#### 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) April 29, 2020

#### **Vertex Pharmaceuticals Incorporated**

(Exact name of registrant as specified in its charter)

Massachusetts

(State or other jurisdiction of incorporation)

**000-19319** (Commission File Number) 04-3039129

(I.R.S. Employer Identification No.)

#### 50 Northern Avenue

#### Boston, Massachusetts 02210

(Address of principal executive offices) (Zip Code)

#### (617) 341-6100

(Registrant's telephone number, including area code)

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

□ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

□ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

□ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

□ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

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

Title of each class	Trading Symbol	Name of each exchange on which registered
Common Stock, \$0.01 Par Value Per Share	VRTX	The Nasdaq Global Select Market

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

Emerging growth company  $\Box$ 

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.

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

On April 29, 2020, we issued a press release in which we reported our consolidated financial results for the three months ended March 31, 2020. A copy of that press release is attached to this Current Report on Form 8-K as Exhibit 99.1 and is incorporated herein by reference.

The information set forth in Exhibit 99.1 shall not be deemed to be "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liability of that section, and shall not be incorporated by reference into any registration statement or other document filed under the Securities Act of 1933, as amended, or the Exchange Act, except as shall be expressly set forth by specific reference in such filing.

#### Item 9.01. Financial Statements and Exhibits.

#### (d) Exhibits

Exhibit Description of Document

99.1 Press Release, dated April 29, 2020

#### 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.

#### VERTEX PHARMACEUTICALS INCORPORATED (Registrant)

Date: April 29, 2020

/s/ Michael Parini

Michael Parini

Executive Vice President, Chief Administrative, Legal and Business Development Officer

#### Vertex Reports First-Quarter 2020 Financial Results

-Product revenues of \$1.52 billion, a 77% increase compared to Q1 2019-

-Company raises revenue guidance; now expects 2020 CF revenues of \$5.3 to \$5.6 billion-

**BOSTON** -- Vertex Pharmaceuticals Incorporated (Nasdaq: VRTX) today reported consolidated financial results for the first quarter ended March 31, 2020 and revised upward its full-year 2020 financial guidance for total cystic fibrosis (CF) product revenues.

"The COVID-19 pandemic has presented unprecedented challenges to societies, communities and businesses around the world, and while these global challenges will continue for some time to come, I am very proud of how Vertex has responded to ensure that we continue to deliver on our mission for patients, keep our employees safe and achieve our business goals," said Reshma Kewalramani, M.D., Chief Executive Officer and President of Vertex. "Importantly, our business continues to grow, and throughout the first quarter, thousands of patients initiated treatment with our medicines worldwide. The U.S. launch of TRIKAFTA has been remarkable, with the majority of eligible patients having now initiated treatment with this medicine. This strong interest reflects TRIKAFTA's substantial benefits for patients and has resulted in significant increases in revenue to support continue to be differentiated by our focus on serial innovation, investment in transformative medicines aimed at the underlying cause of disease, the breadth of our pipeline and capabilities and our financial strength."

<b>First-Quarter</b>	2020	<b>Financial</b>	<u>Highlights</u>
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	Three Months Ended March 31,				%
		2020		2019	Change
		(in millio	ns, except per	share amount	s)
Product revenues, net	\$	1,515	\$	857	77%
TRIKAFTA	\$	895	\$	—	
SYMDEKO/SYMKEVI	\$	173	\$	320	
ORKAMBI	\$	234	\$	293	
KALYDECO	\$	213	\$	244	
GAAP Operating income	\$	720	\$	277	160%
Non-GAAP Operating income	\$	877	\$	377	133%
GAAP Net income	\$	603	\$	269	124%
Non-GAAP Net income	\$	674	\$	296	128%
GAAP Net income per share - diluted	\$	2.29	\$	1.03	122%
Non-GAAP Net income per share - diluted	\$	2.56	\$	1.14	125%

**Total product revenues** increased 77% compared to the first quarter of 2019, primarily driven by the uptake of TRIKAFTA in the U.S. and the uptake of our medicines outside the U.S. following the completion of key reimbursement agreements in 2019.

**GAAP and Non-GAAP net income** each increased more than 120% compared to the first quarter of 2019, largely driven by the strong growth in total product revenues.

**Cash, cash equivalents and marketable securities** as of March 31, 2020 were \$4.2 billion, an increase of approximately \$400 million compared to \$3.8 billion as of December 31, 2019.

#### First-Quarter 2020 Expenses

	Three Months Ended March 31,			
		2020		2019
		(in	millions)	
Combined GAAP R&D and SG&A expenses	\$	631	\$	487
Combined Non-GAAP R&D and SG&A expenses	\$	477	\$	388
GAAP R&D expense	\$	449	\$	339
Non-GAAP R&D expense	\$	337	\$	273
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GAAP SG&A expense	\$	182	\$	147
Non-GAAP SG&A expense	\$	140	\$	114
GAAP income taxes	\$	55	\$	52
Non-GAAP income taxes	\$	184	\$	81
GAAP effective tax rate		8 %		16 %
Non-GAAP effective tax rate		21 %		21 %

**Combined GAAP and Non-GAAP R&D and SG&A expenses** increased compared to the first quarter of 2019, primarily due to the incremental investment to support the global use of Vertex's medicines and the expansion of Vertex's pipeline in CF and other new disease areas.

**GAAP and Non-GAAP income taxes** increased compared to the first quarter of 2019 primarily due to Vertex's increased operating income. Refer to "Supplemental Income Tax Information" for discussion of the cash versus non-cash components of Vertex's provision for income taxes.

#### Full-Year 2020 Financial Guidance

Vertex today revised upward its guidance for full-year 2020 CF product revenues and reiterated its guidance for GAAP and non-GAAP combined R&D and SG&A expenses and for its non-GAAP effective tax rate, as summarized below:

	 Current FY 2020	Previous FY 2020
TOTAL product revenues	\$ 5.3 to 5.6 billion \$	5.1 to 5.3 billion
Combined GAAP R&D and SG&A expenses	Unchanged \$	2.4 to 2.55 billion
Combined Non-GAAP R&D and SG&A expenses	Unchanged \$	1.95 to 2.0 billion
Non-GAAP effective tax rate	Unchanged	21% to 22%

#### Key Business Highlights:

#### **Approved Medicines:**

#### Supply Chain for Approved CF Medicines

- The global COVID-19 outbreak has not had any impact on the continuity of Vertex's supply chain for its approved medicines.
- Vertex remains highly confident in its ability to continue to supply all of its approved medicines to patients around the world.

#### TRIKAFTA (elexacaftor, tezacaftor and ivacaftor)

- Vertex has seen rapid uptake of TRIKAFTA across all groups of eligible patients following approval of this medicine by the U.S. Food and Drug Administration (FDA) in October 2019. The majority of the approximately 18,000 eligible patients have now initiated treatment with TRIKAFTA.
- In Europe, the Marketing Authorization Application for the elexacaftor, tezacaftor and ivacaftor triple combination in patients with at least one *F508del* mutation ages 12 and older continues to be under review with the European Medicines Agency (EMA).
- Vertex also recently submitted applications for approval of the elexacaftor, tezacaftor and ivacaftor triple combination for patients with at least one *F508del* mutation ages 12 and older in Australia and Switzerland.
- Vertex recently completed enrollment for a Phase 3 study evaluating the use of the elexacaftor, tezacaftor and ivacaftor triple combination regimen in children with CF ages 6 through 11 who have two copies of the *F508del* mutation or who have one *F508del* mutation and one minimal function mutation. Pending data from the study, Vertex plans to submit a supplemental New Drug Application (sNDA) to the U.S. FDA in the second half of 2020 for children ages 6-11 with at least one *F508del* mutation, followed by regulatory submissions in other countries.

#### KALYDECO (ivacaftor)

- Vertex recently completed the submission of an sNDA to the U.S. FDA and Type 2 variation to the EMA for the use of KALYDECO in infants ages four to less than six months. KALYDECO is currently approved for use in infants as young as six months of age in both geographies.
  - 3

#### **Development Pipeline:**

Vertex continues to progress its expanding pipeline of programs in the clinic, which span various diseases, modalities and stages of development. To ensure patient safety and reduce the burden on the healthcare system at a time of critical need, Vertex has temporarily paused or delayed enrollment in certain studies.

#### Alpha-1 Antitrypsin (AAT) Deficiency:

• Vertex has temporarily paused screening and enrollment in the Phase 2 study of VX-814; however, the study remains active and Vertex continues to initiate new clinical trial sites to enable future patient enrollment.

#### Beta Thalassemia and Sickle Cell Disease

Vertex and its partner CRISPR Therapeutics remain on track to provide additional data from the two ongoing Phase 1/2 studies of the investigational CRISPR/Cas9 gene-editing therapy CTX001 in patients with transfusion-dependent beta thalassemia and in patients with severe sickle cell disease in 2020. New data expected in 2020 include initial data from additional patients dosed in each of the Phase 1/2 studies and longer duration follow-up data for the first patients dosed in each study. Screening, enrollment and mobilization in these studies is ongoing; however, no additional patients are scheduled to initiate conditioning or dosing at this time.

#### Focal Segmental Glomerulosclerosis (FSGS):

- Vertex recently initiated a Phase 2 proof-of-concept study of VX-147 in people with FSGS.
- The 13-week open-label Phase 2 study is designed to evaluate the reduction in proteinuria in people with FSGS after treatment with VX-147.

#### Type 1 Diabetes:

• Vertex continues to advance its cell therapy program for the treatment of type 1 diabetes and expects to initiate clinical development in patients in late 2020 or early 2021.

#### **Investments in External Innovation**

- Expanding Capabilities in Genetic Therapies: In April, Vertex entered into a collaboration with Affinia Therapeutics to gain access to a novel library of AAV capsids that will bolster Vertex's ongoing research and development work in genetic therapies. The goal of the collaboration will be to develop genetic therapies for people affected by Duchenne muscular dystrophy (DMD), myotonic dystrophy 1 (DM1) and CF.
- mRNA Therapies for CF: Based on promising preclinical data generated to date, Vertex and Moderna recently extended their research collaboration aimed at the discovery and development of mRNA therapeutics for the treatment of CF.

#### **Non-GAAP Financial Measures**

In this press release, Vertex's financial results and financial guidance are provided in accordance with accounting principles generally accepted in the United States (GAAP) and using certain non-GAAP financial measures. In particular, non-GAAP financial results and guidance exclude from Vertex's pre-tax income (i) stock-based compensation expense, (ii) revenues and expenses related to collaboration agreements, (iii) gains or losses related to the fair value of the company's strategic investments, (iv) acquisition-related costs and (v) other adjustments. The company's non-GAAP financial results also exclude from its provision for income taxes the estimated tax impact related to its non-GAAP adjustments to pre-tax income described above and certain discrete items. These results are provided as a complement to results provided in accordance with GAAP because management believes these non-GAAP financial measures help indicate underlying trends in the company's business, are important in comparing current results with prior period results and provide additional information regarding the company's financial position. Management also uses these non-GAAP financial measures to establish budgets and operational goals that are communicated internally and externally and to manage the company's business and to evaluate its performance. The company adjusts, where appropriate, for both revenues and expenses in order to reflect the company's operations. The company provides guidance regarding product revenues in accordance with GAAP and provides guidance regarding combined research and development and sales, general, and administrative expenses on both a GAAP and non-GAAP basis. The company also provides guidance regarding its anticipated income taxes as a percentage of pre-tax income on a non-GAAP basis. The guidance regarding GAAP research and development expenses and sales, general and administrative expenses does not include estimates associated with any potential future business development activities. A reconciliation of the GAAP financial results to non-GAAP financial results is included in the attached financial information.

### Vertex Pharmaceuticals Incorporated First-Quarter Results **Consolidated Statements of Operations** (in thousands, except per share amounts) (unaudited)

	Three Months Ended March 31,			arch 31,
		2020		2019
Revenues:				
Product revenues, net	\$	1,515,107	\$	857,253
Collaboration and royalty revenues		_		1,182
Total revenues		1,515,107		858,435
Costs and expenses:				
Cost of sales		162,497		95,092
Research and development expenses		448,528		339,490
Sales, general and administrative expenses		182,258		147,045
Change in fair value of contingent consideration		1,600		—
Total costs and expenses		794,883		581,627
Income from operations		720,224		276,808
Interest income		12,576		15,615
Interest expense		(14,136)		(14,868)
Other (expense) income, net (1)		(61,130)		42,610
Income before provision for income taxes		657,534		320,165
Provision for income taxes		54,781		51,534
Net income	\$	602,753	\$	268,631
Net income per common share:				
Basic	\$	2.32	\$	1.05
Diluted	\$	2.29	\$	1.03
Shares used in per share calculations:				
Basic		259,815		255,695
Diluted		263,515		260,175

#### **Reconciliation of GAAP to Non-GAAP Net Income First-Quarter Results**

## (in thousands, except per share amounts) (unaudited)

	Three Months Ended March 31,			arch 31,
		2020		2019
GAAP net income	\$	602,753	\$	268,631
Stock-based compensation expense		115,706		93,791
Decrease (increase) in fair value of strategic investments (1)		44,870		(43,551)
Increase in fair value of contingent consideration (2)		1,600		—
Collaborative revenues and expenses (3)		36,250		6,351
Acquisition-related costs (4)		2,883		
Total non-GAAP adjustments to pre-tax income		201,309		56,591
Tax adjustments (5)		(129,608)		(29,392)
Non-GAAP net income	\$	674,454	\$	295,830
Net income per diluted common share:				
GAAP	\$	2.29	\$	1.03
Non-GAAP	\$	2.56	\$	1.14
Shares used in diluted per share calculations:				
GAAP and Non-GAAP		263,515		260,175

#### **Reconciliation of GAAP to Non-GAAP Revenues and Expenses**

**First-Quarter Results** 

(in thousands) (unaudited)

	Three Months Ended March 31,			larch 31,
		2020		2019
GAAP total revenues	\$	1,515,107	\$	858,435
Collaborative revenues		—		(141)
Non-GAAP total revenues	\$	1,515,107	\$	858,294
		Three Months E	nded M	larch 31,
		2020		2019
GAAP cost of sales	\$	162,497	\$	95,092
Stock-based compensation expense		(1,361)		(1,338)
Non-GAAP cost of sales	\$	161,136	\$	93,754
GAAP research and development expenses	\$	448,528	\$	339,490
Stock-based compensation expense		(72,687)		(59,715)
Collaborative expenses (3)		(36,250)		(6,492)
Acquisition-related costs (4)		(2,678)		—
Non-GAAP research and development expenses	\$	336,913	\$	273,283
GAAP sales, general and administrative expenses	\$	182,258	\$	147,045
Stock-based compensation expense		(41,658)		(32,738)
Acquisition-related costs (4)		(205)		—
Non-GAAP sales, general and administrative expenses	\$	140,395	\$	114,307
Combined non-GAAP R&D and SG&A expenses	\$	477,308	\$	387,590
	Three Months Ended March 31,			
		2020		2019
GAAP other (expense) income, net	\$	(61,130)	\$	42,610
Decrease (increase) in fair value of strategic investments (1)		44,870		(43,551)
Non-GAAP other expense, net	\$	(16,260)	\$	(941)
GAAP provision for income taxes	\$	54,781	\$	51,534
Tax adjustments (5)		129,608		29,392

Non-GAAP provision for income taxes (6)

8

\$

184,389 \$

80,926

#### **Condensed Consolidated Balance Sheets**

(in thousands)

(unaudited)

	March 31, 2020		Dece	December 31, 2019		
Assets			· · · · · · · · · · · · · · · · · · ·			
Cash, cash equivalents and marketable securities	\$	4,190,396	\$	3,808,294		
Accounts receivable, net		845,269		633,518		
Inventories		187,087		167,502		
Property and equipment, net		736,303		745,080		
Goodwill and intangible assets		1,402,158		1,402,158		
Deferred tax assets		1,147,705		1,190,815		
Other assets		384,283		371,098		
Total assets	\$	8,893,201	\$	8,318,465		
Liabilities and Shareholders' Equity						
Accounts payable and accrued expenses	\$	1,358,974	\$	1,204,522		
Finance lease liabilities		572,916		577,371		
Contingent consideration		178,100		176,500		
Other liabilities		321,557		274,828		
Shareholders' equity		6,461,654		6,085,244		
Total liabilities and shareholders' equity	\$	8,893,201	\$	8,318,465		
Common shares outstanding		259,079		258,993		

# Supplemental Income Tax Information (in thousands, except percentages) (unaudited)

	Three Months	Ended M	larch 31,
	 2020		2019
Components of provision for income taxes related to:			
Cash paid or accrued for income taxes	\$ 9,370	\$	4,778
Provision for income taxes offset by net operating losses	45,411		46,756
GAAP provision for income taxes (6)	\$ 54,781	\$	51,534
Cash paid or accrued for income taxes	\$ 9,370	\$	4,778
Tax adjustments (5)	129,608		29,392
Provision for income taxes offset by net operating losses	45,411		46,756
Non-GAAP provision for income taxes (6)	\$ 184,389	\$	80,926
Effective tax rate reconciliation:			

GAAP effective tax rate	8%	16%
Impact of GAAP to Non-GAAP adjustments	13%	5%
Non-GAAP effective tax rate	21%	21%

#### **Notes and Explanations**

1: "Other income, net" includes gains and losses related to changes in the fair value of the company's strategic investments.

**2:** During the three months ended March 31, 2020, the increase in the fair value of the contingent consideration relates to potential payments to Exonics' former equity holders.

**3:** "Collaborative revenues and expenses" in the three months ended March 31, 2020 and 2019 primarily related to collaborative milestone payments.

**4:** "Acquisition-related costs" in the three months ended March 31, 2020 related to costs associated with the company's acquisitions of Semma and Exonics. There were no comparable amounts during the three months ended March 31, 2019.

5: In the three months ended March 31, 2020 and 2019, "Tax adjustments" primarily related to the estimated income taxes related to non-GAAP adjustments to pre-tax income including (i) stock-based compensation (including an adjustment for excess tax benefits related to stock-based compensation), (ii) increases or decreases in the fair value of the company's strategic investments and (iii) collaborative payments. In the three months ended March 31, 2020, "Tax adjustments" also included a non-recurring discrete benefit to the company's provision for income taxes of approximately \$50 million that the company excluded from its Non-GAAP measures.

**6:** The company records a provision for income taxes on its pre-tax income using an effective tax rate approximating statutory rates. The provision includes a significant non-cash charge due to the company's ability to offset its pre-tax income against previously benefited net operating losses. The company expects a portion of its tax provision to represent a non-cash expense until its net operating losses have been fully utilized. As of December 31, 2019, the company's federal net operating losses and credits that were available to offset future pre-tax income were approximately \$3.5 billion.

#### **About Vertex**

Vertex is a global biotechnology company that invests in scientific innovation to create transformative medicines for people with serious diseases. The company has multiple approved medicines that treat the underlying cause of cystic fibrosis (CF) — a rare, life-threatening genetic disease — and has several ongoing clinical and research programs in CF. Beyond CF, Vertex has a robust pipeline of investigational small molecule medicines in other serious diseases where it has deep insight into causal human biology, including pain, alpha-1 antitrypsin deficiency and APOL1-mediated kidney diseases. In addition, Vertex has a rapidly expanding pipeline of genetic and cell therapies for diseases such as sickle cell disease, beta thalassemia, Duchenne muscular dystrophy and type 1 diabetes mellitus.

Founded in 1989 in Cambridge, Mass., Vertex's global headquarters is now located in Boston's Innovation District and its international headquarters is in London, UK. Additionally, the company has research and development sites and commercial offices in North America, Europe, Australia and Latin America. Vertex is consistently recognized as one of the industry's top places to work, including 10 consecutive years on Science magazine's Top Employers list and top five on the 2019 Best Employers for Diversity list by Forbes. For company updates and to learn more about Vertex's history of innovation, visit www.vrtx.com or follow us on Facebook, Twitter, LinkedIn, YouTube and Instagram.

#### **Special Note Regarding Forward-Looking Statements**

This press release contains forward-looking statements as defined in the Private Securities Litigation Reform Act of 1995, including, without limitation, Dr. Kewalramani's statements in this press release, the information provided regarding future financial performance, the section captioned "Full-Year 2020 Financial Guidance" and statements regarding (i) regulatory filings, (ii) the development plan and timelines for the company's drug candidates and (iii) the company's expectations regarding the effects COVID-19 will have on its business and operations. While Vertex believes the forward-looking statements contained in this press release are accurate, these forward-looking statements represent the company's beliefs only as of the date of this press release and there are a number of risks and uncertainties that could cause actual events or results to differ materially from those expressed or implied by such forward-looking statements. Those risks and uncertainties include, among other things, that the company's expectations regarding its 2020 CF net product revenues, expenses and effective tax rates may be incorrect (including because one or more of the company's assumptions underlying its expectations may not be realized), that COVID-19 may have different or more significant impacts on the company's business or operations than the company currently expects, that data from the company's development programs may not support registration or further development of its potential medicines due to safety, efficacy or other reasons, and other risks listed under Risk Factors in Vertex's annual report and subsequent quarterly reports filed with the Securities and Exchange Commission and available through the company's website at www.vrtx.com.

#### **Conference Call and Webcast**

The company will host a conference call and webcast today at 5:00 p.m. ET. To access the call, please dial (866) 501-1537 (U.S.) or +1 (720) 545-0001 (International). The conference call will be webcast live and a link to the webcast can be accessed through Vertex's website at www.vrtx.com in the "Investors" section under "Events and Presentations." To ensure a timely connection, it is recommended that users register at least 15 minutes prior to the scheduled webcast. An archived webcast will be available on the company's website.

#### (VRTX-E)

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