 2014 | As of December 31, 2013 |
|--|---|--|
| ASSETS | | |
| Cash, cash equivalents and marketable securities | \$ 1,861,346 | \$ 1,222,729 |
| Accounts receivable, net | 1,091,232 | 824,406 |
| Inventory | 753,063 | 659,003 |
| Other current assets | 666,470 | 478,796 |
| Total current assets | <u>4,372,111</u> | <u>3,184,934</u> |
| Marketable securities | 1,371,431 | 625,772 |
| Property, plant and equipment, net | 1,724,129 | 1,750,710 |
| Intangible assets, net | 4,129,754 | 4,474,653 |
| Goodwill | 1,541,204 | 1,232,916 |
| Investments and other assets | 565,513 | 594,350 |
| TOTAL ASSETS | <u><u>\$ 13,704,142</u></u> | <u><u>\$ 11,863,335</u></u> |
| LIABILITIES AND EQUITY | | |
| Current portion of notes payable | \$ 3,220 | \$ 3,494 |
| Other current liabilities | 1,937,427 | 1,754,785 |
| Notes payable | 583,977 | 592,433 |
| Long-term deferred tax liability | 90,357 | 232,554 |
| Other long-term liabilities | 658,276 | 659,231 |
| Equity | <u>10,430,885</u> | <u>8,620,838</u> |
| TOTAL LIABILITIES AND EQUITY | <u><u>\$ 13,704,142</u></u> | <u><u>\$ 11,863,335</u></u> |

TABLE 3
Biogen Idec Inc. and Subsidiaries
GAAP to Non-GAAP Reconciliation: Net Income and Net Income Per Share
(unaudited, in millions, except per share amounts)

| | For the Three Months Ended September 30, | | For the Nine Months Ended September 30, | |
|--|---|----------------|--|----------------|
| | 2014 | 2013 | 2014 | 2013 |
| EARNINGS PER SHARE | | | | |
| GAAP earnings per share - Diluted | \$ 3.62 | \$ 2.05 | \$ 8.64 | \$ 5.89 |
| Adjustments to net income attributable to Biogen Idec Inc. (as detailed below) | 0.18 | 0.31 | 1.11 | 0.73 |
| Non-GAAP earnings per share - Diluted | <u>\$ 3.80</u> | <u>\$ 2.35</u> | <u>\$ 9.75</u> | <u>\$ 6.62</u> |

An itemized reconciliation between net income attributable to Biogen Idec Inc. on a GAAP basis and on a non-GAAP basis is as follows:

| | | | | |
|---|-----------------|-----------------|-------------------|-------------------|
| GAAP net income attributable to Biogen Idec Inc. | \$ 856.9 | \$ 487.6 | \$ 2,051.3 | \$ 1,405.0 |
| Adjustments: | | | | |
| Amortization of acquired intangible assets | 118.7 | 97.1 | 371.5 | 225.2 |
| (Gain) loss on fair value remeasurement of contingent consideration | (49.4) | (0.1) | (46.2) | (3.0) |
| SG&A: Stock option expense | 1.4 | 1.3 | 5.4 | 4.2 |
| R&D: Stock option expense | 1.2 | 1.2 | 4.8 | 3.6 |
| Donation to Biogen Idec Foundation | — | — | 35.0 | — |
| Income tax effect related to reconciling items | (29.2) | (26.0) | (106.2) | (55.7) |
| Non-GAAP net income attributable to Biogen Idec Inc. | <u>\$ 899.6</u> | <u>\$ 561.1</u> | <u>\$ 2,315.6</u> | <u>\$ 1,579.3</u> |

2014 Full Year Guidance: GAAP to Non-GAAP Reconciliation

An itemized reconciliation between projected net income attributable to Biogen Idec Inc. and diluted earnings per share on a GAAP basis and on a non-GAAP basis is as follows:

| | \$ | Shares | Diluted EPS |
|---|--------------|--------|-------------|
| Projected GAAP net income attributable to Biogen Idec Inc. | 2,860 | 237 | \$ 12.05 |
| Adjustments: | | | |
| Amortization of acquired intangible assets | 470 | | |
| (Gain) loss on fair value remeasurement of contingent consideration | (44) | | |
| Stock option expense | 13 | | |
| Donation to Biogen Idec Foundation | 35 | | |
| Income tax effect related to reconciling items | (130) | | |
| Projected Non-GAAP net income attributable to Biogen Idec Inc. | <u>3,204</u> | 237 | \$ 13.50 |

Numbers may not foot due to rounding.

Use of Non-GAAP Financial Measures

We supplement our consolidated financial statements presented on a GAAP basis by providing additional measures which may be considered “non-GAAP” financial measures under applicable SEC rules. We believe that the disclosure of these non-GAAP financial measures provides additional insight into the ongoing economics of our business and reflects how we manage our business internally, set operational goals and forms the basis of our management incentive programs. These non-GAAP

financial measures are not in accordance with generally accepted accounting principles in the United States and should not be viewed in isolation or as a substitute for reported, or GAAP, net income attributable to Biogen Idec Inc. and diluted earnings per share.

Our “Non-GAAP net income attributable to Biogen Idec Inc.” and “Non-GAAP earnings per share - Diluted” financial measures exclude the following items from GAAP net income attributable to Biogen Idec Inc. and diluted earnings per share:

1. Purchase accounting and merger-related adjustments.

We exclude certain purchase accounting related items associated with the acquisition of businesses, assets and amounts in relation to the consolidation of variable interest entities for which we are the primary beneficiary. These adjustments include charges for in-process research and development, the amortization of certain acquired intangible assets and fair value remeasurements of our contingent consideration obligations. The exclusion of these charges provides management and investors with a supplemental measure of performance which the Company believes better reflects the underlying economics of the business.

2. Stock option expense recorded in accordance with the accounting standard for share-based payments.

We believe that excluding the impact of expensing stock options better reflects the recurring economic characteristics of our business.

3. Other items.

We evaluate other items on an individual basis, and consider both the quantitative and qualitative aspects of the item, including (i) its size and nature, (ii) whether or not it relates to our ongoing business operations, and (iii) whether or not we expect it to occur as part of our normal business on a regular basis. We also include an adjustment to reflect the related tax effect of all reconciling items within our reconciliation of our GAAP to Non-GAAP net income attributable to Biogen Idec Inc.

TABLE 4
Biogen Idec Inc. and Subsidiaries
Product Revenues
(unaudited, in thousands)

| | For the Three Months | | For the Nine Months | |
|-----------------------------|-----------------------------|---------------------|----------------------------|---------------------|
| | Ended September 30, | | Ended September 30, | |
| | 2014 | 2013 | 2014 | 2013 |
| PRODUCT REVENUES | | | | |
| Multiple Sclerosis (MS): | | | | |
| AVONEX | \$ 741,848 | \$ 733,449 | \$ 2,277,094 | \$ 2,253,963 |
| PLEGRIDY | 3,382 | — | 3,382 | — |
| TECFIDERA | 787,122 | 286,366 | 1,993,212 | 478,500 |
| TYSABRI | 501,237 | 400,995 | 1,475,722 | 1,099,906 |
| Hemophilia: | | | | |
| ALPROLIX | 25,305 | — | 35,702 | — |
| ELOCTATE | 21,560 | — | 21,560 | — |
| Other product revenues: | | | | |
| FAMPYRA | 20,384 | 16,691 | 61,660 | 56,705 |
| FUMADERM | 16,528 | 16,053 | 48,091 | 46,177 |
| Total product revenues, net | <u>\$ 2,117,366</u> | <u>\$ 1,453,554</u> | <u>\$ 5,916,423</u> | <u>\$ 3,935,251</u> |