UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, DC 20549

FORM 10-Q

(Mark One)

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

For the quarterly period ended March 31, 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: 001-39696

COMPASS THERAPEUTICS, INC.

(Exact Name of Registrant as Specified in its Charter)

Delaware	**82-4876496**
(State or other jurisdiction of incorporation or organization)	**(I.R.S. Employer Identification No.)**
80 Guest St., Suite 601 Boston, Massachusetts	**02135**
(Address of principal executive offices)	**(Zip Code)**

Registrant's telephone number, including area code: (617) 500-8099

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

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, $0.0001 par value per share	CMPX	NASDAQ Capital Market

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, 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 May 6, 2022, the registrant had 101,283,924 shares of common stock, $0.0001 par value per share, outstanding.

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Item 1. Financial Statements

Compass Therapeutics, Inc. and Subsidiaries
Condensed Consolidated Balance Sheets
(In thousands, except par value)

	March 31, 2022 (unaudited)	December 31, 2021 (Note 1)
Assets		
Current assets:		
Cash and cash equivalents	$ 136,379	$ 144,514
Prepaid expenses and other current assets	3,904	2,591
Total current assets	140,283	147,105
Property and equipment, net	2,142	2,243
Operating lease, right-of-use ("ROU") asset	3,819	4,089
Other assets	320	320
Total assets	$ 146,564	$ 153,757
Liabilities and Stockholders' Equity		
Current liabilities:		
Accounts payable	$ 257	$ 867
Accrued expenses	8,050	8,775
Operating lease obligations, current portion	1,027	989
Total current liabilities	9,334	10,631
Operating lease obligations, long-term portion	2,740	3,048
Total liabilities	12,074	13,679
Commitments and contingencies (Note 6)		
Stockholders' equity:		
Preferred stock, $0.0001 par value; 10,000 shares authorized and no shares issued and outstanding as of March 31, 2022 and December 31, 2021	—	—
Common stock, $0.0001 par value: 300,000 shares authorized; 101,286 and 101,303 shares issued at March 31, 2022 and December 31, 2021, respectively; 100,905 and 100,832 shares outstanding at March 31, 2022 and December 31, 2021, respectively	10	10
Additional paid-in-capital	375,231	373,657
Accumulated deficit	(240,751)	(233,589)
Total stockholders' equity	134,490	140,078
Total liabilities and stockholders' equity	$ 146,564	$ 153,757

The accompanying notes are an integral part of these unaudited condensed consolidated financial statements.

Compass Therapeutics, Inc. and Subsidiaries
Condensed Consolidated Statements of Operations (Unaudited)
(In thousands, except per share data)

	Three Months Ended March 31,	
	2022	**2021**
Operating expenses:		
Research and development	$ 4,415	$ 4,704
General and administrative	2,767	2,635
Total operating expenses	7,182	7,339
Loss from operations	(7,182)	(7,339)
Other income (expense)	20	(83)
Loss before income tax expense	(7,162)	(7,422)
Income tax expense	—	—
Net loss	$ (7,162)	$ (7,422)
Net loss per share - basic and diluted	$ (0.07)	$ (0.14)
Basic and diluted weighted average shares outstanding	100,858	51,313

The accompanying notes are an integral part of these unaudited condensed consolidated financial statements.

Compass Therapeutics, Inc. and Subsidiaries
Condensed Consolidated Statements of Stockholders' Equity (Unaudited)
(In thousands)

	Common Units		Additional Paid-in	Accumulated	Total Stockholders'
	Shares	Amount	Capital	Deficit	Equity
Balance at December 31, 2021	100,832	$ 10	$ 373,657	$ (233,589)	$ 140,078
Vesting of share-based awards	73	—	—	—	—
Stock-based compensation	—	—	1,574	—	1,574
Net loss	—	—	—	(7,162)	(7,162)
Balance at March 31, 2022	100,905	$ 10	$ 375,231	$ (240,751)	$ 134,490

	Common Stock		Additional Paid-in	Accumulated	Total Stockholders'
	Shares	Amount	Capital	Deficit	Equity
Balance at December 31, 2020	51,221	$ 5	$ 191,348	$ (151,408)	$ 39,945
Vesting of share-based awards	92	—	—	—	—
Stock-based compensation	—	—	948	—	948
Net loss	—	—	—	(7,422)	(7,422)
Balance at March 31, 2021	51,313	$ 5	$ 192,296	$ (158,830)	$ 33,471

The accompanying notes are an integral part of these unaudited condensed consolidated financial statements.

Compass Therapeutics, Inc. and Subsidiaries
Condensed Consolidated Statements of Cash Flows (Unaudited)
(In thousands)

	For the Three Months Ended March 31,	
	2022	**2021**
Cash flows from operating activities:		
Net loss	$ (7,162)	$ (7,422)
Adjustments to reconcile net loss to net cash used in operating activities:		
Depreciation and amortization	165	157
Gain on disposal of equipment	—	(44)
Noncash interest expense	—	15
Share-based compensation	1,574	948
ROU asset amortization	270	256
Changes in operating assets and liabilities:		
Prepaid expenses and other current assets	(1,313)	1,173
Accounts payable	(611)	—
Accrued expenses	(595)	(442)
Operating lease liability	(270)	(246)
Net cash used in operating activities	(7,942)	(5,605)
Cash flows from investing activities:		
Purchases of property and equipment	(194)	(128)
Proceeds from sale of equipment	1	115
Net cash used in investing activities	(193)	(13)
Cash flows from financing activities:		
Repayment of borrowings under loan	—	(1,875)
Net cash used in financing activities	—	(1,875)
Net change in cash, cash equivalents and restricted cash	(8,135)	(7,493)
Cash, cash equivalents and restricted cash at beginning of period	144,514	47,339
Cash, cash equivalents and restricted cash at end of period	$ 136,379	$ 39,846
Supplemental disclosure of cash flow information		
Cash paid for interest	$ —	$ 137
Acquisition of equipment included in accrued expenses	$ 57	$ 161
ROU asset acquired through operating leases	$ —	$ 5,148

The accompanying notes are an integral part of these unaudited condensed consolidated financial statements.

1. Nature of Business and Basis of Presentation

Compass Therapeutics, Inc. ("Compass" or the "Company") is a clinical-stage, oncology-focused biopharmaceutical company developing proprietary antibody-based therapeutics to treat multiple human diseases. Our scientific focus is on the relationship between angiogenesis and the immune system. Our pipeline includes novel product candidates that leverage our understanding of the tumor microenvironment, including both angiogenesis-targeted agents and immune-oncology focused agents. These product candidates are designed to optimize critical components required for an effective anti-tumor response to cancer. These include modulation of the microvasculature via angiogenesis-targeted agents; induction of a potent immune response via activators on effector cells in the tumor microenvironment; and alleviation of immunosuppressive mechanisms used by tumors to evade immune surveillance. We plan to advance our product candidates through clinical development as both standalone therapies and in combination with our proprietary drug candidates as long as their continued development is supported by clinical and nonclinical data. References to Compass or the Company herein include Compass Therapeutics, Inc. and its wholly-owned subsidiaries. The Company was incorporated as Olivia Ventures, Inc. ("Olivia") in the State of Delaware on March 20, 2018. Prior to the Company's reverse merger with Compass Therapeutics LLC (the "Merger"), Olivia was a "shell company" (as defined in Rule 12b-2 of the Securities Exchange Act of 1934, as amended).

The Company is subject to risks and uncertainties common to companies in the biotechnology and pharmaceutical industries. There can be no assurance that the Company's research and development will be successfully completed, that adequate protection for the Company's technology will be obtained, that any products developed will obtain necessary government regulatory approval or that any approved products will be commercially viable. The Company operates in an environment of rapid change in technology and substantial competition from pharmaceutical and biotechnology companies. In addition, the Company is dependent upon the services of its employees and consultants.

In the opinion of management, the accompanying unaudited condensed consolidated financial statements include all normal and recurring adjustments (which consist primarily of accruals, estimates and assumptions that impact the financial statements) considered necessary to present fairly the Company's financial position as of March 31, 2022 and its results of operations and changes in stockholders' equity for the three months ended March 31, 2022 and 2021 and cash flows for the three months ended March 31, 2022 and 2021. Operating results for the three months ended March 31, 2022 are not necessarily indicative of the results that may be expected for the year ending December 31, 2022.

The unaudited condensed consolidated financial statements include the accounts of Compass Therapeutics, Inc. and its subsidiaries, and have been prepared by the Company in conformity with accounting principles generally accepted in the United States of America ("GAAP") and pursuant to the rules and regulations of the Securities and Exchange Commission ("SEC") for interim financial statements. Certain information and footnote disclosures normally included in financial statements prepared in accordance with GAAP have been condensed or omitted pursuant to such rules and regulations. The condensed consolidated balance sheet at December 31, 2021 has been derived from the audited consolidated financial statements at that date, but does not include all of the information and footnotes required by GAAP for complete financial statements. Accordingly, these condensed consolidated financial statements should be read in conjunction with the Company's audited financial statements in the Company's Annual Report on Form 10-K for the year ended December 31, 2021 (the "Annual Report").

Liquidity

Since our inception, we have devoted substantially all of our efforts to organizing and staffing our Company, business planning, raising capital, research and development activities, building our intellectual property portfolio and providing general and administrative support for these operations. To date, we have funded our operations primarily with proceeds from the sale of our equity securities and borrowings from debt arrangements. Through March 31, 2022, we have received $329.0 million in gross proceeds from the sale of equity securities. As of March 31, 2022, we had cash and cash equivalents of $136.4 million. Based on our research and development plans, we expect that such cash resources will enable us to fund our operating expenses and capital expenditure requirements into the second half of 2024.

COVID-19 Update

We have been carefully monitoring the COVID-19 pandemic and its potential impact on our business and have taken important steps to help ensure the safety of our employees and to reduce the spread of COVID-19 community-wide. We are ensuring that essential staffing levels at our operations remain in place, including maintaining key personnel in our laboratory facilities. We have implemented stringent safety measures designed to create a safe and clean environment for our employees as we continue to comply with applicable federal, state and local guidelines instituted in response to the COVID-19 pandemic.

There have been delays in sourcing of selected supplies required for the manufacturing of material to be used in our future clinical trials, and these delays have impacted and may continue to impact the timing of our future clinical trials. We expect that COVID-19 may continue to directly or indirectly impact (i) our employees and business operations or personnel at third-party suppliers and other vendors in the U.S. and other countries; (ii) the availability, cost or supply of materials; and (iii) the timeline for our ongoing clinical trial and potential future trials. We are continuing to assess the potential impact of the COVID-19 pandemic on our current and future business and operations, including our expenses and clinical trials, as well as on our industry and the healthcare system.

2. Summary of Significant Accounting Policies

There have been no material changes to the significant accounting policies previously disclosed in the Company's Annual Report, except as noted below.

Recently Adopted Accounting Pronouncements

The Company adopted ASU 2019-12, *Simplifying the Accounting for Income Taxes* on January 1, 2022. The Company accounts for income taxes pursuant to FASB ASC Topic 740, *Income Taxes*. Under FASB ASC Topic 740, deferred tax assets and liabilities are determined based on temporary differences between the bases of certain assets and liabilities for income tax and financial reporting purposes. The deferred tax assets and liabilities are classified according to the financial statement classification of the assets and liabilities generating the differences. The Company maintains a valuation allowance with respect to deferred tax assets. The Company establishes a valuation allowance based upon the potential likelihood of realizing the deferred tax asset and taking into consideration the Company's financial position and results of operations for the current period. Future realization of the deferred tax benefit depends on the existence of sufficient taxable income within the carry-forward period under the federal tax laws. The adoption of ASU 2019-12 did not have any impact on the Company's consolidated financial statement presentation or disclosures.

3. Fair Value Measurements

The following tables represent the Company's financial assets that are measured at fair value on a recurring basis and indicate the level of the fair value hierarchy utilized to determine such fair values (in thousands):

	Fair Value Measurements as of March 31, 2022 Using:			
	Quoted Prices in Active Markets for Identical Assets (Level 1)	Significant Other Observable Inputs (Level 2)	Significant Unobservable Inputs (Level 3)	Fair Value
Assets				
Cash equivalents - money market funds	$ 130,024	$ —	$ —	$ 130,024
Total assets	$ 130,024	$ —	$ —	$ 130,024

	Fair Value Measurements as of December 31, 2021 Using:			
	Quoted Prices in Active Markets for Identical Assets (Level 1)	Significant Other Observable Inputs (Level 2)	Significant Unobservable Inputs (Level 3)	Fair Value
Assets				
Cash equivalents - money market funds	$ 130,005	$ —	$ —	$ 130,005
Total assets	$ 130,005	$ —	$ —	$ 130,005

4. Property and Equipment

Property and equipment consist of the following (in thousands):

	March 31, 2022	December 31, 2021
Equipment	$ 5,345	$ 5,351
Furniture and fixtures	22	22
Leasehold improvements	1,586	1,531
Software	365	365
Total property and equipment–at cost	7,318	7,269
Less: Accumulated depreciation	(5,176)	(5,026)
Property and equipment, net	$ 2,142	$ 2,243

Depreciation expense for each of three months ended March 31, 2022 and 2021 was $0.2 million.

5. Accrued Expenses

Accrued expenses consist of the following (in thousands):

	March 31, 2022	December 31, 2021
Compensation and benefits	$ 490	$ 1,601
Project expenses	1,465	704
Accrued milestone	6,000	6,000
Other	95	470
Total accrued expenses	$ 8,050	$ 8,775

6. Commitments and Contingencies

Leases

The Company adopted Accounting Standards Update 2016-02, *Leases (Topic 842)*, or ASU 2016-02, effective January 1, 2021, using the modified retrospective transition method, in which the new standard is applied as of the date of initial adoption. The Company recognized and measured agreements executed prior to the date of initial adoption that were considered leases on January 1, 2021. No cumulative effect adjustment of initially applying the standard to the opening balance of retained earnings was made upon adoption. The Company elected the package of practical expedients permitted under the transition guidance that will retain the lease classification and initial direct costs for any leases that exist prior to adoption of the standard. In addition, the Company elected the accounting policy of not recording short-term leases with a lease term at the commencement date of 12 months or less on the condensed consolidated balance sheet as permitted by the new standard.

The Company has evaluated its leases and determined that it has one lease that is classified as an operating lease. The classification of this lease is consistent with the Company's determination under the previous accounting standard.

When available, the Company will use the rate implicit in the lease to discount lease payments to present value; however, the Company's current lease does not provide an implicit rate. Therefore, the Company used its incremental borrowing rate to discount the lease payments based on the date of the lease commencement.

The Company has one operating lease for its corporate office and laboratory facility ("Facility") that was signed in December 2020. The Company moved into the Facility in January 2021. The Facility lease has an initial term of four years and five months, beginning on January 1, 2021. The Facility lease contains scheduled rent increases over the lease term. The discount rate used for the Facility lease is 6.25%, and the remaining lease term of the Facility lease is three years and two months as of March 31, 2022.

The table below presents the undiscounted cash flows for the lease term. The undiscounted cash flows are reconciled to the operating lease liabilities recorded on the condensed consolidated balance sheet (000's):

Remainder of 2022	$	875
Years ending December 31,		
2023		1,345
2024		1,379
2025		543
Total minimum lease payments		4,142
Less: amount of lease payments representing interest		(375)
Present value of future minimum lease payments		3,767
Less: operating lease obligations, current portion		(1,027)
Operating lease obligations, long-term portion	$	2,740

Milestone payments

As part of the ABL Bio Agreement, the Company is obligated to pay certain development milestone payments. In the fourth quarter of 2021, the Company was notified of the completion of Phase 1 of the clinical trial for CTX-009. As a result, the Company is obligated to pay a $6.0 million milestone payment to ABL Bio on the deliverance of the final report related to the clinical trial. See Note 10 for additional information on the ABL Bio Agreement.

7. Stock-Based Compensation

Stock-based compensation expense for the three months ended March 31, 2022 and 2021 was classified in the condensed consolidated statement of operations as follows:

	Three Months Ended March 31,	
	2022	2021
	(000's)	
Research and development	$ 375	$ 139
General and administrative	1,199	809
Total	$ 1,574	$ 948

As of March 31, 2022, remaining unrecognized stock-based compensation cost from all plans to be recognized in future periods totaled $14.0 million.

Restricted Stock:

Prior to the adoption of the 2020 Plan, the Company issued restricted stock. A summary of the Company's restricted stock activity during the three months ended March 31, 2022 is as follows:

Weighted Average Fair Value	Shares (000's)	Fair Value Per Share
Unvested, December 31, 2021	471	$ 1.76
Granted	—	$ —
Vested	(73)	$ 1.82
Forfeited or canceled	(17)	$ 1.80
Unvested, March 31, 2022	381	$ 1.75

As of March 31, 2022, the total unrecognized compensation cost related to stock compensation expense for restricted stock is $0.7 million, expected to be recognized over a weighted average period of 1.6 years.

2020 Plan

In June 2020, the Company's board of directors adopted the 2020 Stock Option and Incentive Plan (the "2020 Plan") and reserved 2.9 million shares of common stock for issuance under this plan. The 2020 Plan includes automatic annual increases. The increase on January 1, 2022 was 4.2 million shares. As of March 31, 2022, 2.3 million shares remain available for future grant.

The 2020 Plan authorizes the board of directors or a committee of the board to grant incentive stock options, nonqualified stock options, restricted stock awards and restricted stock units ("RSUs") to eligible officers, employees, consultants and directors of the Company. Options generally vest over a period of four years and have a contractual life of ten years from the date of grant.

Stock Options:

The following table summarizes the stock option activity for the 2020 Plan:

	Number of Unvested Options	Weighted Average Exercise Price Per Share	Weighted Average Remaining Contractual Life (in years)	Aggregate Intrinsic Value (000's)
Outstanding at December 31, 2021	3,659	$ 4.99	8.67	
Granted	1,964	$ 2.31	9.88	
Exercise	—			
Forfeited/cancelled	(100)	$ 4.11		
Outstanding at March 31, 2022	5,523	$ 4.07	9.14	$ 6
Vested at March 31, 2022	1,847	$ 4.93	8.53	$ —

For the three months ended March 31, 2022, the weighted average grant date fair value for options granted was $1.77. There was no intrinsic value for options vested as of March 31, 2022. As of March 31, 2022, the total unrecognized compensation cost related to outstanding options was $9.2 million, to be recognized over a weighted average period of 3.1 years.

For the three months ended March 31, 2021, the weighted average grant date fair value for options granted was $3.58. There was no intrinsic value for options vested as of March 31, 2021.

The weighted average assumptions used in the Black-Scholes pricing model to determine the fair value of stock options granted during the three months ended March 31, 2022 and 2021 were as follows:

	Three Months Ended March 31,	
	2022	2021
Expected term (in years)	6.0	6.1
Risk-free rate	1.85 %	0.66 %
Expected volatility	94.0 %	85.9 %

RSUs:

The following table summarizes the RSU activity for the 2020 Plan:

	Shares (000's)	Weighted Average Price Per Share	Weighted Average Fair Value (000's)
Unvested, December 31, 2021	1,200	$ 3.83	$ 4,596
Granted	—	—	—
Vested	—	—	—
Forfeited or canceled	—	—	—
Unvested, March 31, 2022	1,200	$ 3.83	$ 4,596

Weighted average price per share is the weighted grant price based on the closing market price of each of the stock grants. The weighted average fair value is the weighted average share price times the number of shares.

As of March 31, 2022, remaining unrecognized compensation cost related to RSUs to be recognized in future periods totaled $4.2 million, which is expected to be recognized over a weighted average period of 3.6 years.

8. Related Parties and Related-Party Transactions

In connection with the acquisition of TRIGR Therapeutics, Inc. ("TRIGR") and upon consummation of the merger agreement on June 25, 2021, Miranda Toledano, who previously served as the Chief Financial Officer and Chief Operating Officer of TRIGR, was appointed to Compass Board of Directors as an independent director. Additionally, to facilitate the transition of CTX-009 from TRIGR, the Company entered into a consulting agreement with Ms. Toledano which ended in December 2021.

9. Other Income (Expense)

Other income (expense) consisted of the following:

	March 31,	
	2022	2021
	(000's)	
Interest income	$ 20	$ 15
Interest expense	—	(142)
Realized gain on disposal of equipment	—	44
Total other income (expense)	$ 20	$ (83)

10. License, Research and Collaboration Agreements

Collaboration Agreements

ABL Bio Corporation ("ABL Bio") Agreement

In November 2018, the Company's wholly-owned subsidiary (TRIGR) and ABL Bio, a South Korean biotechnology company, entered into an exclusive global (excluding South Korea) license agreement (the “TRIGR License Agreement”) which granted TRIGR a license to ABL001, ABL Bio’s bispecific antibody targeting DLL4 and VEGF-A (renamed CTX-009). Under the terms of the agreement, ABL Bio and TRIGR would jointly develop CTX-009, with ABL Bio responsible for development of CTX-009 throughout the end of Phase 1 clinical trials and TRIGR responsible for the development of CTX-009 from Phase 2 and onward. ABL Bio received a $5 million upfront payment and is eligible to receive up to $110 million of development and regulatory milestone payments, up to $295 million of commercial milestone payments and tiered single-digit royalties on net sales of CTX-009 in Oncology. ABL Bio is also eligible to receive up to $185 million in development, regulatory and commercial milestone payments and tiered, single-digit royalties on net sales of CTX-009 in Ophthalmology. The financial terms of the agreement were amended in May 2021 but remain substantially similar to the terms in the TRIGR License Agreement. As a result of the TRIGR acquisition in 2021, the TRIGR License Agreement was assigned to the Company and the Company has assumed all the rights and liabilities of the agreement.

In May 2021, TRIGR and ABL Bio terminated license agreements to several preclinical assets. As a result of the return of these assets to ABL Bio and termination of the license agreements, the Company is eligible to receive royalty payments if ABL Bio develops or licenses two bispecific antibodies that were previously licensed to TRIGR.

Adimab Agreement

The Company entered into a collaboration agreement with Adimab, LLC on October 16, 2014. The agreement includes provisions for payment of royalties at rates ranging in the single digits as a percentage of future net sales within a specified term from the first commercial sale. There were no milestone payments made during the first three months of 2022. As of March 31, 2022, future potential milestone payments in connection with this agreement amounted to $2.0 million.

Other License and Research Agreements

FUJIFILM Diosynth Biotechnologies ("Fujifilm") Agreement

The Company entered into a scope of work (“SOW”) under a master services agreement with Fujifilm on July 20, 2020. The Company made cash payments of $46 thousand and recorded $555 thousand in research and

development expense during the three months ended March 31, 2022 related to this agreement. The Company made cash payments of $0.2 million and recorded $1.0 million in research and development expense during the quarter ended March 31, 2021. As of March 31, 2022, future payments in connection with the SOW amounted to $1.2 million and prepayments for future work to be performed amounted to $1.2 million, for a total expense to be recognized of $2.4 million.

11. Subsequent events

None.

Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations.

The following discussion of the financial condition and results of operations of Compass Therapeutics, Inc. should be read in conjunction with the financial statements and the notes to those statements included in this Quarterly Report on Form 10-Q for the period ended March 31, 2022. Some of the information contained in this discussion and analysis, including information with respect to our plans and strategy for our business, includes forward-looking statements that involve risk, uncertainties and assumptions. You should read the "Risk Factors" section of this Quarterly Report on Form 10-Q and the "Risk Factors" section included in our Annual Report on Form 10-K for the fiscal year ended December 31, 2021*, for a discussion of important factors that could cause actual results to differ materially from the results described in or implied by the forward-looking statements contained in the following discussion and analysis.*

Overview

We are a clinical-stage, oncology-focused biopharmaceutical company developing proprietary antibody-based therapeutics to treat multiple human diseases. Our scientific focus is on the relationship between angiogenesis, the immune system, and tumor growth. Our pipeline of novel product candidates is designed to target multiple critical biological pathways required for an effective anti-tumor response. These include modulation of the microvasculature via angiogenesis-targeted agents, induction of a potent immune response via activators on effector cells in the tumor microenvironment, and alleviation of immunosuppressive mechanisms used by tumors to evade immune surveillance. We plan to advance our product candidates through clinical development as both standalone therapies and in combination with proprietary pipeline antibodies based on supportive clinical and nonclinical data.

On June 25, 2021, we consummated a definitive merger agreement (the "Merger Agreement") with Trigr Therapeutics, Inc. ("TRIGR"), a private biotechnology company. Pursuant to the Merger Agreement, through our wholly-owned subsidiaries and a two-step merger structure, we acquired all of the outstanding shares of TRIGR (the "TRIGR Merger"). Consideration payable to TRIGR shareholders at closing totaled an aggregate of 10,265,133 shares of our common stock (after giving effect to elimination of fractional shares that would otherwise be issued). In addition, TRIGR shareholders are eligible to receive up to $9 million, representing earnout payments which are dependent on certain events, including a $5 million earnout which is dependent on biologics license application approval of a product candidate acquired in the transaction, renamed CTX-009. As part of this $9 million in earnout payments to the TRIGR shareholders, $2 million represented a milestone payment that was remitted to TRIGR shareholders in the fourth quarter of 2021, with up to $7 million of eligible earnout payments remaining.

We currently have two product candidates in the clinical stage of development: CTX-009 and CTX-471. In addition, a third product candidate, CTX-8371, is expected to enter the clinic in the first half of 2023. A summary of these product candidates is presented below. We are also developing a portfolio of bispecific and monoclonal antibody product candidates which derive from our in-house antibody discovery and development platforms. For a more detailed description, see our Annual Report on Form 10-K for the fiscal year ended December 31, 2021.

CTX-009 (a.k.a. ABL001) - bispecific antibody targeting DLL4 and VEGF-A

CTX-009 is an investigational bispecific antibody that simultaneously blocks Delta-like ligand 4/Notch ("DLL4") and vascular endothelial growth factor A ("VEGF-A") signaling pathways, which are critical to angiogenesis and tumor vascularization. We have licensed exclusive global rights to CTX-009, outside of South Korea, from ABL Bio, Inc. ("ABL Bio"), a South Korea-based clinical-stage company focused on developing antibody therapeutics. South Korean rights are held by Handok Pharmaceuticals, Inc. ("Handok") and China rights were out-licensed from the Company to Elpiscience Biopharmaceuticals Co., Limited ("Elpiscience").

CTX-009 is undergoing clinical development in patients with advanced solid tumors. A Phase 1 dose escalation and dose expansion monotherapy study and a Phase 1b combination study of CTX-009 in combination with chemotherapy have been completed. In the first quarter of 2021, Handok commenced a Phase 2 study of CTX-009 in combination with paclitaxel in patients with biliary tract cancers ("BTC" or "cholangiocarcinoma") in South Korea. The study has been enrolling patients with unresectable advanced, metastatic, or relapsed BTCs who have received one or two prior systemic therapies. This Phase 2 study has a Simon Two-Stage design where 3 partial responses ("PRs") among the first 21 patients enrolled in the first stage of the study will advance the study to the second stage. As of October 2021, there were five PRs observed among the first 17 evaluable patients (see current status under

Q1 Program Updates below), and therefore, the criteria to advance the study to its second stage was met. The study is currently being conducted at four leading medical centers in South Korea.

We submitted an Investigational New Drug ("IND") application to the U.S. Food and Drug Administration ("FDA") in December 2021 to initiate a global Phase 2 study in the U.S. and South Korea. The FDA cleared our IND application in January 2022, enabling the Company to initiate a global Phase 2 clinical trial for CTX-009 in combination with paclitaxel in patients with advanced BTC in the United States and South Korea. Therefore, patients who were already enrolled in the study in South Korea, and additional patients who will be enrolled in the United States as well as South Korea, will all be included in this global Phase 2 clinical trial.

We plan on opening clinical sites in the United States in the second quarter of 2022 and initiating dosing of patients in these sites in the third quarter of 2022.

Q1 PROGRAM UPDATES

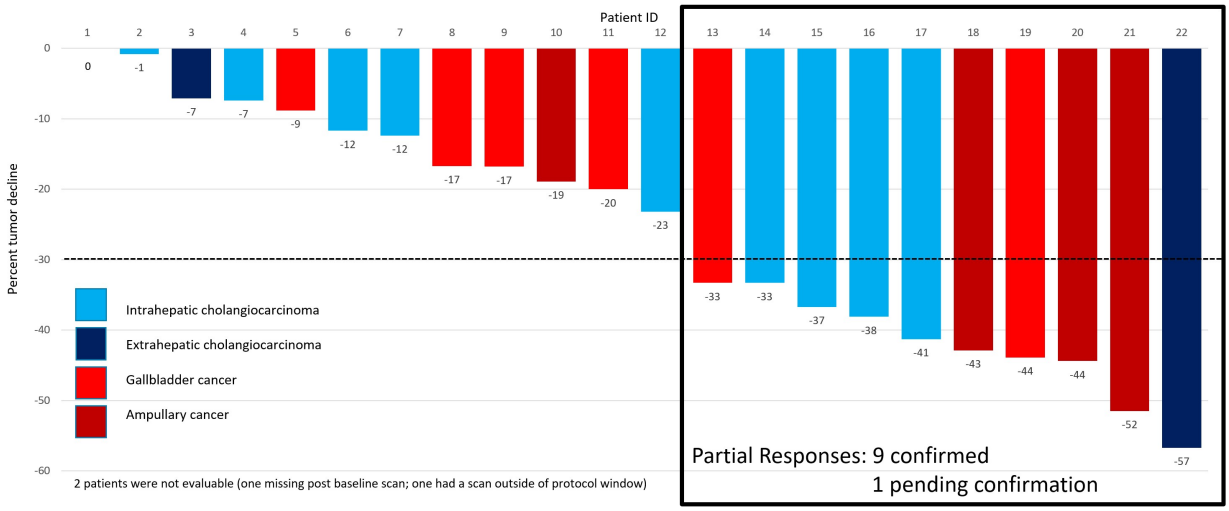
CTX-009 - bispecific antibody targeting DLL4 and VEGF-A

The Company reported the following interim data from its Phase 2 clinical trial of CTX-009 in combination with Paclitaxel in BTC on May 4, 2022. The press release and presentation were included with our current report 8-K filed on May 4, 2022.

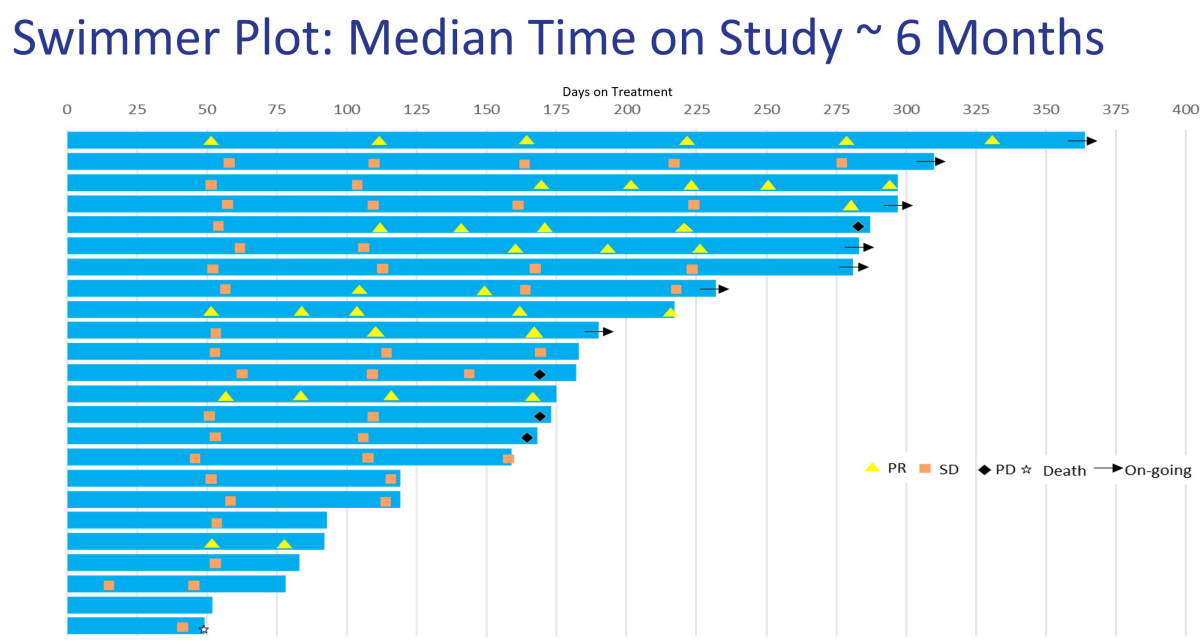
Preliminary Activity Data Summary

As of April 14, 2022, the first stage of the study has been fully enrolled and all 24 patients have been dosed. Of the 24 patients, there were 10 PRs, 9 of which have been confirmed by RECIST 1.1, leading to a preliminary overall response rate ("ORR") of 42%. Two patients are not evaluable for response, and 22 of the 24 patients have had stable disease or better leading to a clinical benefit rate ("CBR") of 92%. The current median time on study is approximately 6 months.

The interim waterfall plot below depicts the best response for 22 patients in the study as of April 14, 2022:



The swimmer plot below depicts the duration that each patient has been on treatment:



Preliminary Safety Data Summary

As of April 14, 2022, no formal safety data analysis has been completed, but CTX-009 in combination with paclitaxel was observed to be generally well-tolerated and the safety data are consistent with the Phase 1 studies with hypertension and neutropenia being the most common events related to CTX-009 and paclitaxel, respectively.

Of the 24 subjects enrolled in the study, all subjects had at least one AE related to CTX-009 and/or paclitaxel. The most common adverse events (all Grades) occurring in at least 3 patients were anemia (12.5%), asthenia (25.0%), fatigue (16.7%), edema (16.7%), pyrexia (16.7%), neutropenia (54.2%), thrombocytopenia (20.8%), headache (16.7%), proteinuria (20.8%), dysphonia (12.5%), dyspnea (25.0%), epistaxis (33.3%), pulmonary hypertension (16.7%, all Grade 1) and hypertension (50.0%).

Grade 3 or greater Adverse Events (“AEs”) that were determined to be probably or possibly related to CTX-009 and paclitaxel treatment included neutropenia (n=12; 50.0%), hypertension (n=4; 16.7%), anemia (n=3; 12.5%) and thrombocytopenia (n=2; 8.3%), which were all attributed to the concomitant chemotherapy agent (paclitaxel) with the exception of hypertension which was attributed to CTX-009. In addition, there were additional Grade 3 or greater events observed in no more than one patient: intestinal perforation, asthenia, catheter site hemorrhage, fatigue, cholangitis, abdominal infection, bacterial gastritis, pneumonia (which was fatal), post-procedure hemorrhage, decreased appetite, cerebral hemorrhage, proteinuria and embolism.

Development Plans for CTX-009

Our development strategy is to develop CTX-009 in all of the indications in which patients have a need for effective and novel therapeutic agents and data that supports the potential therapeutic benefit of CTX-009. We chose BTC as our lead indication due to a number of factors, including CTX-009 activity observed in the Phase 1b and in the ongoing Phase 2 clinical studies, lack of effective therapies for this patient population, and the potential for a straight-forward regulatory route to approval.

We plan to advance CTX-009 into a Phase 2/3 study in the second line setting in the general BTC patient population with ORR and duration of response (“DOR”) as potential endpoints for accelerated approval, with the goal of filing a

biologics license application ("BLA") in 2024. This goal is based on recent approvals of pemigatinib and infigratinib in cholangiocarcinoma patients who harbor the FGFR2 mutation (estimated to be 10-15% of the total BTC patient population - see study from Goyal, L. et al, Targeting FGFR inhibition in cholangiocarcinoma, Cancer Treatment Review, Vol. 95, Apr 2021), the response rate observed with CTX-009 in the general BTC patient population in the Phase 1b and the preliminary response rate observed in the Phase 2 combination trials summarized above.

The second indication we plan on pursuing is advanced colorectal cancer. Colorectal cancer patients who advance on front line therapy have very limited therapeutic options, and this patient population is significant in size and could be well-served by novel and effective therapies. Accordingly, we plan on initiating a Phase 2/3 monotherapy clinical trial of CTX-009 in the third line setting in colorectal cancer. Taking into consideration the activity of regorafenib, which was recently approved in the third line setting in colorectal cancer, and the activity seen with the small molecule KRAS G12C inhibitors, sotorasib and adagrasib in colorectal patients harboring the G12C mutation, and the response rates observed with CTX-009 in the Phase 1a monotherapy trial in colorectal cancer patients, we believe that we can advance CTX-009 into a Phase 2/3 study in the third line setting in colorectal cancer with ORR and DOR as potential endpoints for accelerated approval. We have not yet discussed this plan with regulatory agencies, including the FDA.

The timing of the initiation of the clinical trials in the United States depends, among other things, on the availability of clinical drug product for the studies, communications with the FDA, FDA allowance for each of the proposed studies to proceed and the availability of cash resources to support such trials. These plans remain preliminary and are subject to feedback from regulatory agencies. While our IND has been cleared for CTX-009 in patients with advanced BTC and we had initial communications with the FDA, we have not yet discussed with the FDA or other regulatory agencies the proposed design of the colorectal trial or other potential studies or the regulatory path for BLA submission for CTX-009.

We intend to explore the potential of CTX-009 in additional indications, based on data from pre-clinical models, potential biomarkers such as DLL4 and clinical data from CTX-009 studies providing hints of activity of CTX-009 in additional indications such as gastric cancer, pancreatic cancer, renal cell cancer, prostate cancer and ovarian cancer.

In addition, we are developing a plan to study the combination of CTX-009 with our novel bispecific checkpoint blocker, CTX-8371, and with other checkpoint blockers, such as pembrolizumab and nivolumab. Additionally, we plan to study the combination of CTX-009 with our novel CD137 agonistic antibody, CTX-471, which is currently undergoing a Phase 1b clinical trial in patients with advanced solid tumors.

CTX-471 - monoclonal antibody agonist of CD137

CTX-471, our monoclonal antibody product candidate, is a fully human, IgG4 monoclonal antibody that is an agonist of CD137, a key co-stimulatory receptor on immune cells. Binding of CTX-471 to CD137 has been observed to lead to ligand-stimulated activation of T-cells and NK cells. In tumor models, treatment with CTX-471 as a monotherapy led to recruitment and activation of immune cells in the tumor microenvironment. In the treated mice, dosing with CTX-471 led to extensive reprogramming of the tumor microenvironment, including increased recruitment of immune cells, reversion of exhausted cytotoxic CD8+ T-cells, reductions in immunosuppressive regulatory T-cells, and reductions in immunosuppressive tumor- associated macrophages. Long after the completion of the treatment with CTX-471, a period described as eight half-lives of the antibody, treated mice exhibited immune memory that prevented reestablishment of the same tumor.

In July 2019, we initiated a Phase 1 trial evaluating the safety and tolerability of CTX-471 as a monotherapy in oncology patients who were previously treated with PD-1 or PD-L1 immune checkpoint inhibitors and subsequently relapsed or progressed after a period of stable disease. The design of this trial includes a dose escalation stage (Phase 1a) followed by a dose expansion stage (Phase 1b). The dose-escalation stage of the Phase 1a trial has been completed and CTX-471 was observed to be generally well-tolerated.

The dose expansion stage of the trial is currently ongoing and nearing completion. As of February 25, 2022, 49 patients with 15 different cancers have been enrolled in the study and 38 of those patients are evaluable. Of the 38 evaluable patients, 3 patients had a PR; the first two have been confirmed by RECIST 1.1 and the third PR is unconfirmed. In addition, 19 patients have reached stable disease, leading to a preliminary ORR of 8% and a CBR of 58%. There has been one treatment-related serious adverse event (“SAE”) in the Phase 1b dose expansion stage of the trial. This event was identical to the dose-limiting toxicity seen in the Phase 1a study (thrombocytopenia with elevated liver function tests and elevated C-reactive protein).

The first PR observed in the study was in a patient with advanced small cell lung cancer who had a PR at week 17 and this response has been confirmed at week 25. This patient has now been dosed with CTX-471 for more than one year with a durable PR. In October 2021, a second PR was observed in a patient with metastatic melanoma who was previously treated with nivolumab and progressed on nivolumab. In December 2021, a third PR was observed in a patient with metastatic melanoma of mucosal origin who was previously treated with first-line regimen of ipilimumab plus nivolumab followed by second-line nivolumab as a monotherapy. After progressing on the prior regimens, the patient had joined CTX-471 study with multiple metastases at baseline. This patient reached a PR based on a 58% decline in linear tumor burden at week 17. This PR was not confirmed.

We expect to complete the dose expansion stage of our Phase 1 study in the third quarter of 2022. The results of this study will inform us on the next development steps. There are three studies that are currently been contemplated, and the decision which one(s) to pursue will be informed by the data of our clinical study. The first is a combination study with a PD-1/PD-L1 inhibitor in indications where PD-1 and/or PD-L1 alone are not currently approved. The second is as maintenance treatment for small cell lung cancer. The third is as a monotherapy in the post PD-1/PD-L1 patient population as a salvage therapy across multiple indications.

CTX-8371 - bispecific antibody that targets PD-1 and PD-L1

CTX-8371 is a bispecific antibody that binds to both PD-1 and PD-L1, the targets of well-known and widely used checkpoint inhibitor antibodies. Preclinical studies demonstrate that CTX-8371 has the ability to outperform PD-1, PD-L1, and combinations of the two to activate T-cells in in vitro assays. In mouse xenografts, treatment with CTX-8371 led to significantly greater tumor growth control and longer survival than treatment with a PD-1 inhibitor alone, a PD-L1 inhibitor alone or the combination of PD-1 and PD-L1 inhibitors. IND-enabling studies with CTX-8371 were initiated in August 2020 and are generally progressing well. Our contract development manufacturing organization, Fujifilm Diosynth Biotechnologies (see Note 10 to the financial statements contained in this Form 10-Q for further description of Fujifilm agreement) experienced delays with its supply chain management, leading to a delay in the GMP manufacturing of CTX-8371. In March 2022, the GMP manufacturing campaign started. Based on the timeline provided by Fujifilm for the GMP manufacturing campaign, we are currently targeting an IND submission for CTX-8371 in the first quarter of 2023.

Q1 PROGRAM UPDATE: CTX-8371

In April 2022, the Company presented preclinical data on CTX-8371 involving a unique mechanism of action ("MOA") that involves cleavage of cell surface PD-1, at the 2022 American Association for Cancer Research ("AACR") annual meeting. A summary of the results is as follows:

- Treatment with CTX-8371 leads to PD-1 loss from the surface of intra-tumoral T cells in tumor-bearing transgenic hPD-1/h-PD-L1 mice, and on peripheral blood T cells in cynomolgus monkeys. This unique MOA differentiates CTX-8371 from marketed inhibitors targeting either PD-1 or PD-L1.
- Clearance and half-life of CTX-8371 are within the expected ranges for a human IgG1 antibody in non-human primates (NHP) with a linear pharmacokinetics ("PK").
- Treatment with CTX-8371 in the aggressive MC38-hPD-L1 colorectal mouse model led to a dose-proportional reduction in tumor volume and a complete eradication of tumors at the highest dose.
- Taken together, the murine and cynomolgus monkey PK data, receptor occupancy data, and in vivo efficacy data in murine models will be used to calculate the predicted human efficacious dose range for CTX-8371.

Operating Activities

We have funded our operations primarily with proceeds from the sale of our equity securities. Through March 31, 2022, we have received $329.0 million in gross proceeds from the sale of equity securities.

We have incurred significant operating losses since inception and have not generated any revenue from the sale of products and we do not expect to generate any revenue from the sale of products in the near future, if at all. Our ability to generate product revenue sufficient to achieve profitability will depend heavily on the successful development and eventual commercialization of our treatments and any future product candidates. Our net losses were $7.2 million and $7.4 million for the three months ended March 31, 2022 and 2021, respectively. We had an accumulated deficit of $241 million at March 31, 2022. We expect to continue to incur significant expenses for at least the next several years as we advance through clinical development, develop additional product candidates and seek regulatory approval of any product candidates that complete clinical development. In addition, if we obtain marketing approval for any product candidates, we expect to incur significant commercialization expenses related to product manufacturing, marketing, sales and distribution. We may also incur expenses in connection with the in-licensing or acquisition of additional product candidates.

Until such time as we can generate significant revenue from product sales, if ever, we expect to finance our operations through equity and debt financings, or other capital sources, which may include collaborations with other companies or other strategic transactions. As of March 31, 2022, we had $136 million in cash and cash equivalents. With the proceeds from the underwritten offering in November 2021, and based on our research and development plans, we expect that such cash resources will enable us to fund our operating expenses and capital expenditure requirements into the second half of 2024.

Because of the numerous risks and uncertainties associated with product development, we are unable to predict the timing or amount of increased expenses or when, or if, we will be able to achieve or maintain profitability. Even if we are able to generate product sales, we may not become profitable. If we fail to become profitable or are unable to sustain profitability on a continuing basis, then we may be unable to continue our operations at planned levels and be forced to reduce or terminate our operations. Our financial statements do not include any adjustments that might result from the outcome of this uncertainty.

COVID-19 Update

We have been carefully monitoring the COVID-19 pandemic and its potential impact on our business and have taken important steps to help ensure the safety of our employees and to reduce the spread of COVID-19 community-wide. We are ensuring that essential staffing levels at our operations remain in place, including maintaining key personnel in our laboratory facilities. We have implemented stringent safety measures designed to create a safe and clean environment for our employees as we continue to comply with applicable federal, state and local guidelines instituted in response to the COVID-19 pandemic.

There have been delays in sourcing of selected supplies required for the manufacturing of material to be used in our future clinical trials, and these delays have impacted and may continue to impact the timing of our future clinical trials. We expect that COVID-19 may continue to directly or indirectly impact (i) our employees and business operations or personnel at third-party suppliers and other vendors in the U.S. and other countries; (ii) the availability, cost or supply of materials; and (iii) the timeline for our ongoing clinical trial and potential future trials. We are continuing to assess the potential impact of the COVID-19 pandemic on our current and future business and operations, including our expenses and clinical trials, as well as on our industry and the healthcare system.

Components of Results of Operations

Research and Development

Research and development expenses consist primarily of costs incurred in connection with the development of our product candidates, CTX-471, CTX-8371 and CTX-009, as well as unrelated discovery program expenses. We expense research and development costs as incurred. These expenses include:

- employee-related expenses including salaries, related benefits and equity-based compensation expense for employees engaged in research and development functions;
- expenses incurred under agreements with organizations that support our platform program development;

- Contract Manufacturing Organizations (“CMO”) that are primarily engaged to provide drug substance and product for our clinical trials, research and development programs, as well as investigative sites and consultants that conduct our clinical trials, nonclinical studies and other scientific development services;

- the cost of acquiring and manufacturing nonclinical and clinical trial materials, including manufacturing registration and validation batches;

- costs related to compliance with quality and regulatory requirements; and

- facilities and equipment expenses.

Advance payments that we make for goods or services to be received in the future for use in research and development activities are recorded as prepaid expenses. Such amounts are recognized as an expense as the goods are delivered or the related services are performed, or until it is no longer expected that the goods will be delivered or the services rendered.

Product candidates in later stages of clinical development generally have higher development costs than those in earlier stages of clinical development, primarily due to the increased size and duration of later-stage clinical trials. We expect that our research and development expenses will increase substantially in connection with our planned clinical development activities in the future. At this time, we cannot accurately estimate or know the nature, timing and costs of the efforts that will be necessary to complete the clinical development of any future product candidates.

The successful development and commercialization of product candidates is highly uncertain. This is due to the numerous risks and uncertainties associated with product development and commercialization.

General and Administrative Expenses

General and administrative expenses consist primarily of salaries and related costs for personnel in executive, finance, corporate and business development, and administrative functions. General and administrative expenses also include legal fees relating to patent and corporate matters; professional fees for accounting, auditing, tax and administrative consulting services; insurance costs; administrative travel expenses; marketing expenses and other operating costs.

We anticipate that our general and administrative expenses will increase in the future as we increase our headcount to support our business operations.

Other Income (expense)

Other income (expense) consists of interest expense, interest income and realized gains or losses on sales of furniture and equipment.

Results of Operations

Comparison of the Three Months Ended March 31, 2022 and 2021

The following table summarizes our results of operations for the three months ended March 31, 2022 and 2021:

	Three Months Ended March 31,		
	2022	2021	Change
		(000's)	
Operating expenses:			
Research and development	$ 4,415	$ 4,704	$ (289)
General and administrative	2,767	2,635	132
Total operating expenses	7,182	7,339	(157)
Loss from operations	(7,182)	(7,339)	157
Other income (expense), net	20	(83)	103
Loss before income tax expense	(7,162)	(7,422)	260
Income tax expense	—	—	—
Net loss	$ (7,162)	$ (7,422)	$ 260

Research and Development Expenses

Research and development expenses decreased by $0.3 million, or 6%, for the three months ended March 31, 2022 compared to the three months ended March 31, 2021. The decrease primarily came from a reduction in manufacturing expense of $0.3 million as compared to the same period in 2021.

We track outsourced development, outsourced personnel costs and other research and development costs of specific programs. Research and development expenses are summarized by program in the table below:

	Three Months Ended March 31,	
	2022	2021
	(000's)	
CTX-009	$ 576	$ —
CTX-471	1,296	998
CTX-8371	922	1,494
NKP30 cell engagement platform	—	117
Unallocated research and development expenses	1,621	2,095
Total research and development expenses	$ 4,415	$ 4,704

General and Administrative Expenses

General and administrative expenses increased by $0.1 million, or 5%, to $2.8 million for the three months ended March 31, 2022 as compared to the same period in 2021.

Other Income (Expense)

For the three months ended March 31, 2022, other income (expense) consists exclusively of interest income. For the three months ended March 31, 2021, the primary component was interest expense of $142 thousand related to a term loan facility with Pacific Western Bank, Inc. (the "Credit Facility") which we extinguished in the fourth quarter of 2021 (see our Annual Report on Form 10-K for the fiscal year ended December 31, 2021 for further information on the Credit Facility).

Income Tax Expense

During the three months ended March 31, 2022 and 2021, we recognized no income tax expense.

Liquidity and Capital Resources

Since our inception, we have devoted substantially all of our efforts to organizing and staffing our Company, business planning, raising capital, research and development activities, building our intellectual property portfolio and providing general and administrative support for these operations. We have funded our operations primarily with proceeds from the sale of our equity securities (in addition, we received borrowings from the Credit Facility, which was extinguished in the fourth quarter of 2021). Through March 31, 2022, we have received $329.0 million in gross proceeds from the sale of equity securities. As of March 31, 2022, we had cash and cash equivalents of $136 million.

Funding Requirements

Our primary use of cash is to fund operating expenses, primarily research and development expenditures. Cash used to fund operating expenses is impacted by the timing of when we pay these expenses, as reflected in the change in our outstanding accounts payable, accrued expenses and prepaid expenses. Because of the numerous risks and uncertainties associated with research, development and commercialization of pharmaceutical products, we are unable to estimate the exact amount of our operating capital requirements. Our future funding requirements will depend on many factors, including, but not limited to:

- the scope, timing, progress and results of discovery, preclinical development, laboratory testing and clinical trials for our product candidates;
- the costs of manufacturing our product candidates for clinical trials and in preparation for marketing approval and commercialization;
- the extent to which we enter into collaborations or other arrangements with additional third parties in order to further develop our product candidates;
- the costs of preparing, filing and prosecuting patent applications, maintaining and enforcing our intellectual property rights and defending intellectual property-related claims;
- the costs and fees associated with the discovery, acquisition or in-license of additional product candidates or technologies;
- our ability to establish additional collaborations on favorable terms, if at all;
- the costs required to scale up our clinical, regulatory and manufacturing capabilities;
- the costs of future commercialization activities, if any, including establishing sales, marketing, manufacturing and distribution capabilities, for any of our product candidates for which we receive marketing approval; and
- revenue, if any, received from commercial sales of our product candidates, should any of our product candidates receive marketing approval.

Until such time, if ever, as we can generate substantial product revenue, we expect to finance our operations through a combination of equity offerings, debt financings, collaborations, strategic alliances and marketing, distribution or licensing arrangements. To the extent that we raise additional capital through the sale of equity or convertible debt securities, current stockholders' interest will be diluted, and the terms of these securities may include liquidation or other preferences that adversely affect rights of common stockholders. Debt financing and preferred equity financing, if available, may involve agreements that include covenants limiting or restricting our ability to take specific actions, such as incurring additional debt, making acquisitions or capital expenditures or declaring dividends. If we raise additional funds through collaborations, strategic alliances or marketing, distribution or licensing arrangements with third parties, we may have to relinquish valuable rights to our technologies, future revenue streams, research programs or drug candidates, or grant licenses on terms that may not be favorable to us. If we are unable to raise additional funds through equity or debt financings or other arrangements when needed, we may be required to delay, limit, reduce or terminate our research, product development or future commercialization efforts, or grant rights to develop and market product candidates that we would otherwise prefer to develop and market ourselves.

Cash Flows

The following table shows a summary of our cash flows for the periods indicated:

	Three months Ended March 31,	
	2022	**2021**
	(000's)	
Cash used in operating activities	$ (7,942)	$ (5,605)
Cash used in investing activities	(193)	(13)
Cash used in financing activities	—	(1,875)
Net change in cash, cash equivalents and restricted cash	$ (8,135)	$ (7,493)

Operating Activities

During the three months ended March 31, 2022, we used $7.9 million of cash in operating activities, resulting from our net loss of $7.2 million, offset by non-cash charges of $2.0 million and the change in operating assets and liabilities of $2.8 million. Our non-cash charges are from share-based compensation expense of $1.6 million and depreciation and amortization (including ROU asset amortization) of $0.4 million.

During the three months ended March 31, 2021, we used $5.6 million of cash in operating activities, resulting from our net loss of $7.4 million, offset by non-cash charges of $1.3 million and the change in operating assets and liabilities of $0.5 million. Our non-cash charges were primarily comprised of depreciation and amortization of $0.2 million, share-based compensation expense of $0.9 million, offset by $44 thousand of gain on disposal of equipment.

Investing Activities

During the three months ended March 31, 2022, we used $0.2 million of cash in investing activities which primarily related to leasehold improvements. During the three months ended March 31, 2021, we used $13 thousand of cash in investing activities.

Financing Activities

We had no financing activities during the three months ended March 31, 2022. During the three months ended March 31, 2021, we had $1.9 million in payments under the Credit Facility.

Future Funding Requirements

We expect our expenses to increase substantially in connection with our ongoing activities. The timing and amount of our operating expenditures will depend largely on:

- the initiation, progress, timing, costs and results of clinical trials for our product candidate or any future product candidates we may develop;
- the initiation, progress, timing, costs and results of nonclinical studies for our product candidates or any future product candidates we may develop;

- our ability to maintain our relationships with key collaborators;
- the outcome, timing and cost of seeking and obtaining regulatory approvals from the FDA and comparable foreign regulatory authorities, including the potential for such authorities to require that we perform more nonclinical studies or clinical trials than those that we currently expect or change their requirements on studies that had previously been agreed to;
- the cost to establish, maintain, expand, enforce and defend the scope of our intellectual property portfolio, including the amount and timing of any payments we may be required to make, or that we may receive, in connection with licensing, preparing, filing, prosecuting, defending and enforcing any patents or other intellectual property rights;
- the effect of competing technological and market developments;
- the costs of continuing to grow our business, including hiring key personnel and maintain or acquiring operating space;
- market acceptance of any approved product candidates, including product pricing, as well as product coverage and the adequacy of reimbursement by third-party payors;
- the cost of acquiring, licensing or investing in additional businesses, products, product candidates and technologies;
- the cost and timing of selecting, auditing and potentially validating a manufacturing site for commercial-scale manufacturing;
- the cost of establishing sales, marketing and distribution capabilities for any product candidates for which we may receive regulatory approval and that we determine to commercialize; and
- our need to implement additional internal systems and infrastructure, including financial and reporting systems.

We believe that our existing cash and cash equivalents as of filing of the form 10-Q will enable us to fund our operating expenses and capital expenditure requirements into the second half of 2024 based on our current plans, which may change based on clinical or pre-clinical results. These plans include: initiation of two Phase 2 clinical trials of CTX-009, completion of Phase 1 clinical trial of CTX-471, initiation of a Phase 1b combination trial for CTX-471 and commencement the planned Phase 1 development of CTX-8371, subject to satisfactory completion of IND-enabling activities for that product candidate. We expect that we will require additional funding to complete the clinical development of CTX-009, CTX-471 and CTX-8371, commercialize our product candidates, if we receive regulatory approval, and pursue in-licenses or acquisitions of other product candidates. If we receive regulatory approval for CTX-009, CTX-471 or CTX-8371 or other product candidates, we expect to incur significant commercialization expenses related to product manufacturing, sales, marketing and distribution, depending on where we choose to commercialize ourselves.

Until such time, if ever, as we can generate substantial product revenue, we expect to finance our cash needs through a combination of equity and debt financings, collaborations, strategic alliances, and marketing, distribution or licensing arrangements with third parties. To the extent that we raise additional capital through the sale of equity or convertible debt securities, ownership interest may be materially diluted, and the terms of such securities could include liquidation or other preferences that adversely affect your rights as a common stockholder. Debt financing and preferred equity financing, if available, may involve agreements that include restrictive covenants that limit our ability to take specified actions, such as incurring additional debt, making capital expenditures or declaring dividends. If we raise funds through collaborations, strategic alliances or marketing, distribution 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 grant licenses on terms that may not be favorable to us. If we are unable to raise additional funds through equity or debt financings or other arrangements when needed, we may be required to delay, reduce or eliminate our product development or future commercialization efforts, or grant rights to develop and market product candidates that we would otherwise prefer to develop and market ourselves.

Item 3. Quantitative and Qualitative Disclosures About Market Risk.

Not applicable since we are a smaller reporting company.

Item 4. Controls and Procedures.

Management's Evaluation of our Disclosure Controls and Procedures

Our management, with the participation of our Chief Executive Officer and Chief Operating Officer (Principal Financial Officer), evaluated 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 March 31, 2022. Management recognizes that any controls and procedures, no matter how well designed and operated, can provide only reasonable assurance of achieving their objectives and management necessarily applies its judgment in evaluating the cost-benefit relationship of possible controls and procedures. Based on the evaluation of our disclosure controls and procedures as of March 31, 2022, our Chief Executive Officer and Chief Operating Officer (Principal Financial Officer) concluded that, as of such date, our disclosure controls and procedures were effective.

Changes in Internal Control over Financial Reporting

There was no change in our internal control over financial reporting, as defined in Rules 13a-15(f) and 15d-15(f) under the Exchange Act, that occurred during the quarter ended March 31, 2022 that has materially affected, or is reasonably likely to materially affect, our internal control over financial reporting.

PART II—OTHER INFORMATION

Item 1. Legal Proceedings.

As of the date of this Quarterly Report on Form 10-Q, we are not involved in any material legal proceedings. However, from time to time, we could be subject to various legal proceedings and claims that arise in the ordinary course of our business activities. Regardless of the outcome, legal proceedings can have an adverse impact on us because of defense and settlement costs, diversion of management resources and other factors.

Item 1A. Risk Factors

In addition to the other information set forth in this report, you should carefully consider the risk factors discussed in Part I, Item 1A "Risk Factors" in our Annual Report on Form 10-K for the fiscal year ended December 31, 2021, which could materially affect our business, financial condition, or results of operations.

Item 2. Unregistered Sales of Equity Securities and Use of Proceeds.

None.

Item 3. Defaults Upon Senior Securities.

None.

Item 4. Mine Safety Disclosures.

Not applicable.

Item 5. Other Information.

On May 9, 2022, we issued a press release announcing our financial results for the quarter ended March 31, 2022. A copy of this press release is attached as Exhibit 99.1 to this Quarterly Report. The information regarding this press release in this Item 5 (including Exhibit 99.1) shall not be deemed "filed" for purposes of Section 18 of the Exchange Act or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

Item 6. Exhibits.

Exhibit Number	Description
31.1*	Certification of Principal Executive Officer Pursuant to Rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as Adopted Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
31.2*	Certification of Principal Financial Officer Pursuant to Rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as Adopted Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
32.1**	Certification of Principal Executive Officer Pursuant to 18 U.S.C. Section 1350, as Adopted Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
32.2**	Certification of Principal Financial Officer Pursuant to 18 U.S.C. Section 1350, as Adopted Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
99.1**	Press release titled "Compass Therapeutics Reports First Quarter 2022 Financial Results and Highlights Recent Company Progress"
101.INS	Inline XBRL Instance Document – the instance document does not appear in Interactive Data File because its XBRL tags are embedded within the Inline XBRL Document
101.SCH	Inline XBRL Taxonomy Extension Schema Document
101.CAL	Inline XBRL Taxonomy Extension Calculation Linkbase Document
101.DEF	Inline XBRL Taxonomy Extension Definition Linkbase Document
101.LAB	Inline XBRL Taxonomy Extension Label Linkbase Document
101.PRE	Inline XBRL Taxonomy Extension Presentation Linkbase Document
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

* Filed herewith.

** These exhibits are being furnished and shall not be deemed to be “filed” for purposes of Section 18 of the Exchange Act, or otherwise subject to the liability of that section, nor shall such exhibits be deemed to be incorporated by reference in any registration statement or other document filed under the Securities Act or the Exchange Act, except as otherwise stated in such filing.

SIGNATURES

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

Company Name

Date: May 9, 2022

By: /s/ Thomas Schuetz

Thomas Schuetz, MD
Chief Executive Officer (Principal Executive Officer)

Date: May 9, 2022

By: /s/ Vered Bisker-Leib

Vered Bisker-Leib, PhD
President and Chief Operating Officer (Principal Financial Officer)

Date: May 9, 2022

By: /s/ Neil Lerner

Neil Lerner, CPA
Vice President - Finance

Exhibit 31.1

CERTIFICATION PURSUANT TO
RULES 13a-14(a) AND 15d-14(a) UNDER THE SECURITIES EXCHANGE ACT OF 1934,
AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Thomas Schuetz, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of Compass Therapeutics, 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: May 9, 2022

By: /s/ Thomas Schuetz

Thomas Schuetz, MD

Chief Executive Officer (Principal Executive Officer)

Exhibit 31.2

CERTIFICATION PURSUANT TO
RULES 13a-14(a) AND 15d-14(a) UNDER THE SECURITIES EXCHANGE ACT OF 1934,
AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Vered Bisker-Leib, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of Compass Therapeutics, 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: May 9, 2022

By: /s/ Vered Bisker-Leib

Vered Bisker-Leib, PhD
President and Chief Operating Officer (Principal Financial Officer)

Exhibit 32.1

CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350, AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

In connection with the Quarterly Report of Compass Therapeutics, Inc. (the "Company") on Form 10-Q for the period ended March 31, 2022 as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I certify, pursuant to 18 U.S.C. § 1350, as adopted pursuant to § 906 of the Sarbanes-Oxley Act of 2002, that to the best of my knowledge:

(1) The Report fully complies with the requirements of section 13(a) or 15(d) of the Securities Exchange Act of 1934; and

(2) The information contained in the Report fairly presents, in all material respects, the financial condition and result of operations of the Company.

Date: May 9, 2022

By: /s/ Thomas Schuetz

Thomas Schuetz, MD

Chief Executive Officer (Principal Executive Officer)

Exhibit 32.2

CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350, AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

In connection with the Quarterly Report of Compass Therapeutics, Inc. (the "Company") on Form 10-Q for the period ended March 31, 2022 as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I certify, pursuant to 18 U.S.C. § 1350, as adopted pursuant to § 906 of the Sarbanes-Oxley Act of 2002, that to the best of my knowledge:

(1) The Report fully complies with the requirements of section 13(a) or 15(d) of the Securities Exchange Act of 1934; and

(2) The information contained in the Report fairly presents, in all material respects, the financial condition and result of operations of the Company.

Date: May 9, 2022

By: /s/ Vered Bisker-Leib

Vered Bisker-Leib, PhD
President and Chief Operating Officer (Principal Financial Officer)

Compass Therapeutics Reports First Quarter 2022 Financial Results and Highlights Recent Company Progress

- *The FDA cleared the investigational new drug application for CTX-009 (DLL4 X VEGF-A bispecific) in January allowing the Company to expand the ongoing Phase 2 study in patients with biliary tract cancer (BTC) to a global study and initiate dosing patients in the United States in early Q3 of 2022*
- *The Company reported interim data from the ongoing Phase 2 study of CTX-009 and paclitaxel in patients with advanced BTC demonstrating a 42% overall response rate (ORR) and 92% clinical benefit rate (CBR) among the first 24 patients enrolled and dosed with interim median patient time on study of ~ 6 months*
- *The Company completed enrollment in the CTX-471 (CD137 agonist) Phase 1b monotherapy study and reported 3 partial responses in patients with advanced solid tumors who were dosed with CTX-471 following progression on a prior PD-1/PD-L1 checkpoint blocker*
- *CTX-8371 GMP manufacturing campaign has begun, and the program is on track for an IND in Q1 2023*
- *$136.4 million in cash and cash equivalents at the end of the First Quarter*

Boston, May 9, 2022 - Compass Therapeutics, Inc. (Nasdaq: CMPX), a clinical-stage, oncology-focused biopharmaceutical company developing proprietary antibody-based therapeutics to treat multiple human diseases, today reported first quarter 2022 financial results.

CORPORATE UPDATE

CTX-009 (DLL4 and VEGF-A bispecific antibody)

In January, the Company announced that the FDA cleared its IND application for CTX-009 and, in May the Company released interim results from a Phase 2 study of CTX-009 in combination with paclitaxel in patients with BTC. The data show that:

- CTX-009 demonstrated a 42% overall response rate (ORR) based on 10 patients with Partial Responses (PRs), including 9 PRs confirmed by RECIST 1.1 and 1 PR pending confirmation
- CTX-009 demonstrated anti-tumor activity in previously treated patients with a clinical benefit rate (CBR) of 92% based on 22 patients with a PR or stable disease (SD) out of 24 enrolled patients
- CTX-009 was well-tolerated and preliminary safety profile is consistent with the Phase 1 studies

CTX-471 (CD137 agonist)

- As of February 25, 2022, 49 patients with 15 different cancers have been enrolled in the study and 38 of those patients are evaluable. Of the 38 evaluable patients, 3 patients had a PR; the first two have been confirmed by RECIST 1.1 and the third PR is unconfirmed. In addition, 19 patients have reached stable disease, leading to a preliminary ORR of 8% and a CBR of 58%
- The 3 partial responses observed in the study were in a patient with advanced small cell lung cancer, a patient with metastatic melanoma and a patient with metastatic melanoma of mucosal origin

CTX-8371 (PD-1 and PD-L1 bispecific antibody)

- CTX-8371 GMP manufacturing campaign has begun, and the program is on track for an IND in Q1 2023
- The Company presented preclinical data on CTX-8371 involving a unique mechanism of action (MOA) that involves cleavage of cell surface PD-1, at the 2022 American Association for Cancer Research (AACR) annual meeting

First Quarter 2022 Financial Results

- Cash Position: As of March 31, 2022, cash and cash equivalents were $136.4 million as compared to $39.7 million as of March 31, 2021, providing the Company with an anticipated cash runway into the second half of 2024. The Company used $7.9 million of cash to fund operations in the first quarter of 2022
- Research and development (R&D) Expenses: R&D expenses were $4.4 million for the first quarter ended March 31, 2022, as compared to $4.7 million for the same period in 2021, a decrease of $.3 million or 6%
- General and Administrative (G&A) Expenses: G&A expenses were $2.7 million for the first quarter ended March 31, 2022, as compared to $2.6 million for the same period in 2021, an increase of $.1 million or 5%
- Net Loss: Net loss for the first quarter ended March 31, 2022, was $7.2 million or $0.07 per common share, compared to $7.3 million or $0.14 per common share for the same period in 2021

About Compass Therapeutics

Compass Therapeutics, Inc. is a clinical-stage oncology-focused biopharmaceutical company developing proprietary antibody-based therapeutics to treat multiple human diseases. Compass's scientific focus is on the relationship between angiogenesis, the immune system, and tumor growth. The company pipeline of novel product candidates is designed to target multiple critical biological pathways required for an effective anti-tumor response. These include modulation of the microvasculature via angiogenesis-targeted agents, induction of a potent immune response via activators on effector cells in the tumor microenvironment, and alleviation of immunosuppressive mechanisms used by tumors to evade immune surveillance. Compass plans to advance its product candidates through clinical development as both standalone therapies and in combination with proprietary pipeline antibodies based on supportive clinical and nonclinical data. The company was founded in 2014 and is headquartered in Boston, Massachusetts. For more information, visit the Compass Therapeutics website at https://www.compasstherapeutics.com.

Forward-Looking Statements

This press release contains forward-looking statements. Statements in this press release that are not purely historical are forward-looking statements. Such forward-looking statements include, among other things, references to the Company's financial position to continue advancing its product candidates, expectations about the Company's cash runway, business and development plans, and statements regarding the Company's product candidates, their development, regulatory plans with respect thereto and therapeutic potential thereof, planned interactions with regulatory authorities, and planned clinical development. Actual results could differ from those projected in any forward-looking statements due to numerous factors. Such factors include, among others, the Company's ability to raise the additional funding it will need to continue to pursue its business and product development plans, the inherent uncertainties associated with developing product candidates and operating as a development stage company, the Company's ability to identify additional product candidates for development, the Company's ability to develop, complete clinical trials for, obtain approvals for and commercialize any of our product candidates, competition in the industry in which the Company operates and market conditions. These forward-looking statements are made as of the date of this press release, and the Company assumes no obligation to update the forward-looking statements, or to update the reasons why actual results could differ from those projected in the forward-looking statements, except as required by law. Investors should consult all of the information set forth herein and should also refer to the risk factor disclosure set forth in the reports and other documents we file with the SEC available at www.sec.gov, including without limitation our Form 10-Q for the quarter ended March 31, 2022, and our subsequent filings with the SEC.

Investor Contact
Vered Bisker-Leib, President & Chief Operating Officer
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Media Contact
Anna Gifford, Communications Manager
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617-500-8099

Compass Therapeutics, Inc. and Subsidiaries
Consolidated Statement of Operations (unaudited)
(In thousands, except per share data)

	Three Months Ended March 31,	
	2022	**2021**
Operating expenses:		
Research and development	$ 4,415	$ 4,704
General and administrative	2,767	2,635
Total operating expenses	7,182	7,339
Loss from operations	(7,182)	(7,339)
Other income (expense)	20	(83)
Net loss	$ (7,162)	$ (7,422)
Net loss per share - basic and diluted	$ (0.07)	$ (0.14)
Basic and diluted weighted average shares outstanding	100,858	51,313

Compass Therapeutics, Inc. and Subsidiaries
Consolidated Balance Sheets
(In thousands, except par value)

	March 31, 2022 (unaudited)	December 31, 2021 (audited)
Assets		
Current assets:		
Cash and cash equivalents	$ 136,379	$ 144,514
Prepaid expenses and other current assets	3,904	2,591
Total current assets	140,283	147,105
Property and equipment, net	2,142	2,243
Operating lease, right-of-use ("ROU") asset	3,819	4,089
Other assets	320	320
Total assets	$ 146,564	$ 153,757
Liabilities and Stockholders' Equity		
Current liabilities:		
Accounts payable	$ 257	$ 867
Accrued expenses	8,050	8,775
Operating lease obligations, current portion	1,027	989
Total current liabilities	9,334	10,631
Operating lease obligations, long-term portion	2,740	3,048
Total liabilities	12,074	13,679
Stockholders' equity:		
Common stock, $0.0001 par value: 300,000 shares authorized; 101,286 and 101,303 shares issued at March 31, 2022 and December 31, 2021, respectively; 100,905 and 100,832 shares outstanding at March 31, 2022 and December 31, 2021, respectively	10	10
Additional paid-in-capital	375,231	373,657
Accumulated deficit	(240,751)	(233,589)
Total stockholders' equity	134,490	140,078
Total liabilities and stockholders' equity	$ 146,564	$ 153,757