Verastem, Inc. Form 10-Q May 03, 2018 Table of Contents

UNITED STATES

SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 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, 2018 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 35403

Verastem, Inc.

(Exact name of registrant as specified in its charter)

Delaware 27-3269467 (State or other jurisdiction of incorporation or organization) Identification Number)

117 Kendrick Street, Suite 500

Needham, MA 02494 (Address of principal executive offices) (Zip Code)

(781) 292-4200

(Registrant's telephone number, including area code)

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 and posted on its corporate Web site, if any, every Interactive Data File required to be submitted and posted 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 and post 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 Accelerated filer Non accelerated filer Smaller reporting Emerging growth (Do not check if a company smaller reporting 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 April 30, 2018 there were 56,612,723 shares of Common Stock, \$0.0001 par value per share, outstanding.

Table of Contents

TABLE OF CONTENTS

PARTI—	<u>-FINANCIAL INFORMATIO</u> N	
<u>Item 1.</u>	Condensed Consolidated Financial Statements (unaudited)	4
Item 2.	Management's Discussion and Analysis of Financial Condition and Results of Operations	15
Item 3.	Quantitative and Qualitative Disclosures About Market Risk	19
<u>Item 4.</u>	Controls and Procedures	20
PART II-	<u>–OTHER INFORMATIO</u> N	
<u>Item 1.</u>	<u>Legal Proceedings</u>	21
Item 1A.	Risk Factors	21
<u>Item 2.</u>	<u>Unregistered Sales of Equity Securities and Use of Proceeds</u>	21
<u>Item 3.</u>	<u>Defaults Upon Senior Securities</u>	21
<u>Item 4.</u>	Mine Safety Disclosures	21
<u>Item 5.</u>	Other Information	21
Item 6.	<u>Exhibits</u>	21
EXHIBIT	<u>'INDEX</u>	22
SIGNATU	<u>URES</u>	23

FORWARD LOOKING STATEMENTS

This Quarterly Report on Form 10-Q contains forward-looking statements that involve substantial risks and uncertainties. All statements, other than statements related to present facts or current conditions or historical facts, contained in this Quarterly Report on Form 10-Q, including statements regarding our strategy, future operations, future financial position, future revenues, projected costs, prospects, plans and objectives of management, are forward-looking statements. Such statements relate to, among other things, the development of our product candidates, including duvelisib and defactinib, and our Phosphoinositide 3-kinase (PI3K) and Focal Adhesion Kinase (FAK) programs generally, the timeline for clinical development and regulatory approval of our product candidates, the expected timing for the reporting of data from on-going trials, the structure of our planned or pending clinical trials, additional planned studies, our rights to develop or commercialize our product candidates and our ability to finance contemplated development and commercialization activities and fund operations for a specified period. The words "anticipate," "believe," "estimate," "expect," "intend," "may," "plan," "predict," "project," "target," "potent "would," "could," "should," "continue" and similar expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these identifying words.

Forward-looking statements are not guarantees of future performance and our actual results could differ materially from the results discussed in the forward-looking statements we make. Applicable risks and uncertainties include the risks that approval of our New Drug Application for duvelisib will not occur on the expected timeframe or at all, including by the U.S. Food and Drug Administration's target action date; that a filing of a European Marketing Application may not be achieved in fiscal year 2018 or at all; that we may not enter into any partnerships or collaborations for the potential commercialization of duvelisib outside of the U.S.; that the full data from the Phase 3 DUOTM study will not be consistent with the previously presented results of the study; that the preclinical testing of our product candidates and preliminary or interim data from clinical trials may not be predictive of the results or success of ongoing or later clinical trials; that data may not be available when expected, including for the Phase 3 DUO study; that even if data from clinical trials is positive, regulatory authorities may require additional studies for approval and the product may not prove to be safe and effective; that the degree of market acceptance of product candidates, if approved, may be lower than expected; that the timing, scope and rate of reimbursement for our product candidates is uncertain; that there may be competitive developments affecting our product candidates; that data may not be available when expected; that enrollment of clinical trials may take longer than expected; that our product candidates will cause unexpected safety events or result in an unmanageable safety profile as compared to their level of efficacy; that duvelisib will be ineffective at treating patients with lymphoid malignancies; that we will be unable to successfully initiate or complete the clinical development and eventual commercialization of our product candidates; that the development and commercialization of our product candidates will take longer or cost more than planned; that we may not have sufficient cash to fund our contemplated operations; that we or Infinity Pharmaceuticals, Inc. will fail to fully perform under the duvelisib license agreement; that we may be unable to make additional draws under our debt facility or obtain adequate financing in the future through product licensing, co-promotional arrangements, public or private equity, debt financing or otherwise; that we will not pursue or submit regulatory filings for our product candidates, including for duvelisib in patients with chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL) or indolent non-Hodgkin lymphoma (iNHL); and that our product candidates will not receive regulatory approval, become commercially successful products, or result in new treatment options being offered to patients. Other risks and uncertainties include those identified under the heading "Risk Factors" in our Annual Report on Form 10-K for the year ended December 31, 2017 as filed with the Securities and Exchange Commission (SEC) on March 13, 2018 and in any subsequent filings with the SEC.

As a result of these and other factors, we may not achieve the plans, intentions or expectations disclosed in our forward-looking statements, and you should not place undue reliance on our forward-looking statements. Our forward-looking statements do not reflect the potential impact of any future acquisitions, mergers, dispositions, joint ventures or investments we may make. The forward-looking statements contained in this Quarterly Report on Form 10-Q reflect our views as of the date hereof. We do not assume and specifically disclaim any obligation to update any forward-looking statements, whether as a result of new information, future events or otherwise, except as required by law.

PART I—FINANCIAL INFORMATION

Item 1. Condensed Consolidated Financial Statements (unaudited).

Verastem, Inc.

CONDENSED CONSOLIDATED BALANCE SHEETS

(in thousands, except per share amounts)

Assets Current assets:	March 31, 2018 (unaudited)	December 31, 2017
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Cash and cash equivalents	\$ 64,215	\$ 82,176
Short-term investments	1.015	4,496
Prepaid expenses and other current assets	1,815	1,115
Total current assets	66,030	87,787
Property and equipment, net	1,003	861
Restricted cash	403	162
Other assets	975	981
Total assets	\$ 68,411	\$ 89,791
Liabilities and stockholders' equity		
Current liabilities:		
Accounts payable	\$ 7,267	\$ 9,186
Accrued expenses	7,578	7,942
Total current liabilities	14,845	17,128
Non-current liabilities:		
Long-term debt	14,913	14,828
Other non-current liabilities	101	151
Total liabilities	29,859	32,107
Stockholders' equity:		
Preferred stock, \$0.0001 par value; 5,000 shares authorized, no shares issued		
and outstanding at March 31, 2018 and December 31, 2017, respectively		_
Common stock, \$0.0001 par value; 100,000 shares authorized, 50,968 and 50,801		
shares issued and outstanding at March 31, 2018 and December 31, 2017,		
respectively	5	5
Additional paid-in capital	362,739	360,823
Accumulated other comprehensive loss		(2)
Accumulated deficit	(324,192)	(303,142)
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Total stockholders' equity	38,552	57,684
Total liabilities and stockholders' equity	\$ 68,411	\$ 89,791

See accompanying notes to the condensed consolidated financial statements.

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Verastem, Inc.

CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS AND COMPREHENSIVE LOSS

(unaudited)

(in thousands, except per share amounts)

	Three months	ended March
	31,	
	2018	2017
Operating expenses:		
Research and development	\$ 10,934	\$ 8,385
General and administrative	9,827	4,763
Total operating expenses	20,761	13,148
Loss from operations	(20,761)	(13,148)
Interest income	191	155
Interest expense	(480)	(12)
Net loss	\$ (21,050)	\$ (13,005)
Net loss per share—basic and diluted	\$ (0.41)	\$ (0.35)
Weighted-average number of common shares used in net loss per share—basic and		
diluted	50,835	36,992
Net loss	\$ (21,050)	\$ (13,005)
Unrealized gain (loss) on available-for-sale securities	2	(17)
Comprehensive loss	\$ (21,048)	\$ (13,022)

See accompanying notes to the condensed consolidated financial statements.

Verastem, Inc.

CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS

(unaudited)

(in thousands)

	Three months 31,	ended March
	2018	2017
Operating activities		
Net loss	\$ (21,050)	\$ (13,005)
Adjustments to reconcile net loss to net cash used in operating activities:		
Depreciation	125	146
Stock-based compensation expense	1,328	1,197
Amortization of deferred financing costs, debt discounts and premiums and discounts		
on available-for-sale marketable securities	83	46
Gain on sale of fixed assets	(33)	
Changes in operating assets and liabilities:		
Prepaid expenses, other current assets and other assets	(694)	(1,092)
Accounts payable	(2,078)	4,594
Accrued expenses and other liabilities	(135)	(2,538)
Net cash used in operating activities	(22,454)	(10,652)
Investing activities		
Purchases of property and equipment	(102)	
Sales of property and equipment	37	
Purchases of investments		(6,461)
Maturities of investments	4,500	10,557
Net cash provided by investing activities	4,435	4,096
Financing activities		
Proceeds from long-term debt, net		2,386
Proceeds from the issuance of common stock, net	299	
Net cash provided by financing activities	299	2,386
Decrease in cash, cash equivalents and restricted cash	(17,720)	(4,170)
Cash, cash equivalents and restricted cash at beginning of period	82,338	32,511
Cash, cash equivalents and restricted cash at end of period	\$ 64,618	\$ 28,341
Supplemental disclosure of non-cash financing activities		
Purchases of property and equipment in accounts payable	\$ 169	\$ —
Common stock issuance costs included in accounts payable and accrued expenses	\$ 35	\$ —
Deferred financing costs in accounts payable and accrued expenses	\$ —	\$ 140

See accompanying notes to the condensed consolidated financial statements.

Tabla	of	Contents
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Verastem, Inc.

NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS

(unaudited)

1. Nature of business

Verastem, Inc. (the Company) is a biopharmaceutical company focused on developing and commercializing drugs to improve outcomes for patients with cancer. The Company's operations to date have been limited to organizing and staffing the Company, business planning, raising capital, identifying and acquiring potential product candidates and undertaking preclinical studies and clinical trials of its product candidates.

The Company is subject to a number of risks similar to other life science companies, including, but not limited to, the need to obtain adequate additional funding, possible failure of preclinical testing or clinical trials, inability to obtain marketing approval of product candidates, competitors developing new technological innovations, market acceptance of the Company's products and protection of proprietary technology. If the Company does not successfully commercialize any of its product candidates, it will be unable to generate product revenue or achieve profitability.

As of March 31, 2018, the Company had cash and cash equivalents of \$64.2 million and accumulated deficit of \$324.2 million. The Company has historical losses from operations and anticipates that it will continue to incur losses for the foreseeable future as it continues the research and development and clinical trials of its product candidates, and seeks marketing approval for its lead product candidate, duvelisib. Without additional funding, the Company believes that it will not have sufficient funds to meet its obligations within the next twelve months from the date of issuance of these condensed consolidated financial statements. These factors raise substantial doubt about the Company's ability to continue as a going concern.

The Company plans to continue to fund its operations through proceeds from sales of its common stock under its at-the-market equity offering program, public or private equity offerings, its loan and security agreement with Hercules Capital, Inc. (Hercules), or other strategic transactions. However, adequate additional financing may not be available to the Company on acceptable terms, or at all. If the Company is unable to raise capital when needed or on attractive terms, it may be forced to delay, reduce or eliminate its research and development programs or any commercialization efforts.

2. Summary of significant accounting policies

Basis of Presentation

The accompanying unaudited condensed consolidated financial statements of the Company have been prepared in accordance with generally accepted accounting principles in the United States (GAAP) for interim financial reporting and as required by Regulation S-X, Rule 10-01 under the assumption that the Company will continue as a going concern for the next twelve months. Accordingly, they do not include all of the information and footnotes required by GAAP for complete financial statements, or any adjustments that might result from the uncertainty related to the Company's ability to continue as a going concern. In the opinion of management, all adjustments (including those which are normal and recurring) considered necessary for a fair presentation of the interim financial information have been included. When preparing financial statements in conformity with GAAP, the Company must make estimates and assumptions that affect the reported amounts and related disclosures at the date of the financial statements. Actual results could differ from those estimates. Additionally, operating results for the three months ended March 31, 2018 are not necessarily indicative of the results that may be expected for any other interim period or for the year ending December 31, 2018. For further information, refer to the financial statements and footnotes included in the Company's Annual Report on Form 10-K for the year ended December 31, 2017 as filed with the Securities and Exchange Commission (SEC) on March 13, 2018.

Table of Contents

Recently Issued Accounting Standards Updates

In February 2016, the Financial Accounting Standards Board (FASB) issued Accounting Standards Update (ASU) 2016-02, Leases (Topic 842), which supersedes the guidance under FASB Accounting Standards Codification (ASC) Topic 840, Leases, resulting in the creation of FASB ASC Topic 842, Leases. ASU 2016-02 requires lessees to recognize in the statement of financial position a liability to make lease payments and a right-of-use asset representing its right to use the underlying asset for the lease term for both finance and operating leases. The guidance also eliminates the current real estate-specific provisions for all entities. ASU 2016-02 is effective for fiscal years, and interim periods within those years, beginning after December 15, 2018, with early adoption permitted. The Company has not elected to early adopt this standard and is currently evaluating the impact the adoption of the standard will have on its condensed consolidated financial statements and related disclosures.

Recently Adopted Accounting Standards Updates

In May 2017, the FASB issued ASU 2017-09, Compensation – Stock Compensation (Topic 718): Scope of Modification Accounting. ASU 2017-09 provides guidance about which changes to the terms or conditions of a share-based award require an entity to apply modification accounting under Topic 718. Specifically, an entity would not apply modification accounting if the fair value, vesting conditions and classification of the awards are the same immediately before and after a modification. ASU 2017-09 was effective for annual and interim periods beginning after December 15, 2017, with early adoption permitted. The Company adopted this standard prospectively effective January 1, 2018. The adoption of this ASU did not have an effect on the Company's condensed consolidated financial statements or related disclosures.

In November 2016, the FASB issued ASU 2016-18, Statement of Cash Flows (Topic 230): Restricted Cash. ASU 2016-18 requires that a statement of cash flows explain the change during the period in the total of cash, cash equivalents and amounts generally described as restricted cash or restricted cash equivalents. Amounts generally described as restricted cash and restricted cash equivalents should be included with cash and cash equivalents when reconciling the beginning-of-period and end-of-period total amounts shown on the statement of cash flows. ASU 2016-18 was effective for annual and interim periods beginning after December 15, 2017, with early adoption permitted. The Company adopted this standard effective January 1, 2018. Upon adoption of ASU 2016-18, the Company applied the retrospective transition method for each period presented and included approximately \$162,000 of restricted cash in the beginning-of-period and end-of-period cash, cash equivalents and restricted cash balance reflected in the condensed consolidated statement of cash flows for the three months ended March 31, 2017. A reconciliation of cash, cash equivalents and restricted cash for each period presented is provided in note 3 to the condensed consolidated financial statements.

In August 2016, the FASB issued ASU 2016-15, Statement of Cash Flows (Topic 230): Classification of Certain Cash Receipts and Cash Payments. ASU 2016-15 adds or clarifies guidance on the classification of certain cash receipts and payments in the statement of cash flows. The standard was effective for annual and interim periods beginning after December 15, 2017, with early adoption permitted. The Company adopted this standard effective January 1, 2018. The adoption of this ASU did not have an effect on the Company's condensed consolidated financial statements or related disclosures.

Significant accounting policies

There have been no material changes to the significant accounting policies included in the Company's Annual Report on Form 10-K for the year ended December 31, 2017 as filed with the SEC on March 13, 2018.

3. Cash, cash equivalents and restricted cash

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 (in thousands):

	March 31,	December
	2018	31, 2017
Cash and cash equivalents	\$ 64,215	\$ 82,176
Restricted cash	403	162
Total cash, cash equivalents and restricted cash	\$ 64,618	\$ 82,338

Amounts included in restricted cash represent cash held to collateralize outstanding letters of credit in the amount of approximately \$403,000 and \$162,000 as of March 31, 2018 and December 31, 2017, respectively, provided as a security deposit for the Company's office space located in Needham, Massachusetts.

4. Fair value of financial instruments

The Company determines the fair value of its financial instruments based upon the fair value hierarchy, which prioritizes valuation inputs based on the observable nature of those inputs. The fair value hierarchy applies only to the valuation inputs used in determining the reported fair value of the investments and is not a measure of the investment credit quality. The hierarchy defines three levels of valuation inputs:

Level 1	Quoted prices in active markets for identical assets or liabilities that the Company can access at the
inputs	measurement date.
Level 2	Inputs other than quoted prices included within Level 1 that are observable for the asset or liability, either
inputs	directly or indirectly.
Level 3	Unobservable inputs that reflect the Company's own assumptions about the assumptions market
inputs	participants would use in pricing the asset or liability.

Items Measured at Fair Value on a Recurring Basis

The following table presents information about the Company's financial instruments that are measured at fair value on a recurring basis (in thousands):

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	March 31, 20	018			
Description	Total	Level 1	Level 2	Level 3	
Financial assets					
Cash equivalents	\$ 62,731	\$ 62,731	\$ —	\$ —	
Total financial assets	\$ 62,731	\$ 62,731	\$ —	\$ —	
	December	31, 2017			
Description Financial assets	Total	Level 1	Level 2	Level 3	
Cash equivalents	\$ 80,894	\$ 75,478	\$ 5,416	\$ —	
Short-term investments	4,496	_	4,496	_	
Total financial assets	\$ 85,390	\$ 75,478	\$ 9,912	\$ —	

The Company's cash equivalents and investments are comprised of U.S. Government money market funds and corporate bonds and commercial paper of publicly traded companies. These investments and cash equivalents have been initially valued at the transaction price and subsequently valued, at the end of each reporting period, utilizing third party pricing services or other market observable data. The pricing services utilize industry standard valuation models, including both income and market based approaches and observable market inputs to determine value. These observable

market inputs include reportable trades, benchmark yields, credit spreads, broker/dealer quotes, bids, offers, current spot rates and other industry and economic events. The Company validates the prices provided by third party pricing services by reviewing their pricing methods and matrices, obtaining market values from other pricing sources, analyzing pricing data in certain instances and confirming that the relevant markets are active. After completing its validation procedures, the Company did not adjust or override any fair value measurements provided by the pricing services as of December 31, 2017.

Fair Value of Financial Instruments

The fair value of the Company's long-term debt is determined using current applicable rates for similar instruments as of the condensed consolidated balance sheet dates and an assessment of the credit rating of the Company. The carrying value of the Company's long-term debt at March 31, 2018 approximates fair value because the Company's interest rate yield is near current market rates for comparable debt instruments. The fair value of the Company's long-term debt was determined using Level 3 inputs.

5. Investments

Cash, cash equivalents, and investments consist of the following (in thousands):

	March 31, 20	018		
		Gross	Gross	ъ.
	Amortized	Unrealized	Unrealized	Fair
	Cost	Gains	Losses	Value
Cash and cash equivalents:				
Cash and money market accounts	\$ 64,215	\$ —	\$ —	\$ 64,215
Total cash and cash equivalents	\$ 64,215	\$ —	\$ —	\$ 64,215

	December 31, 2017			
	Amortized Cost	Gross Unrealized Gains	Gross Unrealized Losses	Fair Value
Cash and cash equivalents: Cash and money market accounts	\$ 76,760 5,418	\$ — \$ —	\$ — \$ (2)	\$ 76,760 \$ 5,416

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\$ 82,178	\$	_	\$	(2)	\$ 82,176
\$ 4,496	\$		\$	_	\$ 4,496
\$ 4,496	\$		\$	_	\$ 4,496
\$ 86,674	\$		\$	(2)	\$ 86,672
	\$ 4,496 \$ 4,496	\$ 4,496 \$ \$ 4,496 \$	\$ 4,496 \$ — \$ 4,496 \$ —	\$ 4,496 \$ — \$ \$ 4,496 \$ — \$	\$ 4,496 \$ — \$ — \$ 4,496 \$ — \$ —

There were no realized gains or losses on investments for the three months ended March 31, 2018 or 2017, respectively. There were no investments in an unrealized loss position as of March 31, 2018. There were five investments in an unrealized loss position as of December 31, 2017. None of these investments had been in an unrealized loss position for more than 12 months. The aggregate unrealized loss on these securities as of December 31, 2017 was approximately \$2,000 and the fair value was \$9.9 million.

6. Accrued expenses

Accrued expenses consist of the following (in thousands):

	March	December
	31, 2018	31, 2017
Contract research organization costs	\$ 5,181	\$ 3,774
Compensation and related benefits	1,305	2,622
Professional fees	435	617
Consulting fees	286	579
Deferred rent	194	190
Other	177	160
Total accrued expenses	\$ 7,578	\$ 7,942

7. Long-term debt

On March 21, 2017 (Closing Date), Verastem, Inc. (the Borrower) entered into a term loan facility of up to \$25.0 million with Hercules. The term loan facility is governed by a loan and security agreement, dated March 21, 2017 (the Original Loan Agreement), which was amended on January 4, 2018 and March 6, 2018 (the Amended Loan Agreement) to increase the total borrowing limit under the Original Loan Agreement from up to \$25 million to up to \$50.0 million (the Term Loan), pursuant to certain conditions of funding.

As of March 31, 2018, the Company has borrowed a total of \$15.0 million in term loans. The remaining \$35.0 million of borrowing capacity under the Amended Loan Agreement may be drawn in minimum increments of \$5.0 million in multiple tranches comprised of (i) term loans (each a Term E Loan Advance) in an aggregate principal amount of up to \$10.0 million and (ii) subject to Hercules' sole discretion, term loans (each a Term F Loan Advance) in an aggregate principal amount of up to \$25.0 million. The Amended Loan Agreement permits the Borrower to draw Term E Loan Advances subject to (i) the U.S. Food and Drug Administration accepting on or prior to September 30, 2018 the Company's New Drug Application for duvelisib and (ii) delivery of the Company's financial and business projections to Hercules in form and substance reasonably acceptable to Hercules. In addition, the Amended Loan Agreement allows the Borrower to draw Term F Loan Advances subject to the prior drawing of all other tranches and Hercules' sole discretion.

The Term Loan will mature on December 1, 2020 (Loan Maturity Date). Each advance accrues interest at a floating per annum rate equal to the greater of either (a) 10.5% or (b) the lesser of (i) 12.75% and (ii) the sum of (x) 10.5% plus (y) (A) the prime rate minus (B) 4.5%. The Term Loan provided for interest-only payments until November 1, 2018, which was extended to May 1, 2019 pursuant to the Amended Loan Agreement upon the Borrower's receipt of a minimum of \$20.0 million in cash proceeds from a sale of equity securities in December 2017. Thereafter,

amortization payments will be payable monthly in 20 installments of principal and interest (subject to recalculation upon a change in prime rates).

The Term Loan is secured by a lien on substantially all of the assets of the Borrower, other than intellectual property, and contains customary covenants and representations.

The Company assessed all terms and features of the Original Loan Agreement in order to identify any potential embedded features that would require bifurcation or any beneficial conversion features. As part of this analysis, the Company assessed the economic characteristics and risks of the Original Loan Agreement, including put and call features. The Company determined that all features of the Original Loan Agreement were clearly and closely associated with a debt host and did not require bifurcation as a derivative liability, or the fair value of the feature was immaterial to the Company's financial statements. The Company reassesses the features on a quarterly basis to determine if they require separate accounting. There have been no changes to the Company's original assessment through March 31, 2018.

Table of Contents

The future principal payments under the Loan Agreement are as follows as of March 31, 2018 (in thousands):

Remainder of 2018 \$ —
2019 3,609
2020 11,391
Total principal payments \$ 15,000

8. Net loss per share

Basic and diluted net loss per common share is calculated by dividing net loss applicable to common stockholders by the weighted-average number of common shares outstanding during the period, without consideration for common stock equivalents. The Company's potentially dilutive shares, which include outstanding stock options and restricted stock units (RSUs), are considered to be common stock equivalents and are only included in the calculation of diluted net loss per share when their effect is dilutive.

The following potentially dilutive securities were excluded from the calculation of diluted net loss per share for the periods indicated because including them would have had an anti-dilutive effect:

	Three months ended March		
	31,		
	2018	2017	
Outstanding stock options	10,818,454	7,337,655	
Outstanding restricted stock units	166,250		
Total potentially dilutive securities	10,984,704	7,337,655	

9. Stock based compensation

Stock options

A summary of the Company's stock option activity and related information for the three months ended March 31, 2018 is as follows:

	Shares		Weighted-average remaining contractual term (years)	in	ggregate trinsic value n thousands)
Outstanding at December 31, 2017	8,719,978	\$ 5.19	7.9	\$	6,150
Granted	2,149,000	\$ 3.10			
Exercised	_	\$ 			
Forfeited/cancelled	(50,524)	\$ 2.30			
Outstanding at March 31, 2018	10,818,454	\$ 4.79	8.1	\$	5,734
Vested at March 31, 2018	5,315,689	\$ 6.57	6.8	\$	3,135
Vested and expected to vest at March 31, 2018(1)	10,465,454	\$ 4.85	8.0	\$	5,732

⁽¹⁾ This represents the number of vested options as of March 31, 2018, plus the number of unvested options expected to vest as of March 31, 2018.

Table of Contents

The fair value of each stock option granted during the three months ended March 31, 2018 and 2017 was estimated on the grant date using the Black-Scholes option-pricing model using the following weighted-average assumptions:

	Three months ended				
	March 31	• •			
	2018	2017			
Risk-free interest rate	2.38 %	2.06 %			
Volatility	81 %	79 %			
Dividend yield		_			
Expected term (years)	6.0	6.4			

During the first quarter of 2018, the Company granted stock options to purchase a total of 582,500 shares of common stock to certain executives that vest only upon the achievement of specified performance conditions. The Company determined that a number of the performance conditions are considered probable of achievement as of March 31, 2018, and as a result recognized approximately \$350,000 of stock-based compensation expense related to these awards.

At March 31, 2018, there was \$9.5 million of total unrecognized compensation cost related to unvested stock options and the Company expects to recognize this cost over a remaining weighted-average period of approximately 3 years.

Restricted stock units

The Company awards RSUs to employees under its 2012 Incentive Plan. Each RSU entitles the holder to receive one share of the Company's common stock when the RSU vests. The RSUs vest in four substantially equal installments on each of the first four anniversaries of the vesting commencement date, subject to the employee's continued employment with, or service to, the Company on such vesting date. Compensation expense is recognized on a straight-line basis.

A summary of RSU activity during the three months ended March 31, 2018 is as follows:

Weighted-average grant date fair Shares value per share

Outstanding at December 31, 2017