

CODEXIS INC
Form 10-Q
November 06, 2015
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 September 30, 2015

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: 001-34705

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

Delaware
(State or other jurisdiction of incorporation or organization) 71-0872999
(I.R.S. Employer Identification No.)

200 Penobscot Drive, Redwood City
(Address of principal executive offices) 94063
(Zip Code)
(650) 421-8100
(Registrant's telephone number, including area code)
(Former name, former address and former fiscal year, if changed since last report)

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, or a smaller reporting company. See the definitions of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act. (Check one):
Large accelerated filer Accelerated filer

Non-accelerated filer (Do not check if a smaller reporting company) Smaller reporting company

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 October 30, 2015, there were 40,306,967 shares of the registrant's Common Stock, par value \$0.0001 per share, outstanding.

Codexis, Inc.
 Quarterly Report on Form 10-Q
 For The Three Months Ended September 30, 2015

TABLE OF CONTENTS

	PAGE NUMBER
PART I. FINANCIAL INFORMATION	
ITEM 1: Financial Statements (Unaudited)	
<u>Condensed Consolidated Balance Sheets</u>	<u>3</u>
<u>Condensed Consolidated Statements of Operations</u>	<u>4</u>
<u>Condensed Consolidated Statements of Comprehensive Income (Loss)</u>	<u>5</u>
<u>Condensed Consolidated Statements of Cash Flows</u>	<u>6</u>
<u>Notes to Condensed Consolidated Financial Statements</u>	<u>7</u>
ITEM 2: <u>Management's Discussion and Analysis of Financial Condition and Results of Operations</u>	<u>26</u>
ITEM 3: <u>Quantitative and Qualitative Disclosures about Market Risk</u>	<u>34</u>
ITEM 4: <u>Controls and Procedures</u>	<u>36</u>
 <u>PART II. OTHER INFORMATION</u>	
ITEM 1: <u>Legal Proceedings</u>	<u>37</u>
ITEM 1A: <u>Risk Factors</u>	<u>37</u>
ITEM 2: <u>Unregistered Sales of Equity Securities and Use of Proceeds</u>	<u>37</u>
ITEM 3: <u>Default Upon Senior Securities</u>	<u>37</u>
ITEM 4: <u>Mine Safety Disclosures</u>	<u>37</u>
ITEM 5: <u>Other Information</u>	<u>38</u>
ITEM 6: <u>Exhibits</u>	<u>38</u>
<u>Signatures</u>	

Codexis, Inc.

Condensed Consolidated Balance Sheets

(Unaudited)

(In Thousands, Except Per Share Amounts)

	September 30, 2015	December 31, 2014
Assets		
Current assets:		
Cash and cash equivalents	\$ 16,963	\$ 26,487
Accounts receivable, net of allowances of \$421 at September 30, 2015 and \$428 at December 31, 2014	13,608	3,870
Inventories	678	1,395
Prepaid expenses and other current assets	1,092	1,255
Total current assets	32,341	33,007
Restricted cash	786	711
Marketable securities	1,231	688
Property and equipment, net	2,821	3,995
Intangible assets, net	3,655	6,186
Goodwill	3,241	3,241
Other non-current assets	265	294
Total assets	\$44,340	\$48,122
Liabilities and Stockholders' Equity		
Current liabilities:		
Accounts payable	\$ 1,174	\$ 4,673
Accrued compensation	2,554	2,946
Other accrued liabilities	2,151	2,619
Deferred revenue	6,949	3,497
Total current liabilities	12,828	13,735
Deferred revenue, net of current portion	4,316	3,813
Other long-term liabilities	3,888	4,263
Commitments and contingencies (Note 11)		
Stockholders' equity:		
Preferred stock, \$0.0001 par value; 5,000 shares authorized, none issued and outstanding	—	—
Common stock, \$0.0001 par value; 100,000 shares authorized at September 30, 2015 and December 31, 2014; shares issued and outstanding of 40,300 at September 30, 2015 and 39,563 at December 31, 2014	4	4
Additional paid-in capital	304,561	302,379
Accumulated other comprehensive income (loss)	201	(142)
Accumulated deficit	(281,458)	(275,930)
Total stockholders' equity	23,308	26,311
Total liabilities and stockholders' equity	\$44,340	\$48,122

See accompanying notes to the unaudited condensed consolidated financial statements

Codexis, Inc.
Condensed Consolidated Statements of Operations
(Unaudited)
(In Thousands, Except Per Share Amounts)

	Three Months Ended September		Nine Months Ended September	
	30,		30,	
	2015	2014	2015	2014
Revenues:				
Biocatalyst product sales	\$1,818	\$2,562	\$6,915	\$8,323
Biocatalyst research and development	14,517	3,364	19,247	7,176
Revenue sharing arrangement	1,066	1,546	4,056	5,617
Total revenues	17,401	7,472	30,218	21,116
Costs and operating expenses:				
Cost of biocatalyst product sales	1,302	1,532	4,009	6,179
Research and development	4,994	5,038	15,457	17,708
Selling, general and administrative	5,415	5,157	16,289	16,791
Total costs and operating expenses	11,711	11,727	35,755	40,678
Income (loss) from operations	5,690	(4,255)	(5,537)	(19,562)
Interest income	4	3	12	15
Other expenses	(26)	(57)	(147)	(183)
Income (loss) before income taxes	5,668	(4,309)	(5,672)	(19,730)
Provision for (benefit from) income taxes	274	253	(144)	(314)
Net income (loss)	\$5,394	\$(4,562)	\$(5,528)	\$(19,416)
Net income (loss) per share, basic	\$0.14	\$(0.12)	\$(0.14)	\$(0.51)
Net income (loss) per share, diluted	\$0.13	\$(0.12)	\$(0.14)	\$(0.51)
Weighted average common shares used in computing net income (loss) per share, basic	39,767	38,450	39,340	38,063
Weighted average common shares used in computing net income (loss) per share, diluted	40,970	38,450	39,340	38,063

See accompanying notes to the unaudited condensed consolidated financial statements

Codexis, Inc.

Condensed Consolidated Statements of Comprehensive Income (Loss)

(Unaudited)

(In Thousands)

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2015	2014	2015	2014
Net income (loss)	\$5,394	\$(4,562)) \$(5,528) \$(19,416)
Other comprehensive income (loss):				
Unrealized gain (loss) on marketable securities, net of tax expense of \$263 and \$160 for the three months ended September 30, 2015 and 2014, respectively, and tax benefit of \$200 and \$89 for the nine months ended September 30, 2015 and 2014, respectively.	(449)) (261) 343	145
Other comprehensive income (loss)	(449)) (261) 343	145
Total comprehensive income (loss)	\$4,945	\$(4,823)) \$(5,185) \$(19,271)

See accompanying notes to the unaudited condensed consolidated financial statements

Codexis, Inc.
Condensed Consolidated Statements of Cash Flows
(Unaudited)
(In Thousands)

	Nine Months Ended September	
	30,	
	2015	2014
Operating activities:		
Net loss	\$(5,528)) \$(19,416)
Adjustments to reconcile net loss to net cash used in operating activities:		
Amortization of intangible assets	2,531	2,531
Depreciation and amortization of property and equipment	1,569	2,679
Impairment of property and equipment	—	1,841
Change in the fair value of assets held for sale	—	886
Gain on disposal of property and equipment	(5) (115)
Income tax benefit related to marketable securities	(200) (89)
Gain on sale of Hungarian subsidiary	—	(760)
Stock-based compensation	3,759	3,630
Amortization of premium on marketable securities	—	2
Bad debt expense	—	53
Changes in operating assets and liabilities:		
Accounts receivable, net	(9,738) 2,316
Inventories, net	717	(456)
Prepaid expenses and other current assets	163	(734)
Other assets	29	15
Accounts payable	(3,499) (1,418)
Accrued compensation	(393) (1,100)
Other accrued liabilities	(842) 194
Deferred revenue	3,955	5,288
Net cash used in operating activities	(7,482) (4,653)
Investing activities:		
Purchase of property and equipment	(395) (267)
Proceeds from maturities of marketable securities	—	3,000
Proceeds from sale of Hungarian subsidiary, net of selling costs	—	1,500
Proceeds from the sale of assets held for sale	—	281
Proceeds from sale of property and equipment	5	166
Increase in restricted cash	(75) —
Net cash provided by (used in) investing activities	(465) 4,680
Financing activities:		
Proceeds from exercises of options to purchase common stock	235	180
Taxes paid related to net share settlement of equity awards	(1,812) (815)
Net cash used in financing activities	(1,577) (635)
Net decrease in cash and cash equivalents	(9,524) (608)
Cash and cash equivalents at the beginning of the period	26,487	22,130
Cash and cash equivalents at the end of the period	\$16,963	\$21,522

See accompanying notes to the unaudited condensed consolidated financial statements

Notes to Condensed Consolidated Financial Statements
(Unaudited)

Note 1. Description of Business

In these notes to the condensed consolidated financial statements, the "Company," "we," "us," and "our" refer to Codexis, Inc. and its subsidiaries on a consolidated basis.

We develop biocatalysts for the pharmaceutical and fine chemicals markets. Our proven technologies enable scale-up and implementation of biocatalytic solutions to meet customer needs for rapid, cost-effective and sustainable process development, from research to manufacturing.

Biocatalysts are enzymes that initiate and/or accelerate chemical reactions. Manufacturers have historically used naturally occurring biocatalysts to produce many goods used in everyday life. However, inherent limitations in naturally occurring biocatalysts have restricted their commercial use. Our proprietary CodeEvolver[®] protein engineering technology platform (the "CodeEvolver[®] Platform Technology"), which introduces genetic mutations into microorganisms in order to give rise to changes in enzymes that they produce, is able to overcome many of these limitations, allowing us to evolve and optimize biocatalysts to perform specific and desired chemical reactions at commercial scale.

Once potentially beneficial mutations are identified through this proprietary process, combinations of these mutations can then be tested until variant enzymes have been created that exhibit marketable performance characteristics superior to competitive products. This process allows for continuous, efficient improvements to the performance of enzymes. In the past, we implemented the CodeEvolver[®] Platform Technology through paid collaborations with our customers. In July 2014, we entered into our first license agreement pursuant to which we granted a license to a global pharmaceutical company to use the CodeEvolver[®] Platform Technology for its internal development purposes. In August 2015, we entered into a license agreement involving the CodeEvolver[®] Platform Technology with a second global pharmaceutical company and we continue to pursue licensing opportunities with additional customers. We have commercialized our technology and products in the pharmaceuticals market, which is our primary business focus. Our customers, which include several large global pharmaceutical companies, use our technology, products and services in their manufacturing processes and process development.

We also use our technology to develop biocatalysts for use in the fine chemicals market. The fine chemicals market is similar to our pharmaceutical business and consists of several large market verticals, including food, animal feed, flavors, fragrances, and agricultural chemicals.

We are also using our technology to develop an early stage, novel enzyme therapeutic product candidate for the potential treatment of phenylketonuria ("PKU") in humans. PKU is an inherited metabolic disorder in which the enzyme that converts the essential amino acid phenylalanine into tyrosine is deficient.

Note 2. Basis of Presentation and Summary of Significant Accounting Policies

Basis of Presentation and Principles of Consolidation

The accompanying condensed consolidated financial statements have been prepared in accordance with accounting principles generally accepted in the United States of America ("GAAP") and the applicable rules and regulations of the Securities and Exchange Commission ("SEC") for interim financial information. Accordingly, they do not include all of the information and notes required by GAAP for complete financial statements. These interim condensed consolidated financial statements should be read in conjunction with the consolidated financial statements and notes thereto contained in our Annual Report on Form 10-K for the year ended December 31, 2014. The condensed consolidated balance sheet at December 31, 2014 has been derived from the audited consolidated financial statements at that date, but does not include all disclosures, including notes, required by GAAP for complete financial statements. The unaudited interim condensed consolidated financial statements have been prepared on the same basis as the audited consolidated financial statements and, in the opinion of management, reflect all adjustments of a normal recurring nature considered necessary to present fairly our financial position as of September 30, 2015 and results of our operations and comprehensive income (loss) for the three and nine months ended September 30, 2015 and 2014, and cash flows for the nine months ended September 30, 2015 and 2014. The interim results are not necessarily

indicative of the results for any future interim period or for the entire year. Certain prior period amounts have been reclassified to conform to current period presentation.

7

The unaudited interim condensed consolidated financial statements include Codexis, Inc. and its wholly owned subsidiaries in the United States, Brazil, Hungary (through the sale date of March 13, 2014), India, Mauritius, the Netherlands, and Singapore (dissolved in October 2014). All significant intercompany balances and transactions have been eliminated in consolidation.

Use of Estimates

The preparation of financial statements in conformity with GAAP requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities, disclosures of contingent liabilities at the date of the condensed consolidated financial statements, and the reported amounts of revenue and expenses during the reporting period. We regularly assess these estimates which primarily affect revenue recognition, accounts receivable, inventories, the valuation of investment securities and marketable securities, assets held for sale, intangible assets, goodwill arising out of business acquisitions, accrued liabilities, stock awards and the valuation allowances associated with deferred tax assets. Actual results could differ from those estimates and such differences may be material to the condensed consolidated financial statements.

Segment Reporting

Operating segments are defined as components of an enterprise about which separate financial information is available that is evaluated regularly by the chief operating decision maker, or decision making group, in deciding how to allocate resources and in assessing performance. Our chief operating decision maker is our Chief Executive Officer. The Chief Executive Officer reviews financial information presented on a consolidated basis, accompanied by information about revenues by geographic region, for purposes of allocating resources and evaluating financial performance. We have one business activity and there are no segment managers who are held accountable for operations, operating results beyond revenue goals or plans for levels or components below the consolidated unit level. Accordingly, we have a single reporting segment.

Revenue Recognition

We recognize revenues from the sale of our biocatalyst products, biocatalyst research and development agreements and a revenue sharing arrangement. Revenue is recognized when the related costs are incurred and the four basic criteria of revenue recognition are met: (1) persuasive evidence of an arrangement exists; (2) delivery has occurred or services have been rendered; (3) the fee is fixed or determinable; and (4) collectability is reasonably assured. Where the revenue recognition criteria are not met, we defer the recognition of revenue by recording deferred revenue until such time that all criteria of revenue recognition are met.

We account for revenues from multiple element arrangements, such as license and platform technology transfer agreements in which a licensee may purchase several deliverables, in accordance with Financial Accounting Standards Board ("FASB") Accounting Standards Codification ("ASC") Subtopic 605-25, "Multiple Element Arrangements." For new or materially amended multiple element arrangements, we identify the deliverables at the inception of the arrangement and each deliverable within a multiple deliverable revenue arrangement is accounted for as a separate unit of accounting if both of the following criteria are met: (1) the delivered item or items have value to the customer on a standalone basis and (2) for an arrangement that includes a general right of return relative to the delivered item(s), delivery or performance of the undelivered item(s) is considered probable and substantially in our control. Revenue allocated to each element is then recognized based on when the basic four revenue recognition criteria are met for each element.

Biocatalyst Product Sales

Biocatalyst product sales consist of sales of biocatalyst enzymes, chemical intermediates and Codex[®] Biocatalyst Panels and Kits. Biocatalyst product sales are recognized once passage of title and risk of loss has occurred and contractually specified acceptance criteria, if any, have been met, provided all other revenue recognition criteria have also been met. Shipping and handling costs charged to customers are included in revenue.

Biocatalyst Research and Development

Biocatalyst research and development agreements typically provide us with multiple revenue streams, including research services fees for full time employee ("FTE") research services, up-front licensing fees, technology access, contingent payments upon achievement of contractual criteria, and royalty fees based on the licensees' product sales or cost savings achieved by our customers. We perform biocatalyst research and development activities as specified in each respective customer agreement. Payments for services received are not refundable. Certain research agreements

are based on a contractual reimbursement rate per FTE working on the project. We recognize revenues from research services as those services are performed over the contractual performance periods. When up-front payments are combined with FTE services in a single unit of accounting, we

8

recognize the up-front payments using the proportionate performance method of revenue recognition based upon the actual amount of research labor hours incurred relative to the amount of the total expected labor hours to be incurred by us, up to the amount of cash received. In cases where the planned levels of research services fluctuate substantially over the research term, we are required to make estimates of the total hours required to perform our obligations. We recognize revenues from non-refundable, up-front license fees or technology access payments that are not dependent on any future performance by us when such amounts are earned. If we have continuing obligations to perform under the arrangement, such fees are recorded as deferred revenues and recognized over the estimated period of performance. Estimated performance periods are periodically reviewed based on the progress of the related program. The effect of any change made to an estimated performance period, and therefore to revenue recognized, would occur on a prospective basis in the period that the change was made.

A payment that is contingent upon the achievement of a substantive milestone is recognized in its entirety in the period in which the milestone is achieved. A milestone is an event (i) that can only be achieved based in whole or in part on either our performance or on the occurrence of a specific outcome resulting from our performance, (ii) for which there is, as of the date the arrangement is entered into, substantive uncertainty that the event will be achieved and (iii) results in additional payments being due to us. Milestones are considered substantive when the consideration earned from the achievement of the milestone (i) is commensurate with either our performance to achieve the milestone or the enhancement of the value of the item delivered as a result of a specific outcome resulting from its performance, (ii) relates solely to past performance and (iii) is reasonable relative to all deliverable and payment terms in the arrangement.

We recognize revenues from other contingent payments based on the passage of time or when earned as the result of a customer's performance in accordance with contractual terms and when such payments can be reasonably estimated and collectability of such payments is reasonably assured.

We recognize revenues from royalties based on licensees' sales of our biocatalyst products or products using our technologies.

Royalties are recognized as earned in accordance with the contract terms when royalties from licensees can be reasonably estimated and collectability is reasonably assured. For the majority of our royalty revenues, estimates are made using notification of the sale of licensed products from the licensees.

Revenue Sharing Arrangement

We recognize revenues from a revenue sharing arrangement based upon sales of licensed products by our revenue share partner Exela PharmSci, Inc. ("Exela") (see Note 12, "Related Party Transactions"). We recognize revenues net of product and selling costs upon notification from our revenue share partner of our portion of net profit based on the contractual percentage from the sale of licensed product.

Sales Allowances

Sales allowances primarily relate to product returns and prompt pay sales discounts and are recorded in the same period that the related revenues are recognized, resulting in a reduction in biocatalyst product sales revenue.

Cost of Biocatalyst Product Sales

Cost of biocatalyst product sales comprises both internal and third party fixed and variable costs including amortization of purchased technology, materials and supplies, labor, facilities and other overhead costs associated with our biocatalyst product sales. Shipping costs are included in our cost of biocatalyst product sales. Such shipping costs were not significant in any of the periods presented.

Cost of Research and Development Services

Cost of research and development services related to services under research and development agreements approximate the research funding over the term of the respective agreements and is included in research and development expense.

Research and Development Expenses

Research and development expenses consist of costs incurred for internal projects as well as research and development services as mentioned above. These costs include our direct and research-related overhead expenses, which include salaries and other personnel-related expenses (including stock-based compensation), occupancy-related costs, supplies, depreciation of

facilities and laboratory equipment and amortization of acquired technologies, as well as external costs. Costs to acquire technologies that are utilized in research and development and that have no alternative future use are expensed as incurred.

Stock-Based Compensation

We use the Black-Scholes-Merton option pricing model to estimate the fair value of options granted under our equity incentive plans. The Black-Scholes-Merton option pricing model requires the use of assumptions, including the expected term of the award and the expected stock price volatility. We have, due to insufficient historical data, used the "simplified method," as described in Staff Accounting Bulletin No. 107, "Share-Based Payment," to determine the expected life of all stock options granted from the inception of our equity plans through the first half of 2015. We believe we have sufficient historical data to calculate expected terms for stock options granted beginning in the third quarter of 2015. The expected term was based on historical exercise behavior on similar awards, giving consideration to the contractual terms, vesting schedules and expectations of future employee behavior. We used historical volatility to estimate expected stock price volatility. The risk-free rate assumption was based on United States Treasury instruments whose terms were consistent with the expected term of the stock options. The expected dividend assumption was based on our history and expectation of dividend payouts.

Restricted Stock Units ("RSUs"), Restricted Stock Awards ("RSAs") and performance-contingent restricted stock units ("PSUs") were measured based on the fair market values of the underlying stock on the dates of grant. PSUs awarded may be conditional upon the attainment of one or more performance objectives over a specified period. At the end of the performance period, if the goals are attained, the awards are granted.

Stock-based compensation expense was calculated based on awards ultimately expected to vest and was reduced for estimated forfeitures at the time of grant and revised, if necessary, in subsequent periods if actual forfeitures differed from those estimates. The estimated annual forfeiture rates for stock options, RSUs, PSUs, and RSAs are based on historical forfeiture experience.

The estimated fair value of stock options, RSUs and RSAs is expensed on a straight-line basis over the vesting term of the grant and the estimated fair value of PSUs is expensed using an accelerated method over the term of the award once management has determined that it is probable that the performance objective will be achieved. Compensation expense is recorded over the requisite service period based on management's best estimate as to whether it is probable that the shares awarded are expected to vest. Management assesses the probability of the performance milestones being met on a continuous basis.

We have not recognized, and do not expect to recognize in the near future, any income tax benefit related to employee stock-based compensation expense as a result of the full valuation allowance on our deferred tax assets including deferred tax assets related to net operating loss carryforwards.

Foreign Currency Translation

The United States dollar is the functional currency for our operations outside the United States. Accordingly, nonmonetary assets and liabilities originally acquired or assumed in other currencies are recorded in United States dollars at the exchange rates in effect at the date they were acquired or assumed. Monetary assets and liabilities denominated in other currencies are translated into United States dollars at the exchange rates in effect at the balance sheet date. Translation adjustments are recorded in other expense in the accompanying condensed consolidated statements of operations. Gains and losses realized from non-U.S. dollar transactions, including intercompany balances not considered as permanent investments, denominated in currencies other than an entity's functional currency, are also included in other expense in the accompanying condensed consolidated statements of operations.

Cash and Cash Equivalents

We consider all highly liquid investments with maturity dates of three months or less at the date of purchase to be cash equivalents. Our cash and cash equivalents consist of cash on deposit with banks and money market funds. Most of our cash and cash equivalents are maintained with major financial institutions in North America. Deposits with these financial institutions may exceed the amount of insurance provided on such deposits. Cash and cash equivalents totaled \$17.0 million at September 30, 2015 and were comprised of cash of \$5.9 million and money market funds of \$11.1 million.

Inventories

Inventories are stated at the lower of cost or market value. Cost is determined using a weighted-average approach, assuming full absorption of direct and indirect manufacturing costs, based on our product capacity utilization assumptions. If inventory costs exceed expected market value due to obsolescence or lack of demand, valuation adjustments are recorded for the difference between the cost and the estimated market value. These valuation adjustments are determined based on significant estimates.

Marketable Securities

We invest in equity securities and we classify those investments as available-for-sale. These securities are carried at estimated fair value (see Note 5, "Marketable Securities") with unrealized gains and losses included in accumulated other comprehensive income (loss) in stockholders' equity. Available-for-sale equity securities with remaining maturities of greater than one year or which we currently do not intend to sell are classified as long-term.

We review several factors to determine whether a loss is other-than-temporary. These factors include, but are not limited to, the intent and ability to retain the investment in the issuer for a period of time sufficient to allow for any anticipated recovery in market value, the length of time and the extent to which the market value of the investment has been less than cost and the financial condition and near-term prospects of the issuer. Unrealized losses are charged against "Other expense" when a decline in fair value is determined to be other-than-temporary. Amortization of purchase premiums and accretion of purchase discounts and realized gains and losses of debt securities are included in interest income. The cost of securities sold is based on the specific identification method.

Fair Value Measurements

Fair value is defined as the price that would be received to sell an asset or paid to transfer a liability in an orderly transaction between market participants at the measurement date. In determining fair value, we utilize valuation techniques that maximize the use of observable inputs and minimize the use of unobservable inputs to the extent possible and we consider counterparty credit risk in our assessment of fair value. Carrying amounts of financial instruments, including cash equivalents, short-term investments, marketable investments, accounts receivable, accounts payable and accrued liabilities, approximate their fair values as of the balance sheet dates because of their generally short maturities.

The fair value hierarchy distinguishes between (1) market participant assumptions developed based on market data obtained from independent sources (observable inputs) and (2) an entity's own assumptions about market participant assumptions developed based on the best information available in the circumstances (unobservable inputs). The fair value hierarchy consists of three broad levels, giving the highest priority to unadjusted quoted prices in active markets for identical assets or liabilities (Level 1) and the lowest priority to unobservable inputs (Level 3). The three levels of the fair value hierarchy are described below:

Level 1: Quoted prices (unadjusted) in active markets that are accessible at the measurement date for assets or liabilities.

Level 2: Directly or indirectly observable inputs as of the reporting date through correlation with market data, including quoted prices for similar assets and liabilities in active markets and quoted prices in markets that are not active. Level 2 also includes assets and liabilities that are valued using models or other pricing methodologies that do not require significant judgment since the input assumptions used in the models, such as interest rates and volatility factors, are corroborated by readily observable data from actively quoted markets for substantially the full term of the financial instrument.

Level 3: Unobservable inputs that are supported by little or no market activity and reflect the use of significant management judgment. These values are generally determined using pricing models for which the assumptions utilize management's estimates of market participant assumptions.

Concentrations of Credit Risk

Our financial instruments that are potentially subject to concentration of credit risk primarily consist of cash equivalents, short term investments, accounts receivable, marketable securities and restricted cash. We invest cash that is not required for immediate operating needs principally in money market funds.

Intangible Assets

Our intangible assets are finite-lived and consist of customer relationships, developed core technology, trade names, and the intellectual property rights associated with the acquisition of Maxygen Inc.'s ("Maxygen") directed evolution technology in 2010. Intangible assets were recorded at their fair values at the date we acquired the assets and, for those assets having finite useful lives, are amortized using the straight-line method over their estimated useful lives.

Impairment of Long-Lived Assets

Our long-lived assets include property and equipment and intangible assets. We determined that we have a single entity wide asset group ("Asset Group"). The directed evolution technology patent portfolio acquired from Maxygen ("Core IP") is the most significant component of the Asset Group since it is the base technology for all aspects of our research and development activities, and represents the basis for all of our identifiable cash flow generating capacity. Consequently, we do not believe that identification of independent cash flows associated with long-lived assets is currently possible at any lower level than the Asset Group.

The Core IP is the only finite-lived intangible asset on our condensed consolidated balance sheet as of September 30, 2015. There has been no significant change in the utilization or estimated life of the Core IP since we acquired the technology patent portfolio from Maxygen.

The carrying value of our long-lived assets in the Asset Group may not be recoverable based upon the existence of one or more indicators of impairment which could include: a significant decrease in the market price of our common stock; current period cash flow losses or operating losses combined with a history of losses or a forecast of continuing losses associated with the use of the assets; slower growth rates in our industry; significant adverse changes in the business climate or legal factors; accumulation of costs significantly in excess of the amount originally expected for the acquisition or construction of the assets; loss of significant customers or partners; or the current expectation that the assets will more likely than not be sold or disposed of significantly before the end of their estimated useful life.

We evaluate recoverability of intangible assets based on the sum of the undiscounted cash flows expected to result from the use and the eventual disposal of the Asset Group. We make estimates and judgments about the future undiscounted cash flows over the remaining useful life of the Asset Group. Our anticipated future cash flows include our estimates of existing or in process product sales, production and operating costs, future capital expenditures, working capital needs, and assumptions regarding the ultimate sale of the Asset Group at the end of the life of the primary asset. The useful life of the Asset Group was based on the estimated useful life of the Core IP, the primary asset at the time of acquisition. There has been no change in the estimated useful life of the Asset Group. Although our cash flow forecasts are based on assumptions that are consistent with our plans, there is significant judgment involved in determining the cash flows attributable to the Asset Group over its estimated remaining useful life.

In the fourth quarter of 2014, we determined that there were no events or changes in circumstances that indicated that the carrying value of the Asset Group might not be recoverable. We concluded that the fair value of the reporting unit exceeded its carrying value and no impairment existed. During the nine months ended September 30, 2015, we did not identify any indicators of potential impairment of intangible assets or new information that would have a material impact on the forecast or the impairment analysis prepared as of December 31, 2014.

Goodwill

We determined that we operate in one segment and reporting unit under the criteria in ASC 280, "Segment Reporting." Accordingly, our review of goodwill impairment indicators is performed at the parent level. We review goodwill impairment annually in the fourth quarter of each fiscal year and whenever events or changes in circumstances indicate the carrying value of goodwill may not be recoverable.

The goodwill impairment test consists of a two-step process. The first step of the goodwill impairment test used to identify potential impairment compares the fair value of the reporting unit to carrying value. If the fair value of the reporting unit exceeds its carrying amount, goodwill of the reporting unit is considered not impaired, and the second step of the impairment test is not required.

We use our market capitalization as an indicator of fair value. We believe that because our reporting unit is publicly traded, the ability of a controlling stockholder to benefit from synergies and other intangible assets that arise from control might cause the fair value of our reporting unit as a whole to exceed its market capitalization. However, we believe that the fair value measurement need not be based solely on the quoted market price of an individual share of our common stock, but also can consider the impact of a control premium in measuring the fair value of its reporting

unit.

12

If we were to use an income approach, it would establish a fair value by estimating the present value of our projected future cash flows expected to be generated from our business. The discount rate applied to the projected future cash flows to arrive at the present value would be intended to reflect all risks of ownership and the associated risks of realizing the stream of projected future cash flows. Our discounted cash flow methodology would consider projections of financial performance for a period of several years combined with an estimated residual value. The most significant assumptions we would use in a discounted cash flow methodology are the discount rate, the residual value and expected future revenue, gross margins and operating costs, along with considering any implied control premium. Should our market capitalization be less than total stockholder's equity as of our annual test date or as of any interim impairment testing date, we would also consider market comparables, recent trends in our stock price over a reasonable period and, if appropriate, use an income approach to determine whether the fair value of our reporting unit is greater than the carrying amount.

The second step, if required, compares the implied fair value of the reporting unit goodwill with the carrying amount of that goodwill. If the carrying amount of the reporting unit's goodwill exceeds its implied fair value, an impairment charge is recognized in an amount equal to that excess. Implied fair value is the excess of the fair value of the reporting unit over the fair value of all identified assets and liabilities. We base our fair value estimates on assumptions we believe to be reasonable. Actual future results may differ from those estimates.

Goodwill was tested for impairment in the fourth quarter of 2014. We determined that the fair value of the reporting unit exceeded the carrying value and no impairment existed. Based on the results obtained, we concluded there was no impairment of our goodwill as of December 31, 2014. During the nine months ended September 30, 2015, we did not identify any indicators of potential impairment of goodwill or new information that would have a material impact on the forecast or the impairment analysis prepared as of December 31, 2014.

Income Taxes

We use the liability method of accounting for income taxes, whereby deferred tax assets or liability account balances are calculated at the balance sheet date using current tax laws and rates in effect for the year in which the differences are expected to affect taxable income. Valuation allowances are provided when necessary to reduce deferred tax assets to the amount that will more likely than not be realized.

We make certain estimates and judgments in determining income tax expense for financial statement purposes. These estimates and judgments occur in the calculation of tax credits, benefits and deductions and in the calculation of certain tax assets and liabilities, which arise from differences in the timing of recognition of revenue and expenses for tax and financial statement purposes. Significant changes to these estimates may result in an increase or decrease to our tax provision in a subsequent period.

In assessing the realizability of deferred tax assets, we consider whether it is more likely than not that some portion or all of the deferred tax assets will be realized on a jurisdiction by jurisdiction basis. The ultimate realization of deferred tax assets is dependent upon the generation of taxable income in the future. We have recorded a deferred tax asset in jurisdictions where ultimate realization of deferred tax assets is more likely than not to occur.

We make estimates and judgments about future taxable income that are based on assumptions that are consistent with our plans and estimates. Should the actual amounts differ from our estimates, the amount of our valuation allowance could be materially impacted. Any adjustment to the deferred tax asset valuation allowance would be recorded in the income statement for the periods in which the adjustment is determined to be required. With the sale of the Hungarian subsidiary in the quarter ended March 31, 2014, the related net operating losses and other tax attributes are no longer available to us. The related deferred tax assets had a full valuation allowance and, as a result, their removal did not have a material impact to the financial statements.

We account for uncertainty in income taxes as required by the provisions of ASC Topic 740, "Income Taxes," which clarifies the accounting for uncertainty in income taxes recognized in an enterprise's financial statements. The first step is to evaluate the tax position for recognition by determining if the weight of available evidence indicates that it is more likely than not that the position will be sustained on audit, including resolution of related appeals or litigation processes, if any. The second step is to estimate and measure the tax benefit as the largest amount that is more than 50% likely of being realized upon ultimate settlement. It is inherently difficult and subjective to estimate such amounts, as this requires us to determine the probability of various possible outcomes. We consider many factors when evaluating and estimating our tax positions and tax benefits, which may require periodic adjustments and may

not accurately anticipate actual outcomes.

13

The Tax Reform Act of 1986 and similar state provisions limit the use of net operating loss carryforwards in certain situations where equity transactions result in a change of ownership as defined by Internal Revenue Code Section 382. In the event we should experience such a change of ownership, utilization of our federal and state net operating loss carryforwards could be limited. We maintain a full valuation allowance against net deferred tax assets as we believe that it is more likely than not that the majority of deferred tax assets will not be realized.

We recognized an income tax expense of \$0.3 million for both the three months ended September 30, 2015 and 2014. We recognized an income tax benefit of \$0.1 million for the nine months ended September 30, 2015, as compared to \$0.3 million for the same period in 2014.

Recently Issued and Adopted Accounting Guidance

From time to time, new accounting pronouncements are issued by the FASB or other standards setting bodies that are adopted by us as of the specified effective date. Unless otherwise discussed, we believe that the impact of recently issued standards that are not yet effective will not have a material impact on our consolidated financial statements upon adoption.

In August 2014, the FASB issued Accounting Standards Update ("ASU") 2014-15, "Presentation of Financial Statements - Going Concern (Subtopic 205-40): Disclosure of Uncertainties about an Entity's Ability to Continue as a Going Concern." ASU 2014-15 defines management's responsibility to evaluate whether there is substantial doubt about an organization's ability to continue as a going concern and provide related disclosures. ASU 2014-15 is effective for annual reporting periods ending after December 15, 2016, and interim periods within annual periods beginning after December 15, 2016. The adoption of ASU 2014-15 is not expected to have a material impact on our consolidated financial statements and related disclosures.

In July 2015, the FASB issued ASU 2015-11, "Inventory (Topic 330): Simplifying the Measurement of Inventory," which simplifies the subsequent measurement of inventory by requiring inventory to be measured at the lower of cost and net realizable value. Net realizable value is the estimated selling price of inventory in the ordinary course of business, less reasonably predictable costs of completion, disposal, and transportation. This ASU is effective for public business entities for fiscal years beginning after December 15, 2016, and interim periods within those fiscal years. We are currently evaluating the impact of adopting ASU 2015-11 on our condensed consolidated financial statements and related disclosures.

In August 2015, the FASB issued ASU 2015-14, "Revenue from Contracts with Customers (Topic 606): Deferral of the Effective Date." This ASU defers the effective date of ASU 2014-09, "Revenue from Contracts with Customers (Topic 606)" for all entities by one year. The standard outlines a single comprehensive model for entities to use in accounting for revenue arising from contracts with customers and supersedes most current revenue recognition guidance, including industry-specific guidance. The main principle of ASU 2014-09 is to recognize revenue when promised goods or services are transferred to customers in an amount that reflects the consideration that is expected to be received for those goods or services. ASU 2014-09 provides companies with two implementation methods: (i) apply the standard retrospectively to each prior reporting period presented (full retrospective application); or (ii) apply the standard retrospectively with the cumulative effect of initially applying the standard as an adjustment to the opening balance of retained earnings of the annual reporting period that includes the date of initial application (modified retrospective application). ASU 2014-09 as amended by ASU 2015-14 is effective for annual reporting periods beginning after December 15, 2017, including interim reporting periods within that reporting period. The FASB will permit companies to adopt the new standard early, but not before the original effective date of December 15, 2016. We are currently in the process of evaluating the impact of the pending adoption of this standard on our consolidated financial statements and related disclosures.

Note 3. Net Income (Loss) per Share

Basic net income (loss) per share is computed by dividing net income (loss) by the weighted-average number of shares of common stock outstanding, less RSAs subject to forfeiture. Diluted net income per share is computed by dividing net income by the weighted average number of shares of common stock outstanding, less RSAs subject to forfeiture, plus all additional common shares that would have been outstanding, assuming dilutive potential common shares had been issued for other dilutive securities. For periods of net loss, diluted and basic net loss per share were identical since potential common shares were excluded from the calculation, as their effect would be anti-dilutive.

The following table sets forth the computation of basic and diluted net income per share during the three and nine months ended September 30, 2015 and 2014 (in thousands):

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2015	2014	2015	2014
Numerator				
Net income (loss)	\$5,394	\$(4,562)	\$(5,528)	\$(19,416)
Denominator				
Weighted average common shares used in computing net income (loss) per share, basic	39,767	38,450	39,340	38,063
Effect of dilutive shares	1,203	—	—	—
Weighted average common shares used in computing net income (loss) per share, diluted	40,970	38,450	39,340	38,063
Net income (loss) per share, basic	\$0.14	\$(0.12)	\$(0.14)	\$(0.51)
Net income (loss) per share, diluted	\$0.13	\$(0.12)	\$(0.14)	\$(0.51)
Anti-Dilutive Securities				

In periods of net loss, the weighted average number of shares outstanding related to potentially dilutive securities, prior to the application of the treasury stock method, are excluded from the computation of diluted net loss per common share because including such shares would have an anti-dilutive effect. The following shares were not considered in the computation of diluted net loss per share (in thousands):

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2015	2014	2015	2014
Shares issuable under Equity Incentive Plan	—	6,398	6,121	6,398
Shares issuable upon the conversion of warrants	—	75	75	75
Total shares excluded as anti-dilutive	—	6,473	6,196	6,473

Note 4. Collaborative Arrangements

GSK Platform Technology Transfer, Collaboration and License Agreement

In July 2014, we entered into a CodeEvolver® platform technology transfer, collaboration and license agreement (the "GSK License Agreement") with GlaxoSmithKline ("GSK"). Under the terms of the GSK License Agreement, we granted GSK a non-exclusive license to use the CodeEvolver® Platform Technology to develop novel enzymes for use in the manufacture of GSK's pharmaceutical and health care products.

We received a \$6.0 million up-front licensing fee upon signing the GSK License Agreement and subsequently a \$5.0 million non-creditable, non-refundable milestone payment upon achievement of the first milestone. During the three months ended September 30, 2015, we achieved the second milestone of the agreement earning another milestone payment of \$6.5 million. We are eligible to receive an additional contingent payment of \$7.5 million upon the completion of the technology transfer period. We also have the potential to receive additional contingent payments that range from \$5.75 million to \$38.5 million per project based on GSK's successful application of the licensed technology. The contingent payments are not deemed substantive milestones due to the fact that the achievement of the event underlying the payment predominantly relates to GSK's performance of future development and commercialization activities.

For up to three years following the end of the three-year period during which we will transfer the CodeEvolver® Platform Technology to GSK, GSK can exercise an option, upon payment of certain additional fees, that would extend GSK's license to include certain improvements to the CodeEvolver® Platform Technology that arise during such period. In addition, we will also be eligible to receive royalties based on net sales, if any, of a limited set of products developed by GSK using the CodeEvolver® Platform Technology.

The term of the GSK License Agreement continues, unless earlier terminated, until the expiration of all payment obligations under the GSK License Agreement. At any time following the completion of the first technology transfer stage, GSK can terminate the GSK License Agreement by providing 90 days written notice to us. If GSK exercises this termination right during the three-year technology transfer period, GSK will make a one-time termination

payment to us.

15

Under the GSK License Agreement, the significant deliverables were determined to be the license, platform technology transfer, and contingent obligation to supply GSK with enzymes manufactured by us at GSK's expense. We determined that the license did not have stand-alone value, and we determined that the license and the platform technology transfer and our participation in joint steering committee activities in connection with the platform technology transfer represent a single unit of accounting. We determined that our participation in the joint steering committee does not represent a separate unit of accounting because GSK could not negotiate for and/or acquire these services from other third parties and our participation on the joint steering committee is coterminous with the technology transfer period. Amounts to be received under the supply arrangement described above will be recognized as revenue to the extent GSK purchases enzymes from us.

The up-front license fee of \$6.0 million is being recognized over the technology transfer period of three years. We recognized license fees of \$0.5 million and \$1.5 million for the three and nine months ended September 30, 2015, respectively, and \$0.5 million for the three and nine months ended September 30, 2014 as biocatalyst research and development revenues. We had a deferred revenue balance from GSK related to the upfront license fee of \$3.5 million at September 30, 2015 and \$5.0 million at December 31, 2014.

Merck Supply Agreement

On February 1, 2012, we entered into a five-year Sitagliptin Catalyst Supply Agreement ("Sitagliptin Catalyst Supply Agreement") with Merck Sharp and Dohme Corp., known as MSD outside the United States and Canada ("Merck") whereby Merck may obtain commercial scale substance for use in the manufacture of its products based on the active ingredient sitagliptin, e.g., Januvia®. Merck may extend the term of the Sitagliptin Catalyst Supply Agreement for an additional five years at its sole discretion.

The Sitagliptin Catalyst Supply Agreement requires Merck to pay an annual license fee for the rights to the sitagliptin technology each year for the term of the Sitagliptin Catalyst Supply Agreement. The license fee is being recognized as collaborative research and development revenues ratably over the five year term of the Sitagliptin Catalyst Supply Agreement.

We recognized license fees of \$0.5 million for each of the three months ended September 30, 2015 and 2014 and \$1.5 million for each of the nine months ended September 30, 2015 and 2014 as biocatalyst research and development revenues. We had a deferred revenue balance from Merck related to license fees of \$1.4 million at September 30, 2015 and \$0.9 million at December 31, 2014. In addition, pursuant to the Sitagliptin Catalyst Supply Agreement, Merck may purchase supply from us for a fee based on contractually stated prices.

Merck Platform Technology Transfer and License Agreement

On August 3, 2015 ("Effective Date"), we entered into a CodeEvolver® platform technology transfer and license agreement (the "Merck License Agreement") with Merck. The Agreement allows Merck to use the CodeEvolver® Platform Technology in the field of human and animal healthcare.

We received a \$5.0 million up-front licensing fee upon signing the Merck License Agreement. During the three months ended September 30, 2015, we achieved the first milestone of the Merck License Agreement earning a milestone payment of \$5.0 million. We are eligible to receive an additional \$8.0 million subject to the satisfactory completion of the second milestone of the technology transfer process. We will also be eligible to receive payments of up to a maximum of \$15.0 million for each commercial active pharmaceutical ingredient ("API") that is manufactured by Merck using one or more novel enzymes developed by Merck using the CodeEvolver® Platform Technology. Under the terms of the Merck License Agreement, we granted to Merck a non-exclusive, worldwide license to use the CodeEvolver® Platform Technology to research, develop and manufacture novel enzymes for use by Merck in its internal research programs ("Merck Non-Exclusive Field"). The license to Merck is exclusive for the research, development and manufacture of novel enzymes for use by Merck in the chemical synthesis of therapeutic products owned or controlled by Merck ("Merck Exclusive Field"). Merck has the right to grant sublicenses to affiliates of Merck and, in certain limited circumstances, to third parties. We also granted to Merck a license to make or have made products manufactured using the CodeEvolver® Platform Technology with a right to grant sublicenses solely to affiliates of Merck, contract manufacturing organizations and contract research organizations. The manufacturing license is exclusive in the Merck Exclusive Field and non-exclusive in the Merck Non-Exclusive Field. The licenses are subject to certain limitations based on pre-existing contractual obligations that apply to the technology and intellectual property that are the subject of the license grants. The licenses do not permit the use of the CodeEvolver®

Platform Technology to discover any therapeutic enzyme, diagnostic product or vaccine. In addition, Merck is prohibited from using the CodeEvolver® Platform Technology to develop or produce enzymes or any other compounds for or on behalf of any third parties except in a very limited manner when Merck divests a therapeutic product that is manufactured using an enzyme developed using the CodeEvolver® Platform Technology.

Under the Merck License Agreement, we will transfer the CodeEvolver® Platform Technology to Merck over an approximately 15 to 24 month period starting on the Effective Date (the "Technology Transfer Period"). As part of this technology transfer, we will provide to Merck our proprietary enzymes, proprietary protein engineering protocols and methods, and proprietary software algorithms. In addition, teams of our and Merck scientists will participate in technology training sessions and collaborative research projects at our laboratories in Redwood City, California and at a designated Merck laboratory. Upon completion of technology transfer, Merck will have CodeEvolver® Platform Technology installed at its designated site.

The licenses to Merck are granted under patents, patent applications and know-how that we own or control as of the Effective Date and that cover the CodeEvolver® Platform Technology. Any improvements to the CodeEvolver® Platform Technology during the Technology Transfer Period will also be included in the license grants from Codexis to Merck. At the end of the Technology Transfer Period, Merck can exercise annual options that, upon payment of certain option fees, would extend Merck's license to include certain improvements to the CodeEvolver® Platform Technology that arise during the three-year period that begins at the end of the Technology Transfer Period. During the 15-month period that started on the Effective Date, we will provide additional enzyme evolution services to Merck at our laboratories in Redwood City.

The up-front license fee of \$5.0 million is being recognized ratably over a two-year period. We recognized license fees of \$0.4 million for the three months ended September 30, 2015, as biocatalyst research and development revenues and had a deferred revenue balance from Merck related to the Merck License Agreement license fees of \$4.6 million at September 30, 2015.

Under the Merck License Agreement, we will own any improvements to our protein engineering methods, processes and algorithms that arise and any enzyme technology or process technology that are developed during a technology transfer project, an evolution program or additional services. Merck will own (the "Merck-Owned Technology") (a) any enzyme technology that is developed solely by Merck under the Agreement using the CodeEvolver® Platform Technology (a "Project Enzyme") and (b) the methods of use of any Project Enzyme or any enzyme developed jointly by Merck and us using the CodeEvolver® Platform Technology. Merck granted to us a worldwide, non-exclusive, fully paid-up, royalty-free license, with the right to grant sublicenses, to use the Merck-Owned Technology outside of the Merck Exclusive Field.

For each API that Merck manufactures using an enzyme developed with the CodeEvolver® Platform Technology, we will have a right of first refusal to supply Merck with the enzyme used to manufacture the API if Merck outsources the supply of the enzyme. Our right of first refusal applies during the period that begins on the completion of a Phase III clinical trial for the product containing the API and ends five years following regulatory approval for such product. The Merck License Agreement has a term that begins on the Effective Date and continues, unless earlier terminated, until the expiration of all payment obligations under the agreement. Merck may terminate the Merck License Agreement by providing 90 days written notice to us. If Merck exercises this termination right during the Technology Transfer Period, Merck will make a one-term termination payment to us of \$8.0 million. We can terminate the Merck License Agreement by providing 30 days written notice to Merck if we determine, pursuant to our contractual audit rights under the agreement, that Merck has repeatedly failed to make required payments to us and/or materially underpaid us an amount due under the Merck License Agreement. In the event the Merck License Agreement is terminated earlier by Merck, or by us due to an uncured material breach by Merck, or if Merck sells or transfers to a third party any Merck business or facility that includes any of our proprietary materials, information or technology, we have the right to conduct an audit of Merck's facilities to confirm that all of our proprietary materials, information and technology have been destroyed. The Merck License Agreement contains indemnification provisions under which Merck and we indemnify each other against certain third party claims.

Note 5. Marketable Securities

At September 30, 2015, securities classified as available-for-sale consisted of the following (in thousands):

	September 30, 2015				
	Adjusted Cost	Gross Unrealized Gains	Gross Unrealized Losses	Estimated Fair Value	Average Contractual Maturities (in days)
Money market funds (1)	\$11,115	\$—	\$—	\$11,115	n/a
Common shares of CO2 Solutions (2)	563	668	—	1,231	n/a
Total	\$11,678	\$668	\$—	\$12,346	

(1) Money market funds are classified in cash and cash equivalents on our condensed consolidated balance sheets.

(2) Common shares of CO2 Solutions are classified as marketable securities on our condensed consolidated balance sheets.

There were no marketable securities in an unrealized loss position at September 30, 2015.

At December 31, 2014, securities classified as available-for-sale consisted of the following (in thousands):

	December 31, 2014				
	Adjusted Cost	Gross Unrealized Gains	Gross Unrealized Losses	Estimated Fair Value	Average Contractual Maturities (in days)
Money market funds (1)	\$14,602	\$—	\$—	\$14,602	n/a
Common shares of CO2 Solutions (2)	563	125	—	688	n/a
Total	\$15,165	\$125	\$—	\$15,290	

(1) Money market funds are classified in cash and cash equivalents on our condensed consolidated balance sheets.

(2) Common shares of CO2 Solutions are classified in marketable securities on our condensed consolidated balance sheets.

There were no marketable securities in an unrealized loss position at December 31, 2014.

Note 6. Fair Value Measurements

Fair Value of Financial Instruments

The following table presents the financial instruments that were measured at fair value on a recurring basis at September 30, 2015 by level within the fair value hierarchy (in thousands):

	September 30, 2015			
	Level 1	Level 2	Level 3	Total
Money market funds	\$11,115	\$—	\$—	\$11,115
Common shares of CO2 Solutions	—	1,231	—	1,231
Total	\$11,115	\$1,231	\$—	\$12,346

The following table presents the financial instruments that were measured at fair value on a recurring basis at December 31, 2014 by level within the fair value hierarchy (in thousands):

	December 31, 2014			
	Level 1	Level 2	Level 3	Total
Money market funds	\$14,602	\$—	\$—	\$14,602
Common shares of CO2 Solutions	—	688	—	688
Total	\$14,602	\$688	\$—	\$15,290

We estimated the fair value of our investment in 10,000,000 common shares of CO2 Solutions using the market value of common shares as determined by trading on the TSX Venture Exchange.

Note 7. Balance Sheets Details

Inventories

Inventories consisted of the following (in thousands):

	September 30, 2015	December 31, 2014
Raw materials	\$182	\$84
Work-in-process	11	65
Finished goods	485	1,246
Inventories	\$678	\$1,395

Property and Equipment, net

Property and equipment, net consisted of the following (in thousands):

	September 30, 2015	December 31, 2014
Laboratory equipment	\$21,354	\$23,002
Leasehold improvements	9,782	9,773
Computer equipment	3,271	3,262
Office furniture and equipment	1,227	1,227
Construction in progress (1)	151	24
Property and equipment	35,785	37,288
Less: accumulated depreciation and amortization	(32,964)	(31,452)
Less: impairment of laboratory equipment (2)	—	(1,841)
Property and equipment, net	\$2,821	\$3,995

(1) Construction in progress includes equipment received but not yet placed into service pending installation.

(2) We recorded an impairment charge of \$1.8 million in the second quarter of 2014, reducing the carrying value of certain laboratory equipment related to our Codexol program to zero. The impairment charge was reflected within research and development expenses on the condensed consolidated statements of operations.

Intangible Assets, net

Intangible assets, net consisted of the following (in thousands, except weighted average amortization period):

	September 30, 2015			December 31, 2014			Amortization Period (years)
	Gross Carrying Amount	Accumulated Amortization	Net Carrying Amount	Gross Carrying Amount	Accumulated Amortization	Net Carrying Amount	
Maxygen intellectual property	\$20,244	\$ (16,589)	\$3,655	\$20,244	\$ (14,058)	\$6,186	6

The estimated future amortization expense to be charged to research and development through the year ending December 31, 2016 is as follows (in thousands):

Year ending December 31:	Total
2015 (remaining 3 months)	\$843
2016	2,812
	\$3,655

Goodwill

Goodwill had a carrying value of approximately \$3.2 million at September 30, 2015 and December 31, 2014.

Note 8. Assets Held for Sale and Sale of Former Hungarian Subsidiary

In the fourth quarter of 2013, we announced that we would begin winding down our CodeXyme[®] cellulase enzyme program. As a result of the termination of this research program, we concluded that certain excess research and development equipment, including assets at our Hungarian subsidiary as well as some assets in the United States, were no longer needed and would be sold.

On March 13, 2014, we entered into an agreement with Intrexon Corporation to sell 100% of our equity interests in our Hungarian subsidiary, Codexis Laboratories Hungary Kft, as well as all assets of such subsidiary that were previously classified as held for sale. On March 15, 2014, the sale transaction closed and we received cash proceeds of \$1.5 million from the sale. Accordingly, we reduced the carrying value of assets held for sale by \$0.8 million and recognized a gain of \$0.8 million in connection with the sale which was included in research and development expenses. As part of the purchase, the buyer obtained all of the Hungarian assets held for sale and assumed all employment and facility lease related contract obligations. There were no transaction related costs incurred other than legal fees, which were recorded in selling, general and administrative expenses.

Prior to the sale of our Hungarian subsidiary in the first quarter of 2014, we transferred certain of the subsidiary's equipment to another of our European subsidiaries and incurred a reclaimable VAT liability of approximately \$0.4 million. We paid this VAT amount in July 2014 and recorded a receivable, which is reflected in prepaid expenses and other current assets in our condensed consolidated balance sheets at September 30, 2015 and December 31, 2014.

During the second quarter of 2014, we revised our plan to sell certain U.S. research and development equipment. As part of the revised plan, some equipment was returned to operational use. Additionally, we exchanged certain equipment for more suitable, newer equipment and recognized a loss of approximately \$0.2 million as part of the exchange. We also decided to expedite the disposal of other held for sale assets by selling these assets through auction which resulted in further impairment charges of \$0.6 million for the three months ended June 30, 2014. We disposed of the remaining held for sale equipment in the third quarter of 2014, which resulted in an additional impairment charge of \$0.1 million.

There were no assets classified as held for sale as of September 30, 2015 and as of December 31, 2014.

Note 9. Stock-Based Compensation

Equity Incentive Plans

In March 2010, our board of directors (the "Board") and stockholders approved the 2010 Equity Incentive Award Plan (the "2010 Plan"), which became effective upon the completion of our initial public offering in April 2010. The number of shares of our common stock available for issuance under the 2010 Plan is equal to 1,100,000 shares plus any shares of common stock reserved for future grant or issuance under our 2002 Stock Plan (the "2002 Plan") that remained unissued at the time of completion of the initial public offering. The 2010 Plan also provides for automatic annual increases in the number of shares reserved for future issuance. All grants will reduce the 2010 Plan reserve by one share for every share granted.

The 2010 Plan provides for the grant of incentive stock options, non-statutory stock options, RSUs, RSAs, PSUs, stock appreciation rights, and stock purchase rights to our employees, non-employee directors and consultants. The option exercise price for incentive stock options is at least 100% of the fair value of our common stock on the date of grant and the option exercise price for nonstatutory stock options is at least 85% of the fair value of our common stock on the date of grant, as determined by the Board. If, at the time of a grant, the optionee directly or by attribution owns stock possessing more than 10% of the total combined voting power of all of our outstanding capital stock, the exercise price for these options must be at least 110% of the fair value of the underlying common stock. Stock options granted to employees generally have a maximum term of 10 years and vest over a four year period from the date of grant 25% vest at the end of one year, and 75% vest monthly over the remaining three years. We may grant options with different vesting terms from time to time. Unless an employee's termination of service is due to disability or death, upon termination of service, any unexercised vested options will be forfeited at the end of three months or the expiration of the option, whichever is earlier.

We issue employees RSUs, which generally vest over either a three year period with 33% of the awards vesting on each annual anniversary or a four year period with 25% of the awards vesting on each annual anniversary. We may grant RSUs with different vesting terms from time to time.

Performance-contingent Restricted Stock Units

The compensation committee of the Board has approved grants of PSUs to employees. These awards have dual triggers of vesting based upon the successful achievement of certain corporate operating milestones in specified timelines, as well as a

20

requirement of continued employment. When the performance goals are deemed to be probable of achievement for these types of awards, time-based vesting and, as a result, recognition of stock-based compensation expense commences.

In the first quarter of 2015, we awarded PSUs ("2015 PSUs") based upon the achievement of various weighted performance criteria, including revenue growth, non-GAAP net income growth, new licensing collaborations, and securing a drug development partnership. The 2015 PSUs vest such that one-half of the PSUs subject to the award vest one year following the grant, and the remainder of the PSUs vest two years following the grant, subject to our achievement of the performance goals and the recipient's continued service on each vesting date. If the performance goal is achieved at the threshold level, the number of shares issuable in respect of the 2015 PSUs will be equal to half the number of PSUs granted. If the performance goal is achieved at the target level, the number of shares issuable in respect of the 2015 PSUs will be equal to the number of PSUs granted. If the performance goal is achieved at the superior level, the number of shares issuable in respect of the 2015 PSUs will be equal to two times the number of PSUs granted. The number of shares issuable upon achievement of the performance goal at the levels between the threshold and target levels or target level and superior levels is determined using linear interpolation. Achievement below the threshold level results in no shares being issuable in respect of the 2015 PSUs. During the three and nine months ended September 30, 2015, we evaluated our achievement against the performance criteria for the 2015 PSUs and recognized expense based on the estimated achievement rate.

In 2014 we awarded PSUs ("2014 PSUs") based upon the achievement of certain cash flow performance goals. The 2014 PSUs vest such that one-half of the PSUs subject to the award vest one year following the grant, and the remainder of the PSUs vest two years following the grant, subject to our achievement of the performance goals and the recipient's continued service on each vesting date. If the performance goal is achieved at the threshold level, the number of shares issuable in respect of the 2014 PSUs will be equal to half the number of PSUs granted. If the performance goal is achieved at the target level, the number of shares issuable in respect of the 2014 PSUs will be equal to the number of PSUs granted. If the performance goal is achieved at the superior level, the number of shares issuable in respect of the 2014 PSUs will be equal to two times the number of PSUs granted. The number of shares issuable upon achievement of the performance goal at the levels between the threshold and target levels or target level and superior levels is determined using linear interpolation. Achievement below the threshold level results in no shares being issuable in respect of the 2014 PSUs. During the third quarter of 2014, we concluded that it was not probable that the performance objective would be achieved at the target level of 100%, and we reduced stock-based compensation expense to reflect a lower level of estimated achievement.

Stock-Based Compensation Expense

Stock-based compensation expense is included in the consolidated statements of operations as follows (in thousands):

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2015	2014	2015	2014
Research and development (1)	\$181	\$227	\$710	\$734
Selling, general and administrative	1,042	828	3,049	2,896
Total	\$1,223	\$1,055	\$3,759	\$3,630

(1) Stock-based compensation expense associated with cost of biocatalyst product sales is included in research and development. Amounts were immaterial for all periods presented.

The following table presents total stock-based compensation expense by security types included in the condensed consolidated statements of operations for the three and nine months ended September 30, 2015 and 2014 (in thousands):

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2015	2014	2015	2014
Stock options	\$281	\$247	\$798	\$843
RSUs and RSAs	566	722	2,020	2,383
PSUs	376	86	941	404
Total	\$1,223	\$1,055	\$3,759	\$3,630

As of September 30, 2015, unrecognized stock-based compensation expense, net of expected forfeitures, was \$2.0 million related to unvested employee stock options, \$2.1 million related to unvested RSUs and RSAs and \$0.9 million related to unvested PSUs.

Valuation Assumptions

The weighted-average assumptions used to estimate the fair value of employee stock options granted were as follows:

	Three Months Ended September 30,		Nine Months Ended September 30,		
	2015	2014	2015	2014	
Expected term (in years) (1)	5.2	6.0	6.0	6.0	
Volatility	67	% 68	% 66	% 65	%
Risk-free interest rate	1.64	% 1.91	% 1.70	% 1.915	%
Dividend yield	—	% —	% —	% —	%
Weighted-average estimated fair value of stock options granted	\$2.31	\$1.43	\$2.09	\$1.20	

(1) We have, due to insufficient historical data, used the simplified method to determine the expected term of stock options granted. In the third quarter of 2015, we have applied historical data to calculate an expected term for stock options granted (see Note 2, "Basis of Presentation and Summary of Significant Accounting Policies").

Note 10. Capital Stock

Exercise of options

For the nine months ended September 30, 2015 and 2014, 128,921 and 136,796 shares were exercised at a weighted-average exercise price of \$1.82 and \$1.34 per share, respectively, with net cash proceeds of \$0.2 million for both periods.

Warrants

Our outstanding warrants are exercisable for common stock at any time during their respective terms. As of September 30, 2015, the following warrants remain outstanding:

September 30, 2015

Issue Date	Shares Subject to Warrants	Exercise Price per Share	Expiration
July 17, 2007	2,384	\$12.45	February 9, 2016
September 28, 2007	72,727	\$8.25	September 28, 2017

Note 11. Commitments and Contingencies

Operating Leases

Our headquarters are located in Redwood City, California, where we occupy approximately 107,000 square feet of office and laboratory space in four buildings within the same business park of Metropolitan Life Insurance Company ("Met-Life"). We entered into the initial lease with Met-Life for a portion of this space in 2004 and the lease has been amended multiple times since then to adjust space and amend the terms of the lease, with the latest amendment being in 2012. The various terms for the spaces under the lease have expiration dates that range from January 2017 through January 2020.

We incurred \$3.6 million of capital improvement costs related to the facilities leased from Met-Life through December 31, 2012. During 2011 and 2012, we requested and received \$3.1 million of reimbursements from the landlord from the tenant improvement and HVAC allowances for the completed construction. The reimbursements were recorded once cash was received and are amortized on a straight line basis over the term of the lease as a reduction in rent expense. The remaining lease incentive obligation was \$1.4 million at September 30, 2015, and is reflected in other liabilities on the consolidated balance sheet. Rent expense for the Redwood City properties is recognized on a straight-line basis over the term of the lease.

We are required to restore certain of the Redwood City facilities that we are renting to their original form. We are expensing the asset retirement obligation over the terms of the respective leases. We review the estimated obligation each reporting period and make adjustments if our estimates change. In 2014, we entered into a sublease agreement whereby certain changes were made to our facility by our sublessor. As such, on December 31, 2014, we revised our estimated asset retirement obligation to restore the sublet facility to its original form and recognized an asset retirement obligation of \$0.3 million and correspondingly increased our related estimated cash payments. Accretion expense related to our asset retirement obligations

was nominal in each of the three and nine months ended September 30, 2015 and nil in each of the three and nine months ended September 30, 2014.

In accordance with the terms of the amended lease agreement, we exercised our right to deliver a letter of credit in lieu of a security deposit. The letters of credit are collateralized by deposit balances held by the bank in the amount of \$0.7 million as of September 30, 2015 and December 31, 2014. These deposits are recorded as restricted cash on the consolidated balance sheets.

Prior to March 2014, we also rented facilities in Hungary. Rent expense was being recognized on a straight-line basis over the respective terms of the leases. The facility lease was transferred to Intrexon Corporation in connection with the sale of Codexis Laboratories Hungary Kft (see Note 8, "Assets Held for Sale and Sale of Former Hungarian Subsidiary").

Rent expense was \$0.9 million and \$2.6 million in the three and nine months ended September 30, 2015, respectively, partially offset by sublease income of \$0.2 million and \$0.5 million, respectively. Rent expense was \$0.9 million and \$2.5 million in the three and nine months ended September 30, 2014, respectively, partially offset by sublease income of \$0.1 million and \$0.3 million during the respective periods in 2014.

Future minimum payments under noncancellable operating leases are as follows at September 30, 2015 (in thousands):

	Lease payments
Years ending December 31, 2015 (3 months remaining)	\$689
2016	2,827
2017	2,677
2018	2,736
2019 and beyond	3,054
Total	\$11,983

Legal Proceedings

From time to time we are involved in various legal proceedings related to matters that have arisen during the ordinary course of business. Although there can be no assurance as to the ultimate disposition of these matters, we have determined, based upon the information available, that the expected outcome of these matters, individually or in the aggregate, will not have a material adverse effect on our consolidated financial position, results of operations or cash flows.

Other Contingencies

In November 2009, one of our foreign subsidiaries sold intellectual property to Codexis, Inc. Under the local laws, the sale of intellectual property to a nonresident legal entity is deemed an export and is not subject to VAT. However, there is uncertainty regarding whether the items sold represented intellectual property or research and development services, which would subject the sale to VAT. We believe that the uncertainty results in an exposure to pay VAT that is more than remote but less than likely to occur and, accordingly, we have not recorded an accrual for this exposure. If the sale is deemed a sale of research and development services, we could be obligated to pay an estimated amount of \$0.6 million.

Indemnifications

We are required to recognize a liability for the fair value of any obligations we assume upon the issuance of a guarantee. We have certain agreements with licensors, licensees and collaborators that contain indemnification provisions. In such provisions, we typically agree to indemnify the licensor, licensee and collaborator against certain types of third party claims. The maximum amount of the indemnifications is not limited. We accrue for known indemnification issues when a loss is probable and can be reasonably estimated. There were no accruals for expenses related to indemnification issues for any periods presented.

Note 12. Related Party Transactions

Exela PharmSci, Inc.

We signed a commercialization agreement with Exela in 2007, whereby Exela agreed to pay to us a contractual percentage share of Exela's net profit from the sales of licensed products.

Thomas R. Baruch, one of our directors, serves on the board of directors of Exela and is a limited partner in Presidio Partners 2007, L.P., which owns more than 10% of Exela's outstanding capital stock. Consequently, Mr. Baruch has an indirect pecuniary interest in the shares of Exela held by Presidio Partners 2007, L.P. Mr. Baruch is also a limited partner in CMEA Ventures, which owned 7.4% of our common stock until November 10, 2014, at which time the shares were transferred to Presidio Partners 2014, L.P. Mr. Baruch has no direct or indirect pecuniary interest in the shares of our common stock owned by Presidio Partners 2014, L.P.

We recognized \$1.1 million and \$4.1 million for the three and nine months ended September 30, 2015, respectively, and \$1.5 million and \$5.6 million for the three and nine months ended September 30, 2014, respectively, shown in the consolidated statement of operations as revenue sharing arrangement. We had receivables of \$0.3 million at September 30, 2015 and no receivables at December 31, 2014 from Exela.

Alexander A. Karsner

Alexander A. Karsner was a member of Board until the expiration of his term at the close of our Annual Meeting of Stockholders on June 11, 2014. In addition, Mr. Karsner provided consulting services to us beginning in 2011 through June 30, 2014. Amounts paid to Mr. Karsner for consulting services were nil for the three and nine months ended September 30, 2015 and nil and \$60,000 for the three and nine months ended September 30, 2014, respectively.

Note 13. Significant Customer and Geographic Information

Significant Customers

Customers that each contributed 10% or more of our total revenues were as follows:

	Percentage of Total Revenues for the Three Months Ended September 30,		Nine Months Ended September 30,		
	2015	2014	2015	2014	
Customer A	39	% 27	% 34	% 26	%
Customer B	40	% *	26	% *	
Customer C (related party)	*	21	% 13	% 27	%

* Less than 10% in period presented

Of the customers that contributed 10% or more of our total revenues, the following had accounts receivable balances for the periods presented:

	Percentage of Accounts Receivables at		
	September 30, 2015	December 31, 2014	
Customer A	42	% 63	%
Customer B	47	% 2	%
Customer C (related party)	*	—	%

* Revenue percentage was less than 10%, accounts receivable balance not applicable

Geographic Information

Geographic revenues are identified by the location of the customer and consist of the following (in thousands):

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2015	2014	2015	2014
Revenues:				
United States	\$8,755	\$4,747	\$16,516	\$12,518
Asia				
India	369	225	519	636
Others	619	658	1,305	1,338
Europe				
Ireland	160	—	160	2,744
Others	7,498	1,842	11,712	3,864
Other	—	—	6	16
Total revenues	\$17,401	\$7,472	\$30,218	\$21,116

Identifiable long-lived assets were all in the United States as follows (in thousands):

	September 30,	December 31,
	2015	2014
Long-lived assets		
United States	\$6,741	\$10,475

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

The following discussion and analysis of our financial condition and results of operations should be read in conjunction with the unaudited condensed consolidated financial statements and the related notes thereto included elsewhere in this Quarterly Report on Form 10-Q and the audited consolidated financial statements and notes thereto and management's discussion and analysis of financial condition and results of operations for the year ended December 31, 2014 included in our Annual Report on Form 10-K filed with the SEC on March 6, 2015. This Quarterly Report on Form 10-Q contains forward-looking statements within the meaning of Section 21E of the Securities Exchange Act of 1934, as amended. These statements are often identified by the use of words such as may, will, expect, believe, anticipate, intend, could, should, estimate, or continue, and similar expressions or variations. Such forward-looking statements are subject to risks, uncertainties and other factors that could cause actual results and the timing of certain events to differ materially from future results expressed or implied by such forward-looking statements. Factors that could cause or contribute to such differences include, but are not limited to, those set forth in Part II, Item 1A of this Quarterly Report on Form 10-Q and in Part I, Item 1A of our Annual Report on Form 10-K for the year ended December 31, 2014 filed with the SEC on March 6, 2015, as incorporated herein and referenced in Part II, Item 1A of this Quarterly Report on Form 10-Q and elsewhere in this report. The forward-looking statements in this Quarterly Report on Form 10-Q represent our views as of the date of this Quarterly Report on Form 10-Q. We anticipate that subsequent events and developments will cause our views to change. However, while we may elect to update these forward-looking statements at some point in the future, we have no current intention of doing so except to the extent required by applicable law. You should, therefore, not rely on these forward-looking statements as representing our views as of any date subsequent to the date of this Quarterly Report on Form 10-Q.

Business Overview

We develop biocatalysts for the pharmaceutical and fine chemicals markets. Our proven technologies enable scale-up and implementation of biocatalytic solutions to meet customer needs for rapid, cost-effective and sustainable process development, from research to manufacturing.

Biocatalysts are enzymes that initiate and/or accelerate chemical reactions. Manufacturers have historically used naturally occurring biocatalysts to produce many goods used in everyday life. However, inherent limitations in naturally occurring biocatalysts have restricted their commercial use. Our proprietary CodeEvolver[®] protein engineering technology platform (the "CodeEvolver[®] Platform Technology"), which introduces genetic mutations into microorganisms in order to give rise to changes in enzymes that they produce, is able to overcome many of these limitations, allowing us to evolve and optimize biocatalysts to perform specific and desired chemical reactions at commercial scale. Once potentially beneficial mutations are identified through this proprietary process, combinations of these mutations can then be tested until variant enzymes have been created that exhibit marketable performance characteristics superior to competitive products. This process allows for continuous, efficient improvements to the performance of enzymes. In the past, we implemented the CodeEvolver[®] Platform Technology through paid collaborations with our customers. In July 2014, we entered into our first license agreement pursuant to which we granted a license to a global pharmaceutical company to use the CodeEvolver[®] Platform Technology for their internal development purposes. In August 2015, we entered into a license agreement involving the CodeEvolver[®] Platform Technology with a second global pharmaceutical company and we continue to pursue licensing opportunities with additional customers.

We have commercialized our technology and products in the pharmaceuticals market, which is our primary business focus. Our customers, which include several large global pharmaceutical companies, use our technology, products and services in their manufacturing processes and process development.

We also use our technology to develop biocatalysts for use in the fine chemicals market. The fine chemicals market is similar to our pharmaceutical business and consists of several large market verticals, including food, animal feed, flavors, fragrances, and agricultural chemicals.

We are also using our technology to develop an early stage, novel enzyme therapeutic product candidate for the potential treatment of phenylketonuria ("PKU") in humans. PKU is an inherited metabolic disorder in which the enzyme that converts the essential amino acid phenylalanine into tyrosine is deficient.

We are actively collaborating with new and existing customers in the pharmaceutical and other markets and we believe that we can utilize our products and services, and develop new products and services, to increase our revenue and gross margins in future periods.

Results of Operations Overview

Revenues were \$17.4 million for the third quarter of 2015, an increase of 133% from \$7.5 million for the third quarter of 2014. The increase in revenues was due to the achievement of milestones of \$5.0 million and \$6.5 million under our collaborative arrangements with Merck Sharp & Dohme Corp., known as MSD outside the United States and Canada ("Merck"), and GlaxoSmithKline ("GSK"), respectively, during the third quarter of 2015, partially offset by lower biocatalyst product sales and lower revenues from our revenue sharing arrangement with Exela PharmSci, Inc. ("Exela").

Revenues from our revenue sharing arrangement with Exela decreased by \$0.5 million, or 31%, to \$1.1 million for the third quarter of 2015 compared to the same period in 2014. The decrease was due to lower sales of the argatroban injectable drug, which resulted from the expiration of the formulation patent for argatroban in June 2014, allowing for generic competition in the subsequent quarters.

Biocatalyst product sales decreased by \$0.7 million, or 29%, to \$1.8 million for the third quarter of 2015 compared to the corresponding period in 2014, primarily due to the timing of customer demands.

Biocatalyst research and development revenues were \$14.5 million for the third quarter of 2015, an increase of \$11.2 million, or 332%, compared to the third quarter of 2014, driven by the achievement of milestones under our collaborative arrangements with Merck and GSK.

Cost of biocatalyst product sales was \$1.3 million for the third quarter of 2015, a decrease of 15% from \$1.5 million for the third quarter of 2014, due primarily to the decrease of biocatalyst product sales. Biocatalyst product gross margin in the third quarter of 2015 was 28% compared to 40% in the same period in 2014, due to changes in the product sales mix.

Research and development expenses remained flat at \$5.0 million for the third quarter of 2015 compared to the same period in 2014. Research and development expenses in the third quarter of 2015 included higher lab supply expenses offset by lower depreciation expenses and lower contractor charges.

Selling, general and administrative expenses were \$5.4 million for the third quarter of 2015, an increase of 5% from \$5.2 million for the third quarter of 2014. The increase in general and administrative expenses for the third quarter of 2015 was primarily driven by higher personnel-related expenses (including stock-based compensation), partially offset by lower legal expenses and contractor charges.

Net income for the third quarter of 2015 was \$5.4 million, representing basic net income of \$0.14 per share. This compares favorably to a net loss of \$4.6 million, or net loss of \$0.12 per share, for the third quarter of 2014. The net income position is primarily related to the achievement of the milestones from the Merck License Agreement and the GSK License Agreement.

Cash and cash equivalents decreased to \$17.0 million as of September 30, 2015 compared to \$26.5 million as of December 31, 2014. Net cash used in operating activities increased to \$7.5 million in the nine months ended September 30, 2015 compared to \$4.7 million in the nine months ended September 30, 2014. We are actively collaborating with new and existing customers in the pharmaceutical and other markets and we believe that we can utilize our products and services, and develop new products and services, to increase our revenue and gross margins in future periods. We believe that based on our current level of operations, our existing cash, cash equivalents, and marketable securities will provide adequate funds for ongoing operations, planned capital expenditures and working capital requirements for at least the next 12 months.

GSK Platform Technology Transfer, Collaboration and License Agreement

In July 2014, we entered into a CodeEvolver[®] platform technology transfer, collaboration and license agreement (the "GSK License Agreement") with GSK. Under the terms of the GSK License Agreement, we granted GSK a non-exclusive license to use the CodeEvolver[®] Platform Technology to develop novel enzymes for use in the manufacture of GSK's pharmaceutical and health care products.

We received a \$6.0 million up-front licensing fee upon signing the GSK License Agreement and subsequently a \$5.0 million non-creditable, non-refundable milestone payment upon achievement of the first milestone. During the three months ended September 30, 2015, we achieved the second milestone of the agreement earning another milestone payment of \$6.5 million. We are eligible to receive an additional contingent payment of \$7.5 million upon the completion of the technology transfer period. We also have the potential to receive additional contingent payments that range from \$5.75 million to \$38.5 million per project based on GSK's successful application of the licensed

technology. The contingent payments are not deemed substantive milestones due to the fact that the achievement of the event underlying the payment predominantly relates to GSK's performance of future development and commercialization activities. We do not expect to begin receiving these additional contingent payments, if any, during the first three years of the GSK License Agreement.

For up to three years following the end of the three-year period during which we will transfer the CodeEvolver® Platform Technology to GSK, GSK can exercise an option, upon payment of certain additional fees, that would extend GSK's license to include certain improvements to the CodeEvolver® Platform Technology that arise during such period. In addition, we will also be eligible to receive royalties based on net sales, if any, of a limited set of products developed by GSK using the CodeEvolver® Platform Technology.

The term of the GSK License Agreement continues, unless earlier terminated, until the expiration of all payment obligations under the GSK License Agreement. At any time following the completion of the first technology transfer stage, GSK can terminate the GSK License Agreement by providing 90 days written notice to us. If GSK exercises this termination right during the three-year technology transfer period, GSK will make a one-time termination payment to us.

Under the GSK License Agreement, the significant deliverables were determined to be the license, platform technology transfer, and contingent obligation to supply GSK with enzymes manufactured by us at GSK's expense. We determined that the license did not have stand-alone value, and we determined that the license and the platform technology transfer and our participation in joint steering committee activities in connection with the platform technology transfer represent a single unit of accounting. We determined that our participation in the joint steering committee does not represent a separate unit of accounting because GSK could not negotiate for and/or acquire these services from other third parties and our participation on the joint steering committee is coterminous with the technology transfer period. Amounts to be received under the supply arrangement described above will be recognized as revenue to the extent GSK purchases enzymes from us.

The up-front license fee of \$6.0 million is being recognized over the technology transfer period of three years. We recognized license fees of \$0.5 million and \$1.5 million for the three and nine months ended September 30, 2015, respectively, and \$0.5 million for the three and nine months ended September 30, 2014 as biocatalyst research and development revenues. We had a deferred revenue balance from GSK related to the upfront license fee of \$3.5 million at September 30, 2015 and \$5.0 million at December 31, 2014.

Merck Platform Technology Transfer and License Agreement

On August 3, 2015 ("Effective Date"), we entered into a CodeEvolver® platform technology transfer and license agreement (the "Merck License Agreement") with Merck. The Agreement allows Merck to use the CodeEvolver® Platform Technology in the field of human and animal healthcare.

We received a \$5.0 million up-front licensing fee upon signing the Merck License Agreement. During the three months ended September 30, 2015, we achieved the first milestone of the Merck License Agreement earning a milestone payment of \$5.0 million. We are eligible to receive an additional \$8.0 million subject to the satisfactory completion of the second milestone of the technology transfer process. We will also be eligible to receive payments of up to a maximum of \$15.0 million for each commercial active pharmaceutical ingredient ("API") that is manufactured by Merck using one or more novel enzymes developed by Merck using the CodeEvolver® Platform Technology. Under the terms of the Merck License Agreement, we granted to Merck a non-exclusive, worldwide license to use the CodeEvolver® Platform Technology to research, develop and manufacture novel enzymes for use by Merck in its internal research programs ("Merck Non-Exclusive Field"). The license to Merck is exclusive for the research, development and manufacture of novel enzymes for use by Merck in the chemical synthesis of therapeutic products owned or controlled by Merck ("Merck Exclusive Field"). Merck has the right to grant sublicenses to affiliates of Merck and, in certain limited circumstances, to third parties. We also granted to Merck a license to make or have made products manufactured using the CodeEvolver® Platform Technology with a right to grant sublicenses solely to affiliates of Merck, contract manufacturing organizations and contract research organizations. The manufacturing license is exclusive in the Merck Exclusive Field and non-exclusive in the Merck Non-Exclusive Field. The licenses are subject to certain limitations based on pre-existing contractual obligations that apply to the technology and intellectual property that are the subject of the license grants. The licenses do not permit the use of the CodeEvolver® Platform Technology to discover any therapeutic enzyme, diagnostic product or vaccine. In addition, Merck is prohibited from using the CodeEvolver® Platform Technology to develop or produce enzymes or any other compounds for or on behalf of any third parties except in a very limited manner when Merck divests a therapeutic product that is manufactured using an enzyme developed using the CodeEvolver® Platform Technology.

Under the Merck License Agreement, we will transfer the CodeEvolver® Platform Technology to Merck over an approximately 15 to 24 month period starting on the Effective Date (the "Technology Transfer Period"). As part of this technology transfer, we will provide to Merck our proprietary enzymes, proprietary protein engineering protocols and methods, and proprietary software algorithms. In addition, teams of our and Merck scientists will participate in technology training sessions and collaborative research projects at our laboratories in Redwood City, California and at a designated Merck laboratory. Upon completion of technology transfer, Merck will have CodeEvolver® Platform Technology installed at its designated site.

The licenses to Merck are granted under patents, patent applications and know-how that we own or control as of the Effective Date and that cover the CodeEvolver® Platform Technology. Any improvements to the CodeEvolver® Platform Technology during the Technology Transfer Period will also be included in the license grants from Codexis to Merck. At the end of the Technology Transfer Period, Merck can exercise annual options that, upon payment of certain option fees, would extend Merck's license to include certain improvements to the CodeEvolver® Platform Technology that arise during the three-year period that begins at the end of the Technology Transfer Period. During the 15-month period that started on the Effective Date, we will provide additional enzyme evolution services to Merck at our laboratories in Redwood City.

The up-front license fee of \$5.0 million is being recognized ratably over a two-year period. We recognized license fees of \$0.4 million for the three months ended September 30, 2015 as biocatalyst research and development revenue and had a deferred revenue balance from Merck related to the Merck License Agreement license fees of \$4.6 million at September 30, 2015.

Under the Merck License Agreement, we will own any improvements to our protein engineering methods, processes and algorithms that arise and any enzyme technology or process technology that are developed during a technology transfer project, an evolution program or additional services. Merck will own (the "Merck-Owned Technology") (a) any enzyme technology that is developed solely by Merck under the Agreement using the CodeEvolver® Platform Technology (a "Project Enzyme") and (b) the methods of use of any Project Enzyme or any enzyme developed jointly by Merck and us using the CodeEvolver® Platform Technology. Merck granted to us a worldwide, non-exclusive, fully paid-up, royalty-free license, with the right to grant sublicenses, to use the Merck-Owned Technology outside of the Merck Exclusive Field.

For each API that Merck manufactures using an enzyme developed using the CodeEvolver® Platform Technology, we will have a right of first refusal to supply Merck with the enzyme used to manufacture the API if Merck outsources the supply of the enzyme. Our right of first refusal applies during the period that begins on the completion of a Phase III clinical trial for the product containing the API and ends five years following regulatory approval for such product. The Merck License Agreement has a term that begins on the Effective Date and continues, unless earlier terminated, until the expiration of all payment obligations under the agreement. Merck may terminate the Merck License Agreement by providing 90 days written notice to us. If Merck exercises this termination right during the Technology Transfer Period, Merck will make a one-term termination payment to us of \$8.0 million. We can terminate the Merck License Agreement by providing 30 days written notice to Merck if we determine, pursuant to our contractual audit rights under the agreement, that Merck has repeatedly failed to make required payments to us and/or materially underpaid us an amount due under the Merck License Agreement. In the event the Merck License Agreement is terminated earlier by Merck, or by us due to an uncured material breach by Merck, or if Merck sells or transfers to a third party any Merck business or facility that includes any of our proprietary materials, information or technology, we have the right to conduct an audit of Merck's facilities to confirm that all of our proprietary materials, information and technology have been destroyed. The Merck License Agreement contains indemnification provisions under which Merck and we indemnify each other against certain third party claims.

Results of Operations

The following table shows the amounts from our consolidated statements of operations for the periods presented (in thousands):

	Three months ended September 30,		Change		Nine months ended September 30,		Change	
	2015	2014	\$	%	2015	2014	\$	%
Revenues:								
Biocatalyst product sales	\$1,818	\$2,562	\$(744)	(29)%	\$6,915	\$8,323	\$(1,408)	(17)%
Biocatalyst research and development	14,517	3,364	11,153	332%	19,247	7,176	12,071	168%
Revenue sharing arrangement	1,066	1,546	(480)	(31)%	4,056	5,617	(1,561)	(28)%
Total revenues	17,401	7,472	9,929	133%	30,218	21,116	9,102	43%
Costs and operating expenses:								
Cost of biocatalyst product sales	1,302	1,532	(230)	(15)%	4,009	6,179	(2,170)	(35)%
Research and development	4,994	5,038	(44)	(1)%	15,457	17,708	(2,251)	(13)%
Selling, general and administrative	5,415	5,157	258	5%	16,289	16,791	(502)	(3)%
Total costs and operating expenses	11,711	11,727	(16)	—%	35,755	40,678	(4,923)	(12)%
Income (loss) from operations	5,690	(4,255)	9,945	(234)%	(5,537)	(19,562)	14,025	(72)%
Interest income	4	3	1	33%	12	15	(3)	(20)%
Other expenses	(26)	(57)	31	(54)%	(147)	(183)	36	(20)%
Income (loss) before income taxes	5,668	(4,309)	9,977	(232)%	(5,672)	(19,730)	14,058	(71)%
Provision for (benefit from) income taxes	274	253	21	8%	(144)	(314)	170	(54)%
Net income (loss)	\$5,394	\$(4,562)	\$9,956	(218)%	\$(5,528)	\$(19,416)	\$13,888	(72)%

Our revenues are comprised of biocatalyst product sales, biocatalyst research and development arrangements, and a revenue sharing arrangement.

- Biocatalyst product sales revenues consist of sales of biocatalyst enzymes, chemical intermediates, and CodeX® Biocatalyst Panels and Kits.

- Biocatalyst research and development revenues include license, technology access and exclusivity fees, research services FTE, contingent payments, royalties, and optimization and screening fees.

- Revenue sharing arrangement revenues are recognized based upon sales of licensed products by Exela.

(In Thousands)	Three months ended September 30,		Change		Nine months ended September 30,		Change	
	2015	2014	\$	%	2015	2014	\$	%
Biocatalyst product sales	\$1,818	\$2,562	\$(744)	(29)%	\$6,915	\$8,323	\$(1,408)	(17)%
Biocatalyst research and development	14,517	3,364	11,153	332%	19,247	7,176	12,071	168%
Revenue sharing arrangement	1,066	1,546	(480)	(31)%	4,056	5,617	(1,561)	(28)%
Total revenues	\$17,401	\$7,472	\$9,929	133%	\$30,218	\$21,116	\$9,102	43%

Typically, revenues fluctuate on a quarterly basis due to the variability in our customers' manufacturing schedules and the timing of our customers' clinical trials. In addition, we have limited internal capacity to manufacture enzymes, and as a result, we are dependent upon the performance and capacity of third party manufacturers for the commercial scale manufacturing of the enzymes used in our pharmaceutical and fine chemicals business.

We accept purchase orders for deliveries covering periods from one day up to approximately one year from the date on which the order is placed. However, purchase orders can generally be revised or cancelled by the customer without penalty. Considering these industry practices and our experience, we do not believe the total of customer purchase orders outstanding (backlog) provides meaningful information that can be relied on to predict actual sales for future periods.

Total revenues increased \$9.9 million and \$9.1 million in the three and nine months ended September 30, 2015, respectively, compared to the same periods in 2014 as a result of the increase in biocatalyst research and development revenues, partially offset by decreases in biocatalyst product sales and revenues from our revenue-sharing arrangement with Exela.

Biocatalyst product sales decreased \$0.7 million and \$1.4 million in the three and nine months ended September 30, 2015, respectively, compared to the same periods in 2014. The decreases were primarily due to the timing of customer demand during the three and nine months ended September 30, 2015 compared to the same periods in 2014.

Biocatalyst research and development revenues increased approximately \$11.2 million and \$12.1 million in the three and nine months ended September 30, 2015 as a result of the achievement of milestones of \$5.0 million and \$6.5 million from the Merck License Agreement and the GSK License Agreement, respectively, during the third quarter of 2015.

Revenues from the revenue-sharing arrangement with Exela for the sales of argatroban injectable drug decreased \$0.5 million and \$1.6 million during the three and nine months ended September 30, 2015, respectively, compared to the same periods in 2014. The decrease is the result of the expiration of the formulation patent for argatroban in June 2014, allowing for generic competition in the subsequent quarters. We believe that revenues from our revenue-sharing arrangement with Exela may continue to decline in future quarters due to increasing generic competition.

Cost and Operating Expenses

(In Thousands)	Three months ended September 30,		Change		Nine months ended September 30,		Change	
	2015	2014	\$	%	2015	2014	\$	%
Cost of biocatalyst product sales	\$1,302	\$1,532	\$(230)	(15)%	\$4,009	\$6,179	\$(2,170)	(35)%
Research and development	4,994	5,038	(44)	(1)%	15,457	17,708	(2,251)	(13)%
Selling, general and administrative	5,415	5,157	258	5%	16,289	16,791	(502)	(3)%
Total operating expenses	\$11,711	\$11,727	\$(16)	—%	\$35,755	\$40,678	\$(4,923)	(12)%

Cost of Biocatalyst Product Sales and Product Gross Margin

(In Thousands)	Three months ended September 30,		Change		Nine months ended September 30,		Change	
	2015	2014	\$	%	2015	2014	\$	%
Biocatalyst product sales	\$1,818	\$2,562	\$(744)	(29)%	\$6,915	\$8,323	\$(1,408)	(17)%
Cost of biocatalyst product sales	1,302	1,532	(230)	(15)%	4,009	6,179	(2,170)	(35)%
Biocatalyst product gross profit	\$516	\$1,030	\$(514)	(50)%	\$2,906	\$2,144	\$762	36%
Product gross margin (%)	28%	40%			42%	26%		

Cost of biocatalyst product sales comprises both internal and third-party fixed and variable costs, including the amortization of purchased technology, materials and supplies, labor, facilities and other overhead costs associated with our biocatalyst product sales.

Our cost of biocatalyst product sales decreased by \$0.2 million, or 15%, during the three months ended September 30, 2015 and \$2.2 million, or 35%, during the nine months ended September 30, 2015 due to lower biocatalyst product sales compared to the same periods in 2014. Product gross margins were 28% and 42% in the three and nine months ended September 30, 2015, compared to 40% and 26% in the same periods in 2014 due to changes in the product sales mix.

Research and Development Expenses

Research and development expenses consist of costs incurred for internal projects as well as partner-funded collaborative research and development activities. These costs primarily consist of (i) employee-related costs, which include salaries and other personnel-related expenses (including stock-based compensation), (ii) various allocable expenses, which include occupancy-related costs, supplies, depreciation of facilities and laboratory equipment and amortization of acquired technologies, and (iii) external costs. Research and development expenses are expensed when incurred. We budget total

31

research and development expenses on an internal department level because we do not have project or program level reporting capabilities.

Research and development expenses remained flat for the third quarter of 2015 and decreased by \$2.3 million during the nine months ended September 30, 2015 compared to the same periods in 2014. Research and development expenses for the nine months ended 2014 included non-recurring non-cash impairment charges of \$2.7 million, of which \$1.8 million was related to the write down of assets associated with our CodeXol program and the remainder due to changes in fair value of assets held for sale. Additionally, research and development expenses for the first quarter of 2014 included a \$0.8 million gain from the sale of our former Hungarian subsidiary. Excluding such non-recurring charges, research and development expenses decreased \$0.4 million during the nine months ended September 30, 2015 compared to the same periods in 2014, primarily due to a decrease in depreciation expenses resulting from the aforementioned impairments and the sale of our Hungarian subsidiary in 2014, partially offset by an increase in employee-related expenses of \$0.4 million in the nine months ended September 30, 2015 compared to the same period in 2014.

Selling, General and Administrative Expenses

Selling, general and administrative expenses consist of employee-related costs, which include salaries and other personnel related expenses (including stock-based compensation), hiring and training costs, consulting and outside services expenses (including audit and legal counsel related costs), marketing costs, building lease costs, and depreciation and amortization expenses.

Selling, general and administrative expenses increased marginally by \$0.3 million, or 5%, during the third quarter of 2015 and decreased by \$0.5 million, or 3%, for the nine months ended September 30, 2015, compared to the same periods in 2014. The increase during the third quarter of 2015 was driven by increases in personnel-related expenses, including an increase in stock-based compensation expense of \$0.2 million, partially offset by decreases in legal, insurance and contractor expenses. The decrease during the nine months ended September 30, 2015 related primarily to a decrease in legal fees.

Interest income and other expenses

(In Thousands)	Three months ended September 30,		Change		Nine months ended September 30,		Change		
	2015	2014	\$	%	2015	2014	\$	%	
Interest income	\$4	\$3	\$1	33	% \$12	\$15	\$(3)	(20)	%
Other expenses	(26)	(57)	31	(54))% (147)	(183)	36	(20))%
Total other expenses	\$(22)	\$(54)	\$32	(59))% \$(135)	\$(168)	\$33	(20))%

Changes to our interest income were marginal during the three and nine months ended September 30, 2015 compared to the same periods in 2014, driven primarily by changes in our cash and cash equivalent balances.

Other expenses decreased for the three and nine months ended September 30, 2015 compared to the same periods in 2014. These changes were primarily related to fluctuations in foreign currency.

Provision for (benefit from) income taxes

We recognized an income tax expense of \$0.3 million for both the three months ended September 30, 2015 and 2014. We recognized an income tax benefit of \$0.1 million for the nine months ended September 30, 2015, as compared to \$0.3 million for the same period in 2014. The decrease was primarily due to the release of reserves related to uncertain tax positions that was recorded in 2014 whereas no release was recorded in 2015.

The tax benefit for the nine months ended September 30, 2015 primarily related to unrecognized gains from changes in the fair value of our investment in CO2 Solutions. The tax benefit for the nine months ended September 30, 2014 primarily consisted of income tax benefit attributable to foreign operations offset by the tax effect of the unrecognized gain from our investment in CO2 Solutions. Additionally, for the nine months ended September 30, 2014, we recognized approximately \$0.4 million of previously unrecognized tax benefits related to our operations in Singapore.

We continue to recognize a full valuation allowance against our net deferred tax assets as we believe that it is more likely than not that the majority of our deferred tax assets will not be realized.

Liquidity and Capital Resources

Liquidity is the measurement of our ability to meet working capital needs and to fund capital expenditures. Our sources of cash include operations and stock option exercises. We actively manage our cash usage and investment of liquid cash to ensure the maintenance of sufficient funds to meet our daily needs. The majority of our cash and investments are held in U.S. banks, and our foreign subsidiaries maintain a limited amount of cash in their local banks to cover their short-term operating expenses.

(In Thousands)	September 30, 2015	December 31, 2014
Cash and cash equivalents	\$ 16,963	\$ 26,487
Working capital	\$ 19,513	\$ 19,272

(In Thousands)	Nine months ended September 30,	
	2015	2014
Net cash used in operating activities	\$(7,482)	\$(4,653)
Net cash provided by (used in) investing activities	(465)	4,680
Net cash used in financing activities	(1,577)	(635)
Net decrease in cash and cash equivalents	\$(9,524)	\$(608)

We have historically experienced negative cash flows from operations as we continue to invest in key technology development projects and improvements to our biocatalysis technology platform, and expand our business development and collaboration with new customers. Our cash flows from operations will continue to be affected principally by sales and gross margins from biocatalyst product sales and research and development services provided to customers, as well as our headcount costs, primarily in research and development. Our primary source of cash flows from operating activities is cash receipts from our customers for purchases of biocatalyst products and/or biocatalyst research and development services. Our largest uses of cash from operating activities are for employee-related expenditures, rent payments, inventory purchases to support our product sales and non-payroll research and development costs.

We are actively collaborating with new and existing customers in the pharmaceutical and food industries and we believe that we can utilize our current products and services, and develop new products and services, to increase our revenue and gross margins in future periods.

We expect to receive payments totaling \$11.5 million during the fourth quarter of 2015 from the achievement of milestones under our collaborative arrangements with Merck and GSK. We believe that based on our current level of operations, our existing cash and cash equivalents will provide adequate funds for ongoing operations, planned capital expenditures and working capital requirements for at least the next 12 months. However, we may need additional capital if our current plans and assumptions change. Our need for additional capital will depend on many factors, including the financial success of our business, the spending required to develop and commercialize new and existing products, the effect of any acquisitions of other businesses, technologies or facilities that we may make or develop in the future, our spending on new market opportunities, and the potential costs for the filing, prosecution, enforcement and defense of patent claims, if necessary.

If our capital resources are insufficient to meet our capital requirements, and we are unable to enter into or maintain collaborations with partners that are able or willing to fund our development efforts or commercialize any products that we develop or enable, we will have to raise additional funds to continue the development of our technology and products and complete the commercialization of products, if any, resulting from our technologies. If future financings involve the issuance of equity securities, our existing stockholders would suffer dilution. If we raise debt financing, we may be subject to restrictive covenants that limit our ability to conduct our business. We may not be able to raise sufficient additional funds on terms that are favorable to us, if at all. If we fail to raise sufficient funds and fail to generate sufficient revenue to achieve planned gross margins and to control operating costs, our ability to fund our operations, take advantage of strategic opportunities, develop products or technologies, or otherwise respond to competitive pressures could be significantly limited. If this happens, we may be forced to delay or terminate research or development programs or the commercialization of products resulting from our technologies, curtail or cease operations or obtain funds through collaborative and licensing arrangements that may require us to relinquish commercial rights, or grant licenses on terms that are not favorable to us. If adequate funds are not available, we will not be able to successfully execute our business plan or continue our business.

Cash Flows from Operating Activities

Cash used in operating activities was \$7.5 million for the nine months ended September 30, 2015, which resulted from a net loss of \$5.5 million for the nine months ended September 30, 2015 adjusted for non-cash depreciation and amortization of \$4.1 million and stock-based compensation of \$3.8 million, as well as changes in operating assets and liabilities. Such changes included an increase in accounts receivable of \$9.7 million related to the achievement of milestones from collaborative arrangements with Merck and GSK during the third quarter of 2015 and a decrease in accounts payable of \$3.5 million, which were partially offset by an increase in deferred revenues of \$4.0 million.

Cash used in operating activities was \$4.7 million for the nine months ended September 30, 2014, which resulted from a net loss of \$19.4 million for the nine months ended September 30, 2014, adjusted for non-cash depreciation and amortization of \$5.2 million, stock-based compensation of \$3.6 million and impairment and changes in fair values of assets held for sale totaling \$2.7 million. Cash used in operating activities was offset by the receipt of \$6.0 million in up-front fees under the collaborative arrangement with GSK and a \$0.8 million gain on the sale of our Hungarian subsidiary.

Cash Flows from Investing Activities

Cash used in investing activities was \$0.5 million for the nine months ended September 30, 2015 primarily due to the purchase of property and equipment.

Cash provided by investing activities was \$4.7 million for the nine months ended September 30, 2014 mainly resulting from \$3.0 million in proceeds from the maturity of our investments in marketable securities and \$1.5 million in proceeds from the sale of our Hungarian subsidiary.

Cash Flows from Financing Activities

Cash used in financing activities was \$1.6 million and \$0.6 million for the nine months ended September 30, 2015 and 2014, respectively. Cash used in financing activities consisted of the payment of taxes related to the net share settlement of equity awards, partially offset by the proceeds from exercises of employee stock options.

Off-Balance Sheet Arrangements

We did not have any off-balance sheet arrangements as of September 30, 2015.

Contractual Obligations

Our contractual obligations principally arise from operating leases primarily related to our leased facilities in Redwood City, California. There have been no significant changes in our payments due under contractual obligations, compared to those disclosed in our Annual Report on Form 10-K for the year ended December 31, 2014.

Critical Accounting Policies and Estimates

The preparation of financial statements in conformity with generally accepted accounting principles requires management to make judgments, estimates and assumptions in the preparation of our consolidated financial statements and accompanying notes. Actual results could differ from those estimates. There have been no significant changes to our critical accounting policies as discussed in our Annual Report on Form 10-K for the year ended December 31, 2014.

ITEM 3. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK

Market Risk Management

Our cash flows and earnings are subject to fluctuations due to changes in foreign currency exchange rates, interest rates and other factors. There were no significant changes in our market risk exposures for the three and nine months ended September 30, 2015. This is discussed in further detail in our Annual Report on Form 10-K for the year ended December 31, 2014 filed with the SEC on March 6, 2015.

Equity Price Risk

As described in Note 5, "Marketable Securities" and Note 6, "Fair Value Measurements" to the condensed consolidated financial statements, we have an investment in common shares of CO2 Solutions, whose shares are publicly traded in Canada on the TSX Venture Exchange. As of September 30, 2015, the fair value of our investment in CO2 Solutions' common stock was \$1.2 million and our carrying cost for the investment was \$0.6 million.

This investment is exposed to fluctuations in both the market price of CO2 Solutions' common shares and changes in the exchange rate between the U.S. dollar and the Canadian dollar. The effect of a 10% adverse change in the market price of CO2 Solution's common shares as of September 30, 2015 would have been an unrealized loss of approximately \$0.1 million, recognized as a component of our condensed consolidated statements of comprehensive income (loss). The effect of a 10% adverse change in the exchange rate between the U.S. dollar and the Canadian dollar as of September 30, 2015 would have been an unrealized loss of approximately \$0.1 million, recognized as a component of our condensed consolidated statements of comprehensive income (loss).

ITEM 4. CONTROLS AND PROCEDURES

Evaluation of Disclosure Controls and Procedures

We maintain disclosure controls and procedures and internal controls that are designed to provide reasonable assurance that information required to be disclosed in our Exchange Act reports is recorded, processed, summarized and reported within the time periods specified in the SEC's rules and forms and that such information is accumulated and communicated to our management, including our principal executive officer and our principal financial and accounting officer, as appropriate, to allow timely decisions regarding required disclosure.

Our management, including our principal executive officer and our principal financial and accounting officer, evaluated the effectiveness of our disclosure controls and procedures as required by Rules 13a-15(e) and 15d-15(e) of the Securities Exchange Act of 1934, as amended. Based on this review, our principal executive officer and our principal financial and accounting officer concluded that these disclosure controls and procedures were effective as of September 30, 2015 at the reasonable assurance level.

Changes in Internal Control over Financial Reporting

There were no changes in our internal control over financial reporting identified in connection with the evaluation required by paragraph (d) of Exchange Act Rules 13a-15 or 15d-15 that occurred during our last fiscal quarter that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

Inherent Limitations on Effectiveness of Controls

In designing and evaluating the disclosure controls and procedures, management recognizes that any controls and procedures, even if determined effective and no matter how well designed and operated, can provide only reasonable assurance of achieving the desired control objectives to prevent or detect misstatements. In addition, the design of disclosure controls and procedures must reflect the fact that there are resource constraints and that management is required to apply judgment in evaluating the benefits of possible controls and procedures relative to their costs. Also, projections of any evaluation of effectiveness to future periods are subject to the risk that controls may become inadequate because of changes in conditions, or that the degree of compliance with the policies or procedures may deteriorate.

PART II. OTHER INFORMATION

ITEM 1. LEGAL PROCEEDINGS

We are not currently a party to any material litigation or other material legal proceedings.

ITEM 1A. RISK FACTORS

We have included in Part I, Item 1A of our Annual Report on Form 10-K for the year ended December 31, 2014, a description of certain risks and uncertainties that could affect our business, future performance or financial condition (the "Risk Factors"). Except as set forth below, there are no material changes from the disclosure provided in the Form 10-K for the year ended December 31, 2014 with respect to the Risk Factors. Investors should consider the Risk Factors, as updated below, prior to making an investment decision with respect to our stock.

Our results of operations may be adversely affected by the results of regulatory tax examinations.

We are subject to value added tax, customs tax, sales and use tax, withholding tax, payroll tax, income tax and other taxes in connection with the operation of our business. The regulators from the various jurisdictions in which we operate periodically perform audits, and we are regularly subject to, and are currently undergoing, audits and assessments by tax authorities in the United States and foreign jurisdictions for prior tax years. Although we believe our tax estimates are reasonable, and we intend to defend our positions if necessary, the final outcome of tax audits and related proceedings is inherently uncertain and could be materially different than that reflected in our historical income tax provisions and accruals. Moreover, we could be subject to assessments of substantial additional taxes and/or fines or penalties relating to ongoing or future audits. The adverse resolution of any audits or related proceedings could have an adverse effect on our financial position and results of operations.

Compliance with European Union chemical regulations could be costly and adversely affect our business and results of operations.

Some of our products are subject to the European Union regulatory regime known as The Registration, Evaluation and Authorization of Chemicals ("REACH"). REACH mandates that certain chemicals manufactured in, or imported into, the European Union be registered and evaluated for their potential effects on human health and the environment.

Under REACH, we and our contract manufacturers located in the European Union are required to register certain of our products based on the quantity of such product imported into or manufactured in the European Union and on the product's intended end-use. The registration, evaluation and authorization process under REACH can be costly and time consuming. Problems or delays in the registration, evaluation or authorization process under REACH could delay or prevent the manufacture of some of our products in, or the importation of some of our products into, the European Union, which could adversely affect our business and results of operations. In addition, if we or our contract manufacturers fail to comply with REACH, we may be subject to penalties or other enforcement actions, which could have a material adverse effect on our business and results of operations.

ITEM 2. UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS

Not applicable.

ITEM 3. DEFAULTS UPON SENIOR SECURITIES

Not applicable.

ITEM 4. MINE SAFETY DISCLOSURES

Not applicable.

ITEM 5. OTHER INFORMATION

Not applicable.

ITEM 6. EXHIBITS

See the Exhibit Index on the page immediately following the signature page to this Quarterly Report on Form 10-Q for a list of exhibits filed as part of this Quarterly Report, which Exhibit Index is incorporated herein by reference.

38

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

Codexis, Inc.

Date: November 5, 2015

By: /s/ John Nicols
John Nicols
President and Chief Executive Officer
(principal executive officer)

Date: November 5, 2015

By: /s/ Gordon Sangster
Gordon Sangster
Chief Financial Officer
(principal financial and accounting officer)

EXHIBIT INDEX

Listed and indexed below are all Exhibits filed as part of this report.

ITEM 6. Exhibits

- 3.1 Amended and Restated Certificate of Incorporation of Codexis, Inc. filed with the Secretary of the State of the State of Delaware on April 27, 2010 and effective as of April 27, 2010 (incorporated by reference to Exhibit 3.1 to the Company's Quarterly Report on Form 10-Q for the quarter ended March 31, 2010, filed on May 28, 2010).
- 3.2 Certificate of Designations of Series A Junior Participating Preferred Stock of Codexis, Inc., filed with the Secretary of State of the State of Delaware on September 4, 2012 (incorporated by reference to Exhibit 3.1 to the Company's Current Report on Form 8-K, filed on September 4, 2012).
- 3.3 Amended and Restated Bylaws of Codexis, Inc. effective as of April 27, 2010 (incorporated by reference to Exhibit 3.2 to the Company's Quarterly Report on Form 10-Q for the quarter ended March 31, 2010, filed on May 28, 2010).
- 4.1 Form of the Company's Common Stock Certificate (incorporated by reference to Exhibit 4.1 to the Company's Quarterly Report on Form 10-Q for the quarter ended June 30, 2012, filed on August 9, 2012).
- 10.1† Platform Technology Transfer and License Agreement by and between the Company and Merck Sharp & Dohme Corp., dated as of August 3, 2015.
- 31.1 Certification of Principal Executive Officer Required Under Rule 13a-14(a) and 15d-14(a) of the Securities Exchange Act of 1934, as amended.
- 31.2 Certification of Principal Financial Officer Required Under Rule 13a-14(a) and 15d-14(a) of the Securities Exchange Act of 1934, as amended.
- 32.1 Certification of Principal Executive Officer and Principal Financial Officer Required Under Rule 13a-14(b) of the Securities Exchange Act of 1934, as amended, and 18 U.S.C. §1350.
- 101 The following materials from the Company's Quarterly Report on Form 10-Q for the quarter ended September 30, 2015, formatted in Extensible Business Reporting Language (XBRL) includes:
 (i) Condensed Consolidated Balance Sheets at September 30, 2015 and December 31, 2014,
 (ii) Condensed Consolidated Statements of Income for the Three and Nine Months Ended September 30, 2015 and 2014, (iii) Condensed Consolidated Statements of Comprehensive Income (Loss) for the Three and Nine Months Ended September 30, 2015 and 2014, (iv) Condensed Consolidated Statements of Cash Flows for the Nine Months Ended September 30, 2015 and 2014, and (v) Notes to Condensed Consolidated Financial Statements.

† Certain portions have been omitted pursuant to a confidential treatment request. Omitted information has been filed separately with the Securities and Exchange Commission.