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

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**FORM 10-Q**

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(Mark One)

**QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the quarterly period ended June 30, 2022

or

**TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the transition period from \_\_\_\_\_ to \_\_\_\_\_

Commission File Number: 001-36274

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**INTRA-CELLULAR THERAPIES, INC.**

(Exact name of registrant as specified in its charter)

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**Delaware**  
(State or other jurisdiction of  
incorporation or organization)

**430 East 29th Street**  
**New York, New York**  
(Address of principal executive offices)

**36-4742850**  
(I.R.S. Employer  
Identification No.)

**10016**  
(Zip Code)

**(646) 440-9333**

(Registrant's telephone number, including area code)

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

<u>Title of each class</u>	<u>Trading Symbol(s)</u>	<u>Name of each exchange on which registered</u>
Common Stock	ITCI	The Nasdaq Global Select Market

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Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. Yes  No

Indicate by check mark whether the registrant has submitted electronically every Interactive Data File required to be submitted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit such files). Yes  No

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, a smaller reporting company, or an emerging growth company. See the definitions of “large accelerated filer,” “accelerated filer,” “smaller reporting company,” and “emerging growth company” in Rule 12b-2 of the Exchange Act.

Large accelerated filer

Accelerated filer

Non-accelerated filer

Smaller reporting company

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.

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). Yes  No

As of August 5, 2022, the registrant had 94,401,745 shares of common stock outstanding.

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In this Quarterly Report on Form 10-Q, the terms "we," "us," "our," and the "Company" mean Intra-Cellular Therapies, Inc. and our subsidiaries. "ITI" refers to our wholly-owned subsidiary ITI, Inc. and "ITI Limited" refers to our wholly-owned subsidiary ITI Limited.

**PART I: FINANCIAL INFORMATION****Item 1. FINANCIAL STATEMENTS**

Intra-Cellular Therapies, Inc. and Subsidiaries

Condensed Consolidated Balance Sheets (in thousands except share and per share amounts)

	June 30, 2022 (Unaudited)	December 31, 2021
<b>Assets</b>		
Current assets:		
Cash and cash equivalents	\$ 77,235	\$ 92,365
Investment securities, available-for-sale	600,594	319,968
Restricted cash	1,400	1,400
Accounts receivable, net	46,976	20,156
Inventory	25,022	7,948
Prepaid expenses and other current assets	37,979	25,444
Total current assets	789,206	467,281
Property and equipment, net	2,137	1,791
Right of use assets, net	20,477	20,764
Other assets	86	86
Total assets	<u>\$ 811,906</u>	<u>\$ 489,922</u>
<b>Liabilities and stockholders' equity</b>		
Current liabilities:		
Accounts payable	\$ 10,749	\$ 8,691
Accrued and other current liabilities	17,274	11,073
Accrued customer programs	16,483	5,964
Accrued employee benefits	18,385	20,897
Lease liabilities, short-term	7,743	6,732
Total current liabilities	70,634	53,357
Lease liabilities	19,075	18,675
Total liabilities	89,709	72,032
Stockholders' equity:		
Common stock, \$0.0001 par value: 175,000,000 shares authorized at June 30, 2022 and December 31, 2021; 94,367,233 and 81,886,965 shares issued and outstanding at June 30, 2022 and December 31, 2021, respectively	9	8
Additional paid-in capital	2,106,942	1,639,476
Accumulated deficit	(1,379,952)	(1,221,230)
Accumulated comprehensive loss	(4,802)	(364)
Total stockholders' equity	722,197	417,890
Total liabilities and stockholders' equity	<u>\$ 811,906</u>	<u>\$ 489,922</u>

*See accompanying notes to these condensed consolidated financial statements.*

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## Intra-Cellular Therapies, Inc. and Subsidiaries

## Condensed Consolidated Statements of Operations (in thousands except share and per share amounts) (Unaudited)

	<u>Three Months Ended June 30,</u>		<u>Six Months Ended June 30,</u>	
	<u>2022</u>	<u>2021</u>	<u>2022</u>	<u>2021</u>
Revenues				
Product sales, net	\$ 55,074	\$ 19,007	\$ 89,829	\$ 34,586
Grant revenue	505	1,040	746	1,339
Total revenues	55,579	20,047	90,575	35,925
Operating expenses:				
Cost of product sales	4,650	2,040	7,805	3,495
Selling, general and administrative	100,316	69,851	175,776	122,435
Research and development	38,536	17,297	67,579	32,355
Total operating expenses	143,502	89,188	251,160	158,285
Loss from operations	(87,923)	(69,141)	(160,585)	(122,360)
Interest income	1,320	421	1,868	905
Loss before provision for income taxes	(86,603)	(68,720)	(158,717)	(121,455)
Income tax expense	—	24	5	29
Net loss	\$ (86,603)	\$ (68,744)	\$ (158,722)	\$ (121,484)
Net loss per common share:				
Basic & Diluted	\$ (0.92)	\$ (0.85)	\$ (1.70)	\$ (1.50)
Weighted average number of common shares:				
Basic & Diluted	94,285,117	81,229,788	93,449,424	81,088,900

See accompanying notes to these condensed consolidated financial statements.

## Intra-Cellular Therapies, Inc. and Subsidiaries

## Condensed Consolidated Statements of Comprehensive Loss (in thousands) (Unaudited)

	Three Months Ended June 30,		Six Months Ended June 30,	
	2022	2021	2022	2021
Net loss	<b>\$ (86,603)</b>	\$ (68,744)	<b>\$ (158,722)</b>	\$ (121,484)
Other comprehensive income:				
Unrealized (loss) on investment securities	<b>(1,474)</b>	(70)	<b>(4,438)</b>	(304)
Comprehensive loss	<b><u>\$ (88,077)</u></b>	<u>\$ (68,814)</u>	<b><u>\$ (163,160)</u></b>	<u>\$ (121,788)</u>

See accompanying notes to these condensed consolidated financial statements.

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## Intra-Cellular Therapies, Inc. and Subsidiaries

## Condensed Consolidated Statements of Stockholders' Equity (in thousands except share and per share amounts) (Unaudited)

	Common Stock		Additional Paid-in Capital	Accumulated Deficit	Accumulated Comprehensive (Loss) Income	Total Stockholders' Equity
	Shares	Amount				
Balance at March 31, 2021	81,133,849	\$ 8	\$ 1,601,739	\$ (989,844)	\$ 247	\$ 612,150
Exercise of stock options and issuances of restricted stock	176,757	—	1,582	—	—	1,582
Stock issued for services	1,272	—	52	—	—	52
Share-based compensation	—	—	8,617	—	—	8,617
Net loss	—	—	—	(68,744)	—	(68,744)
Other comprehensive loss	—	—	—	—	(70)	(70)
Balance at June 30, 2021	81,311,878	\$ 8	\$ 1,611,990	\$(1,058,588)	\$ 177	\$ 553,587
	Common Stock		Additional Paid-in Capital	Accumulated Deficit	Accumulated Comprehensive (Loss) Income	Total Stockholders' Equity
	Shares	Amount				
Balance at December 31, 2020	80,463,089	\$ 8	\$ 1,593,475	\$ (937,104)	\$ 481	\$ 656,860
Exercise of stock options and issuances of restricted stock	845,939	—	3,014	—	—	3,014
Stock issued for services	2,850	—	106	—	—	106
Share-based compensation	—	—	15,395	—	—	15,395
Net loss	—	—	—	(121,484)	—	(121,484)
Other comprehensive loss	—	—	—	—	(304)	(304)
Balance at June 30, 2021	81,311,878	\$ 8	\$ 1,611,990	\$(1,058,588)	\$ 177	\$ 553,587
	Common Stock		Additional Paid-in Capital	Accumulated Deficit	Accumulated Comprehensive Loss	Total Stockholders' Equity
	Shares	Amount				
Balance at March 31, 2022	94,020,425	\$ 9	\$ 2,089,418	\$ (1,293,349)	\$ (3,328)	\$ 792,750
Exercise of stock options and issuances of restricted stock	346,411	—	5,504	—	—	5,504
Stock issued for services	397	—	22	—	—	22
Share-based compensation	—	—	11,998	—	—	11,998
Net loss	—	—	—	(86,603)	—	(86,603)
Other comprehensive loss	—	—	—	—	(1,474)	(1,474)
Balance at June 30, 2022	94,367,233	\$ 9	\$ 2,106,942	\$(1,379,952)	\$ (4,802)	\$ 722,197
	Common Stock		Additional Paid-in Capital	Accumulated Deficit	Accumulated Comprehensive Loss	Total Stockholders' Equity
	Shares	Amount				
Balance at December 31, 2021	81,886,965	\$ 8	\$ 1,639,476	\$ (1,221,230)	\$ (364)	\$ 417,890
Common shares issued January 7, 2022	10,952,381	1	433,724	—	—	433,725
Exercise of stock options and issuances of restricted stock	1,527,107	—	13,593	—	—	13,593
Stock issued for services	780	—	46	—	—	46
Share-based compensation	—	—	20,103	—	—	20,103
Net loss	—	—	—	(158,722)	—	(158,722)
Other comprehensive loss	—	—	—	—	(4,438)	(4,438)
Balance at June 30, 2022	94,367,233	\$ 9	\$ 2,106,942	\$(1,379,952)	\$ (4,802)	\$ 722,197

See accompanying notes to these condensed consolidated financial statements.

Intra-Cellular Therapies, Inc. and Subsidiaries

Condensed Consolidated Statements of Cash Flows (in thousands) (Unaudited)

	<b>Six Months Ended June 30,</b>	
	<b>2022</b>	<b>2021</b>
<b>Cash flows used in operating activities</b>		
Net loss	\$ (158,722)	\$ (121,484)
Adjustments to reconcile net loss to net cash used in operating activities:		
Depreciation	343	253
Share-based compensation	20,103	15,395
Stock issued for services	46	106
Amortization of premiums and discounts on investment securities, net	(947)	(1,708)
Changes in operating assets and liabilities:		
Accounts receivable, net	(26,820)	(4,423)
Inventory	(17,074)	(712)
Prepaid expenses and other assets	(12,535)	(7,954)
Accounts payable	2,058	10,439
Accrued liabilities and other	3,689	1,680
Accrued customer programs	10,519	1,395
Lease liabilities, net	1,698	(80)
Net cash used in operating activities	(177,642)	(107,093)
<b>Cash flows provided by (used in) investing activities</b>		
Purchases of investments	(552,782)	(119,899)
Maturities of investments	268,665	284,977
Purchases of property and equipment	(689)	(15)
Net cash provided by (used in) investing activities	(284,806)	165,063
<b>Cash flows provided by financing activities</b>		
Proceeds of public offering, net	433,725	—
Proceeds from exercise of stock options	13,593	3,014
Net cash provided by financing activities	447,318	3,014
Net increase (decrease) in cash, cash equivalents, and restricted cash	(15,130)	60,984
Cash, cash equivalents, and restricted cash at beginning of period	93,765	61,446
Cash, cash equivalents, and restricted cash at end of period	<u>\$ 78,635</u>	<u>\$ 122,430</u>
<b>Non-cash investing and financing activities</b>		
<b>Right of use assets under operating leases</b>	\$ 4,167	\$ 271

The following table provides a reconciliation of cash, cash equivalents and restricted cash reported within the condensed consolidated balance sheets that sum to the total of the same such amounts shown in the condensed consolidated statements of cash flows:

	<b>June 30,</b>	
	<b>2022</b>	<b>2021</b>
Cash and cash equivalents	\$77,235	\$121,030
Restricted cash	1,400	1,400
Total cash, cash equivalents and restricted cash	<u>\$78,635</u>	<u>\$122,430</u>

See accompanying notes to these condensed consolidated financial statements.



Intra-Cellular Therapies, Inc.

Notes to Condensed Consolidated Financial Statements (Unaudited)

June 30, 2022

**1. Organization**

Intra-Cellular Therapies, Inc. (the “Company”), through its wholly-owned operating subsidiaries, ITI, Inc. (“ITI”) and ITI Limited, is a biopharmaceutical company focused on the discovery, clinical development and commercialization of innovative, small molecule drugs that address underserved medical needs primarily in neuropsychiatric and neurological disorders by targeting intracellular signaling mechanisms within the central nervous system (“CNS”). In December 2019, the U.S. Food and Drug Administration (“FDA”) approved CAPLYTA<sup>®</sup> (lumateperone) for the treatment of schizophrenia in adults (42mg/day). The Company initiated the commercial launch of CAPLYTA in late March 2020. In December 2021, the FDA approved CAPLYTA for the treatment of depressive episodes associated with bipolar I or II disorder in adults, as monotherapy and as adjunctive therapy with lithium or valproate (42mg/day). Additionally, in April 2022, the FDA approved two new dosage strengths of CAPLYTA, 10.5 mg and 21 mg capsules, to provide dosage recommendations for patients concomitantly taking strong or moderate CYP3A4 inhibitors, and 21 mg for patients with moderate or severe hepatic impairment (Child-Pugh class B or C). As used in these Notes to Condensed Consolidated Financial Statements, “CAPLYTA” refers to lumateperone approved by the FDA for the treatment of schizophrenia and bipolar depression in adults, and “lumateperone” refers to, where applicable, CAPLYTA as well as lumateperone for the treatment of indications beyond schizophrenia and bipolar depression. Lumateperone is in Phase 3 clinical development as a novel treatment for major depressive disorder.

On January 7, 2022, the Company completed a public offering of common stock in which the Company sold 10,952,381 shares of common stock at a public offering price of \$42.00 per share for aggregate gross proceeds of \$460.0 million. After deducting underwriting discounts, commissions and offering expenses, the net proceeds to the Company were approximately \$433.7 million. In order to further its commercial activities and research projects and support its collaborations, the Company may require additional financing until such time, if ever, that revenue streams are sufficient to generate consistent positive cash flow from operations. The Company currently projects that its cash, cash equivalents and investments will be sufficient to fund operating expenses and capital expenditures for at least one year from the date that these financial statements are filed with the Securities and Exchange Commission (the “SEC”). Possible sources of funds include public or private sales of the Company’s equity securities, sales of debt or convertible debt securities, the incurrence of debt from commercial lenders, strategic collaborations, licensing a portion or all of the Company’s products, product candidates and technology and, to a much lesser extent, grant funding.

On January 6, 2020, the Company filed an automatic shelf registration statement on Form S-3 with the SEC, which became effective upon filing, on which the Company registered for sale an unlimited amount of any combination of its common stock, preferred stock, debt securities, warrants, rights, and/or units from time to time and at prices and on terms that the Company may determine, so long as the Company continues to satisfy the requirements of a “well-known seasoned issuer” under SEC rules. This registration statement will remain in effect for up to three years from the date it became effective.

**2. Summary of Significant Accounting Policies**

**Basis of Presentation**

The accompanying condensed consolidated financial statements of Intra-Cellular Therapies, Inc. and its wholly own subsidiaries have been prepared in conformity with accounting principles generally accepted in the United States of America (“GAAP”). Any reference in these notes to applicable guidance is meant to refer to the authoritative U.S. GAAP set forth in the Accounting Standards Codification (“ASC”) and Accounting Standards Update (“ASU”) of the Financial Accounting Standards Board (“FASB”). All intercompany accounts and transactions have been eliminated in consolidation. The Company currently operates in one operating segment. Operating segments are defined as components of an enterprise about which separate discrete information is available for the chief operating decision maker, or decision making group, in deciding how to allocate resources and assessing performance. The Company views its operations and manages its business in one segment, which is discovering, developing and commercializing drugs primarily for the treatment of neurological and psychiatric disorders.

**Recent Accounting Pronouncements**

Management has evaluated all accounting pronouncements issued through the date of the condensed consolidated financial statements and does not believe that any recently issued, but not yet effective, pronouncements, if currently adopted, would have a material effect on the Company’s financial statements.

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### Use of Estimates

The preparation of financial statements in conformity with GAAP requires management to make estimates and assumptions that affect the amounts reported in the financial statements and accompanying notes. Although actual results could differ from those estimates, management does not believe that such differences would be material.

### Significant Accounting Policies

The accounting policies used by the Company in its presentation of interim financial results are consistent with those presented in Note 2 to the consolidated financial statements included in the Company's Annual Report on Form 10-K for the year ended December 31, 2021.

### Concentration of Credit Risk

Financial instruments which potentially subject the Company to concentrations of credit risk consist of accounts receivable, net from customers and cash, cash equivalent and investments held at financial institutions. As of June 30, 2022, all of the Company's accounts receivable, net arose from product sales in the United States. For the six-month periods ended June 30, 2022 and 2021, respectively, 97% and 97% of sales were from three major wholesalers accounting for approximately 38%, 30%, and 29%, and 40%, 30%, and 27% of product sales, for the same periods, respectively. The percentage of the total net product sales by individual customers has not significantly changed since inception. All customers have standard payment terms which generally require payment within 60 days. As of June 30, 2022, the Company continues to believe that such customers are of high credit quality.

### 3. Investment Securities

Investment securities consisted of the following (in thousands):

	June 30, 2022			Estimated Fair Value
	Amortized Cost	Unrealized Gains	Unrealized (Losses)	
			(Unaudited)	
U.S. Government Agency Securities	\$ 197,886	\$ 10	\$ (1,611)	\$ 196,285
Certificates of Deposit	10,434	—	(38)	10,396
Commercial Paper	109,634	—	(476)	109,158
Corporate Notes/Bonds	288,441	1	(2,688)	285,754
	<u>\$606,395</u>	<u>\$ 11</u>	<u>\$ (4,813)</u>	<u>\$601,593</u>
			December 31, 2021	
	Amortized Cost	Unrealized Gains	Unrealized (Losses)	Estimated Fair Value
U.S. Government Agency Securities	\$ 71,752	\$ —	\$ (100)	\$ 71,652
Certificates of Deposit	26,232	1	—	26,233
Commercial Paper	55,955	—	(36)	55,919
Corporate Notes/Bonds	166,394	19	(249)	166,164
	<u>\$320,333</u>	<u>\$ 20</u>	<u>\$ (385)</u>	<u>\$319,968</u>

The Company has classified all of its investment securities as available-for-sale, including those with maturities beyond one year, as current assets on the condensed consolidated balance sheets based on the highly liquid nature of the investment securities and because these investment securities are considered available for use in current operations. As of June 30, 2022 and December 31, 2021, the Company held \$109.3 million and \$67.2 million, respectively, of available-for-sale investment securities with contractual maturity dates more than one year and less than two years.

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The aggregate related fair value of investments with unrealized losses as of June 30, 2022 was \$585.3 million, which consisted of \$191.4 million from U.S. government agency securities, \$1.9 million of certificates of deposit, \$109.2 million of commercial paper, and \$282.8 million of corporate notes/bonds. \$20.1 million of the aggregate fair value of investments with unrealized losses as of June 30, 2022 has been held in a continuous unrealized loss position for over 12 months, with the remaining \$565.1 million held in a continuous unrealized loss position for less than 12 months. As of December 31, 2021, the aggregate related fair value of investments with unrealized losses was \$263.5 million. \$9.5 million of the aggregate fair value of investments with unrealized losses as of December 31, 2021 has been held in a continuous unrealized loss position for over than 12 months, with the remaining \$254.1 million held in a continuous unrealized loss position for less than 12 months.

The Company reviewed all of the investments which were in a loss position at the respective balance sheet dates, as well as the remainder of the portfolio. The Company has analyzed the unrealized losses and determined that market conditions were the primary factor driving these changes. After analyzing the securities in an unrealized loss position, the portion of these losses that relate to changes in credit quality is insignificant. The Company does not intend to sell these securities, nor is it more likely than not that the Company will be required to sell them prior to the end of their contractual terms. Furthermore, the Company does not believe that these securities expose the Company to undue market risk or counterparty credit risk.

#### 4. Fair Value Measurements

The Company applies the fair value method under ASC Topic 820, *Fair Value Measurements and Disclosures*. The ASC Topic 820 hierarchy ranks the quality and reliability of inputs, or assumptions, used in the determination of fair value and requires assets and liabilities carried at fair value to be classified and disclosed in one of the following categories based on the lowest level input used that is significant to a particular fair value measurement:

- Level 1—Fair value is determined by using unadjusted quoted prices that are available in active markets for identical assets and liabilities.
- Level 2—Fair value is determined by using inputs other than Level 1 quoted prices that are directly or indirectly observable. Inputs can include quoted prices for similar assets and liabilities in active markets or quoted prices for identical assets and liabilities in inactive markets. Related inputs can also include those used in valuation or other pricing models, such as interest rates and yield curves that can be corroborated by observable market data.
- Level 3—Fair value is determined by inputs that are unobservable and not corroborated by market data. Use of these inputs involves significant and subjective judgments to be made by a reporting entity—e.g., determining an appropriate adjustment to a discount factor for illiquidity associated with a given security.

The Company has no assets or liabilities that were measured using quoted prices for significant unobservable inputs (Level 3 assets and liabilities) as of June 30, 2022 or December 31, 2021. The carrying value of cash held in money market funds of approximately \$55.8 million as of June 30, 2022 and \$56.5 million as of December 31, 2021 is included in cash and cash equivalents on the condensed consolidated balance sheets and approximates market value based on quoted market prices or Level 1 inputs. The carrying value of cash held in corporate bonds of approximately \$1.0 million and \$0 as of June 30, 2022 and December 31, 2021, respectively, is included in cash and cash equivalents.

The fair value measurements of the Company's cash equivalents and available-for-sale investment securities are identified in the following tables (in thousands):

	Fair Value Measurements at Reporting Date Using			
	June 30, 2022	Quoted Prices in Active Markets for Identical Assets (Level 1)	Significant Other Observable Inputs (Level 2)	Significant Unobservable Inputs (Level 3)
Money Market Funds	\$ 55,775	\$ 55,775	\$ —	\$ —
U.S. Government Agency Securities	196,285	—	196,285	—
Certificates of Deposit	10,396	—	10,396	—
Commercial Paper	109,158	—	109,158	—
Corporate Notes/Bonds	285,754	—	285,754	—
	<u>\$657,368</u>	<u>\$ 55,775</u>	<u>\$ 601,593</u>	<u>\$ —</u>

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	Fair Value Measurements at Reporting Date Using			
	December 31, 2021	Quoted Prices in Active Markets for Identical Assets (Level 1)	Significant Other Observable Inputs (Level 2)	Significant Unobservable Inputs (Level 3)
Money Market Funds	\$ 56,539	\$ 56,539	\$ —	\$ —
U.S. Government Agency Securities	71,652	—	71,652	—
Certificates of Deposit	26,233	—	26,233	—
Commercial Paper	55,919	—	55,919	—
Corporate Notes/Bonds	166,164	—	166,164	—
	<u>\$ 376,507</u>	<u>\$ 56,539</u>	<u>\$ 319,968</u>	<u>\$ —</u>

### 5. Inventory

Inventory consists of the following (in thousands):

	June 30, 2022	December 31, 2021
Raw materials	<u>\$18,160</u>	\$ 2,484
Work in process	<u>1,741</u>	2,407
Finished goods	<u>5,121</u>	3,057
	<u>\$25,022</u>	<u>\$ 7,948</u>

Product costs incurred prior to receipt of the FDA approval on December 20, 2019 for CAPLYTA was expensed as research and development as incurred.

### 6. Right of Use Assets and Lease Liabilities

In 2014, the Company entered into a long-term lease with a related party, which, as amended, provided for a lease of useable laboratory and office space located in New York, New York. A member of the Company's board of directors is the Executive Chairman of the parent company to the landlord under this lease. Concurrent with this lease, the Company entered into a license agreement to occupy certain vivarium-related space in the same facility for the same term and rent escalation provisions as the lease. This license has the primary characteristics of a lease and is characterized as a lease. In September 2018, the Company further amended the lease to obtain an additional office space beginning October 1, 2018 and to extend the term of the lease for previously acquired space. The lease, as amended, has a term of 14.3 years ending in May 2029.

On May 17, 2019, the Company entered into an agreement (the "Vehicle Lease") with a company (the "Lessor") to acquire motor vehicles for certain employees. The Vehicle Lease provides for individual leases for the vehicles, which at each lease commencement was determined to qualify for operating lease treatment. The Company began leasing vehicles under the Vehicle Lease in March 2020. The contractual period of each lease is 12 months, followed by month-to-month renewal periods. The Company estimates the lease term for each vehicle to be 30 months based on industry standards. The Company is required to maintain an irrevocable \$1.4 million letter of credit that the Lessor may draw upon in the event the Company defaults on the Vehicle Lease, which has been recorded as restricted cash on the condensed consolidated balance sheet.

The Company has no other significant leases. In addition, no identified leases require allocations between lease and non-lease components.

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Maturity analyses under the lease agreements are as follows (in thousands):

	<u>Real Estate</u>	<u>Fleet</u>
Six months ending December 31, 2022	\$ 1,792	\$ 2,352
Year ending December 31, 2023	3,680	2,687
Year ending December 31, 2024	3,792	2,130
Year ending December 31, 2025	3,907	—
Year ending December 31, 2026	3,974	—
Thereafter	9,937	—
Total	27,082	7,169
Less: Present value discount	(7,194)	(239)
Total Lease liability	19,888	6,930
Less: Current portion	(3,451)	(4,292)
Long-term lease liabilities	<u>\$ 16,437</u>	<u>\$ 2,638</u>

## 7. Commitments and Contingencies

### *License and Royalty Commitments*

On May 31, 2005, the Company entered into a worldwide, exclusive License Agreement with Bristol-Myers Squibb Company (“BMS”), pursuant to which the Company holds a license to certain patents and know-how of BMS relating to lumateperone and other specified compounds. The agreement was amended on November 3, 2010. The licensed rights are exclusive, except BMS retains rights in specified compounds in the fields of obesity, diabetes, metabolic syndrome and cardiovascular disease. However, BMS has no right to use, develop or commercialize lumateperone and other specified compounds in any field of use. The Company has the right to grant sublicenses of the rights conveyed by BMS. The Company is obliged under the agreement to use commercially reasonable efforts to develop and commercialize the licensed technology. The Company is also prohibited from engaging in the clinical development or commercialization of specified competitive compounds.

Under the agreement, the Company has made payments of \$10.75 million to BMS related to milestones achieved to date for lumateperone. Possible milestone payments remaining total \$5.0 million. Under the agreement, the Company may be obliged to make other milestone payments to BMS for each licensed product of up to an aggregate of approximately \$14.75 million. The Company is also obliged to make tiered single digit percentage royalty payments ranging between 5 – 9% on sales of licensed products. The Company is obliged to pay to BMS a percentage of non-royalty payments made in consideration of any sublicense.

The agreement extends, and royalties are payable, on a country-by-country and product-by-product basis, through the later of 10 years after first commercial sale of a licensed product in such country, expiration of the last licensed patent covering a licensed product, its method of manufacture or use, or the expiration of other government grants providing market exclusivity, subject to certain rights of the parties to terminate the agreement on the occurrence of certain events. On termination of the agreement, the Company may be obliged to convey to BMS rights in developments relating to a licensed compound or licensed product, including regulatory filings, research results and other intellectual property rights.

In September 2016, the Company transferred certain of its rights under the BMS agreement to its wholly owned subsidiary, ITI Limited. In connection with the transfer, the Company guaranteed ITI Limited’s performance of its obligations under the BMS agreement. The Company expensed approximately \$4.5 million and \$1.7 million, respectively, for the six-month periods ended June 30, 2022 and 2021, in cost of product sales to satisfy its obligation under the BMS agreement.

### *Research and Other Commitments*

As of June 30, 2022, the Company has committed to purchasing production campaigns for various raw materials and API from each of its supply vendors – Siegfried Evionnaz SA (“Siegfried”) and Lonza Ltd. (“Lonza”). The campaigns are expected to be received into inventory during 2022 through 2024. The Company had paid deposits of \$15.0 million and \$9.5 million as of June 30, 2022 and December 31, 2021, respectively, for the Siegfried and Lonza campaigns, which are recorded within prepaid expenses and other current assets. Over the course of the vendors’ manufacturing period, the Company will remit payments to each vendor based on the payment plan within the executed agreements.

## 8. Share-Based Compensation

In 2020, the Company’s stockholders approved the Amended and Restated 2018 Equity Incentive Plan (the “Amended 2018 Plan”), which provides for the granting of up to 6,500,000 additional share-based awards, such as

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stock options, restricted common stock, RSUs and stock appreciation rights to employees, directors and consultants as determined by the Board of Directors. In December 2019, the Company adopted the 2019 Inducement Award Plan (the “2019 Inducement Plan”) for the grant of equity awards of up to 1,000,000 shares of common stock to newly hired employees.

Information regarding the stock options activity, including with respect to grants to employees, directors and consultants as of June 30, 2022, and changes during the six-month period then ended, are summarized as follows:

	<u>Number of Shares</u>	<u>Weighted-Average Exercise Price</u>	<u>Weighted-Average Contractual Life</u>
Outstanding at December 31, 2021	5,451,398	\$ 21.09	6.1 years
Options granted 2022	592,332	\$ —	
Options exercised 2022	(778,916)	\$ —	
Options canceled or expired 2022	(23,010)	\$ —	
Outstanding at June 30, 2022	<u>5,241,804</u>	<u>\$ 25.54</u>	6.2 years
Vested and expected to vest at June 30, 2022	<u>5,241,804</u>	<u>\$ 25.54</u>	
Exercisable at June 30, 2022	<u>4,042,330</u>	<u>\$ 20.04</u>	5.4 years

The fair value of the time based RSUs is based on the closing price of the Company’s common stock on the date of grant. Information regarding the time based RSU activity and changes during the six-month period ended June 30, 2022 is summarized as follows:

	<u>Number of Shares</u>	<u>Weighted-Average Grant Date Fair Value Per Share</u>	<u>Weighted-Average Contractual Life</u>
Outstanding at December 31, 2021	1,529,652	\$ 26.95	1.5 years
Time based RSUs granted in 2022	582,768	\$ —	
Time based RSUs vested in 2022	(751,695)	\$ —	
Time based RSUs cancelled in 2022	(37,919)	\$ —	
Outstanding at June 30, 2022	<u>1,322,806</u>	<u>\$ 42.39</u>	1.3 years

The Company recognized non-cash share-based compensation expense related to time based RSUs for the six months ended June 30, 2022 and 2021 of approximately \$10.4 million and \$7.9 million, respectively. As of June 30, 2022, there was approximately \$48.0 million of unrecognized compensation costs estimated related to unvested time based RSUs.

## 9. Loss Per Share

The following awards were excluded in the calculation of diluted loss per share because their effect could be anti-dilutive as applied to the loss from operations for the three and six months ended June 30, 2022 and 2021:

	<u>Three and Six Months Ended June 30,</u>	
	<u>2022</u>	<u>2021</u>
Stock options	5,241,804	6,014,017
RSUs	1,420,313	1,631,478
TSR RSUs	97,507	68,598

**Item 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS**

*You should read the following in conjunction with our unaudited condensed consolidated financial statements and the related notes thereto that appear elsewhere in this Quarterly Report on Form 10-Q and the audited consolidated financial statements and notes thereto and under the heading "Management's Discussion and Analysis of Financial Condition and Results of Operations" in our Annual Report on Form 10-K filed on March 1, 2022. In addition to historical information, the following discussion and analysis includes forward-looking information that involves risks, uncertainties and assumptions. Our actual results and the timing of events could differ materially from those anticipated by these forward-looking statements as a result of many factors, including those discussed under "Risk Factors" in our Annual Report on Form 10-K filed on March 1, 2022, as updated from time to time in our subsequent periodic and current reports filed with the SEC.*

**Overview**

We are a biopharmaceutical company focused on the discovery, clinical development and commercialization of innovative, small molecule drugs that address underserved medical needs primarily in neuropsychiatric and neurological disorders by targeting intracellular signaling mechanisms within the central nervous system, or CNS. In December 2019, CAPLYTA (lumateperone) was approved by the U.S. Food and Drug Administration, or FDA, for the treatment of schizophrenia in adults (42mg/day) and we initiated the commercial launch of CAPLYTA in late March 2020. In support of our commercialization efforts, we employ a national sales force. In December 2021, CAPLYTA was approved by the FDA for the treatment of bipolar depression in adults (42mg/day). CAPLYTA is the only FDA-approved treatment for depressive episodes associated with bipolar I or II disorder (bipolar depression) in adults as monotherapy and as adjunctive therapy with lithium or valproate. We initiated the commercial launch of CAPLYTA for the treatment of bipolar depression in late December 2021. In support of the commercial launch of lumateperone for the treatment of bipolar depression, we have expanded our sales force from approximately 240 sales representatives to approximately 320 sales representatives. In addition, in April 2022, the FDA approved two new dosage strengths of CAPLYTA, 10.5 mg and 21 mg capsules, to provide dosage recommendations for patients concomitantly taking strong or moderate CYP3A4 inhibitors, and 21 mg for patients with moderate or severe hepatic impairment (Child-Pugh class B or C). As used in this report, "CAPLYTA" refers to lumateperone approved by the FDA for the treatment of schizophrenia in adults and for the treatment of bipolar depression in adults, and "lumateperone" refers to, where applicable, CAPLYTA as well as lumateperone for the treatment of indications beyond schizophrenia and bipolar depression.

Lumateperone is in Phase 3 clinical development as a novel treatment for major depressive disorder, or MDD. Patient enrollment in Study 501 and Study 502, our global Phase 3 clinical trials evaluating lumateperone 42 mg as an adjunctive therapy to antidepressants for the treatment of MDD is ongoing. We expect to file an sNDA with the FDA for approval of lumateperone as an adjunctive therapy to antidepressants for the treatment of MDD in 2024. In the first quarter of 2020, as part of our lumateperone bipolar depression clinical program, we initiated our third monotherapy Phase 3 study, Study 403, evaluating lumateperone as monotherapy in the treatment of major depressive episodes associated with bipolar I or bipolar II disorder. Following the positive results in our adjunctive study that was part of our bipolar depression clinical program, Study 402, we amended Study 403 to evaluate major depressive episodes with mixed features in bipolar disorder in patients with bipolar I or bipolar II disorder and mixed features in patients with MDD. We expect to complete clinical conduct in Study 403 in late 2022 and, following completion of data analysis, we expect to report topline results in the first quarter of 2023. Following completion of Study 403, we intend to discuss the results with the FDA to determine whether this study will provide supportive data for a potential future regulatory filing for this indication.

We have also initiated a Phase 3 study evaluating lumateperone for the prevention of relapse in patients with schizophrenia. The study is being conducted in five phases consisting of a screening phase, a 6-week, open-label run-in phase during which all patients will receive 42 mg of lumateperone per day, a 12-week, open-label stabilization phase during which all patients will receive 42 mg of lumateperone per day; a double-blind treatment phase 26 weeks in duration during which patients receive either 42 mg of lumateperone per day or placebo (1:1 ratio) and a 2-week safety follow-up phase. This study is being conducted in accordance with our post approval marketing commitment to the FDA in connection with the approval of CAPLYTA for the treatment of schizophrenia as is typical for antipsychotics.

Within the lumateperone portfolio, we are also developing a long-acting injectable, or LAI, formulation to provide more treatment options to patients suffering from mental illness. We have completed the preclinical development of an LAI formulation, and we have conducted a Phase 1 single ascending dose study with this formulation. This study evaluated the pharmacokinetics, safety and tolerability of lumateperone LAI in patients with stable symptoms of schizophrenia. We are now exploring alternate sites of injection with this formulation as well as progressing other formulations. This will assist us in evaluating dosing strategies and formulation for our efficacy studies. The goal of our program is to develop LAI formulations that are effective, safe and well-tolerated with treatment durations of one month and longer. Given the encouraging tolerability data to date with oral lumateperone, we believe that an LAI option, in particular, may lend itself to being an important formulation choice for certain patients.

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We are developing ITI-1284-ODT-SL for the treatment of behavioral disturbances in patients with dementia, the treatment of dementia-related psychosis and for the treatment of certain depressive disorders in the elderly. ITI-1284-ODT-SL is a deuterated form of lumateperone, a new molecular entity formulated as an oral disintegrating tablet for sublingual administration. ITI-1284-ODT-SL is formulated as an oral solid dosage form that dissolves almost instantly when placed under the tongue, allowing for ease of use in the elderly and may be particularly beneficial for patients who have difficulty swallowing conventional tablets. Phase 1 single and multiple ascending dose studies in healthy volunteers and healthy elderly volunteers (> than 65 years of age) evaluated the safety, tolerability and pharmacokinetics of ITI-1284-ODT-SL. In these studies, there were no reported serious adverse events in either age group. In the elderly cohort, reported adverse events were infrequent with the most common adverse event being transient dry mouth (mild). Based on these results, we have initiated our program evaluating ITI-1284-ODT-SL for the treatment of agitation in patients with probable Alzheimer's disease. We are in discussions with the FDA regarding the non-clinical toxicological profile of ITI-1284-ODT-SL. The FDA has informed us that they do not believe the deuterated and undeuterated forms of lumateperone are identical. As a result, the non-clinical data from lumateperone may not be broadly applied to ITI-1284-ODT-SL and we are conducting additional toxicology studies and we expect to commence clinical conduct in a Phase 2 study in agitation in patients with probable Alzheimer's disease in 2023. Additional studies in dementia-related psychosis and certain depressive disorders in the elderly are also planned for 2023. We are continuing with Phase 1 studies with ITI-1284-ODT-SL including drug-drug interaction studies.

We have another major program that has yielded a portfolio of compounds that selectively inhibit the enzyme phosphodiesterase type 1, or PDE1. PDE1 enzymes are highly active in multiple disease states and our PDE1 inhibitors are designed to reestablish normal function in these disease states. Abnormal PDE1 activity is associated with cellular proliferation and activation of inflammatory cells. Our PDE1 inhibitors ameliorate both of these effects in animal models. We intend to pursue the development of our phosphodiesterase, or PDE, program, for the treatment of aberrant immune system activation in several CNS and non-CNS conditions with a focus on diseases where excessive PDE1 activity has been demonstrated and increased inflammation is an important contributor to disease pathogenesis. Our potential disease targets include heart failure, immune system regulation, neurodegenerative diseases, cancers and other non-CNS disorders. Lenrispodun (ITI-214) is our lead compound in this program. Following the favorable safety and tolerability results in our Phase 1 program, we initiated our development program for lenrispodun for Parkinson's disease and conducted a Phase 1/2 clinical trial of lenrispodun in patients with Parkinson's disease to evaluate safety and tolerability in this patient population, as well as motor and non-motor exploratory endpoints. In this study lenrispodun was generally well-tolerated with a favorable safety profile and clinical signs consistent with improvements in motor symptoms and dyskinesias. We have initiated our Phase 2 clinical program with lenrispodun for Parkinson's disease and expect to commence patient enrollment in the second half of 2022. In addition, we have conducted a Phase 1/2 translational study of single ascending doses of lenrispodun in patients with chronic systolic heart failure with reduced ejection fraction. In this study, lenrispodun improved cardiac output by increasing heart contractility and decreasing vascular resistance. Agents that both increase heart contractility (inotropism) and decrease vascular resistance (vasodilation) are called inodilators. Inodilators in current clinical use are associated with the development of arrhythmias, which are abnormal heart rhythms that, when serious, can impair heart function and lead to mortality. Lenrispodun, which acts through a novel mechanism of action, was not associated with arrhythmias in this study and was generally well-tolerated in all patients.

We also have a development program with our ITI-333 compound as a potential treatment for substance use disorders, pain and psychiatric comorbidities including depression and anxiety. There is a pressing need to develop new drugs to treat opioid addiction and for safe, effective, non-addictive treatments to manage pain. ITI-333 is a novel compound that uniquely combines activity as an antagonist at serotonin 5-HT<sub>2A</sub> receptors and a partial agonist at  $\mu$ -opioid receptors. These combined actions support the potential utility of ITI-333 in the treatment of opioid use disorder and associated comorbidities (e.g., depression, anxiety, sleep disorders) without opioid-like safety and tolerability concerns. We have conducted a Phase 1 single ascending dose study evaluating the safety, tolerability and pharmacokinetics of ITI-333 in healthy volunteers. In this study ITI-333 achieved plasma exposures at or above those required for efficacy and was generally safe and well-tolerated. We have commenced a neuroimaging study to investigate brain occupancy for receptors that play a role in substance use disorder and also have applicability for pain. The results of this study will support the dose selection for future studies. We have received a grant from the National Institute on Drug Abuse under the Helping to End Addiction Long-term Initiative, or NIH HEAL Initiative, that we expect will fund a significant portion of the early stage clinical development costs associated with this program.

We have assembled a management team with significant industry experience to lead the commercialization of our product and the discovery, development and potential commercialization of our product candidates. We complement our management team with a group of scientific and clinical advisors that includes recognized experts in the fields of schizophrenia, bipolar depression and other CNS disorders.



## COVID-19

In December 2019, a novel strain of coronavirus, SARS-CoV-2, which causes coronavirus disease 2019, or COVID-19, surfaced in Wuhan, China. Since then, SARS-CoV-2 and COVID-19 have spread to countries worldwide, including the United States. The COVID-19 pandemic continues to evolve, and to date has led to the implementation of various responses, including government-imposed quarantines, travel restrictions and other public health safety measures. As a result of the COVID-19 pandemic, we expect to continue to experience disruptions, which could severely impact our business, including our ability to successfully commercialize our only commercial product, CAPLYTA, in the United States, and could negatively impact our sales of CAPLYTA. Our commercial organization, sales force and medical organization have had, and, depending on the severity and duration of the pandemic, may continue to have significantly reduced personal interactions with physicians and customers and may need to continue to conduct many promotional activities virtually, and we may elect to cease in-person interactions with physicians and customers entirely for some period of time in the interest of employee and community safety. In addition, the pandemic may continue to impact the willingness of patients to visit their healthcare provider. Business interruptions from the current or future pandemics may also adversely impact the third parties we rely on to sufficiently manufacture CAPLYTA and to produce our product candidates in quantities we require, which may impair the commercialization and our research and development activities.

We conduct clinical trials for our product candidates in many countries and regions, including the United States and Europe, and may expand to other geographies. Timely enrollment of, completion of and reporting on our clinical trials is dependent upon these global clinical trial sites which are, or in the future may be, adversely affected by the COVID-19 pandemic or other pandemics. Some factors from the COVID-19 pandemic that have or may adversely affect the timing and conduct of our clinical trials and adversely impact our business generally, include but are not limited to, delays or difficulties in clinical site initiation, patient enrollment, diversion of healthcare resources away from clinical trials to pandemic concerns, limitations on travel, regulatory delays and supply chain disruptions.

During the global response to the COVID-19 pandemic, moreover, the responses of the federal, international, state and regional governments to the public health emergency, including but not limited to the shelter in place orders, the allocation of healthcare resources to treating those infected with the virus, and the strategic redeployment of FDA and EMA resources and staff to priority projects, could have an impact on the timeline for review and approval of new marketing applications. Over the course of the pandemic, FDA's new drug review programs continued to meet key performance goals related to working with applicants and approving NDAs and NDA supplements, although the agency has also stated that the uncertainty of the COVID-19 situation may make it difficult to sustain that level of performance indefinitely. The FDA may not be able to maintain its normal pace with respect to new drug applications and delays or setbacks are possible in the future.

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The COVID-19 pandemic continues to rapidly evolve, and the severity and duration of the pandemic remain uncertain. The extent to which the pandemic impacts our business, including our commercial results, clinical trials, and preclinical studies will depend on future developments, which are highly uncertain.

### **Results of Operations**

The following discussion summarizes the key factors our management believes are necessary for an understanding of our financial statements.

#### ***Revenues***

Net revenues from product sales consist of sales of CAPLYTA, which was approved by the FDA for the treatment of schizophrenia in adults in December 2019 and for the treatment of bipolar depression in adults in December 2021. We initiated the commercial launch of CAPLYTA in late March 2020. During the three and six months ended June 30, 2022, net sales increased to approximately \$55.1 million and \$89.8 million, respectively, from approximately \$19.0 million and \$34.6 million for the three and six months ended June 30, 2021, respectively.

#### ***Expenses***

The process of researching, developing and commercializing drugs for human use is lengthy, unpredictable and subject to many risks. We are unable, with certainty, to estimate either the costs or the timelines in which those costs will be incurred. The costs associated with the commercialization of CAPLYTA are substantial and will be incurred prior to our generating sufficient revenue to offset these costs. Costs for the clinical development of lumateperone-related projects, including for the treatment of MDD, consumes and, together with our other anticipated clinical development programs, will continue to consume a large portion of our current, as well as projected, resources. We intend to pursue other disease indications that lumateperone may address, but there are significant costs associated with pursuing FDA approval for those indications, which would include the cost of additional clinical trials.

Our PDE, ITI-1284 and ITI-333 development programs are currently in clinical stage development. Our other programs are still in the preclinical stages and will require extensive funding not only to complete preclinical testing, but also to commence and complete clinical trials. Expenditures that we incur on these programs will be subject to availability of funding in addition to the funding required for the advancement of lumateperone. Any failure or delay in the advancement of lumateperone could require us to re-allocate resources from our other programs to the advancement of lumateperone, which could have a material adverse impact on the advancement of these other programs and on our results of operations.

Our operating expenses are comprised of (i) costs of product sales; (ii) selling expenses; (iii) general and administrative expenses; and (iv) research and development expenses.

Costs of product sales are comprised of:

- direct costs of formulating, manufacturing and packaging drug product;
- overhead costs consisting of labor, customs, share-based compensation, shipping, outside inventory management and other miscellaneous operating costs; and
- royalty payments on product sales.

Selling expenses are incurred in three major categories:

- salaries and related benefit costs of a dedicated sales force;
- sales operation costs; and
- marketing, promotional and advertising expenses.

General and administrative expenses are incurred in three major categories:

- salaries and related benefit costs;

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- patent, legal, and professional costs; and
- office and facilities overhead.

Research and development costs are comprised of:

- fees paid to external parties who provide us with contract services, such as pre-clinical testing, manufacturing and related testing, clinical trial activities and license milestone payments; and
- internal recurring costs, such as costs relating to labor and fringe benefits, materials, supplies, facilities and maintenance.

Product sold through June 30, 2022 consisted of drug product that was previously charged to research and development expense prior to FDA approval of CAPLYTA and other direct, indirect, and overhead costs required to make final product for sale. Because the Company's policy does not allow for the capitalization of pre-approval product, the cost of drug product sold is lower than it would have been and has a positive impact on our cost of product sales for the three and six-month periods ended June 30, 2022 and 2021. We expect to continue to have this favorable impact on cost of product sales and related product gross margins until our sales of CAPLYTA include drug product that is manufactured entirely after the FDA approval. We expect that this will be the case for the near-term and, as a result, our cost of product sales will be less than we anticipate it will be in future periods.

We expect that research and development expenses will increase as we proceed with our clinical trials including, increased manufacturing of drug product for clinical trials and pre-clinical development activities. We also expect that our selling, general and administrative costs will increase from prior periods primarily due to costs associated with promotional activities to support the commercial sales of CAPLYTA as well as costs associated with building and maintaining infrastructure, which will include hiring additional personnel and increasing technological capabilities. We granted significant share-based awards in 2021 and 2022. We expect to continue to grant share-based awards in the future due to our growing employee base, which will increase our share-based compensation expense in future periods.

The following table sets forth our revenues, operating expenses, interest income and income tax expense for the three and six-month periods ended June 30, 2022 and 2021 (in thousands):

	For the Three Months Ended June 30,		For the Six Months Ended June 30,	
	2022	2021	2022	2021
	(Unaudited)		(Unaudited)	
<b>Revenues</b>				
Product sales, net	\$ 55,074	\$ 19,007	\$ 89,829	\$ 34,586
Grant revenue	505	1,040	746	1,339
<b>Total revenues, net</b>	<b>55,579</b>	<b>20,047</b>	<b>90,575</b>	<b>35,925</b>
<b>Expenses</b>				
Cost of product sales	4,650	2,040	7,805	3,495
Selling, general and administrative	100,316	69,851	175,776	122,435
Research and development	38,536	17,297	67,579	32,355
<b>Total costs and expenses</b>	<b>143,502</b>	<b>89,188</b>	<b>251,160</b>	<b>158,285</b>
<b>Loss from operations</b>	<b>(87,923)</b>	<b>(69,141)</b>	<b>(160,585)</b>	<b>(122,360)</b>
Interest income	1,320	421	1,868	905
Income tax expense	—	(24)	(5)	(29)
<b>Net loss</b>	<b>\$(86,603)</b>	<b>\$(68,744)</b>	<b>\$(158,722)</b>	<b>\$(121,484)</b>

### Comparison of Three and Six-Month Periods Ended June 30, 2022 and June 30, 2021

#### Total Revenues, Net

Total revenues, net for the three and six-month periods ended June 30, 2022 were approximately \$55.6 million and \$90.6 million, respectively, compared to \$20.0 million and \$35.9 million for the three and six-month periods ended June 30, 2021, respectively. Net product sales were approximately \$55.1 million and \$89.8 million, respectively, for the three and six-month periods ended June 30, 2022 and \$19.0 million and \$34.6 million for the three and six-month periods ended June 30, 2021, respectively. Net product revenue in 2021 was comprised of sales of CAPLYTA for the treatment of schizophrenia, while net product revenue in 2022 was comprised of sales of CAPLYTA for the treatment of schizophrenia and bipolar depression.

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### *Cost of Product Sales*

Cost of product sales was approximately \$4.7 million and \$7.8 million, respectively, for the three and six-month periods ended June 30, 2022, compared to \$2.0 million and \$3.5 million, respectively, for the three and six-month periods ended June 30, 2021. Cost of product sales consisted primarily of product royalty fees, overhead and direct costs. Drug product costs, including certain direct, indirect, and overhead costs, incurred through June 30, 2022 were previously charged to research and development expense prior to FDA approval and are not a component of cost of product sales. This minimal cost drug product had a positive impact on our cost of product sales and related product gross margins for the three and six-month periods ended June 30, 2022 and 2021.

We will continue to have a lower cost of product sales that excludes the cost of the drug product that was incurred prior to FDA approval until our sales of CAPLYTA include drug product that is entirely manufactured after the FDA approval. We expect that this will be the case for the near-term and, as a result, our cost of product sales will be less than we anticipate it will be in future periods.

### *Selling, General and Administrative Expenses*

Selling, general and administrative costs for the three-month period ended June 30, 2022 were \$100.3 million as compared to \$69.9 million in the three-month period ended June 30, 2021, which represents an increase of 44%, which was due to an increase in selling costs and an increase in general and administrative expenses as discussed below.

Selling costs were \$81.3 million for the three-month period ended June 30, 2022 as compared to selling costs of \$52.1 million in the same period in 2021, which represents an increase of 56%. This increase is primarily due to increases of marketing and advertising expenses of approximately \$20.9 million, sales related labor costs of approximately \$6.8 million, and approximately \$1.5 million in travel and other costs. Salaries, bonuses and related benefit costs for our sales and marketing functions for the three months ended June 30, 2022 and 2021 constituted approximately 29% and 33%, respectively, of our selling costs.

General and administrative expenses were \$19.0 million in the three-month period ended June 30, 2022 as compared to \$17.7 million for the same period in 2021, an increase of 7%. This increase is due to increases in labor related costs of approximately \$0.9 million, stock-based compensation of \$0.7 million, and \$0.3 million for insurance and other expenses, partially offset by a decrease in professional fees of \$0.6 million. Salaries, bonuses and related benefit costs for our general and administrative functions for the three months ended June 30, 2022 and 2021 constituted approximately 58% and 54%, respectively, of our general and administrative costs.

Selling, general and administrative costs for the six-month period ended June 30, 2022 were \$175.8 million as compared to \$122.4 million in the six month period ended June 30, 2021, which represents an increase of 44%, which was due to an increase in selling, marketing, and advertising expenses and an increase in general and administrative expenses as discussed below.

Selling costs were \$137.3 million for the six-month period ended June 30, 2022 as compared to \$90.4 million in the same period in 2021, or an increase of 52%. This increase is primarily due to increases in commercialization costs of \$27.4 million, sales related labor costs of approximately \$14.4 million and approximately \$5.2 million in travel and other sales related expenses. Salaries, bonuses and related benefit costs for our sales and marketing functions for the six months ended June 30, 2022 and 2021 constituted approximately 35% and 37%, respectively, of our selling costs.

General and administrative expenses for the six months ended June 30, 2022 were \$38.4 million as compared to \$32.0 million for the same period in 2021, an increase of 20%. This increase is due to increases in stock compensation expense of \$2.0 million, labor and bonus expense of \$1.7 million, and the remainder for insurance, lease expense, and other administrative expenses. Salaries, bonuses and related benefit costs for our general and administrative functions for the six months ended June 30, 2022 and 2021 constituted approximately 56% and 51%, respectively, of our general and administrative costs.

We expect selling, general and administrative costs to increase moderately in 2022 as compared to 2021. We are expanding marketing, promotional, and advertising costs and increasing efforts to educate physicians, expand market access, and enhance our administrative infrastructure.

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### *Research and Development Expenses*

The following tables set forth our research and development expenses for the three and six-month periods ended June 30, 2022 and 2021 (in thousands):

	<b>Three Months Ended June 30,</b>		<b>Six Months Ended June 30,</b>	
	<b>2022</b>	<b>2021</b>	<b>2022</b>	<b>2021</b>
External costs	25,863	8,733	45,156	16,480
Internal costs	12,673	8,564	22,423	15,875
<b>Total research and development expenses</b>	<b>\$ 38,536</b>	<b>\$ 17,297</b>	<b>\$ 67,579</b>	<b>\$ 32,355</b>

  

	<b>Three Months Ended June 30,</b>		<b>Six Months Ended June 30,</b>	
	<b>2022</b>	<b>2021</b>	<b>2022</b>	<b>2021</b>
Lumateperone costs	21,978	10,364	39,451	18,891
Non-lumateperone project costs	10,885	3,723	19,892	7,411
Stock based compensation	5,507	2,862	7,878	5,184
Overhead	166	348	358	869
<b>Total research and development expenses</b>	<b>\$ 38,536</b>	<b>\$ 17,297</b>	<b>\$ 67,579</b>	<b>\$ 32,355</b>

Research and development expenses increased to \$38.5 million for the three-month period ended June 30, 2022 as compared to \$17.3 million for the three-month period ended June 30, 2021, representing an increase of approximately \$21.2 million, or 123%. This increase is due primarily to an increase of \$11.6 million for lumateperone clinical trial and other costs, and approximately \$7.2 million for other projects including the ITI-1284, ITI-214, and ITI-333 programs, among others, and an increase of approximately \$2.6 million for stock based compensation, partially offset by a decrease of approximately \$0.2 million of overhead costs. Internal costs increased by approximately \$4.1 million for the period due primarily to labor related costs and share-based compensation.

Research and development expenses increased to \$67.6 million for the six-month period ended June 30, 2022 as compared to \$32.4 million for the six-month period ended June 30, 2021, representing an increase of approximately 109%. This increase is due primarily to an increase of approximately \$20.6 million for lumateperone costs, an increase of approximately \$12.5 million for other projects including the ITI-1284, ITI-214, and ITI-333 programs, among others, and an increase of approximately \$2.7 million for stock based compensation, partially offset by a decrease of approximately \$0.5 million of overhead costs. Internal costs increased by approximately \$6.5 million for the period due primarily to labor related costs and stock based compensation.

As the development of lumateperone and non-lumateperone programs progress, we anticipate research and development costs to increase moderately due primarily to pre-clinical testing and conducting ongoing and planned clinical trials during the next several years. We are also required to complete non-clinical testing to obtain FDA approval and manufacture materials needed for clinical trial use, which includes non-clinical testing of the drug product, and manufacturing of drug product in anticipation of possible additional FDA approvals of lumateperone for indications beyond schizophrenia and bipolar depression.

We currently have several projects, in addition to lumateperone, that are in the research and development stages. We have used internal resources and incurred expenses not only in relation to the development of lumateperone, but also in connection with these additional projects as well, including our PDE program. We have not, however, reported these costs on a project-by-project basis, as these costs are broadly spread among these projects. The external costs for these projects have been modest and are reflected in the table above in this section “—*Research and Development Expenses*.”

The research and development process necessary to develop a pharmaceutical product for commercialization is subject to extensive regulation by numerous governmental authorities in the United States and other countries. This process typically takes years to complete and requires the expenditure of substantial resources. The steps required before a drug may be marketed in the United States generally include the following:

- completion of extensive pre-clinical laboratory tests, animal studies, and formulation studies in accordance with the FDA’s Good Laboratory Practice, or GLP, regulations;
- submission to the FDA of an Investigational New Drug application, or IND, for human clinical testing, which must become effective before human clinical trials may begin;
- performance of adequate and well-controlled human clinical trials to establish the safety and efficacy of the drug for each proposed indication;
- submission to the FDA of a New Drug Application, or NDA, after completion of all clinical trials;

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- satisfactory completion of an FDA pre-approval inspection of the manufacturing facility or facilities at which the active pharmaceutical ingredient, or API, and finished drug product are produced and tested to assess compliance with current Good Manufacturing Practices, or cGMPs;
- satisfactory completion of FDA inspections of clinical trial sites to assure that data supporting the safety and effectiveness of product candidates has been generated in compliance with Good Clinical Practices; and
- FDA review and approval of the NDA prior to any commercial marketing or sale of the drug in the United States.

The successful development of our product candidates and the approval process requires substantial time, effort and financial resources, and is uncertain and subject to a number of risks. We cannot be certain that any of our product candidates will prove to be safe and effective, will meet all of the applicable regulatory requirements needed to receive and maintain marketing approval, or will be granted marketing approval on a timely basis, if at all. Data from pre-clinical studies and clinical trials are susceptible to varying interpretations that could delay, limit or prevent regulatory approval or could result in label warnings related to or recalls of approved products. We, the FDA, or other regulatory authorities may suspend clinical trials at any time if we or they believe that the subjects participating in such trials are being exposed to unacceptable risks or if such regulatory agencies find deficiencies in the conduct of the trials or other problems with our product candidates. Other risks associated with our product candidates are described in the section entitled “Risk Factors” in our Annual Report on Form 10-K for the year ended December 31, 2021, as updated by the section entitled “Risk Factors” in this Quarterly Report on Form 10-Q and from time to time in our other periodic and current reports filed with the SEC.

### **Liquidity and Capital Resources**

From inception through June 30, 2022, we have financed the Company primarily through the issuance of public and private offerings of our common stock and other securities, and to a far lesser extent, through proceeds from grants from government agencies and foundations. From the date of approval of CAPLYTA through June 30, 2022, we have collected approximately \$192.6 million from product sales, which we believe will increase going forward. We do not believe that grant revenue will be a significant source of funding in the future.

On January 7, 2022, we completed a public offering of common stock in which we sold 10,952,381 shares of common stock at a public offering price of \$42.00 per share for aggregate gross proceeds of \$460.0 million. After deducting underwriting discounts, commissions and offering expenses, the net proceeds to the Company were approximately \$433.7 million.

As of June 30, 2022, we had a total of approximately \$679.2 million in cash and cash equivalents, available-for-sale investment securities and restricted cash, and approximately \$70.6 million of short-term liabilities consisting entirely of liabilities from operations, including approximately \$7.7 million of short-term lease obligations. In the six months ended June 30, 2022, we spent approximately \$260 million in cash for operations and equipment. During this period, we collected \$88 million from product sales and \$1.8 million of interest income and cost of product sales was \$7.8 million which resulted in \$178.3 million of net cash used for operations and equipment. The use of cash was primarily for selling and marketing costs in connection with our commercialization of CAPLYTA, conducting clinical trials and non-clinical testing, funding recurring operating expenses, and product manufacturing.

Based on our current operating plans, we expect that our existing cash, cash equivalents and marketable securities will enable us to fund our operating expenses and capital expenditure requirements for at least the next 12 months from the filing date of this quarterly report. During that time, we expect that our expenses will increase, primarily due to the commercialization of CAPLYTA for the treatment of schizophrenia and bipolar depression; the development of lumateperone in our late-stage clinical programs; the development of our other product candidates, including ITI-214; the continuation of manufacturing activities for anticipated future product sales and in connection with the development of lumateperone; and infrastructure expansion and general operations.

For the first six months of 2022, we used approximately \$260 million for operating costs. We expect to spend up to \$240 million in the second half of 2022, primarily for the marketing and commercialization of CAPLYTA, lumateperone clinical development including clinical trial conduct, regulatory activities, manufacturing and inventory production, expansion of our administrative infrastructure and other development activities. Our other development activities will include efforts relating to our ITI-1284, lenrispodun and ITI-333 programs, among others. However, the COVID-19 pandemic may negatively impact our commercialization of CAPLYTA, our ability to complete our ongoing or planned nonclinical and clinical trials, our ability to obtain approval of any product candidates from the FDA or other regulatory authorities, and our workforce and therefore our research, development and commercialization activities. This may ultimately have a material adverse effect on our liquidity, although we are unable to make any prediction with certainty given the rapidly changing nature of the pandemic and governmental and other responses to it.

We may require additional financing in the future to continue to fund our operations. We believe that we have the funding in place to commercialize CAPLYTA in patients with schizophrenia and bipolar depression. We also plan to fund additional clinical

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trials of lumateperone for the treatment of depressive disorders and other CNS disorders; preclinical and clinical development of our ITI-007 long acting injectable development program; clinical development of ITI-1284; continued clinical development of our PDE product candidates, including lenrispodun; research and preclinical development of our other product candidates. We have incurred losses in every year since inception with the exception of 2011, when we received an up-front fee and a milestone payment related to a license agreement that has been terminated. These losses have resulted in significant cash used in operations.

We seek to balance the level of cash, cash equivalents and investments on hand with our projected needs and to allow us to withstand periods of uncertainty relative to the availability of funding on favorable terms. Subject to our ability to generate significant revenues from operations, we may need to satisfy our future cash needs through public or private sales of our equity securities, sales of debt securities, incurrence of debt from commercial lenders, strategic collaborations, licensing a portion or all of our product candidates and technology and, to a lesser extent, grant funding. On January 6, 2020, we filed an automatic shelf registration statement on Form S-3 with the SEC, which became effective upon filing, on which we registered for sale an unlimited amount of any combination of its common stock, preferred stock, debt securities, warrants, rights, and/or units from time to time and at prices and on terms that we may determine, so long as we continue to satisfy the requirements of a “well-known seasoned issuer” under SEC rules. This registration statement will remain in effect for up to three years from the date it became effective.

We cannot be sure that future funding will be available to us when we need it on terms that are acceptable to us, or at all. We sell securities and incur debt when the terms of such transactions are deemed favorable to us and as necessary to fund our current and projected cash needs. The amount of funding we raise through sales of our common stock or other securities depends on many factors, including, but not limited to, the magnitude of sales of CAPLYTA, the status and progress of our product development programs, projected cash needs, availability of funding from other sources, our stock price and the status of the capital markets. Due to the volatile nature of the financial markets, equity and debt financing may be difficult to obtain. Additionally, the uncertain market conditions and continued spread of COVID-19 may limit our ability to access any financing. In addition, any unfavorable results in the commercialization of CAPLYTA and unfavorable development or delay in the progress of our lumateperone program could have a material adverse impact on our ability to raise additional capital.

To the extent that we raise additional capital through the sale of equity or convertible debt securities, the ownership interest of our existing stockholders will be diluted, and the terms may include liquidation or other preferences that adversely affect the rights of our stockholders. Debt financing, if available, may involve agreements that include covenants limiting or restricting our ability to take specific actions, such as incurring debt, making capital expenditures or declaring dividends. If we raise additional funds through government or other third-party funding, marketing and distribution arrangements or other collaborations, strategic alliances or licensing arrangements with third parties, we may have to relinquish valuable rights to our technologies, future revenue streams, research programs or product candidates or to grant licenses on terms that may not be favorable to us.

If adequate funds are not available to us on a timely basis, we may be required to: (1) delay, limit, reduce or terminate nonclinical studies, clinical trials or other clinical development activities for one or more of our product candidates, including our lead product candidate lumateperone, lenrispodun, and our other product candidates; (2) delay, limit, reduce or terminate our discovery research or pre-clinical development activities; (3) enter into licenses or other arrangements with third parties on terms that may be unfavorable to us or sell, license or relinquish rights to develop or commercialize our product candidates, technologies or intellectual property at an earlier stage of development and on less favorable terms than we would otherwise agree; or (4) limit or reduce commercialization efforts related to CAPLYTA.

Our cash is maintained in checking accounts, money market accounts, money market mutual funds, U.S. government agency securities, certificates of deposit, commercial paper, corporate notes and corporate bonds at major financial institutions. Due to the recent history of low interest rates available for these instruments, we have been earning limited interest income. During the six months ended June 30, 2022, interest rates have risen. This trend has resulted in approximately \$4.8 million of unrealized loss on investments during the six months ended June 30, 2022. Due to the short-term nature of these investments, we do not expect to recognize these losses. Even with the rise or further potential rise in interest rates, we do not expect interest income to be a significant source of funding over the next several quarters. In addition, our investment portfolio historically has not been adversely impacted by problems in the credit markets, but there can be no assurance that our investment portfolio will not be adversely affected in the future.

In 2014, we entered into a lease of 16,753 square feet of useable laboratory and office space located at 430 East 29th Street, New York, New York 10016. In September 2018, we further amended the lease to obtain an additional 15,534 square feet of office space beginning October 1, 2018 and to extend the term of the lease for previously acquired space. The lease, as amended, has a term of 14.3 years ending in May 2029. On May 17, 2019, we entered into a vehicle fleet lease with a company to acquire motor vehicles for certain employees. The vehicle fleet lease provides for individual leases for the vehicles, which at each lease commencement was determined to qualify for operating lease treatment. We began leasing vehicles under the vehicle fleet lease in March 2020. Restricted cash of \$1.4 million on our condensed consolidated balance sheet as of June 30, 2022 and December 31, 2021 relates to a letter of credit issued as part of the vehicle fleet lease.

### **Critical Accounting Policies and Estimates**

Our critical accounting policies are those policies which require the most significant judgments and estimates in the preparation of our condensed consolidated financial statements. We evaluate our estimates, judgments, and assumptions on an ongoing basis. Actual results may differ from these estimates under different assumptions or conditions. A summary of our critical accounting policies is presented in Part II, Item 7, of our Annual Report on Form 10-K for the year ended December 31, 2021. There have been no material changes to our critical accounting policies during the six months ended June 30, 2022.

The discussion and analysis of our financial condition and results of operations are based on our financial statements, which have been prepared in accordance with accounting principles generally accepted in the United States, or U.S. GAAP. The preparation of these financial statements requires management to make estimates and assumptions that affect reported amounts of assets and liabilities as of the date of the balance sheet and reported amounts of revenues and expenses for the periods presented. Judgments must also be made about the disclosure of contingent liabilities. We base our estimates on historical experience and on various other assumptions that we believe to be reasonable under the circumstances. These estimates and assumptions form the basis for making judgments about the carrying values of assets and liabilities that are not readily apparent from other sources. Management makes estimates and exercises judgment in research and development, including clinical trial accruals. Actual results may differ from those estimates and under different assumptions or conditions.

### **Recently Issued Accounting Pronouncements**

We review new accounting standards to determine the expected financial impact, if any, that the adoption of each such standard will have. For the recently issued accounting standards that we believe may have an impact on our financial statements, see “Recent Accounting Pronouncements” in Note 2 to our unaudited condensed consolidated financial statements included elsewhere in this Quarterly Report on Form 10-Q, and “Recently Issued Accounting Standards” in Note 2 to our audited consolidated financial statements and “Recently Issued Accounting Pronouncements” in Part II, Item 7, in our Annual Report on Form 10-K for the year ended December 31, 2021 filed on March 1, 2022.

### **Certain Factors That May Affect Future Results of Operations**

The SEC encourages companies to disclose forward-looking information so that investors can better understand a company’s future prospects and make informed investment decisions. This Quarterly Report on Form 10-Q contains such “forward-looking statements” within the meaning of the Private Securities Litigation Reform Act of 1995. These statements involve known and unknown risks, uncertainties and other important factors which may cause our actual results, performance or achievements to be materially different from any future results, performances or achievements expressed or implied by the forward-looking statements. Forward-looking statements include, but are not limited to, statements about: the accuracy of our estimates regarding expenses, revenues, uses of cash, cash equivalents and investment securities, capital requirements and the need for additional financing; our expectations regarding our commercialization of CAPLYTA, including the impact of COVID-19 on the commercialization of CAPLYTA and our ability to adapt our approach as appropriate; the duration and severity of the COVID-19 pandemic and its impact on our business; the supply and availability of and demand for our product; the initiation, cost, timing, progress and results of our development activities, non-clinical studies and clinical trials; the timing of and our ability to obtain and maintain regulatory approval, or submit an application for regulatory approval, of lumateperone and our other existing product candidates, any product candidates that we may develop, and any related restrictions, limitations, and/or warnings in the label of any approved product candidates; our plans to research, develop and commercialize lumateperone and our other current and future product candidates; the election by any collaborator to pursue research, development and commercialization activities; our ability to obtain future reimbursement and/or milestone payments from our collaborators; our ability to attract collaborators with development, regulatory and commercialization expertise; our ability to obtain and maintain intellectual property protection for our product candidates; our ability to successfully commercialize lumateperone and our other product candidates; the performance of our third-party suppliers and manufacturers and our ability to obtain alternative sources of raw materials; our ability to obtain additional financing; our use of the proceeds from our securities offerings; our exposure to investment risk, interest rate risk, inflation risk and capital market risk; and our ability to attract and retain key scientific, management, or sales and marketing personnel.

Words such as “may,” “anticipate,” “estimate,” “expect,” “may,” “project,” “intend,” “plan,” “believe,” “potential,” “predict,” “project,” “likely,” “will,” “would,” “could,” “should,” “continue” and words and terms of similar substance used in connection with any discussion of future operating or financial performance, identify forward-looking statements. All forward-looking statements are management’s present expectations of future events and are subject to a number of risks and uncertainties that could cause actual results to differ materially and adversely from those described in the forward-looking statements. These risks and uncertainties



include, but are not limited to, the following: there are no guarantees that CAPLYTA will be commercially successful; we may encounter issues, delays or other challenges in commercializing CAPLYTA; the COVID-19 pandemic may negatively impact our commercial plans and sales for CAPLYTA; the COVID-19 pandemic may negatively impact the conduct of, and the timing of enrollment, completion and reporting with respect to, our clinical trials; whether CAPLYTA receives adequate reimbursement from third-party payors; the degree to which CAPLYTA receives acceptance from patients and physicians for its approved indications; challenges associated with execution of our sales activities, which in each case could limit the potential of our product; results achieved in CAPLYTA in the treatment of schizophrenia and bipolar depression following commercial launch of the product may be different than observed in clinical trials, and may vary among patients; any other impacts on our business as a result of or related to the COVID-19 pandemic; challenges associated with supply and manufacturing activities, which in each case could limit our sales and the availability of our product; impacts on our business, including on the commercialization of CAPLYTA and our clinical trials, as a result of the conflict in Ukraine; risks associated with our current and planned clinical trials; we may encounter unexpected safety or tolerability issues with CAPLYTA following commercial launch for the treatment of schizophrenia or bipolar depression or in ongoing or future trials and other development activities; our other product candidates may not be successful or may take longer and be more costly than anticipated; product candidates that appeared promising in earlier research and clinical trials may not demonstrate safety and/or efficacy in larger-scale or later clinical trials or in clinical trials for other indications; our proposals with respect to the regulatory path for our product candidates may not be acceptable to the FDA; our reliance on collaborative partners and other third parties for development, commercialization, manufacturing or supply of our product and product candidates; and the other risk factors detailed under the heading “Risk Factors” in our most recent Annual Report on Form 10-K, as updated under the heading “Risk Factors” from time to time in our subsequent periodic and current reports filed with the SEC.

In light of these assumptions, risks and uncertainties, the results and events discussed in the forward-looking statements contained in this Quarterly Report on Form 10-Q or in any document incorporated by reference might not occur. Stockholders are cautioned not to place undue reliance on the forward-looking statements, which speak only as of the date of this Quarterly Report on Form 10-Q. We are not under any obligation, and we expressly disclaim any obligation, to update or alter any forward-looking statements, whether as a result of new information, future events or otherwise. All subsequent forward-looking statements attributable to the Company or to any person acting on its behalf are expressly qualified in their entirety by the cautionary statements contained or referred to in this section.

### **Item 3. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK**

*Interest Rate Sensitivity.* As of June 30, 2022, we had cash, cash equivalents, marketable securities and restricted cash of approximately \$679.2 million consisting of cash deposited in a highly rated financial institution in the United States and in a short-term U.S. Treasury money market fund, as well as high-grade corporate bonds and commercial paper. The primary objective of our investment activities is to preserve our capital for the purpose of funding operations and we do not enter into investments for trading or speculative purposes. We believe that we do not have material exposure to high-risk investments such as mortgage-backed securities, auction rate securities or other special investment vehicles within our money-market fund investments. We believe that we do not have any material exposure to changes in fair value as a result of changes in interest rates, although the recent increase in interest rates has resulted in our unrealized loss on investments, net, as of June 30, 2022 of approximately \$4.8 million and an unrealized loss on investments, net, as of December 2021 totaling approximately \$0.4 million. We plan on holding those investments to maturity, and should interest rates rise, there would be no recognition of impairment required. Declines in interest rates, however, would reduce future investment income.

*Inflation Risk.* Inflation generally may affect us by increasing clinical trial and other operational costs. To date, inflation has not had a material impact on our business, but if the global inflationary trends continue, we expect appreciable increases in clinical trial, selling, labor, and other operating costs. If our costs were to become subject to significant inflationary pressures, we may not be able to fully offset such higher costs through price increases of our product. Our inability or failure to do so could adversely affect our business, financial condition and results of operations.

*Capital Market Risk.* Although we receive product revenues from commercial sales of CAPLYTA, we continue to depend on funds raised through other sources. One possible source of funding is through further equity offerings. Our ability to raise funds in this manner depends upon capital market forces affecting our stock price.

### **Item 4. CONTROLS AND PROCEDURES**

(a) *Evaluation of Disclosure Controls and Procedures.* Our principal executive officer and principal financial officer, after evaluating the effectiveness of our disclosure controls and procedures (as defined in Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934, as amended, or the Exchange Act) as of the end of the period covered by this Form 10-Q, have concluded that, based on such evaluation, our disclosure controls and procedures were effective at a reasonable assurance level to ensure that information required to be disclosed by us in the reports that we file or submit under the Exchange Act is recorded,

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processed, summarized and reported within the time periods specified in the SEC's rules and forms, and is accumulated and communicated to our management, including our principal executive and principal financial officers, or persons performing similar functions, as appropriate, to allow timely decisions regarding required disclosure.

(b) *Changes in Internal Controls.* There were no changes in our internal control over financial reporting identified in connection with the evaluation of such internal control that occurred during the three months ended June 30, 2022 that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

## PART II: OTHER INFORMATION

### Item 1. LEGAL PROCEEDINGS

On July 8, 2022, a purported shareholder derivative complaint was filed in the Supreme Court for the State of New York against the directors serving on the Company's board of directors and the Company as a nominal defendant alleging breach of fiduciary duty and unjust enrichment alleging that compensation awarded to the director defendants for service on the board of directors was excessive. We believe that the complaint is without merit. At present, we are unable to estimate potential losses, if any, related to the lawsuit.

### Item 1A. RISK FACTORS

There have been no material changes to the risk factors discussed in Item 1A. Risk Factors in our Annual Report on Form 10-K for the year ended December 31, 2021, filed with the Securities and Exchange Commission on March 1, 2022, except as set forth below for the following risk factor which was previously included in our Quarterly Report on Form 10-Q for the quarter ended March 31, 2022.

***We rely on third parties to conduct our clinical trials and perform data collection and analysis, which may result in costs and delays that prevent us from successfully commercializing our product candidates.***

Although we design and manage our current preclinical studies and clinical trials, we do not now have the ability to conduct clinical trials for our product candidates on our own. In addition to our collaborators, we rely on contract research organizations, medical institutions, clinical investigators, and contract laboratories to perform data collection and analysis and other aspects of our clinical trials. In addition, we also rely on third parties to assist with our preclinical studies, including studies regarding biological activity, safety, absorption, metabolism, and excretion of product candidates.

Our preclinical activities or clinical trials may be delayed, suspended, or terminated if: the quality or accuracy of the data obtained by the third parties on whom we rely is compromised due to their failure to adhere to our clinical protocols or regulatory requirements or if for other reasons, these third parties do not successfully carry out their contractual duties or fail to meet regulatory obligations or expected deadlines, or these third parties need to be replaced.

If the third parties on whom we rely fail to perform, our development costs may increase, our ability to obtain regulatory approval, and consequently, to commercialize our product candidates may be delayed or prevented altogether. We currently use several contract research organizations to perform services for our preclinical studies and clinical trials. While we believe that there are numerous alternative sources to provide these services, in the event that we seek such alternative sources, we may not be able to enter into replacement arrangements without delays or incurring additional expenses.

Further, we are currently conducting clinical trials for our product candidates in many countries, including the United States and Europe, including Russia and Ukraine, and may expand to other geographies. Timely enrollment of, completion of and reporting on our clinical trials is dependent upon these global clinical trial sites which are, or in the future may be, adversely affected by political instability or conflict. For example, in connection with Studies 501, 502, and 503 for MDD and Study 403 for major depressive episodes with mixed features in bipolar disorder, we have third-party clinical sites located in Ukraine and Russia, which are experiencing significant conflict. Political instability and conflict in Ukraine and Russia, and any other areas in the world where we have clinical operations, may delay our trials and negatively affect our business and operations in those regions.

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**Item 2. UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS**

**Unregistered Sales of Equity Securities**

Not applicable.

**Issuer Purchases of Equity Securities**

We did not repurchase any of our equity securities during the quarter ended June 30, 2022.

**Item 3. DEFAULTS UPON SENIOR SECURITIES**

Not applicable.

**Item 4. MINE SAFETY DISCLOSURES**

Not applicable.

**Item 5. OTHER INFORMATION**

Not applicable.

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### Item 6. EXHIBITS

<u>Exhibit Number</u>	<u>Exhibit Description</u>	<u>Filed Herewith</u>	<u>Incorporated by Reference herein from Form or Schedule</u>	<u>Filing Date</u>	<u>SEC File/Reg. Number</u>
31.1	<a href="#">Certification of the Registrant's Principal Executive Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.</a>	X			
31.2	<a href="#">Certification of the Registrant's Principal Financial Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.</a>	X			
32	<a href="#">Certification of Principal Executive Officer and Principal Financial Officer pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.</a>	X			
101	The following materials from the Registrant's Quarterly Report on Form 10-Q for the quarter ended June 30, 2022, formatted in Inline XBRL (eXtensible Business Reporting Language): (i) Condensed Consolidated Balance Sheets as of June 30, 2022 (unaudited) and December 31, 2021 (audited), (ii) Condensed Consolidated Statements of Operations (unaudited) for the three and six months ended June 30, 2022 and 2021, (iii) Condensed Consolidated Statements of Comprehensive Loss (unaudited) for the three and six months ended June 30, 2022 and 2021, (iv) Condensed Consolidated Statements of Stockholders' Equity (unaudited) for the three and six months ended June 30, 2022 and 2021, (v) Condensed Consolidated Statements of Cash Flows (unaudited) for the six months ended June 30, 2022 and 2021, and (vi) Notes to Condensed Consolidated Financial Statements (unaudited).	X			
104	Cover Page Interactive Data File (formatted as Inline XBRL and contained in Exhibit 101).	X			

**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 thereunto duly authorized.

INTRA-CELLULAR THERAPIES, INC.

Date: August 9, 2022

By: /s/ Sharon Mates, Ph.D.

Sharon Mates, Ph.D.

Chairman, President and Chief Executive Officer

Date: August 9, 2022

By: /s/ Lawrence J. Hinline

Lawrence J. Hinline

Senior Vice President of Finance and Chief Financial Officer

## CERTIFICATIONS UNDER SECTION 302

I, Sharon Mates, Ph.D., certify that:

1. I have reviewed this quarterly report on Form 10-Q of Intra-Cellular Therapies, Inc.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
  - a) designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
  - b) designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
  - c) evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
  - d) disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
5. The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
  - a) all significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
  - b) any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: August 9, 2022

/s/ Sharon Mates, Ph.D.

Sharon Mates, Ph.D.

Chairman, President and Chief Executive Officer  
(principal executive officer)

## CERTIFICATIONS UNDER SECTION 302

I, Lawrence J. Hinline, certify that:

1. I have reviewed this quarterly report on Form 10-Q of Intra-Cellular Therapies, Inc.;

2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;

4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:

a) designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

b) designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

c) evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and

d) disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

5. The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):

a) all significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and

b) any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: August 9, 2022

/s/ Lawrence J. Hinline

Lawrence J. Hinline  
Senior Vice President of Finance and Chief Financial Officer  
(principal financial officer)

## CERTIFICATIONS UNDER SECTION 906

Pursuant to section 906 of the Sarbanes-Oxley Act of 2002 (subsections (a) and (b) of section 1350, chapter 63 of title 18, United States Code), each of the undersigned officers of Intra-Cellular Therapies, Inc., a Delaware corporation (the “Company”), does hereby certify, to such officer’s knowledge, that:

The Quarterly Report for the quarter ended June 30, 2022 (the “Form 10-Q”) of the Company fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934, and the information contained in the Form 10-Q fairly presents, in all material respects, the financial condition and results of operations of the Company.

Dated: August 9, 2022

/s/ Sharon Mates, Ph.D.

Sharon Mates, Ph.D.

Chairman, President and Chief Executive Officer  
(principal executive officer)

Dated: August 9, 2022

/s/ Lawrence J. Hinline

Lawrence J. Hinline

Senior Vice President of Finance and Chief Financial Officer  
(principal financial officer)