

**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): July 1, 2022

Intercept Pharmaceuticals, Inc.
(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation)	001-35668 (Commission File Number)	22-3868459 (IRS Employer Identification No.)
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305 Madison Avenue, Morristown, NJ 07960
(Address of principal executive offices) (Zip Code)

Registrant's telephone number, including area code: (646) 747-1000

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
Common Stock, par value \$0.001 per share	ICPT	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

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 1.01 Entry into a Material Definitive Agreement.

On July 1, 2022, Intercept Pharmaceuticals, Inc. (“ICPT Inc.”) entered into an amendment of its Share Purchase Agreement with Mercury Pharma Group Limited (“Mercury”), an affiliate of Advanz Pharma Corp. Limited (together with its affiliates, “Advanz”). In the amendment, ICPT Inc. agreed to provide for continuity of cash collateral that is securing letters of credit issued for the benefit of former affiliates of ICPT Inc. that have now been sold to Advanz. Advanz agreed to work to replace the letters of credit reasonably promptly. Advanz will pay for the expenses of ICPT Inc. in maintaining the letters of credit. Advanz will also pay interest to ICPT Inc. at a rate of 5% for the first three months following the sale closing date of July 1, 2022, and 6.5% for the next three months, on the balance of cash collateral required to be posted by ICPT Inc. to the letter of credit bank to continue to support the remaining letters of credit. After six months, Advanz will deliver cash collateral to ICPT Inc. in the amount of the remaining cash collateral posted by ICPT Inc., which will be returned following the release by the bank of cash collateral to ICPT Inc.

On July 1, 2022, ICPT Inc. and Mercury entered into an amendment of their Safety Data Exchange Agreement. The original agreement specified that ICPT Inc. would retain responsibility for performance of pharmacovigilance obligations ex-U.S. until ex-U.S. marketing authorizations have been transferred to Mercury or a designated third party, not longer than a period of three months. In the amendment, this period was changed to six months.

The descriptions above of these amendments do not purport to be complete, and are qualified in their entirety by reference to the text of the amendments, which will be filed as exhibits to the Company’s Quarterly Report on Form 10-Q for the quarter ending June 30, 2022.

Item 2.01 Completion of Acquisition or Disposition of Assets.

As previously disclosed, on May 5, 2022, ICPT Inc. and affiliates entered into a series of agreements selling to Advanz their ex-U.S. commercial operations, and sublicensing the right to commercialize Ocaliva outside of the United States. On July 1, 2022, the sale closed. Consideration under the agreements totaled \$405 million up front, plus a \$45 million earnout, payable upon Advanz’s receipt of extensions of orphan drug exclusivity in Europe. The consideration was comprised of:

- (1) \$364.5 million (plus the earnout) under the Sublicense Agreement, with ICPT’s wholly owned subsidiary Intercept Pharma Europe Ltd. (“IPEL”) sublicensing to Mercury the right to commercialize Ocaliva outside of the United States.
 - (2) \$38.5 million under the Share Purchase Agreement, with ICPT Inc. selling to Mercury the shares of its foreign subsidiaries (excluding IPEL).
 - (3) \$1 million under the Business Transfer Agreement, with IPEL transferring certain business assets to Advanz Pharma Services (UK) Limited.
 - (4) \$1 million under the Master Trademark Assignment Agreement, with ICPT Inc. and its wholly owned subsidiary RXF Technologies, Inc. assigning to Mercury certain trademarks.
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ICPT Inc. and IPEL together received aggregate cash consideration of \$366.5 million. The consideration under the Share Purchase Agreement will be settled in connection with the completion statements, which will also include adjustments including for cash, working capital, and assumed liabilities. As previously disclosed, the transaction also included a Supply and Manufacture Agreement, a Safety Data Exchange Agreement, and a Transitional Services Agreement, which did not include upfront consideration.

Item 9.01 Financial Statements and Exhibits.

(b)(1) *Pro forma financial information.*

Attached as Exhibit 99.1 and incorporated by reference is the pro forma financial information required by Article 11 of Regulation S-X.

(d) *Exhibits.*

<u>Exhibit Number</u>	<u>Description</u>
99.1	Pro Forma Financial Information
104	Cover Page Interactive Data File (embedded as Inline XBRL document)

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.

INTERCEPT PHARMACEUTICALS, INC.

By: /s/ Rocco Venezia

Name: Rocco Venezia

Title: Chief Accounting Officer

Date: July 8, 2022

UNAUDITED PRO FORMA FINANCIAL INFORMATION*(In millions, except share information)*

As previously disclosed, on May 5, 2022, Intercept Pharmaceuticals, Inc., a Delaware corporation (“ICPT Inc.”), and its wholly owned subsidiary Intercept Pharma Europe Ltd., a UK private limited company (“IPEL” and, together with ICPT Inc. and affiliates, the “Company”), entered into a series of agreements to sell the Company’s ex-U.S. commercial operations, and sublicense the right to commercialize Ocaliva® (obeticholic acid) outside of the United States, to Advanz Pharma and its affiliates (collectively, “Advanz”).

Total consideration to the Company under the agreements is \$405 million up front (subject to working capital, closing costs, France reimbursement liability, and other adjustments), consisting of \$38.5 million under the Share Purchase Agreement (“SPA”) between ICPT Inc. as seller and Mercury Pharma as purchaser, pursuant to which ICPT Inc. will sell the shares of its foreign subsidiaries (excluding IPEL) to Mercury Pharma, \$364.5 million under the Sublicense Agreement (the “Sublicense Agreement”) among IPEL, ICPT Inc., and Mercury Pharma, \$1 million under the Business Transfer Agreement (“BTA”) between IPEL and Advanz Services, and \$1 million under the Master Trademark Assignment Agreement among ICPT Inc., its wholly owned subsidiary RXF Technologies, Inc., and Mercury Pharma. The Sublicense Agreement also includes a \$45 million earnout, payable upon Advanz’s receipt of extensions of orphan drug exclusivity in Europe. These transactions are referred to collectively as the “Disposition Transactions”.

On July 1, 2022, upon the terms and subject to the conditions set forth in the SPA and the other transaction documents, the Disposition Transactions contemplated thereby were consummated.

The following unaudited pro forma condensed consolidated financial statements are intended to show how the Disposition Transactions might have affected the historical financial statements of the Company had the transaction been completed at an earlier time as indicated herein. The unaudited pro forma condensed consolidated financial statements have been prepared in accordance with Article 11 of Regulation S-X, were derived from the Company’s historical condensed consolidated (for interim periods) or consolidated (for annual periods) financial statements, and are being presented to give effect to the Disposition Transactions. The historical results of the Company will present the international operations disposed in the Disposition Transactions on a discontinued operations basis in accordance with ASC 205, *Discontinued Operations*, beginning in the second quarter of 2022. The unaudited pro forma condensed consolidated financial statements should be read with:

- i. The accompanying notes to the unaudited pro forma condensed consolidated financial statements
- ii. The Current Report on Form 8-K filed with the Securities & Exchange Commission (the “SEC”) on July 8, 2022, reporting the Disposition Transactions
- iii. The audited consolidated financial statements of the Company and its subsidiaries and the accompanying notes included in the Company’s Annual Report on Form 10-K filed with the SEC on March 2, 2022.
- iv. The unaudited interim historical financial statements of the Company and its subsidiaries, the accompanying notes and “Management’s Discussion and Analysis of Financial Condition and Results of Operations” included in the Company’s Quarterly Report on Form 10-Q for the three months ended March 31, 2022 filed with the SEC on May 6, 2022.

The unaudited pro forma condensed consolidated financial statements are based on available information and assumptions that the Company’s management believes are reasonable as of the date of this filing. The unaudited pro forma condensed consolidated statements of operations for the years ended December 31, 2021, 2020 and 2019, and the three months ended March 31, 2022, present the Company’s results as if the transaction had occurred on January 1, 2019, the beginning of the earliest period presented. The unaudited pro forma condensed consolidated balance sheet as of March 31, 2022 reflects the Company’s assets, liabilities and equity as if the transaction had occurred on March 31, 2022.

The unaudited pro forma condensed consolidated financial statements are not necessarily indicative of what the Company's actual results of operations would have been had the transaction been consummated on the dates assumed nor does it purport to represent the Company's results of operations or financial condition for any future period. Actual amounts could differ materially from these estimates.

The transaction accounting adjustments to reflect the Disposition Transactions in the unaudited pro forma condensed consolidated financial statements include:

- The sale of the assets and liabilities pursuant to the Disposition Transactions agreements.
- Estimated impact of the cash proceeds received in connection with the Disposition Transactions, net of closing costs and income taxes.

There were no autonomous entity adjustments included in the pro form financial information because the Company currently operates, and will continue to operate, as an independent, standalone entity after the completion of the Disposition Transactions. The pro forma financial statements do not include management adjustments to reflect synergies or dis-synergies because we do not believe presenting such adjustments would enhance an understanding of the pro forma effects of the transaction.

On May 5, 2022, the parties entered into a Transition Services Agreement ("TSA") for the Company and Advanz to provide certain transition services for continuity purposes. The TSA details the services to be provided, the terms over which the services will be provided, and the compensation to be paid for providing those services. The unaudited pro forma condensed consolidated financial statements do not include any compensation or costs related to the TSA as they are not material. Similarly, on May 5, 2022, the parties entered into a Supply and Manufacture Agreement and a Safety Data Exchange Agreement. We do not consider the amounts under those agreements to be material for the periods presented.

INTERCEPT PHARMACEUTICALS, INC.
Pro Forma Condensed Consolidated Balance Sheet
As of March 31, 2022
(In thousands, except per share amounts)
(Unaudited)

	<u>Historical Financial Statements as Reported</u>	<u>Disposition Transactions (b)</u>	<u>Pro Forma</u>
Assets			
Current assets:			
Cash and cash equivalents	\$ 48,320	\$ 341,819(a)	\$ 390,139
Restricted cash	11,153	(1,549)	9,604
Investment debt securities, available for sale	347,382	—	347,382
Accounts receivable, net of allowance for credit losses of \$255	46,144	(20,740)	25,404
Prepaid expenses and other current assets	24,354	34,553(f)	58,907
Total current assets	477,353	354,083	831,436
Fixed assets, net	3,059	(61)	2,998
Inventory	8,257	(639)	7,618
Security deposits	6,926	(2,285)	4,641
Other assets	7,762	(2,001)	5,761
Total assets	<u>\$ 503,357</u>	<u>\$ 349,097</u>	<u>\$ 852,454</u>
Liabilities and Stockholders' Deficit			
Current liabilities:			
Accounts payable, accrued expenses and other liabilities	\$ 147,060	\$ (54,295)(c)	\$ 92,765
Short-term interest payable	3,976	—	3,976
Total current liabilities	151,036	(54,295)	96,741
Long-term liabilities:			
Long-term debt	716,885	—	716,885
Long-term other liabilities	7,197	(1,084)	6,113
Total liabilities	<u>\$ 875,118</u>	<u>\$ (55,379)</u>	<u>\$ 819,739</u>
Stockholders' deficit			
Common stock par value \$0.001 per share, 90,000,000 shares authorized, 29,708,846 shares issued and outstanding	30	—	30
Additional paid in capital	2,007,684	—	2,007,684
Accumulated other comprehensive loss, net	(3,487)	1,589	(1,898)
Accumulated deficit	(2,375,988)	402,887(d)	(1,973,101)
Total stockholders' deficit	<u>(371,761)</u>	<u>404,476</u>	<u>32,715</u>
Total liabilities and stockholders' deficit	<u>\$ 503,357</u>	<u>\$ 349,097</u>	<u>\$ 852,454</u>

INTERCEPT PHARMACEUTICALS, INC.
Pro Forma Condensed Consolidated Statement of Operations
For the three months ended March 31, 2022
(In thousands, except per share amounts)
(Unaudited)

	<u>Historical Financial Statements as Reported</u>	<u>Disposition Transactions (e)</u>	<u>Pro Forma</u>
Revenue:			
Product revenue, net	\$ 88,582	\$ (29,436)	\$ 59,146
Total revenue	<u>88,582</u>	<u>(29,436)</u>	<u>59,146</u>
Operating expenses:			
Cost of sales	758	(368)	390
Selling, general and administrative	50,007	(14,909)	35,098
Research and development	48,089	(196)	47,893
Restructuring	—	—	—
Total operating expenses	<u>98,854</u>	<u>(15,473)</u>	<u>83,381</u>
Operating loss	<u>(10,272)</u>	<u>(13,963)</u>	<u>(24,235)</u>
Other (expense) income:			
Interest expense	(6,673)	—	(6,673)
Other (expense) income, net	(339)	312	(27)
Total other (expense), net	<u>(7,012)</u>	<u>312</u>	<u>(6,700)</u>
Net loss	<u>\$ (17,284)</u>	<u>\$ (13,651)</u>	<u>\$ (30,935)</u>
Net loss per common and potential common share:			
Basic and diluted	\$ (0.58)		\$ (1.04)
Weighted average common and potential common shares outstanding:			
Basic and diluted	29,696		29,696

INTERCEPT PHARMACEUTICALS, INC.
Pro Forma Condensed Consolidated Statement of Operations
For the year ended December 31, 2021
(In thousands, except per share amounts)
(Unaudited)

	<u>Historical Financial Statements as Reported</u>	<u>Disposition Transactions (e)</u>	<u>Pro Forma</u>
Revenue:			
Product revenue, net	\$ 363,468	\$ (102,718)	\$ 260,750
Total revenue	<u>363,468</u>	<u>(102,718)</u>	<u>260,750</u>
Operating expenses:			
Cost of sales	3,100	(1,585)	1,515
Selling, general and administrative	230,855	(48,068)	182,787
Research and development	185,272	(876)	184,396
Restructuring	(86)	—	(86)
Total operating expenses	<u>419,141</u>	<u>(50,529)</u>	<u>368,612</u>
Operating loss	<u>(55,673)</u>	<u>(52,189)</u>	<u>(107,862)</u>
Other (expense) income:			
Interest expense	(54,419)	—	(54,419)
Gain on extinguishment of debt	16,511	—	16,511
Other (expense) income, net	2,155	(1,388)	767
Total other (expense), net	<u>(35,753)</u>	<u>(1,388)</u>	<u>(37,141)</u>
Net loss	<u>\$ (91,426)</u>	<u>\$ (53,577)</u>	<u>\$ (145,003)</u>
Net loss per common and potential common share:			
Basic and diluted	\$ (2.87)		\$ (4.55)
Weighted average common and potential common shares outstanding:			
Basic and diluted	31,894		31,894

INTERCEPT PHARMACEUTICALS, INC.
Pro Forma Condensed Consolidated Statement of Operations
For the year ended December 31, 2020
(In thousands, except per share amounts)
(Unaudited)

	<u>Historical Financial Statements as Reported</u>	<u>Disposition Transactions (e)</u>	<u>Pro Forma</u>
Revenue:			
Product revenue, net	\$ 312,690	\$ (78,720)	\$ 233,970
Total revenue	<u>312,690</u>	<u>(78,720)</u>	<u>233,970</u>
Operating expenses:			
Cost of sales	5,322	(1,340)	3,982
Selling, general and administrative	332,493	(62,380)	270,113
Research and development	191,485	(595)	190,890
Restructuring	14,630	—	14,630
Total operating expenses	<u>543,930</u>	<u>(64,315)</u>	<u>479,615</u>
Operating loss	<u>(231,240)</u>	<u>(14,405)</u>	<u>(245,645)</u>
Other (expense) income:			
Interest expense	(48,054)	—	(48,054)
Other (expense) income, net	4,414	2,187	6,601
Total other (expense), net	<u>(43,640)</u>	<u>2,187</u>	<u>(41,453)</u>
Net loss	<u>\$ (274,880)</u>	<u>\$ (12,218)</u>	<u>\$ (287,098)</u>
Net loss per common and potential common share:			
Basic and diluted	\$ (8.34)		\$ (8.71)
Weighted average common and potential common shares outstanding:			
Basic and diluted	32,970		32,970

INTERCEPT PHARMACEUTICALS, INC.
Pro Forma Condensed Consolidated Statement of Operations
For the year ended December 31, 2019
(In thousands, except per share amounts)
(Unaudited)

	<u>Historical Financial Statements as Reported</u>	<u>Disposition Transactions (e)</u>	<u>Pro Forma</u>
Revenue:			
Product revenue, net	\$ 249,570	\$ (62,134)	\$ 187,436
Licensing revenue	2,432	—	2,432
Total revenue	<u>252,002</u>	<u>(62,134)</u>	<u>189,868</u>
Operating expenses:			
Cost of sales	4,212	(1,049)	3,163
Selling, general and administrative	317,418	(66,354)	251,064
Research and development	242,799	(283)	242,516
Total operating expenses	<u>564,429</u>	<u>(67,686)</u>	<u>496,743</u>
Operating loss	<u>(312,427)</u>	<u>5,552</u>	<u>(306,875)</u>
Other (expense) income:			
Interest expense	(41,144)	—	(41,144)
Other (expense) income, net	8,890	3,966	12,856
Total other (expense), net	<u>(32,254)</u>	<u>3,966</u>	<u>(28,288)</u>
Net loss	<u>\$ (344,681)</u>	<u>\$ 9,518</u>	<u>\$ (335,163)</u>
Net loss per common and potential common share:			
Basic and diluted	\$ (10.89)		\$ (10.59)
Weighted average common and potential common shares outstanding:			
Basic and diluted	31,654		31,654

Notes to Pro forma Condensed Consolidated Financial Statements

The unaudited pro forma condensed consolidated financial statements reflect the following transaction accounting adjustments:

- a) Represents the estimated cash proceeds received, net of closing costs and taxes related to the gain on the Disposition Transactions, based on total consideration of \$405 million. Actual cash proceeds received upon closing prior to any estimated closing costs and taxes were \$366.5 million, based on the timing of payment under the SPA, which includes a post-closing mechanism for settling the completion statements, including the \$38.5 million consideration under the SPA, and adjustments for working capital, France reimbursement liability, etc. The cash proceeds do not reflect the \$45 million due contingent upon receipt of extensions of orphan drug exclusivity from the European Medicines Agency (“EMA”) and Medicines and Healthcare products Regulatory Agency (“MHRA”), which if realized, would result in an incremental gain.
- i) Represents total consideration from the transaction of \$366.5 million, less (a) estimated closing costs of \$9.5 million that are likely to be incurred as part of the consummation of the transaction and (b) the expected tax effects on the estimated U.S. federal, U.S. state and foreign income taxes paid of \$15.2 million related to the gain on the transaction. The estimated expected tax effects are calculated based on the amount of the taxable gain considering the use of historical net operation losses and other carryforwards in place to reduce taxable income, and using the applicable income tax rates in the respective jurisdictions. The estimates, including the jurisdiction income tax effects, are subject to change and actual amounts may differ from the results reported herein. The estimated gain is reflected in the unaudited pro forma condensed consolidated balance sheet within accumulated deficit. The estimated gain is not reflected in the unaudited pro forma condensed consolidated statement of operations as there is no continuing impact of the gain on the Company’s operating results.
- b) Adjustments represent the elimination of assets and liabilities attributable to the Disposition Transactions and the disposed operations.
- c) Adjustment includes the transfer of the payback liability of \$42.0 million USD currently being negotiated between the Company and the French government.
- d) Represents the estimated gain on sale (in millions) calculated as follows:

Total consideration	\$	405.0
Estimated closing costs		9.5
Total consideration (net of estimated closing costs)		395.5
Company’s carrying value in disposed operations		24.2
Release of accumulated other comprehensive loss		(1.6)
Pro forma gain before income taxes	\$	418.1
(Provision) for income taxes		(15.2)
Pro forma net gain on disposition of operations	\$	402.9

The pro forma net gain of \$402.9 million is reflected as an adjustment to retained deficit. This amount is based on historical information as of March 31, 2022 for the Company’s carrying value of the disposed operations and the related amount of accumulated other comprehensive loss. The actual net gain will be based on the Company’s carrying value in the disposed operations as of July 1, 2022 and may differ materially from the information presented.

- e) Adjustments represent the elimination of revenues and costs and expenses attributable to the Disposition Transactions and the disposed operations.
- f) Adjustment includes consideration of \$38.5 million to be settled post-closing.
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