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**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
WASHINGTON, D.C. 20549**

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**FORM 8-K**

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**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 7, 2019**

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**Horizon Therapeutics Public Limited Company**  
(Exact name of registrant as specified in its charter)

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**Ireland**  
(State or other jurisdiction  
of incorporation)

**001-35238**  
(Commission  
File No.)

**Not Applicable**  
(IRS Employer  
Identification No.)

**Connaught House, 1<sup>st</sup> Floor, 1 Burlington Road, Dublin 4, D04 C5Y6, Ireland**  
(Address of principal executive offices)

**Registrant's telephone number, including area code: 011-353-1-772-2100**

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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))

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

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
<b>Ordinary shares, nominal value \$0.0001 per share</b>	<b>HZNP</b>	<b>The Nasdaq Global Select Market</b>

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

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.

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**Item 2.02 Results of Operations and Financial Condition.**

On August 7, 2019, Horizon Therapeutics plc (“the Company”) issued a press release announcing its financial results for the second quarter ended June 30, 2019. A copy of this press release is attached hereto as Exhibit 99.1.

As previously disclosed in its interim financial statements on Form 10-Q for the three months ended March 31, 2019, the Company changed its accounting for business combinations in respect of intangible assets acquired and their related third-party contingent royalties. When accounting for business combinations under ASC Topic 805, Business Combinations, the Company previously separately identified and recorded at fair value intangible assets acquired and their related third-party contingent royalties at the date of acquisition. Third-party contingent royalties are royalties payable to parties other than sellers of the businesses. Effective January 1, 2019, the Company retrospectively changed its accounting for business combinations and is now recording acquired intangible assets and their related third-party contingent royalties on a net basis (“New Method”). The Company changed its accounting principle on the basis that the use of the New Method is preferable, primarily due to improved comparability with the Company’s peers. The “Adjusted Historical Financial Information” that is attached hereto as Exhibit 99.2 presents selected line items from the Company’s annual and quarterly consolidated financial statements for certain historical periods illustrating the effect of the change in accounting principle.

The information in this Item 2.02 and Exhibits 99.1 and 99.2 hereto are being furnished and shall not be deemed to be “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liability of that section, nor shall they be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits.

<u>Exhibit No.</u>	<u>Description</u>
99.1	<a href="#"><u>Press Release of Horizon Therapeutics plc, dated August 7, 2019.</u></a>
99.2	<a href="#"><u>Horizon Therapeutics plc Adjusted Historical Financial Information (Unaudited), dated August 7, 2019.</u></a>

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**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.

Date: August 7, 2019

**HORIZON THERAPEUTICS PUBLIC LIMITED COMPANY**

By: /s/ Paul W. Hoelscher  
Paul W. Hoelscher  
Executive Vice President and Chief Financial Officer



**Horizon Therapeutics plc Reports Strong Second-Quarter 2019 Results;  
Increases Full-Year 2019 Net Sales and Adjusted EBITDA Guidance**

— *Second-Quarter 2019 Net Sales of \$320.6 Million Increased 6 Percent;  
Second-Quarter 2019 GAAP Net Loss of \$5.1 Million; Adjusted EBITDA of \$124.1 Million* —

— *Quarterly Orphan and Rheumatology Segment Net Sales Increased 11 Percent to \$223.5 Million;  
KRYSTEXXA® Second-Quarter 2019 Net Sales Growth of 36 Percent* —

— *Increasing Full-Year 2019 Net Sales Guidance Range to \$1.28 Billion to \$1.30 Billion and  
Adjusted EBITDA Guidance Range to \$460 Million to \$475 Million;  
KRYSTEXXA Full-Year 2019 Net Sales Growth Expected to Be Greater Than 20 Percent* —

— *Submitted Teprotumumab U.S. Biologics License Application (BLA) for the  
Treatment of Active Thyroid Eye Disease (TED)* —

— *Initiated Registrational Clinical Trial MIRROR, Evaluating KRYSTEXXA in Combination with  
Methotrexate to Potentially Improve Patient Response* —

— *Cash Position of \$866 Million; Net Leverage of 1.1 Times as of June 30, 2019* —

**DUBLIN** – Aug. 7, 2019 – Horizon Therapeutics plc (Nasdaq: HZNP) today announced its second-quarter 2019 financial results and increased its full-year 2019 net sales and adjusted EBITDA guidance.

“The second quarter was another quarter of outstanding execution and strategic progress,” said Timothy Walbert, chairman, president and chief executive officer, Horizon. “We generated double-digit net sales growth in our orphan and rheumatology segment, driven by continued momentum from KRYSTEXXA, our medicine for uncontrolled gout and our main growth driver. In addition, we recently submitted teprotumumab for U.S. FDA approval, another milestone toward delivering the first FDA-approved treatment to people living with active thyroid eye disease.”

**Financial Highlights**

(in millions except for per share amounts and percentages)	Q2 19	Q2 18	% Change	YTD 19	YTD 18	% Change
Net sales	\$ 320.6	\$ 302.8	6	\$ 601.0	\$ 526.7	14
Net loss	(5.1)	(24.8)	79	(38.0)	(173.4)	78
Non-GAAP net income	95.6	80.5	19	149.6	85.3	75
Adjusted EBITDA	124.1	116.8	6	212.5	150.4	41
Loss per share - diluted	(0.03)	(0.15)	80	(0.21)	(1.05)	80
Non-GAAP earnings per share - diluted	0.49	0.48	2	0.80	0.51	57

**Horizon Therapeutics plc**

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## Second-Quarter and Recent Company Highlights

- **Submitted BLA for Teprotumumab for Active TED:** In early July, the Company submitted a BLA for its investigational medicine teprotumumab for the treatment of active TED to the U.S. Food and Drug Administration (FDA). The submission included results from the Phase 3 clinical trial, OPTIC (Treatment of Graves' Orbitopathy (Thyroid Eye Disease) to Reduce Proptosis with Teprotumumab Infusions in a Randomized, Placebo-Controlled, Clinical Study), as well as the positive Phase 2 results.  
  
Teprotumumab has Breakthrough Therapy, Orphan Drug and Fast Track designations from the FDA. Horizon has requested Priority Review for the application, which, if granted, could result in a six-month review process. The FDA has a 60-day filing review period to determine whether the BLA is complete and acceptable for filing. If approved, teprotumumab would be the first and only approved treatment for active TED.  
  
In April, additional results from OPTIC were presented at the American Association of Clinical Endocrinologists (AACE) Scientific and Clinical Congress, which included measurements of improvement in proptosis, the major driver of morbidity in TED. These data showed that after the full course of treatment for 24 weeks, patients treated with teprotumumab demonstrated a mean proptosis reduction of 3.32 mm compared with 0.53 mm for patients on placebo ( $p < 0.001$ ).
- **Announced Teprotumumab Expanded Access Program (EAP):** The Company recently announced the availability of an expanded access program for teprotumumab. The expanded access program will be available for people living with active TED while the FDA reviews the teprotumumab BLA.
- **Initiated KRYSTEXXA Immunomodulation Trial:** In June, the Company initiated its registrational clinical trial MIRROR (Methotrexate to Increase Response Rates in Patients with Uncontrolled GOut Receiving KRYSTEXXA). The trial is evaluating administration of KRYSTEXXA in combination with methotrexate to determine the potential for dampening anti-drug antibody formation and increasing response rates with KRYSTEXXA, allowing more patients living with uncontrolled gout to fully benefit from treatment. The randomized placebo-controlled study is expected to enroll approximately 135 patients to receive either KRYSTEXXA and methotrexate or KRYSTEXXA and placebo. The primary endpoint will assess the proportion of serum uric acid (sUA) responders (sUA  $< 6$  mg/dL) at Month 6.
- **FDA Accepted New Drug Application (NDA) for PROCYSBI® Oral Granules:** In July, the FDA accepted the NDA for PROCYSBI Delayed-Release Oral Granules in Packets. If approved, this new dosage form would provide another administration option for patients, in addition to the currently available PROCYSBI delayed-release capsules, which are FDA-approved for children one year of age and older and adults living with nephropathic cystinosis. The submission is part of the Company's ongoing investment in the cystinosis community.



- **Appointed Sue Mahony to the Board of Directors:** The Company recently appointed Sue Mahony, Ph.D., MBA, to its board of directors. Dr. Mahony brings more than 30 years of diverse industry experience to the Board, including an 18-year tenure at Eli Lilly and Company, where she served in a variety of global and domestic leadership roles of increasing responsibility, including helping oversee the development of an innovative pipeline. Before Lilly, Dr. Mahony spent five years at Bristol-Myers Squibb Company.
- **Changed Company Name to Horizon Therapeutics plc:** In May, shareholders approved the change of the Company's name to Horizon Therapeutics Public Limited Company at the Annual General Meeting. The new name captures the Company's long-term strategy to develop and commercialize innovative new medicines that address rare and rheumatic diseases with very few effective treatment options. The Company believes the new name also better reflects its work with patients, caregivers, physicians and communities that goes well beyond its medicines.
- **Improved the Company's Capital Structure:** In May, the Company repaid \$250 million of its outstanding debt, reducing it to \$1.443 billion as of June 30, 2019. In May, the Company also refinanced its senior secured term loans, lowering the interest rate by 25 basis points and extending the final maturity date to May 22, 2026. Additionally, in July, the Company issued \$600 million of 5.5 percent Senior Notes due 2027 and is using the proceeds along with cash on hand to repay \$625 million of its outstanding debt. These actions serve to reduce interest expense and extend the maturity of the debt, furthering the Company's strategy to improve its capital structure.

## Research and Development Programs

### *Orphan Disease Candidate and Program:*

- **Teprotumumab:** Teprotumumab is a fully human monoclonal antibody insulin-like growth factor-1 receptor (IGF-1R) inhibitor candidate for the treatment of active TED, a serious, progressive, vision-threatening autoimmune disease in which the muscles and fatty tissue behind the eye become inflamed and expand. This can lead to proptosis (eye bulging) and diplopia (double vision) and impact activities of daily living and quality of life. The development program for teprotumumab in TED includes positive Phase 2 results published in *The New England Journal of Medicine*, as well as positive results from the confirmatory Phase 3 OPTIC clinical trial, announced in February 2019. The OPTIC study met its primary endpoint of a  $\geq 2$  mm reduction in proptosis ( $p < 0.001$ ), the main cause of morbidity in TED, with 82.9 percent of patients treated with teprotumumab demonstrating a significant improvement in proptosis compared to 9.5 percent of placebo patients. In addition, all secondary endpoints were met ( $p \leq 0.001$ ), and the safety profile was consistent with the Phase 2 study.

### *Rheumatology Pipeline Candidates and Programs:*

- **KRYSTEXXA Immunomodulation Trial:** The Company is evaluating the use of methotrexate to increase the response rate with KRYSTEXXA through its MIRROR study. Methotrexate is the immunomodulator most used by rheumatologists, and has been shown to reduce anti-drug antibody formation to biologic therapies when combined with these therapies. The MIRROR trial is designed to support the potential for registration and commenced in June.



- **KRYSTEXXA Study in Kidney Transplant Patients with Uncontrolled Gout:** The Company plans to initiate a clinical trial in the second half of 2019 evaluating the effect of KRYSTEXXA on serum uric acid levels in kidney transplant patients with uncontrolled gout. Kidney transplant patients have more than a tenfold increase in the prevalence of gout when compared to the general population, and literature suggests that persistently high serum uric acid levels can be associated with organ rejection. Managing uncontrolled gout is one of the most common and significant unmet needs of kidney transplant patients.
- **Next-generation Biologic Programs for Uncontrolled Gout:** The Company is pursuing several development programs for next-generation biologics for uncontrolled gout to support and sustain the Company's market leadership in this area. These include HZN-003, HZN-007 and a discovery and development collaboration with HemoShear Therapeutics, LLC.

### Second-Quarter Financial Results

*Note: For additional detail and reconciliation of non-GAAP financial measures to the most directly comparable GAAP financial measures, please refer to the tables at the end of this release.*

- **Net Sales:** Second-quarter 2019 net sales were \$320.6 million, an increase of 6 percent.
- **Gross Profit:** Under U.S. GAAP, the second-quarter 2019 gross profit ratio was 72.2 percent compared to 69.8 percent in the second quarter of 2018. The non-GAAP gross profit ratio in the second quarter of 2019 was 90.9 percent compared to 90.2 percent in the second quarter of 2018.
- **Operating Expenses:** Research and development (R&D) expenses were 8.8 percent of net sales and selling, general and administrative (SG&A) expenses were 52.1 percent of net sales. Non-GAAP R&D expenses were 6.9 percent of net sales, and non-GAAP SG&A expenses were 45.4 percent of net sales.
- **Income Tax Rate:** In the second quarter of 2019, the income tax benefit rate on a GAAP basis was 48.8 percent and the income tax expense rate on a non-GAAP basis was 11.3 percent.
- **Net Income (Loss):** On a GAAP basis in the second quarter of 2019, net loss was \$5.1 million. Second-quarter 2019 non-GAAP net income was \$95.6 million.
- **Adjusted EBITDA:** Second-quarter 2019 adjusted EBITDA was \$124.1 million.
- **Earnings (Loss) per Share:** On a GAAP basis diluted loss per share in the second quarter of 2019 and 2018 was \$0.03 and \$0.15, respectively. Non-GAAP diluted earnings per share in the second quarter of 2019 and 2018 was \$0.49 and \$0.48, respectively. Weighted average shares outstanding used for calculating GAAP and non-GAAP diluted earnings per share in the second quarter of 2019 were 185.3 million and 193.2 million, respectively.



## Second-Quarter Segment Results

Management uses net sales and segment operating income to evaluate the performance of the Company's two segments. While segment operating income contains certain adjustments to the directly comparable GAAP figures in the Company's consolidated financial results, it is considered to be prepared in accordance with GAAP for purposes of presenting the Company's segment operating results.

### Orphan and Rheumatology Segment

(in millions except for percentages)	Q2 19	Q2 18	% Change	YTD 19	YTD 18	% Change
KRYSTEXXA	79.8	58.6	36	132.1	105.3	25
RAVICTI®(1)	50.4	57.0	(11)	100.3	106.1	(5)
PROCYSBI	41.2	38.4	7	80.7	73.4	10
ACTIMMUNE®	29.3	27.4	7	51.0	52.2	(2)
RAYOS®	20.3	13.5	51	39.7	24.1	64
BUPHENYL®(1)	2.3	5.2	(55)	5.2	11.0	(53)
QUINSAIR™	0.2	0.1	75	0.4	0.2	55
LODOTRA®(1)	—	1.5	NM	—	1.7	NM
<b>Orphan and Rheumatology Net Sales</b>	<b>\$ 223.5</b>	<b>\$ 201.7</b>	<b>11</b>	<b>\$ 409.4</b>	<b>\$ 374.0</b>	<b>9</b>

<b>Orphan and Rheumatology Segment Operating Income</b>	<b>\$ 74.5</b>	<b>\$ 70.6</b>	<b>6</b>	<b>\$ 121.2</b>	<b>\$ 113.7</b>	<b>7</b>
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(1) Beginning in 2019, the Company no longer recognizes revenue from RAVICTI and AMMONAPS sales outside of North America and Japan, nor from sales of LODOTRA. On Dec. 28, 2018, the Company divested the rights to RAVICTI and AMMONAPS outside of North America and Japan. AMMONAPS is known as BUPHENYL in the United States. In addition, effective Jan. 1, 2019, the RAYOS and LODOTRA license and supply agreements were amended, including the transfer of LODOTRA to Vectura Group plc. LODOTRA is known as RAYOS in the United States.

- Second-quarter 2019 net sales of the orphan and rheumatology segment, the Company's strategic growth segment, were \$223.5 million, an increase of 11 percent over the prior year's quarter, driven by growth of KRYSTEXXA, RAYOS, PROCYSBI and ACTIMMUNE.
- Second-quarter 2019 orphan and rheumatology segment operating income was \$74.5 million, which includes the impact of investment in teprotumumab pre-launch activities.

### Inflammation Segment<sup>(1)</sup>

(in millions except for percentages)	Q2 19	Q2 18	% Change	YTD 19	YTD 18	% Change
PENNSAID® 2%	51.5	47.6	8	101.7	74.4	37
DUEXIS®	30.0	30.7	(2)	59.5	46.4	28
VIMOVO®	14.6	21.9	(33)	28.6	30.2	(5)
MIGERGOT®(2)	1.0	0.9	5	1.8	1.7	8
<b>Inflammation Net Sales</b>	<b>\$ 97.1</b>	<b>\$ 101.1</b>	<b>(4)</b>	<b>\$ 191.6</b>	<b>\$ 152.7</b>	<b>25</b>

<b>Inflammation Segment Operating Income</b>	<b>\$ 49.7</b>	<b>\$ 45.9</b>	<b>8</b>	<b>\$ 91.1</b>	<b>\$ 36.3</b>	<b>151</b>
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(1) Previously known as the primary care segment.

(2) In June 2019, the Company divested the rights to MIGERGOT.





- Second-quarter 2019 net sales of the inflammation segment were \$97.1 million and segment operating income was \$49.7 million.

#### **Cash Flow Statement and Balance Sheet Highlights**

- On a GAAP basis in the second quarter of 2019, operating cash flow was \$91.3 million. Non-GAAP operating cash flow was \$95.7 million.
- The Company had cash and cash equivalents of \$866.0 million as of June 30, 2019.
- As of June 30, 2019, the total principal amount of debt outstanding was \$1.443 billion. As of June 30, 2019, net debt was \$577 million and net-debt-to-last-12-months adjusted EBITDA leverage ratio was 1.1 times, compared to 3.6 times at June 30, 2018.

In May, the Company repaid \$250 million of its outstanding debt, reducing it to \$1.443 billion as of June 30, 2019. In May, the Company also refinanced its senior secured term loans, lowering the interest rate by 25 basis points and extending the final maturity date to May 22, 2026. In July, the Company issued \$600 million of 5.5 percent Senior Notes due 2027 and is using the proceeds along with cash on hand to repay \$625 million of its outstanding debt. Following the refinancing transactions, the Company expects the total principal amount of debt outstanding to be \$1.418 billion, consisting of \$418 million in senior secured term loans due 2026, \$600 million of Senior Notes due 2027 and \$400 million of Exchangeable Senior Notes due 2022.

#### **New 2019 Guidance**

The Company now expects full-year 2019 net sales to range between \$1.28 billion to \$1.30 billion, an increase from the previous guidance range of \$1.26 billion to \$1.28 billion. Full-year 2019 adjusted EBITDA is now expected to range between \$460 million to \$475 million, an increase from the previous guidance range of \$450 million to \$465 million.

#### **Webcast**

At 8 a.m. EDT / 1 p.m. IST today, the Company will host a live webcast to review its financial and operating results and provide a general business update. The live webcast and a replay may be accessed at <http://ir.horizontherapeutics.com>. Please connect to the Company's website at least 15 minutes prior to the live webcast to ensure adequate time for any software download that may be needed to access the webcast. A replay of the webcast will be available approximately two hours after the live webcast.

#### **About Horizon**

Horizon is focused on researching, developing and commercializing medicines that address critical needs for people impacted by rare and rheumatic diseases. Our pipeline is purposeful: we apply scientific expertise and courage to bring clinically meaningful therapies to patients. We believe science and compassion must work together to transform lives. For more information on how we go to incredible lengths to impact lives, please visit <https://www.horizontherapeutics.com/>, follow us [@HorizonNews](#) on Twitter, like us on [Facebook](#) or explore career opportunities on [LinkedIn](#).

#### **Note Regarding Use of Non-GAAP Financial Measures**

*EBITDA, or earnings before interest, taxes, depreciation and amortization, and adjusted EBITDA are used and provided by Horizon as non-GAAP financial measures. Horizon provides certain other financial measures such as non-GAAP net income, non-GAAP diluted earnings per share, non-GAAP gross profit and gross profit ratio, non-GAAP operating expenses, non-GAAP operating income, non-GAAP tax rate, non-GAAP operating cash flow, net leverage ratio and net debt, each of which include adjustments to GAAP figures. These non-GAAP measures are intended to provide additional information on Horizon's performance, operations, expenses, profitability and cash flows. Adjustments to Horizon's GAAP figures as well as EBITDA exclude acquisition and/or divestiture-related expenses, charges related to the discontinuation of ACTIMMUNE development for Friedreich's ataxia, gain or loss from divestiture, gain or loss from sale of assets, upfront, progress and milestone payments related to license and collaboration agreements, litigation settlements, loss on debt extinguishment, costs of debt refinancing, drug manufacturing harmonization costs, restructuring and realignment costs, as well as non-cash items such as share-based compensation, depreciation and amortization, non-cash interest expense, long-lived asset impairment charges and other non-cash adjustments. Certain other special items or substantive events may also be included in the non-GAAP adjustments periodically when their magnitude is significant within the periods incurred. Horizon maintains an established non-GAAP cost policy that guides the determination of what costs will be excluded in non-GAAP measures. Horizon believes that these non-GAAP financial measures, when considered together with the GAAP figures, can enhance an overall understanding of Horizon's financial and operating performance. The non-GAAP financial measures are included with the intent of providing investors with a more complete understanding of the Company's historical and expected 2019 financial results and trends and to facilitate comparisons between periods and with respect to projected information. In addition, these non-GAAP financial measures are among the indicators Horizon's management uses for planning and forecasting purposes and measuring the Company's performance. For example, adjusted EBITDA is used by Horizon as one measure of management performance under certain incentive compensation arrangements. These non-GAAP financial measures should be considered in addition to, and not as a substitute for, or superior to, financial measures calculated in accordance with GAAP. The non-GAAP financial measures used by the Company may be calculated differently from, and therefore may not be comparable to, non-GAAP financial measures used by other companies. Horizon has not provided a reconciliation of its full-year 2019 adjusted EBITDA outlook to an expected net income (loss) outlook because certain items such as acquisition/divestiture-related expenses and share-based compensation that are a component of net income (loss) cannot be reasonably projected due to the significant impact of changes in Horizon's stock price, the variability associated with the size or timing of acquisitions/divestitures and other factors. These components of net income (loss) could significantly impact Horizon's actual net income (loss).*

#### **Forward-Looking Statements**

*This press release contains forward-looking statements, including, but not limited to, statements related to Horizon's full-year 2019 net sales and adjusted EBITDA guidance; expected financial performance and operating results in future periods, including potential growth in net sales of certain of Horizon's medicines; expected impact of refinancing transactions; expected timing of clinical trials and regulatory submissions and decisions, including related to the BLA submission for teprotumumab and the NDA for PROCYSBI Delayed-Release Oral Granules in Packets; expected expansion of Horizon's rare disease medicine pipeline and the impact thereof; potential market opportunity for Horizon's medicines and medicine candidates; and business and other statements that are not historical facts. These forward-looking statements are based on Horizon's current expectations and inherently involve significant risks and uncertainties. Actual results and the timing of events could differ materially from those anticipated in such forward-looking statements as a result of these risks and uncertainties, which include, without limitation, risks that Horizon's*



*actual future financial and operating results may differ from its expectations or goals; Horizon's ability to grow net sales from existing medicines; the availability of coverage and adequate reimbursement and pricing from government and third-party payers; risks relating to Horizon's ability to successfully implement its business strategies; risks inherent in developing novel medicine candidates, such as teprotumumab, and existing medicines for new indications; risks associated with regulatory approvals; risks in the ability to recruit, train and retain qualified personnel; competition, including potential generic competition; the ability to protect intellectual property and defend patents; regulatory obligations and oversight, including any changes in the legal and regulatory environment in which Horizon operates and those risks detailed from time-to-time under the caption "Risk Factors" and elsewhere in Horizon's filings and reports with the SEC. Horizon undertakes no duty or obligation to update any forward-looking statements contained in this press release as a result of new information.*

**Contacts:**

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**Horizon Therapeutics plc**

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**Horizon Therapeutics plc**  
**Condensed Consolidated Statements of Operations (Unaudited)**  
(in thousands, except share and per share data)

	Three Months Ended June 30,		Six Months Ended June 30,	
	2019	2018	2019	2018
Net sales	\$ 320,647	\$ 302,835	\$ 601,018	\$ 526,716
Cost of goods sold	89,163	91,337	177,305	201,625
<b>Gross profit</b>	<b>231,484</b>	<b>211,498</b>	<b>423,713</b>	<b>325,091</b>
<b>OPERATING EXPENSES:</b>				
Research and development	28,314	24,265	50,039	41,910
Selling, general and administrative	167,095	176,674	339,394	356,273
Loss on sale of assets	10,963	—	10,963	—
Impairment of long-lived assets	—	—	—	33,647
<b>Total operating expenses</b>	<b>206,372</b>	<b>200,939</b>	<b>400,396</b>	<b>431,830</b>
<b>Operating income (loss)</b>	<b>25,112</b>	<b>10,559</b>	<b>23,317</b>	<b>(106,739)</b>
<b>OTHER EXPENSE, NET:</b>				
Interest expense, net	(22,033)	(31,030)	(49,563)	(61,484)
Loss on debt extinguishment	(11,878)	—	(17,464)	—
Foreign exchange gain (loss)	76	(5)	15	(115)
Other (expense) income, net	(1,272)	346	(1,083)	497
<b>Total other expense, net</b>	<b>(35,107)</b>	<b>(30,689)</b>	<b>(68,095)</b>	<b>(61,102)</b>
<b>Loss before (benefit) expense for income taxes</b>	<b>(9,995)</b>	<b>(20,130)</b>	<b>(44,778)</b>	<b>(167,841)</b>
(Benefit) expense for income taxes	(4,875)	4,621	(6,795)	5,566
<b>Net loss</b>	<b>\$ (5,120)</b>	<b>\$ (24,751)</b>	<b>\$ (37,983)</b>	<b>\$ (173,407)</b>
Loss per ordinary share - basic and diluted	\$ (0.03)	\$ (0.15)	\$ (0.21)	\$ (1.05)
Weighted average ordinary shares outstanding - basic and diluted	185,327,383	165,536,826	178,866,391	164,921,722



**Horizon Therapeutics plc**  
**Condensed Consolidated Balance Sheets (Unaudited)**  
(in thousands, except share data)

	As of	
	June 30, 2019	December 31, 2018
<b>ASSETS</b>		
<b>CURRENT ASSETS:</b>		
Cash and cash equivalents	\$ 865,997	\$ 958,712
Restricted cash	3,739	3,405
Accounts receivable, net	395,018	464,730
Inventories, net	51,019	50,751
Prepaid expenses and other current assets	85,728	68,218
<b>Total current assets</b>	<b><u>1,401,501</u></b>	<b><u>1,545,816</u></b>
Property and equipment, net	24,808	20,101
Developed technology, net	1,813,950	1,945,639
Other intangible assets, net	4,229	4,630
Goodwill	413,669	413,669
Deferred tax assets, net	6,080	3,148
Other assets	43,767	8,959
<b>Total assets</b>	<b><u>\$ 3,708,004</u></b>	<b><u>\$ 3,941,962</u></b>
<b>LIABILITIES AND SHAREHOLDERS' EQUITY</b>		
<b>CURRENT LIABILITIES:</b>		
Accounts payable	\$ 42,672	\$ 30,284
Accrued expenses	188,192	215,739
Accrued trade discounts and rebates	398,657	457,763
Deferred revenues, current portion	7,311	4,901
<b>Total current liabilities</b>	<b><u>636,832</u></b>	<b><u>708,687</u></b>
<b>LONG-TERM LIABILITIES:</b>		
Exchangeable notes, net	341,682	332,199
Long-term debt, net of current	1,025,096	1,564,485
Deferred tax liabilities, net	109,443	107,768
Other long-term liabilities	74,078	38,717
<b>Total long-term liabilities</b>	<b><u>1,550,299</u></b>	<b><u>2,043,169</u></b>
<b>COMMITMENTS AND CONTINGENCIES</b>		
<b>SHAREHOLDERS' EQUITY:</b>		
Ordinary shares, \$0.0001 nominal value; 600,000,000 and 300,000,000 shares authorized at June 30, 2019 and December 31, 2018, respectively; 186,470,230 and 169,244,520 shares issued at June 30, 2019 and December 31, 2018, respectively, and 186,085,864 and 168,860,154 shares outstanding at June 30, 2019 and December 31, 2018, respectively	19	17
Treasury stock, 384,366 ordinary shares at June 30, 2019 and December 31, 2018	(4,585)	(4,585)
Additional paid-in capital	2,743,793	2,374,966
Accumulated other comprehensive loss	(1,666)	(1,523)
Accumulated deficit	(1,216,688)	(1,178,769)
<b>Total shareholders' equity</b>	<b><u>1,520,873</u></b>	<b><u>1,190,106</u></b>
<b>Total liabilities and shareholders' equity</b>	<b><u>\$ 3,708,004</u></b>	<b><u>\$ 3,941,962</u></b>



**Horizon Therapeutics plc**  
**Condensed Consolidated Statements of Cash Flows (Unaudited)**  
(in thousands)

	<u>Three Months Ended June 30,</u>		<u>Six Months Ended June 30,</u>	
	<u>2019</u>	<u>2018</u>	<u>2019</u>	<u>2018</u>
<b>CASH FLOWS FROM OPERATING ACTIVITIES:</b>				
Net loss	\$ (5,120)	\$ (24,751)	\$ (37,983)	\$ (173,407)
<b>Adjustments to reconcile net loss to net cash provided by operating activities:</b>				
Depreciation and amortization expense	59,126	62,031	118,017	124,467
Equity-settled share-based compensation	21,367	30,721	48,915	58,554
Impairment of long-lived assets	—	—	—	33,647
Loss on debt extinguishment	11,878	—	17,464	—
Amortization of debt discount and deferred financing costs	5,771	5,690	11,622	11,185
Loss on sale of assets	10,963	—	10,963	—
Deferred income taxes	(2,759)	(3,433)	(1,257)	(1,753)
Foreign exchange and other adjustments	84	580	493	459
Changes in operating assets and liabilities:				
Accounts receivable	9,019	678	69,787	1,742
Inventories	343	(2,741)	(504)	11,549
Prepaid expenses and other current assets	(17,807)	(11,934)	(17,696)	(21,738)
Accounts payable	5,138	(10,120)	11,554	(3,592)
Accrued trade discounts and rebates	(8,247)	19,982	(59,151)	(52,138)
Accrued expenses	(6,736)	(5,371)	(28,071)	13,654
Deferred revenues	2,477	1,817	2,410	333
Other non-current assets and liabilities	5,770	(1,361)	873	(1,988)
<b>Net cash provided by operating activities</b>	<b>91,267</b>	<b>61,788</b>	<b>147,436</b>	<b>974</b>
<b>CASH FLOWS FROM INVESTING ACTIVITIES:</b>				
Payment related to license agreement	—	—	—	(12,000)
Proceeds from sale of assets	6,000	—	6,000	—
Purchases of property and equipment	(5,009)	(96)	(6,858)	(762)
<b>Net cash provided by (used in) investing activities</b>	<b>991</b>	<b>(96)</b>	<b>(858)</b>	<b>(12,762)</b>
<b>CASH FLOWS FROM FINANCING ACTIVITIES:</b>				
Net proceeds from the issuance of ordinary shares	(957)	—	326,793	—
Repayment of term loans	(518,026)	(25,598)	(818,026)	(27,722)
Repayment of senior notes	(258,282)	—	(258,282)	—
Net proceeds from the term loans	517,378	—	517,378	—
Proceeds from the issuance of ordinary shares in conjunction with ESPP program	5,465	4,720	5,465	4,734
Proceeds from the issuance of ordinary shares in connection with stock option exercises	1,987	2,727	12,029	3,672
Payment of employee withholding taxes relating to share-based awards	(7,203)	(5,668)	(24,374)	(9,185)
<b>Net cash used in financing activities</b>	<b>(259,638)</b>	<b>(23,819)</b>	<b>(239,017)</b>	<b>(28,501)</b>
<b>Effect of foreign exchange rate changes on cash, cash equivalents and restricted cash</b>	<b>576</b>	<b>(1,988)</b>	<b>58</b>	<b>(1,003)</b>
Net (decrease) increase in cash, cash equivalents and restricted cash	(166,804)	35,885	(92,381)	(41,292)
Cash, cash equivalents and restricted cash, beginning of the period <sup>(1)</sup>	1,036,540	680,720	962,117	757,897
<b>Cash, cash equivalents and restricted cash, end of the period<sup>(1)</sup></b>	<b>\$ 869,736</b>	<b>\$ 716,605</b>	<b>\$ 869,736</b>	<b>\$ 716,605</b>

(1) Amounts include restricted cash balance in accordance with ASU No. 2016-18. Cash and cash equivalents excluding restricted cash are shown on the balance sheet.



**Horizon Therapeutics plc**  
**GAAP to Non-GAAP Reconciliations**  
**Net Income and Earnings Per Share (Unaudited)**  
**(in thousands, except share and per share data)**

	Three Months Ended June 30,		Six Months Ended June 30,	
	2019	2018	2019	2018
<b>GAAP net loss</b>	<b>\$ (5,120)</b>	<b>\$ (24,751)</b>	<b>\$ (37,983)</b>	<b>\$ (173,407)</b>
Non-GAAP adjustments:				
Acquisition/divestiture-related costs	1,200	1,078	2,546	5,803
Restructuring and realignment costs	13	7,039	33	10,307
Amortization and step-up:				
Intangible amortization expense	57,683	60,480	115,100	121,364
Inventory step-up expense	(25)	53	90	17,129
Amortization of debt discount and deferred financing costs	5,710	5,691	11,622	11,187
Impairment of long-lived assets	—	—	—	33,647
Loss on sale of assets	10,963	—	10,963	—
Share-based compensation	21,367	30,721	48,915	58,554
Depreciation	1,443	1,551	2,916	3,104
Litigation settlements	1,000	4,250	1,000	4,250
Upfront, progress and milestone payments related to license and collaboration agreements	4,000	—	6,000	90
Fees related to refinancing activities	1,033	15	1,175	42
Loss on debt extinguishment	11,878	—	17,464	—
Drug substance harmonization costs	234	475	314	1,279
Charges relating to discontinuation of Friedreich's ataxia program	1,300	272	1,221	1,222
<b>Total of pre-tax non-GAAP adjustments</b>	<b>117,799</b>	<b>111,625</b>	<b>219,359</b>	<b>267,978</b>
Income tax effect of pre-tax non-GAAP adjustments	(15,621)	(6,356)	(30,372)	26,638
Other non-GAAP income tax adjustments	(1,452)	—	(1,452)	(35,893)
<b>Total of non-GAAP adjustments</b>	<b>100,726</b>	<b>105,269</b>	<b>187,535</b>	<b>258,723</b>
<b>Non-GAAP Net Income</b>	<b>\$ 95,606</b>	<b>\$ 80,518</b>	<b>\$ 149,552</b>	<b>\$ 85,316</b>
<b>Non-GAAP Earnings Per Share:</b>				
<b>Weighted average ordinary shares - Basic</b>	<b>185,327,383</b>	<b>165,536,826</b>	<b>178,866,391</b>	<b>164,921,722</b>
<b>Non-GAAP Earnings Per Share - Basic:</b>				
<b>GAAP loss per share - Basic</b>	<b>\$ (0.03)</b>	<b>\$ (0.15)</b>	<b>\$ (0.21)</b>	<b>\$ (1.05)</b>
Non-GAAP adjustments	0.55	0.64	1.05	1.57
<b>Non-GAAP earnings per share - Basic</b>	<b>\$ 0.52</b>	<b>\$ 0.49</b>	<b>\$ 0.84</b>	<b>\$ 0.52</b>
<b>Weighted average ordinary shares - Diluted</b>				
Weighted average ordinary shares - Basic	185,327,383	165,536,826	178,866,391	164,921,722
Ordinary share equivalents	7,897,507	3,820,913	7,658,133	3,678,249
<b>Weighted average shares - Diluted</b>	<b>193,224,890</b>	<b>169,357,739</b>	<b>186,524,524</b>	<b>168,599,971</b>
<b>Non-GAAP Earnings Per Share - Diluted</b>				
<b>GAAP loss per share - Diluted</b>	<b>\$ (0.03)</b>	<b>\$ (0.15)</b>	<b>\$ (0.21)</b>	<b>\$ (1.05)</b>
Non-GAAP adjustments	0.55	0.64	1.05	1.57
Diluted earnings per share effect of ordinary share equivalents	(0.03)	(0.01)	(0.04)	(0.01)
<b>Non-GAAP earnings per share - Diluted</b>	<b>\$ 0.49</b>	<b>\$ 0.48</b>	<b>\$ 0.80</b>	<b>\$ 0.51</b>



**Horizon Therapeutics plc**  
**GAAP to Non-GAAP Reconciliations**  
**EBITDA (Unaudited)**  
**(in thousands)**

	<u>Three Months Ended June 30,</u>		<u>Six Months Ended June 30,</u>	
	<u>2019</u>	<u>2018</u>	<u>2019</u>	<u>2018</u>
<b>GAAP net loss</b>	<b>\$ (5,120)</b>	<b>\$ (24,751)</b>	<b>\$ (37,983)</b>	<b>\$ (173,407)</b>
Depreciation	1,443	1,551	2,916	3,104
Amortization, accretion and step-up:				
Intangible amortization expense	57,683	60,480	115,100	121,364
Inventory step-up expense	(25)	53	90	17,129
Interest expense, net (including amortization of debt discount and deferred financing costs)	22,033	31,030	49,563	61,484
(Benefit) expense for income taxes	(4,875)	4,621	(6,795)	5,566
<b>EBITDA</b>	<b>\$ 71,139</b>	<b>\$ 72,984</b>	<b>\$ 122,891</b>	<b>\$ 35,240</b>
Other non-GAAP adjustments:				
Acquisition/divestiture-related costs	1,200	1,078	2,546	5,803
Restructuring and realignment costs	13	7,039	33	10,307
Impairment of long-lived assets	—	—	—	33,647
Loss on sale of assets	10,963	—	10,963	—
Share-based compensation	21,367	30,721	48,915	58,554
Litigation settlements	1,000	4,250	1,000	4,250
Upfront, progress and milestone payments related to license and collaboration agreements	4,000	—	6,000	90
Fees related to refinancing activities	1,033	15	1,175	42
Loss on debt extinguishment	11,878	—	17,464	—
Drug substance harmonization costs	234	475	314	1,279
Charges relating to discontinuation of Friedreich's ataxia program	1,300	272	1,221	1,222
<b>Total of other non-GAAP adjustments</b>	<b>52,988</b>	<b>43,850</b>	<b>89,631</b>	<b>115,194</b>
<b>Adjusted EBITDA</b>	<b>\$ 124,127</b>	<b>\$ 116,834</b>	<b>\$ 212,522</b>	<b>\$ 150,434</b>

Horizon Therapeutics plc





**Horizon Therapeutics plc**  
**GAAP to Non-GAAP Reconciliations**  
**EBITDA (Unaudited) – 2018**  
**(in thousands)**

	<u>Twelve Months</u> <u>Ended December 31,</u> <u>2018</u>
<b>GAAP net loss</b>	<b>\$ (38,380)</b>
Depreciation	6,126
Amortization, accretion and step-up:	
Intangible amortization expense	243,634
Inventory step-up expense	17,312
Interest expense, net (including amortization of debt discount and deferred financing costs)	121,692
Benefit for income taxes	(44,752)
<b>EBITDA</b>	<b>\$ 305,632</b>
Other non-GAAP adjustments:	
Acquisition/divestiture-related costs	4,396
Restructuring and realignment costs	15,350
Share-based compensation	114,860
Impairment of long-lived assets	46,096
Litigation settlements	5,750
Upfront, progress and milestone payments related to license and collaboration agreements	(10)
Fees related to refinancing activities	937
Drug substance harmonization costs	2,855
Charges relating to discontinuation of Friedreich's ataxia program	(1,464)
Gain on sale of assets	(42,985)
Total of other non-GAAP adjustments	145,785
<b>Adjusted EBITDA</b>	<b>\$ 451,417</b>



**Horizon Therapeutics plc**  
**GAAP to Non-GAAP Reconciliations**  
**Operating Income (Unaudited)**  
**(in thousands)**

	<u>Three Months Ended June 30,</u>		<u>Six Months Ended June 30,</u>	
	2019	2018	2019	2018
<b>GAAP operating income (loss)</b>	<b>\$ 25,112</b>	<b>\$ 10,559</b>	<b>\$ 23,317</b>	<b>\$ (106,739)</b>
Non-GAAP adjustments:				
Acquisition/divestiture-related costs	73	1,077	1,275	5,775
Restructuring and realignment costs	13	7,039	33	10,307
Amortization and step-up:				
Intangible amortization expense	57,683	60,480	115,100	121,364
Inventory step-up expense	(25)	53	90	17,129
Impairment of long-lived assets	—	—	—	33,647
Loss on sale of assets	10,963	—	10,963	—
Share-based compensation	21,367	30,721	48,915	58,554
Depreciation	1,443	1,551	2,916	3,104
Litigation settlements	1,000	4,250	1,000	4,250
Upfront, progress and milestone payments related to license and collaboration agreements	4,000	—	6,000	90
Fees related to refinancing activities	1,033	15	1,175	42
Drug substance harmonization costs	234	475	314	1,279
Charges relating to discontinuation of Friedreich's ataxia program	1,300	272	1,221	1,222
<b>Total of non-GAAP adjustments</b>	<b>99,084</b>	<b>105,933</b>	<b>189,002</b>	<b>256,763</b>
<b>Non-GAAP operating income</b>	<b>\$ 124,196</b>	<b>\$ 116,492</b>	<b>\$ 212,319</b>	<b>\$ 150,024</b>
Orphan and Rheumatology segment operating income	74,502	70,609	121,180	113,713
Inflammation segment operating income	49,694	45,883	91,139	36,311
<b>Total segment operating income</b>	<b>\$ 124,196</b>	<b>\$ 116,492</b>	<b>\$ 212,319</b>	<b>\$ 150,024</b>
Foreign exchange gain (loss)	76	(5)	15	(115)
Other income, net	(145)	347	188	525
<b>Adjusted EBITDA</b>	<b>\$ 124,127</b>	<b>\$ 116,834</b>	<b>\$ 212,522</b>	<b>\$ 150,434</b>



**Horizon Therapeutics plc**  
**GAAP to Non-GAAP Reconciliations**  
**Gross Profit and Operating Cash Flow (Unaudited)**  
(in thousands, except percentages)

	<u>Three Months Ended June 30,</u>		<u>Six Months Ended June 30,</u>	
	<u>2019</u>	<u>2018</u>	<u>2019</u>	<u>2018</u>
<b>Non-GAAP Gross Profit:</b>				
<b>GAAP gross profit</b>	<b>\$ 231,484</b>	<b>\$ 211,498</b>	<b>\$ 423,713</b>	<b>\$ 325,091</b>
Non-GAAP gross profit adjustments:				
Acquisition/divestiture-related costs	—	(664)	1,114	68
Intangible amortization expense	57,481	60,277	114,699	120,961
Inventory step-up expense	(25)	53	90	17,129
Share-based compensation	951	1,110	1,990	1,893
Depreciation	158	176	317	353
Drug substance harmonization costs	234	475	314	1,279
Charges relating to discontinuation of Friedreich's ataxia program	1,300	185	1,221	1,135
<b>Total of Non-GAAP adjustments</b>	<b>60,099</b>	<b>61,612</b>	<b>119,745</b>	<b>142,818</b>
<b>Non-GAAP gross profit</b>	<b>\$ 291,583</b>	<b>\$ 273,110</b>	<b>\$ 543,458</b>	<b>\$ 467,909</b>
<b>GAAP gross profit %</b>	72.2%	69.8%	70.5%	61.7%
<b>Non-GAAP gross profit %</b>	90.9%	90.2%	90.4%	88.8%
<b>GAAP cash provided by operating activities</b>	<b>\$ 91,267</b>	<b>\$ 61,788</b>	<b>\$ 147,436</b>	<b>\$ 974</b>
Cash payments for acquisition/divestiture-related costs	142	1,597	495	5,555
Cash payments for restructuring and realignment costs	839	4,230	2,882	4,677
Cash payments for litigation settlements	—	1,500	—	1,500
Cash payments for upfront, progress and milestone payments related to license and collaboration agreement	—	—	2,000	275
Cash payments drug substance harmonization costs	25	5,960	672	5,960
Cash payments for discontinuation of Friedreich's ataxia program	1,659	108	2,589	3,507
Cash payments relating to refinancing activities	1,797	13	1,806	31
<b>Non-GAAP operating cash flow</b>	<b>\$ 95,729</b>	<b>\$ 75,196</b>	<b>\$ 157,880</b>	<b>\$ 22,479</b>



**Horizon Therapeutics plc**  
**Net Debt Reconciliation (Unaudited)**  
**(in thousands)**

	As of		
	June 30, 2019	December 31, 2018	June 30, 2018
Long-term debt, net of current Exchangeable notes, net	\$1,025,096 341,682	\$ 1,564,485 332,199	\$1,562,013 323,105
<b>Total Debt</b>	<b>1,366,778</b>	<b>1,896,684</b>	<b>1,885,118</b>
Debt discount	70,754	87,038	97,737
Deferred financing fees	5,494	9,304	10,171
<b>Total Principal Amount Debt</b>	<b>1,443,026</b>	<b>1,993,026</b>	<b>1,993,026</b>
Less: cash and cash equivalents	865,997	958,712	710,211
<b>Net Debt</b>	<b>\$ 577,029</b>	<b>\$ 1,034,314</b>	<b>\$1,282,815</b>

Horizon Therapeutics plc

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**Horizon Therapeutics plc**  
**GAAP to Non-GAAP Tax Rate Reconciliation (Unaudited)**  
(in millions, except percentages and per share amounts)

**Q2 2019**

	Pre-tax Net (Loss) Income	Income Tax (Benefit) Expense	Tax Rate	Net (Loss) Income	Diluted (Loss) Earnings Per Share
As reported - GAAP	\$ (10.0)	\$ (4.9)	48.8%	\$ (5.1)	\$ (0.03)
Non-GAAP adjustments	117.8	17.1		100.7	
<b>Non-GAAP</b>	<b>\$ 107.8</b>	<b>\$ 12.2</b>	<b>11.3%</b>	<b>\$ 95.6</b>	<b>\$ 0.49</b>

**Q2 2018**

	Pre-tax Net (Loss) Income	Income Tax (Benefit) Expense	Tax Rate	Net (Loss) Income	Diluted (Loss) Earnings Per Share
As reported - GAAP	\$ (20.1)	\$ 4.6	(23.0)%	\$ (24.8)	\$ (0.15)
Non-GAAP adjustments	111.6	6.4		105.3	
<b>Non-GAAP</b>	<b>\$ 91.5</b>	<b>\$ 11.0</b>	<b>12.0%</b>	<b>\$ 80.5</b>	<b>\$ 0.48</b>

**YTD 2019**

	Pre-tax Net (Loss) Income	Income Tax (Benefit) Expense	Tax Rate	Net (Loss) Income	Diluted (Loss) Earnings Per Share
As reported - GAAP	\$ (44.8)	\$ (6.8)	15.2%	\$ (38.0)	\$ (0.21)
Non-GAAP adjustments	219.4	31.8		187.5	
<b>Non-GAAP</b>	<b>\$ 174.6</b>	<b>\$ 25.0</b>	<b>14.3%</b>	<b>\$ 149.5</b>	<b>\$ 0.80</b>

**YTD 2018**

	Pre-tax Net (Loss) Income	Income Tax (Benefit) Expense	Tax Rate	Net (Loss) Income	Diluted (Loss) Earnings Per Share
As reported - GAAP	\$ (167.8)	\$ 5.6	(3.3)%	\$ (173.4)	\$ (1.05)
Non-GAAP adjustments	268.0	9.3		258.7	
<b>Non-GAAP</b>	<b>\$ 100.2</b>	<b>\$ 14.9</b>	<b>14.9%</b>	<b>\$ 85.3</b>	<b>\$ 0.51</b>



**Horizon Therapeutics plc**  
**Certain Income Statement Line Items - Non-GAAP Adjusted**  
**For the Three Months Ended June 30, 2019**  
**(Unaudited)**

	COGS	Research & Development	Selling, General & Administrative	Loss on Sale of Assets	Loss on Debt Extinguishment	Interest Expense	Other Expense	Income Tax Benefit (Expense)
<b>GAAP as reported</b>	<b>\$(89,163)</b>	<b>\$ (28,314)</b>	<b>\$ (167,095)</b>	<b>\$ (10,963)</b>	<b>\$ (11,878)</b>	<b>\$(22,033)</b>	<b>\$ (1,272)</b>	<b>\$ 4,875</b>
<b>Non-GAAP Adjustments (in thousands):</b>								
Acquisition/divestiture-related costs(1)	—	—	73	—	—	—	1,127	—
Restructuring and realignment costs(2)	—	—	13	—	—	—	—	—
<b>Amortization and step-up:</b>								
Intangible amortization expense(3)	57,481	—	202	—	—	—	—	—
Inventory step-up expense(4)	(25)	—	—	—	—	—	—	—
Amortization of debt discount and deferred financing costs(5)	—	—	—	—	—	5,710	—	—
Loss on sale of assets(7)	—	—	—	10,963	—	—	—	—
Share-based compensation(8)	951	2,343	18,073	—	—	—	—	—
Depreciation(9)	158	—	1,285	—	—	—	—	—
Litigation settlements(10)	—	—	1,000	—	—	—	—	—
<b>Upfront, progress and milestone payments related to license and collaboration agreements(11)</b>								
Fees related to refinancing activities (12)	—	4,000	—	—	—	—	—	—
Loss on debt extinguishment(13)	—	—	—	—	11,878	—	—	—
Drug substance harmonization costs(14)	234	—	—	—	—	—	—	—
Charges relating to discontinuation of Friedreich's ataxia program(15)	1,300	—	—	—	—	—	—	—
Income tax effect on pre-tax non-GAAP adjustments(16)	—	—	—	—	—	—	—	(15,621)
Other non-GAAP income tax adjustments(17)	—	—	—	—	—	—	—	(1,452)
<b>Total of non-GAAP adjustments</b>	<b>60,099</b>	<b>6,343</b>	<b>21,679</b>	<b>10,963</b>	<b>11,878</b>	<b>5,710</b>	<b>1,127</b>	<b>(17,073)</b>
<b>Non-GAAP</b>	<b>\$(29,064)</b>	<b>\$ (21,971)</b>	<b>\$ (145,416)</b>	<b>\$ —</b>	<b>\$ —</b>	<b>\$(16,323)</b>	<b>\$ (145)</b>	<b>\$ (12,198)</b>

**Horizon Therapeutics plc**  
**Certain Income Statement Line Items - Non-GAAP Adjusted**  
**For the Three Months Ended June 30, 2018**  
**(Unaudited)**

	COGS	Research & Development	Selling, General & Administrative	Interest Expense	Income Tax Benefit (Expense)
<b>GAAP as reported</b>	<b>\$(91,337)</b>	<b>\$ (24,265)</b>	<b>\$ (176,674)</b>	<b>\$(31,030)</b>	<b>\$ (4,621)</b>
<b>Non-GAAP Adjustments (in thousands):</b>					
Acquisition/divestiture-related costs(1)	(664)	18	1,724	—	—
Restructuring and realignment costs(2)	—	1,733	5,306	—	—
<b>Amortization and step-up:</b>					
Intangible amortization expense(3)	60,277	—	202	—	—
Inventory step-up expense(4)	53	—	—	—	—
Amortization of debt discount and deferred financing costs(5)	—	—	—	5,691	—
Share-based compensation(8)	1,110	2,209	27,402	—	—
Depreciation(9)	176	—	1,375	—	—
Litigation settlements(10)	—	—	4,250	—	—
Fees related to refinancing activities (12)	—	—	15	—	—
Drug substance harmonization costs(14)	475	—	—	—	—
Charges relating to discontinuation of Friedreich's ataxia program(15)	185	87	—	—	—
Income tax effect on pre-tax non-GAAP adjustments(16)	—	—	—	—	(6,356)
<b>Total of non-GAAP adjustments</b>	<b>61,612</b>	<b>4,047</b>	<b>40,274</b>	<b>5,691</b>	<b>(6,356)</b>
<b>Non-GAAP</b>	<b>\$(29,725)</b>	<b>\$ (20,218)</b>	<b>\$ (136,400)</b>	<b>\$(25,339)</b>	<b>\$ (10,977)</b>



**Horizon Therapeutics plc**  
**Certain Income Statement Line Items - Non-GAAP Adjusted**  
**For the Six Months Ended June 30, 2019**  
**(Unaudited)**

	COGS	Research & Development	Selling, General & Administrative	Loss on Sale of Assets	Interest Expense	Other Expense	Loss on Debt Extinguishment	Income Tax Benefit (Expense)
<b>GAAP as reported</b>	<b>\$(177,305)</b>	<b>\$ (50,039)</b>	<b>\$ (339,394)</b>	<b>\$ (10,963)</b>	<b>\$(49,563)</b>	<b>\$ (1,083)</b>	<b>\$ (17,464)</b>	<b>\$ 6,795</b>
<b>Non-GAAP Adjustments (in thousands):</b>								
Acquisition/divestiture-related costs(1)	1,114	—	164	—	—	1,268	—	—
Restructuring and realignment costs(2)	—	—	33	—	—	—	—	—
<b>Amortization and step-up:</b>								
Intangible amortization expense(3)	114,699	—	401	—	—	—	—	—
Inventory step-up expense(4)	90	—	—	—	—	—	—	—
Amortization of debt discount and deferred financing costs(5)	—	—	—	—	11,622	—	—	—
Loss on sale of assets(7)	—	—	—	10,963	—	—	—	—
Share-based compensation(8)	1,990	4,979	41,946	—	—	—	—	—
Depreciation(9)	317	—	2,599	—	—	—	—	—
Litigation settlements(10)	—	—	1,000	—	—	—	—	—
Upfront, progress and milestone payments related to license and collaboration agreements(11)	—	6,000	—	—	—	—	—	—
Fees related to refinancing activities (12)	—	—	1,175	—	—	—	—	—
Loss on debt extinguishment(13)	—	—	—	—	—	—	17,464	—
Drug substance harmonization costs(14)	314	—	—	—	—	—	—	—
Charges relating to discontinuation of Friedrich's ataxia program(15)	1,221	—	—	—	—	—	—	—
Income tax effect on pre-tax non-GAAP adjustments(16)	—	—	—	—	—	—	—	(30,372)
Other non-GAAP income tax adjustments(17)	—	—	—	—	—	—	—	(1,452)
<b>Total of non-GAAP adjustments</b>	<b>119,745</b>	<b>10,979</b>	<b>47,318</b>	<b>10,963</b>	<b>11,622</b>	<b>1,268</b>	<b>17,464</b>	<b>(31,824)</b>
<b>Non-GAAP</b>	<b>\$ (57,560)</b>	<b>\$ (39,060)</b>	<b>\$ (292,076)</b>	<b>\$ —</b>	<b>\$(37,941)</b>	<b>\$ 185</b>	<b>\$ —</b>	<b>\$ (25,029)</b>

**Horizon Therapeutics plc**  
**Certain Income Statement Line Items - Non-GAAP Adjusted**  
**For the Six Months Ended June 30, 2018**  
**(Unaudited)**

	COGS	Research & Development	Selling, General & Administrative	Impairment of Long-Lived Assets	Interest Expense	Income Tax Benefit (Expense)
<b>GAAP as reported</b>	<b>\$(201,625)</b>	<b>\$ (41,910)</b>	<b>\$ (356,273)</b>	<b>\$ (33,647)</b>	<b>\$(61,484)</b>	<b>\$ (5,566)</b>
<b>Non-GAAP Adjustments (in thousands):</b>						
Acquisition/divestiture-related costs(1)	68	(67)	5,800	—	—	—
Restructuring and realignment costs(2)	—	1,733	8,574	—	—	—
<b>Amortization and step-up:</b>						
Intangible amortization expense(3)	120,961	—	402	—	—	—
Inventory step-up expense(4)	17,129	—	—	—	—	—
Amortization of debt discount and deferred financing costs(5)	—	—	—	—	11,187	—
Impairment of long lived assets(6)	—	—	—	33,647	—	—
Share-based compensation(8)	1,893	4,649	52,012	—	—	—
Depreciation(9)	353	—	2,751	—	—	—
Litigation settlements(10)	—	—	4,250	—	—	—
Upfront, progress and milestone payments related to license and collaboration agreements(11)	—	90	—	—	—	—
Fees related to refinancing activities (12)	—	—	42	—	—	—
Drug substance harmonization costs(14)	1,279	—	—	—	—	—
Charges relating to discontinuation of Friedrich's ataxia program(15)	1,135	87	—	—	—	—
Income tax effect on pre-tax non-GAAP adjustments(16)	—	—	—	—	—	26,638
Other non-GAAP income tax adjustments(17)	—	—	—	—	—	(35,893)
<b>Total of non-GAAP adjustments</b>	<b>142,818</b>	<b>6,492</b>	<b>73,831</b>	<b>33,647</b>	<b>11,187</b>	<b>(9,255)</b>
<b>Non-GAAP</b>	<b>\$ (58,807)</b>	<b>\$ (35,418)</b>	<b>\$ (282,442)</b>	<b>\$ —</b>	<b>\$(50,297)</b>	<b>\$ (14,821)</b>



#### NOTES FOR CERTAIN INCOME STATEMENT LINE ITEMS - NON-GAAP

1. Represents expenses, including legal and consulting fees, incurred in connection with our acquisitions and divestitures.
2. Represents expenses, including severance costs and consulting fees, related to restructuring and realignment activities.
3. Intangible amortization expenses are associated with our intellectual property rights, developed technology and customer relationships related to ACTIMMUNE, BUPHENYL, KRYSTEXXA, LODOTRA, MIGERGOT, PENNSAID 2%, PROCYSBI, RAVICTI, VIMOVO and RAYOS.
4. During the six months ended June 30, 2018, we recognized in cost of goods sold \$17.1 million for inventory step-up expense primarily related to KRYSTEXXA inventory sold.
5. Represents amortization of debt discount and deferred financing costs associated with our debt.
6. Impairment of long-lived assets during the six months ended June 30, 2018, relates to the write-off of the book value of developed technology related to PROCYSBI in Canada and Latin America.
7. During the six months ended June 30, 2019, we recorded a loss of \$10.9 million on the sale of our rights to MIGERGOT.
8. Represents share-based compensation expense associated with our stock option, restricted stock unit and performance stock unit grants to our employees and non-employees and our employee share purchase plan.
9. Represents depreciation expense related to our property, equipment, software and leasehold improvements.
10. The company recorded \$1.0 million and \$4.3 million of expense during the three months ended June 30, 2019, and June 30, 2018, respectively, for litigation settlements.
11. During the six months ended June 30, 2019, we recorded an upfront cash payment of \$2.0 million and a \$4.0 million progress payment in relation to the collaboration agreement with HemoShear.
12. Represents arrangement and other fees relating to our refinancing activities.
13. During the six months ended June 30, 2019, we recorded a loss on debt extinguishment of \$17.5 million in the condensed consolidated statements of comprehensive loss, which reflected the write-off of the deferred financing fees and debt discount fees related to the prepayment of \$225.0 million of 2023 Senior Notes and term loan repayments of \$300.0 million.



14. During the year ended December 31, 2016, we entered into a definitive agreement to acquire certain rights to interferon gamma-1b, marketed as IMUKIN in an estimated thirty countries primarily in Europe and the Middle East, or the IMUKIN purchase agreement. We already owned the rights to interferon gamma-1b marketed as ACTIMMUNE in the United States, Canada and Japan. In connection with the IMUKIN purchase agreement, we also committed to pay our contract manufacturer certain amounts related to the harmonization of the manufacturing processes for ACTIMMUNE and IMUKIN drug substance, or the harmonization program. At the time we entered into the IMUKIN purchase agreement and the harmonization program commitment was made, we had anticipated achieving certain benefits should the Phase 3 clinical trial evaluating ACTIMMUNE for the treatment of Friedreich's ataxia, or FA, be successful. If the study had been successful and if U.S. marketing approval had subsequently been obtained, we had forecasted significant increases in demand for the medicine and the harmonization program would have resulted in significant benefits for us. Following our discontinuation of the FA program, we determined that certain assets, including an upfront payment related to the IMUKIN purchase agreement, were impaired, and the costs under the harmonization program would no longer have benefit to us and should be expensed as incurred.
15. Represents expenses incurred relating to discontinuation of Friedreich's ataxia program and a reduction to previous charges recorded.
16. Income tax adjustments on pre-tax non-GAAP adjustments represent the estimated income tax impact of each pre-tax non-GAAP adjustment based on the statutory income tax rate of the applicable jurisdictions for each non-GAAP adjustment.
17. Following Notice 2018-28, issued by the U.S. Treasury Department and the U.S. Internal Revenue Service on April 2, 2018 and in accordance with the measurement period provisions under Staff Accounting Bulletin No. 118, or SAB 118, during the six months ended June 30, 2019 we reinstated the deferred tax asset previously written off during the year ended December 31, 2017, related to our U.S. interest expense carry forwards under Section 163(j) of the Internal Revenue Code of 1986, as amended, based on the revised U.S. federal tax rate of 21 percent. The impact of the deferred tax asset reinstatement in accordance with SAB 118 was a \$35.9 million increase to our benefit for income taxes and a corresponding decrease to the U.S. group net deferred tax liability position.

During the three months ended June 30, 2019 the Company released a reserve related to an uncertain tax position in connection with an acquisition resulting in a non-GAAP tax adjustment of \$1.5 million.

**HORIZON THERAPEUTICS PLC**  
**ADJUSTED HISTORICAL FINANCIAL INFORMATION (UNAUDITED)**  
**DATED AUGUST 7, 2019**

*Change in Accounting Principle*

As previously disclosed in its interim financial statements on Form 10-Q for the three months ended March 31, 2019, Horizon Therapeutics plc (“the Company”) changed its accounting for business combinations in respect of intangible assets acquired and their related third-party contingent royalties. When accounting for business combinations under ASC Topic 805, Business Combinations, the Company previously separately identified and recorded at fair value intangible assets acquired and their related third-party contingent royalties at the date of acquisition. Third-party contingent royalties are payable to parties other than sellers of the businesses. Effective January 1, 2019, the Company retrospectively changed its accounting for business combinations and is now recording acquired intangible assets and their related third-party contingent royalties on a net basis (“New Method”). The Company changed its accounting principle on the basis that the use of the New Method is preferable, primarily due to improved comparability with the Company’s peers.

The change in accounting principle resulted in the Company re-performing its purchase price allocations as of the respective acquisition dates for prior business combinations. The adjustments to the purchase price allocations primarily resulted in a decrease in developed technology intangible assets and the elimination of liabilities for accrued contingent royalties due to recording these items on a net basis. The re-performance of purchase price allocations also impacted goodwill and deferred tax liabilities. In addition, the change in accounting principle resulted in the elimination of royalty reimbursement assets and accrued contingent royalty liabilities that were recorded in connection with divestitures, impacting prepaid expenses and other current assets, other assets, accrued expenses and other long-term liabilities captions as shown in the tables below. In addition, under the New Method of accounting, the Company is presenting accrued royalties based on each period’s net sales as part of the accrued expenses line item on its consolidated balance sheets.

Furthermore, the adjustments to the purchase price allocations primarily resulted in a net decrease in cost of goods sold reflecting lower intangible asset amortization and the elimination of royalty accretion and remeasurement expenses, partially offset by the royalty expense based on the periods’ net sales. The re-performance of purchase price allocations also directly impacted impairments of long-lived assets and benefit/expense for income taxes, as shown in the tables below. In addition, the elimination of royalty reimbursement assets and accrued contingent royalty liabilities that were recorded in connection with divestitures resulted in adjustments to gain on sale of assets, gain on divestiture and other income, net.

*Revision of Prior Period Financial Statements*

As previously disclosed in the Company’s Annual Report on Form 10-K for the year ended December 31, 2018, during the course of preparing its consolidated financial statements for the year ended December 31, 2018, the Company identified an error in the measurement of the contingent royalty liability calculation pertaining to the royalty end date for one of its medicines. The royalty end date for KRYSTEXXA is approximately two and one half years earlier than the date originally assumed in the calculations. As a result of the error, accrued royalties, net of current, and cost of goods sold had been overstated and shareholders’ equity had been understated. The Company concluded that the amounts were not material to any of its previously issued consolidated financial statements.

The revision did not impact the consolidated balance sheet and consolidated statement of comprehensive loss as of and for the year ended December 31, 2018, as the error had been identified prior to the reporting of these consolidated financial statements. The comparative amounts included in the consolidated financial statements included in the Company’s Annual Report on Form 10-K for the year ended December 31, 2018 reflected the impact of the revision; therefore, the ‘As Previously Reported’ columns in the tables below for the consolidated balance sheet and consolidated statement of comprehensive loss as of and for the year ended December 31, 2017, and the consolidated statement of comprehensive loss for the year ended December 31, 2016, reflect the impact of the revision. The revision did not impact the consolidated financial statements as of and for the years ended December 31, 2013, 2014 and 2015 as the error related to the accounting for a business combination that occurred in January 2016.

The ‘As Previously Reported’ column in the table below for the consolidated statement of comprehensive loss for the three months ended December 31, 2018 was derived from Note 23 to the consolidated financial statements included in the Company’s Annual Report on Form 10-K for the year ended December 31, 2018, which reflected the impact of the revision. The ‘As Previously Reported’ columns in the tables below for the consolidated statements of comprehensive loss for the three-month periods ended September 30, 2018, June 30, 2018 and March 31, 2018, were derived from the previously reported interim financial statements on Form for 10-Q for each of the respective periods.

*Adjusted Financial Information*

The impact of the change in accounting principle and the revision on the consolidated statements of cash flows consisted of adjustments to reconcile net (loss) income to net cash provided by operating activities and changes in operating assets and liabilities for all periods presented. There was no impact on total operating, investing or financing cash flows for any prior period. In addition, there was no impact from the change in accounting principle and the revision on the Company’s previously reported adjusted EBITDA, non-GAAP net income, non-GAAP diluted earnings per share, non-GAAP gross profit and gross profit ratio, non-GAAP operating expenses, non-GAAP operating income, non-GAAP tax rate, non-GAAP operating cash flow and net debt for any prior period.

The following tables present selected line items from the Company's consolidated financial statements for the years ended December 31, 2018, 2017, 2016, 2015, 2014 and 2013 illustrating the effect of the change in accounting principle, in addition to the effect of the revision, where applicable:

<b>Consolidated Balance Sheet</b>			
<b>As of December 31, 2018</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Prepaid expenses and other current assets	\$ 70,828	\$ (2,610)	\$ 68,218
Total current assets	1,548,426	(2,610)	1,545,816
Developed technology, net	2,120,596	(174,957)	1,945,639
Goodwill	426,441	(12,772)	413,669
Other assets	23,029	(14,070)	8,959
Total assets	4,146,371	(204,409)	3,941,962
Accrued expenses	205,593	10,146	215,739
Accrued royalties - current portion	63,363	(63,363)	—
Total current liabilities	761,904	(53,217)	708,687
Accrued royalties - net of current	285,374	(285,374)	—
Deferred tax liabilities, net	93,630	14,138	107,768
Other long-term liabilities	54,622	(15,905)	38,717
Total long-term liabilities	2,330,310	(287,141)	2,043,169
Accumulated deficit	(1,314,718)	135,949	(1,178,769)
Total shareholders' equity	1,054,157	135,949	1,190,106
Total liabilities and shareholders' equity	4,146,371	(204,409)	3,941,962

<b>Consolidated Statement of Comprehensive Loss</b>			
<b>For the Twelve Months Ended December 31, 2018</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Cost of goods sold	\$ 422,317	\$ (31,016)	\$ 391,301
Gross profit	785,253	31,016	816,269
Impairment of long-lived assets	50,302	(4,206)	46,096
Gain on sale of assets	(42,688)	(297)	(42,985)
Total operating expenses	782,861	(4,503)	778,358
Operating income	2,392	35,519	37,911
Other income, net	346	495	841
Total other expenses, net	(121,538)	495	(121,043)
Loss before benefit for income taxes	(119,146)	36,014	(83,132)
Benefit for income taxes	(44,959)	207	(44,752)
Net loss	(74,187)	35,807	(38,380)
Net loss per ordinary share - basic and diluted	(0.45)	0.22	(0.23)
Comprehensive loss	(74,727)	35,807	(38,920)

**Consolidated Balance Sheet**

**As of December 31, 2017**

	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Prepaid expenses and other current assets	\$ 43,402	\$ (2,643)	\$ 40,759
Total current assets	1,268,168	(2,643)	1,265,525
Developed technology, net	2,442,292	(205,426)	2,236,866
Goodwill	426,441	(12,772)	413,669
Other assets	36,081	(19,985)	16,096
Total assets	4,202,298	(240,826)	3,961,472
Accrued expenses	175,697	8,979	184,676
Accrued royalties - current portion	65,328	(65,328)	—
Total current liabilities	794,969	(56,349)	738,620
Accrued royalties - net of current	279,316	(279,316)	—
Deferred tax liabilities, net	157,945	13,932	171,877
Other long-term liabilities	68,015	(19,235)	48,780
Total long-term liabilities	2,406,019	(284,619)	2,121,400
Accumulated deficit	(1,242,117)	100,142	(1,141,975)
Total shareholders' equity	1,001,310	100,142	1,101,452
Total liabilities and shareholders' equity	4,202,298	(240,826)	3,961,472

**Consolidated Statement of Comprehensive  
Loss**

**For the Twelve Months Ended  
December 31, 2017**

	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Cost of goods sold	\$ 537,334	\$ (43,966)	\$ 493,368
Gross profit	518,897	43,966	562,863
Operating loss	(383,428)	43,966	(339,462)
Gain on divestiture	6,267	1,698	7,965
Other income, net	588	(141)	447
Total other expenses, net	(120,906)	1,557	(119,349)
Loss before benefit for income taxes	(504,334)	45,523	(458,811)
Benefit for income taxes	(102,749)	(5,937)	(108,686)
Net loss	(401,585)	51,460	(350,125)
Net loss per ordinary share - basic and diluted	(2.46)	0.31	(2.15)
Comprehensive loss	(399,482)	51,460	(348,022)

**Consolidated Balance Sheet**  
**As of December 31, 2016**

	<u>As Previously Reported</u>	<u>Revision</u>	<u>As Revised</u>	<u>Impact of Accounting Change</u>	<u>As Adjusted</u>
Developed technology, net	\$ 2,767,184	\$(1,829)	\$2,765,355	\$(237,216)	\$2,528,139
Goodwill	445,579	—	445,579	(13,364)	432,215
Total assets	4,307,306	(1,829)	4,305,477	(250,580)	4,054,897
Accrued expenses	198,012	—	198,012	10,698	208,710
Accrued royalties - current portion	61,981	—	61,981	(61,981)	—
Total current liabilities	621,099	—	621,099	(51,283)	569,816
Accrued royalties - net of current	272,293	(3,100)	269,193	(269,193)	—
Deferred tax liabilities, net	296,568	—	296,568	19,869	316,437
Other long-term liabilities	46,061	—	46,061	1,412	47,473
Total long-term liabilities	2,422,428	(3,100)	2,419,328	(247,912)	2,171,416
Accumulated deficit	(848,021)	1,271	(846,750)	48,615	(798,135)
Total shareholders' equity	1,263,779	1,271	1,265,050	48,615	1,313,665
Total liabilities and shareholders' equity	4,307,306	(1,829)	4,305,477	(250,580)	4,054,897

**Consolidated Statement of Comprehensive  
Loss**  
**For the Twelve Months Ended  
December 31, 2016**

	<u>As Previously Reported</u>	<u>Impact of Accounting Change</u>	<u>As Adjusted</u>
Cost of goods sold	\$ 392,001	\$ (25,596)	\$ 366,405
Gross profit	589,119	25,596	614,715
Impairment of long-lived assets	66,000	(1,300)	64,700
Total operating expenses	735,015	(1,300)	733,715
Operating loss	(145,896)	26,896	(119,000)
Other income, net	6,697	—	6,697
Total other expenses, net	(80,918)	—	(80,918)
Loss before benefit for income taxes	(226,814)	26,896	(199,918)
Benefit for income taxes	(61,251)	8,425	(52,826)
Net loss	(165,563)	18,471	(147,092)
Net loss per ordinary share - basic and diluted	(1.03)	0.11	(0.92)
Comprehensive loss	(165,998)	18,471	(147,527)

<b>Consolidated Balance Sheet</b>			
<b>As of December 31, 2015</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Developed technology, net	\$ 1,609,049	\$ (130,353)	\$1,478,696
In-process research and development	66,000	(1,300)	64,700
<b>Total assets</b>	<b>3,073,060</b>	<b>(131,653)</b>	<b>2,941,407</b>
Accrued expenses	114,518	9,314	123,832
Accrued royalties - current portion	51,700	(51,700)	—
Total current liabilities	372,024	(42,386)	329,638
Accrued royalties - net of current	123,519	(123,519)	—
Deferred tax liabilities, net	113,400	4,108	117,508
Total long-term liabilities	1,387,891	(119,411)	1,268,480
Accumulated deficit	(681,187)	30,144	(651,043)
Total shareholders' equity	1,313,145	30,144	1,343,289
<b>Total liabilities and shareholders' equity</b>	<b>3,073,060</b>	<b>(131,653)</b>	<b>2,941,407</b>

<b>Consolidated Statement of Comprehensive Income</b>			
<b>For the Twelve Months Ended December 31, 2015</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Cost of goods sold	\$ 219,502	\$ (24,986)	\$ 194,516
Gross profit	537,542	24,986	562,528
Operating income	55,372	24,986	80,358
Loss before benefit for income taxes	(132,712)	24,986	(107,726)
Benefit for income taxes	(172,244)	4,107	(168,137)
Net income	39,532	20,879	60,411
Net income per ordinary share - basic	0.27	0.14	0.41
Net income per ordinary share - diluted	0.25	0.14	0.39
Comprehensive income	41,244	20,879	62,123

<b>Consolidated Balance Sheet</b>			
<b>As of December 31, 2014</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Developed technology, net	\$ 696,963	\$ (57,484)	\$ 639,479
In-process research and development	66,000	(1,300)	64,700
Deferred tax assets, net, non-current	20,291	2,106	22,397
Total assets	1,126,302	(56,678)	1,069,624
Accrued expenses	49,794	6,163	55,957
Accrued royalties - current portion	25,325	(25,325)	—
Total current liabilities	221,840	(19,162)	202,678
Accrued royalties - net of current	48,887	(48,887)	—
Deferred tax liabilities, net, non-current	20,291	2,106	22,397
Total long-term liabilities	364,258	(46,781)	317,477
Accumulated deficit	(720,719)	9,265	(711,454)
Total shareholders' equity	540,204	9,265	549,469
Total liabilities and shareholders' equity	1,126,302	(56,678)	1,069,624

<b>Consolidated Statement of Comprehensive Loss</b>			
<b>For the Twelve Months Ended December 31, 2014</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Cost of goods sold	\$ 78,753	\$ (8,589)	\$ 70,164
Gross profit	218,202	8,589	226,791
Operating loss	(8,491)	8,589	98
Loss before benefit for income taxes	(269,687)	8,589	(261,098)
Net loss	(263,603)	8,589	(255,014)
Net loss per ordinary share - basic and diluted	(3.15)	0.11	(3.04)
Comprehensive loss	(265,563)	8,589	(256,974)

<b>Consolidated Balance Sheet</b>			
<b>As of December 31, 2013</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Developed technology, net	\$ 131,094	\$ (32,316)	\$ 98,778
Deferred tax assets, net, non-current	—	254	254
<b>Total assets</b>	<b>246,328</b>	<b>(32,062)</b>	<b>214,266</b>
Accrued royalties - current portion	8,010	(8,010)	—
<b>Total current liabilities</b>	<b>43,310</b>	<b>(8,010)</b>	<b>35,300</b>
Accrued royalties - net of current	24,982	(24,982)	—
Deferred tax liabilities, net, non-current	3,362	254	3,616
<b>Total long-term liabilities</b>	<b>252,100</b>	<b>(24,728)</b>	<b>227,372</b>
Accumulated deficit	(457,116)	676	(456,440)
Total shareholders' equity	(49,082)	676	(48,406)
<b>Total liabilities and shareholders' equity</b>	<b>246,328</b>	<b>(32,062)</b>	<b>214,266</b>

<b>Consolidated Statement of Comprehensive Loss</b>			
<b>For the Twelve Months Ended December 31, 2013</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Cost of goods sold	\$ 14,625	\$ (676)	\$ 13,949
Gross profit	59,391	676	60,067
Operating loss	(42,854)	676	(42,178)
Loss before benefit for income taxes	(150,126)	676	(149,450)
Net loss	(149,005)	676	(148,329)
Net loss per ordinary share - basic and diluted	(2.34)	0.01	(2.33)
Comprehensive loss	(148,036)	676	(147,360)



The following tables present selected line items from the Company's consolidated statement of comprehensive loss for the four quarters of 2018 illustrating the effect of the change in accounting principle, in addition to the effect of the revisions, where applicable:

<b>Consolidated Statement of Comprehensive Loss</b>			
<b>For the Three Months Ended December 31, 2018</b>			
	<b>As Previously Reported</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Cost of goods sold	\$ 109,520	\$ (10,921)	\$ 98,599
Gross profit	246,023	10,921	256,944
Gain on sale of assets	(30,385)	(297)	(30,682)
Total operating expenses	174,773	(297)	174,476
Operating income	71,250	11,218	82,468
Other income, net	(632)	640	8
Total other expenses, net	(30,514)	640	(29,874)
Income before benefit for income taxes	40,736	11,858	52,594
Benefit for income taxes	(46,822)	(2,232)	(49,054)
Net income	87,558	14,090	101,648
Net income per ordinary share - basic	0.52	0.08	0.60
Net income per ordinary share - diluted	0.50	0.08	0.58
Comprehensive income	87,296	14,090	101,386

<b>Consolidated Statement of Comprehensive Loss</b>					
<b>For the Three Months Ended September 30, 2018</b>					
	<b>As Previously Reported</b>	<b>Revision</b>	<b>As Revised</b>	<b>Impact of Accounting Change</b>	<b>As Adjusted</b>
Cost of goods sold	\$ 99,011	\$ (840)	\$ 98,171	\$ (7,094)	\$ 91,077
Gross profit	226,300	840	227,140	7,094	234,234
Operating income	54,246	840	55,086	7,094	62,180
Other income, net	453	—	453	(116)	337
Total other expenses, net	(29,949)	—	(29,949)	(116)	(30,065)
Income before benefit for income taxes	24,297	840	25,137	6,978	32,115
Benefit for income taxes	(1,733)	—	(1,733)	467	(1,266)
Net income	26,030	840	26,870	6,511	33,381
Net income per ordinary share - basic	0.16	—	0.16	0.04	0.20
Net income per ordinary share - diluted	0.15	0.01	0.16	0.03	0.19
Comprehensive income	25,897	840	26,737	6,511	33,248

**Consolidated Statement of Comprehensive Loss**  
**For the Three Months Ended June 30, 2018**

	<u>As Previously Reported</u>	<u>Revision</u>	<u>As Revised</u>	<u>Impact of Accounting Change</u>	<u>As Adjusted</u>
Cost of goods sold	\$ 100,082	\$ (795)	\$ 99,287	\$ (7,950)	\$ 91,337
Gross profit	202,753	795	203,548	7,950	211,498
Operating income	1,814	795	2,609	7,950	10,559
Other income, net	347	—	347	(1)	346
Total other expenses, net	(30,688)	—	(30,688)	(1)	(30,689)
Loss before expense for income taxes	(28,874)	795	(28,079)	7,949	(20,130)
Expense for income taxes	3,962	—	3,962	659	4,621
Net loss	(32,836)	795	(32,041)	7,290	(24,751)
Net loss per ordinary share - basic and diluted	(0.20)	0.01	(0.19)	0.04	(0.15)
Comprehensive loss	(33,444)	795	(32,649)	7,290	(25,359)

**Consolidated Statement of Comprehensive Loss**  
**For the Three Months Ended March 31, 2018**

	<u>As Previously Reported</u>	<u>Revision</u>	<u>As Revised</u>	<u>Impact of Accounting Change</u>	<u>As Adjusted</u>
Cost of goods sold	\$ 116,092	\$ (753)	\$ 115,339	\$ (5,051)	\$ 110,288
Gross profit	107,789	753	108,542	5,051	113,593
Impairment of long-lived assets	37,853	—	37,853	(4,206)	33,647
Total operating expenses	235,097	—	235,097	(4,206)	230,891
Operating loss	(127,308)	753	(126,555)	9,257	(117,298)
Other income, net	178	—	178	(27)	151
Total other expenses, net	(30,386)	—	(30,386)	(27)	(30,413)
Loss before (benefit) expense for income taxes	(157,694)	753	(156,941)	9,230	(147,711)
(Benefit) expense for income taxes	(367)	—	(367)	1,312	945
Net loss	(157,327)	753	(156,574)	7,918	(148,656)
Net loss per ordinary share - basic and diluted	(0.96)	0.01	(0.95)	0.05	(0.90)
Comprehensive loss	(156,864)	753	(156,111)	7,918	(148,193)