

ORAMED PHARMACEUTICALS INC.

Form 10-Q

January 14, 2019

**UNITED STATES**

**SECURITIES AND EXCHANGE COMMISSION**

**WASHINGTON, D.C. 20549**

**FORM 10-Q**

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

**For the quarterly period ended November 30, 2018**

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

**Commission file number: 000-50298**

**ORAMED PHARMACEUTICALS INC.**

(Exact Name of Registrant as Specified in Its Charter)

**Delaware**

(State or Other Jurisdiction of  
Incorporation or Organization)

**98-0376008**

(I.R.S. Employer  
Identification No.)

**142 W. 57th Street**

**New York, New York**

(Address of Principal Executive Offices)

**10019**

(Zip Code)



As of January 11, 2019, there were 17,378,359 shares of the issuer's common stock, \$0.012 par value per share, outstanding.

**ORAMED PHARMACEUTICALS INC.**

**FORM 10-Q**

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As used in this Quarterly Report on Form 10-Q, the terms "we," "us," "our" and the "Company" mean Oramed Pharmaceuticals Inc. and our wholly-owned Israeli subsidiary, Oramed Ltd., unless otherwise indicated. All dollar amounts refer to U.S. Dollars unless otherwise indicated.

On November 30, 2018, the exchange rate between the New Israeli Shekel, or NIS, and the dollar, as quoted by the Bank of Israel, was NIS 3.701 to \$1.00. Unless indicated otherwise by the context, statements in this Quarterly Report on Form 10-Q that provide the dollar equivalent of NIS amounts or provide the NIS equivalent of dollar amounts are based on such exchange rate.

**PART I – FINANCIAL INFORMATION**

**ITEM 1 - FINANCIAL STATEMENTS**

**ORAMED PHARMACEUTICALS INC.**

**CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

AS OF November 30, 2018

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**ORAMED PHARMACEUTICALS INC.****CONDENSED CONSOLIDATED BALANCE SHEETS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

	November 30, 2018	August 31, 2018
Assets		
<b>CURRENT ASSETS:</b>		
Cash and cash equivalents	\$ 3,861	\$4,996
Short-term deposits	19,920	20,875
Marketable securities	5,143	4,592
Prepaid expenses and other current assets	727	574
Total current assets	29,651	31,037
<b>LONG-TERM ASSETS:</b>		
Long-term deposits	11,613	13,542
Marketable securities	2,290	2,785
Amounts funded in respect of employee rights upon retirement	16	16
Property and equipment, net	23	17
Total long-term assets	13,942	16,360
Total assets	\$ 43,593	\$47,397
Liabilities and stockholders' equity		
<b>CURRENT LIABILITIES:</b>		
Accounts payable and accrued expenses	\$ 2,982	\$2,058
Contract liabilities	1,131	2,449
Payable to related parties	51	46
Total current liabilities	4,164	4,553
<b>LONG-TERM LIABILITIES:</b>		
Contract liabilities	10,259	11,388
Employee rights upon retirement	20	20
Provision for uncertain tax position	11	11
Other liabilities	281	313
Total long-term liabilities	10,571	11,732
<b>COMMITMENTS</b> (note 2)		
<b>STOCKHOLDERS' EQUITY:</b>		

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Common stock, \$0.012 par value (30,000,000 authorized shares; 17,378,359 and 17,369,875 shares issued and outstanding as of November 30, 2018 and August 31, 2018, respectively)	207	207
Additional paid-in capital	99,701	99,426
Accumulated other comprehensive income	-	702
Accumulated deficit	(71,050	) (69,223)
Total stockholders' equity	28,858	31,112
Total liabilities and stockholders' equity	\$ 43,593	\$47,397

**The accompanying notes are an integral part of the condensed consolidated financial statements.**



**ORAMED PHARMACEUTICALS INC.****CONDENSED CONSOLIDATED STATEMENTS OF COMPREHENSIVE LOSS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

	Three months ended	
	November 30, 2018	November 30, 2017
REVENUES	\$674	\$ 611
COST OF REVENUES	35	-
RESEARCH AND DEVELOPMENT EXPENSES	4,347	2,327
GENERAL AND ADMINISTRATIVE EXPENSES	932	1,016
OPERATING LOSS	4,640	2,732
FINANCIAL INCOME	286	222
FINANCIAL EXPENSES	8	21
INCOME FROM CHANGES IN FAIR VALUE OF INVESTMENTS	60	-
NET LOSS FOR THE PERIOD	\$4,302	\$ 2,531
UNREALIZED INCOME ON AVAILABLE FOR SALE SECURITIES	-	326
TOTAL OTHER COMPREHENSIVE INCOME	-	326
TOTAL COMPREHENSIVE LOSS FOR THE PERIOD	\$4,302	\$ 2,205
LOSS PER SHARE OF COMMON STOCK:		
BASIC AND DILUTED LOSS PER SHARE OF COMMON STOCK	\$0.25	\$ 0.18
WEIGHTED AVERAGE NUMBER OF SHARES OF COMMON STOCK USED IN COMPUTING BASIC AND DILUTED LOSS PER SHARE OF COMMON STOCK	17,448,744	14,239,346

**The accompanying notes are an integral part of the condensed consolidated financial statements.**

**ORAMED PHARMACEUTICALS INC.****CONDENSED CONSOLIDATED STATEMENTS OF CHANGES IN STOCKHOLDERS' EQUITY**

U.S. Dollars in thousands

(UNAUDITED)

	Common Stock Shares	\$	Additional paid-in capital	Accumulated other comprehensive income	Accumulated deficit	Total stockholders' equity
BALANCE AS OF AUGUST 31, 2018	17,369	\$ 207	\$ 99,426	\$ 702	\$ (69,223 )	\$ 31,112
INITIAL ADOPTION OF ASC 606					1,773	1,773
INITIAL ADOPTION OF ASU 2016-01				(702 )	702	-
CHANGES DURING THE THREE-MONTH PERIOD ENDED NOVEMBER 30, 2018:						
SHARES ISSUED FOR SERVICES	8	*	36	-	-	36
STOCK-BASED COMPENSATION	-	*	239	-	-	239
NET LOSS	-	-	-	-	(4,302 )	(4,302 )
BALANCE AS OF NOVEMBER 30, 2018	17,377	\$ 207	\$ 99,701	-	\$ (71,050 )	\$ 28,858

	Common Stock Shares	\$	Additional paid-in capital	Accumulated other comprehensive income	Accumulated deficit	Total stockholders' equity
BALANCE AS OF AUGUST 31, 2017	13,668	\$ 163	\$ 75,170	\$ 401	\$ (56,496 )	\$ 19,238
CHANGES DURING THE THREE-MONTH PERIOD ENDED NOVEMBER 30, 2017:						
SHARES ISSUED FOR SERVICES	3	*	24	-	-	24
ISSUANCE OF COMMON STOCK, NET	454	5	4,225	-	-	4,230
EXERCISE OF WARRANTS AND OPTIONS	178	2	928	-	-	930
STOCK-BASED COMPENSATION	5	*	524	-	-	524
NET LOSS	-	-	-	-	(2,531 )	(2,531 )
OTHER COMPREHENSIVE INCOME	-	-	-	326	-	326
BALANCE AS OF NOVEMBER 30, 2017	14,308	\$ 170	\$ 80,871	\$ 727	\$ (59,027 )	\$ 22,741

\* Represents an amount of less than \$1.

**The accompanying notes are an integral part of the condensed consolidated financial statements.**

**ORAMED PHARMACEUTICALS INC.****CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS**

U.S. dollars in thousands

(UNAUDITED)

	Three months ended November 30,	
	2018	2017
<b>CASH FLOWS FROM OPERATING ACTIVITIES:</b>		
Net loss	\$(4,302)	\$(2,531)
Adjustments required to reconcile net loss to net cash used in operating activities:		
Depreciation	2	1
Exchange differences and interest on deposits and held to maturity bonds	(116 )	(71 )
Changes at fair value of investments	(60 )	
Stock-based compensation	239	524
Shares issued for services	36	24
Changes in operating assets and liabilities:		
Prepaid expenses and other current assets	(153 )	(4 )
Accounts payable, accrued expenses and related parties	929	(41 )
Contract liabilities	(674 )	(611 )
Liability for employee rights upon retirement	-	1
Other liabilities	(32 )	(20 )
Total net cash used in operating activities	(4,131)	(2,728)
<b>CASH FLOWS FROM INVESTING ACTIVITIES:</b>		
Purchase of short-term deposits	-	(2,039)
Purchase of long-term deposits	-	(3,540)
Purchase of held to maturity securities	(397 )	(2,879)
Proceeds from sale of short-term deposits	3,000	2,455
Proceeds from maturity of held to maturity securities	400	857
Purchase of property and equipment	(8 )	-
Total net cash provided by (used in) investing activities	2,995	(5,146)
<b>CASH FLOWS FROM FINANCING ACTIVITIES:</b>		
Proceeds from issuance of common stock, net of issuance costs	-	4,230
Proceeds from exercise of warrants and options	-	930
Total net cash provided by financing activities	-	5,160
EFFECT OF EXCHANGE RATE CHANGES ON CASH	1	3
DECREASE IN CASH AND CASH EQUIVALENTS	(1,135)	(2,711)
CASH AND CASH EQUIVALENTS AT BEGINNING OF PERIOD	4,996	3,969
CASH AND CASH EQUIVALENTS AT END OF PERIOD	\$3,861	\$1,258

SUPPLEMENTARY DISCLOSURE ON CASH FLOWS -

Interest received	\$159	\$133
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**The accompanying notes are an integral part of the condensed consolidated financial statements.**

**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 1 - SIGNIFICANT ACCOUNTING POLICIES:**

**a. General:**

**1) Incorporation and operations**

Oramed Pharmaceuticals Inc. (collectively with its subsidiary, the “Company”, unless the context indicates otherwise) was incorporated on April 12, 2002, under the laws of the State of Nevada. From incorporation until March 3, 2006, the Company was an exploration stage company engaged in the acquisition and exploration of mineral properties. On February 17, 2006, the Company entered into an agreement with Hadasit Medical Services and Development Ltd. to acquire the provisional patent related to an orally ingestible insulin capsule to be used for the treatment of individuals with diabetes.

On May 14, 2007, the Company incorporated a wholly-owned subsidiary in Israel, Oramed Ltd. (the “Subsidiary”), which is engaged in research and development.

On March 11, 2011, the Company was reincorporated from the State of Nevada to the State of Delaware.

On November 30, 2015, the Company entered into a Technology License Agreement with Hefei Tianhui Incubator of Technologies Co. Ltd. (“HTIT”) and on December 21, 2015, the parties entered into an Amended and Restated Technology License Agreement that was further amended by the parties on June 3, 2016 and July 24, 2016 (the “License Agreement”). According to the License Agreement, the Company granted HTIT an exclusive commercialization license in the territory of the People’s Republic of China, Macau and Hong Kong (the “Territory”), related to the Company’s oral insulin capsule, ORMD-0801 (the “Product”). Pursuant to the License Agreement, HTIT will conduct, at its own expense, certain pre-commercialization and regulatory activities with respect to the Subsidiary’s technology and ORMD-0801 capsule, and will pay to the Subsidiary (i) royalties of 10% on net sales of the related commercialized products to be sold by HTIT in the Territory (“Royalties”), and (ii) an aggregate of \$37,500,

of which \$3,000 was payable immediately, \$8,000 will be paid subject to the Company entering into certain agreements with certain third parties, and \$26,500 will be paid upon achievement of certain milestones and conditions. In the event that the Company does not meet certain conditions, the Royalties rate may be reduced to a minimum of 8%. Following the final expiration of the Company's patents covering the technology in the Territory in 2033, the Royalties rate may be reduced, under certain circumstances, to 5%.

The royalty payment obligation shall apply during the period of time beginning upon the first commercial sale of the Product in the Territory, and ending upon the later of (i) the expiration of the last-to-expire licensed patents in the Territory; and (ii) 15 years after the first commercial sale of the Product in the Territory (the "Royalty Term").

The License Agreement shall remain in effect until the expiration of the Royalty Term. The License Agreement contains customary termination provisions.

**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 1 - SIGNIFICANT ACCOUNTING POLICIES** (continued):

Among others, the Company's involvement through the product submission date will include consultancy for the pre-commercialization activities in the Territory, as well as advisory services to HTIT on an ongoing basis.

The initial payment of \$3,000 was received in January 2016. Following the achievement of certain milestones, the second and third payments of \$6,500 and \$4,000, respectively, were received in July 2016, the fourth milestone payment of \$4,000 was received in October 2016 and the fifth milestone payment of \$3,000 was received in January 2019. Milestone payments received as of January 2019 totaled \$20,500.

In addition, on November 30, 2015, the Company entered into a Stock Purchase Agreement with HTIT (the "SPA"). According to the SPA, the Company issued 1,155,367 shares of common stock to HTIT for \$12,000. The transaction closed on December 28, 2015.

In July 2015, according to the letter of intent signed between the parties or their affiliates, HTIT's affiliate paid the Subsidiary a non-refundable amount of \$500 as a no-shop fee. The no-shop fee was deferred and the related revenue is recognized over the estimated term of the License Agreement.

For revenue recognition policy see note 1c.

**2) Development and liquidity risks**

The Company is engaged in research and development in the biotechnology field for innovative pharmaceutical solutions, including an orally ingestible insulin capsule to be used for the treatment of individuals with diabetes, and



the use of orally ingestible capsules for delivery of other polypeptides, and has not generated significant revenues from its operations. Based on our current cash resources and commitments, we believe we will be able to maintain our current planned development activities and the corresponding level of expenditures for at least the next 12 months and beyond, although no assurance can be given that we will not need additional funds prior to such time. If there are unexpected increases in our operating expenses, we may need to seek additional financing during the next 12 months. Successful completion of the Company's development programs and its transition to normal operations is dependent upon obtaining necessary regulatory approvals from the U.S. Food and Drug Administration prior to selling its products within the United States, obtaining foreign regulatory approvals to sell its products internationally, or entering into licensing agreements with third parties. There can be no assurance that the Company will receive regulatory approval of any of its product candidates, and a substantial amount of time may pass before the Company achieves a level of revenues adequate to support its operations, if at all. The Company also expects to incur substantial expenditures in connection with the regulatory approval process for each of its product candidates during their respective developmental periods. Obtaining marketing approval will be directly dependent on the Company's ability to implement the necessary regulatory steps required to obtain marketing approval in the United States and in other countries. The Company cannot predict the outcome of these activities.

**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 1 - SIGNIFICANT ACCOUNTING POLICIES** (continued):

**b. Loss per common share**

Basic and diluted net loss per common share are computed by dividing the net loss for the period by the weighted average number of shares of common stock outstanding for each period. Outstanding stock options, warrants and restricted stock units (“RSUs”) have been excluded from the calculation of the diluted loss per share because all such securities are anti-dilutive for all periods presented. The weighted average number of common stock options, warrants and RSUs excluded from the calculation of diluted net loss was 4,352,798 and 1,424,029 for the three-month periods ended November 30, 2018 and 2017, respectively.

**c. Revenue recognition**

The License Agreement and the SPA were considered a single arrangement with multiple deliverables. The Company allocated the total consideration of \$49,500 between the License Agreement and the SPA according to their fair value, as follows: \$10,617 was allocated to the issuance of common stock (less issuance expenses of \$23), based on the quoted price of the Company’s shares on the closing date of the SPA on December 28, 2015, and \$38,883 was allocated to the License Agreement.

Under Accounting Standards Codification (“ASC”) 605 (which was the authoritative revenue recognition guidance applied for all periods prior to September 1, 2018) given the Company’s continuing involvement through the expected product submission in June 2023, amounts received relating to the License Agreement were recognized over the period from which the Company was entitled to the respective payment, and the expected product submission date using a time-based model approach over the periods that the fees were earned.

On September 1, 2018, the Company adopted Accounting Standards Update (“ASU”) 2014-09 “Revenue from Contracts with Customers (Topic 606)” (“ASC 606”), using the modified retrospective method of adoption. Under this

method, the Company applied ASC 606 to the License Agreement at the adoption date and was required to make an adjustment to the September 1, 2018 opening accumulated deficit balance and all prior periods continue to be presented under ASC 605. The most significant impact from adopting ASC 606 was the impact of the timing of recognition of revenue associated with the milestone payment. Under ASC 605 (which was the authoritative revenue recognition guidance applied for all periods prior to September 1, 2018) given the Company's continuing involvement through the expected product submission in June 2023, amounts received relating to the License Agreement were recognized over the period from which the Company was entitled to the respective payment, and the expected product submission date using a time-based model approach over the periods that the fees were earned. However, under ASC 606, the Company is required to recognize the total transaction price (which includes consideration related to milestones once the criteria for recognition have been satisfied) using the input method over the period the performance obligation is fulfilled. Accordingly, once the consideration associated with a milestone is included in the transaction price, incremental revenue is recognized immediately based on the period of time that has elapsed towards complete satisfaction of the performance obligation. This method results in the recognition of revenue earlier than under ASC 605 and the resulting impact was recorded as a reduction of the opening balance of accumulated deficit at September 1, 2018 as further described below.

Under ASC 606, the Company identified a single performance obligation in the agreement and determined that the license and services are not distinct as the license and services are highly dependent on each other. In other words, HTIT cannot benefit from the license without the related services, and vice versa.

Since the customer benefits from the services as the entity performs, revenue is recognized over time through the expected product submission date in June 2023, using the input method. The Company used the input method to measure the process for the purpose of recognizing revenue, which approximates the straight line attribution. The Company used significant judgment when it determined the product submission date.

Under ASC 606, the consideration that the Company would be entitled to upon the achievement of contractual milestones, which are contingent upon the occurrence of future events, are a form of variable consideration. When assessing the portion, if any, of such milestones-related consideration to be included in the transaction price, the Company first assesses the most likely outcome for each milestone and excludes the consideration related to milestones of which the occurrence is not considered the most likely outcome.

The Company then evaluates if any of the variable consideration determined in the first step is constrained by including in the transaction price variable consideration to the extent that it is probable that a significant reversal in the amount of cumulative revenue recognized will not occur when the uncertainty associated with the variable consideration is subsequently resolved. The Company used significant judgment when it determined the first step of variable consideration.

**ORAMED PHARMACEUTICALS INC.**

NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 1 - SIGNIFICANT ACCOUNTING POLICIES** (continued):

The potential future royalty consideration is also considered a form of variable consideration under ASC 606 as it is based on a percentage of potential future sales of the Company's products. However, the Company applies the sales-based royalty exception and accordingly will recognize the sales-based royalty amounts at the earlier of the time (a) when the related sale has occurred and (b) the company has fulfilled the related performance obligation. To date, the Company has not recognized any royalty-related revenue.

As of the adoption date, the Company adjusted its accumulated deficit by \$1,773 against contract liabilities due to the effect of variable consideration.

Amounts that were allocated to the License Agreement as of November 30, 2018 aggregated \$20,557, \$19,383 of which was received through the balance sheet date. Through November 30, 2018, the Company has recognized revenue associated with this agreement in the aggregate amount of \$7,993 (of which \$674 was recognized in the quarter ended November 30, 2018 and \$1,773 was recognized as an increase to the September 1, 2018 opening balance of stockholders' equity associated with the impact of the adoption of ASC 606 under the modified retrospective method of adoption), and deferred the remaining amount of \$11,390 which is presented as a contract liability on the condensed consolidated balance sheet.

In accordance with ASC 606, the disclosure of the impact of adoption to the Company's consolidated balance sheet as of August 31, 2018 was as follows:

<b>As reported  August 31,</b>	Updated September 1, 2018	Effect of Change
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**2018**

Contract liabilities (short term)	2,449	1,230	(1,219 )
Contract liabilities (long term)	11,388	10,834	(554 )
Accumulated deficit	69,223	67,450	(1,773 )

The impact of adoption of ASC 606 on the condensed consolidated balance sheet as of November 30, 2018 and on the condensed consolidated statement of operations for the three months ended November 30, 2018 was as follows:

	<b>As reported</b>	Balances without Adoption of ASC 606	Effect of Change
	<b>November 30, 2018</b>		
Revenues	\$ 674	\$ 611	\$63
Cost of revenues	35	-	35
Contract liabilities (short term)	1,131	2,449	(1,318 )
Contract liabilities (long term)	10,259	10,777	(518 )
Accumulated deficit	71,050	72,851	(1,801 )

**d. Financial instruments**

In January 2016, the Financial Accounting Standards Board (“FASB”) issued guidance which updates certain aspects of recognition, measurement, presentation and disclosure of financial assets and financial liabilities (“ASU 2016-01”). The guidance requires entities to recognize changes in fair value in net income rather than in accumulated other comprehensive income. The Company adopted the provisions of this update in the first quarter of fiscal year 2019. Following the adoption, as of September 1, 2018, the Company classified the available for sale securities (investments in equity securities of D.N.A Biomedical Solutions Ltd. (“D.N.A”) and Entera Bio Ltd. (“Entera”)) to financial assets measured in fair value through profit or loss. The Company adopted the standard using the modified retrospective method and, accordingly, reclassified the cumulative unrealized gain from accumulated other comprehensive income to a reduction of its accumulated deficit in an amount of \$702.

**e. Condensed Consolidated Financial Statements Preparation**

The condensed consolidated financial statements included herein have been prepared in accordance with United States generally accepted accounting principles (“U.S. GAAP”) and, except as described in note 1f, on the same basis as the audited consolidated financial statements included in the Company’s Annual Report on Form 10-K for the fiscal year ended August 31, 2018 (the “2018 Form 10-K”). These condensed consolidated financial statements reflect all adjustments that are of a normal recurring nature and that are considered necessary for a fair statement of the results of the periods presented. Certain information and disclosures normally included in annual consolidated financial statements have been omitted in this interim period report pursuant to the rules and regulations of the Securities and Exchange Commission. Because the condensed consolidated interim financial statements do not include all of the

information and disclosures required by U.S. GAAP for annual financial statements, they should be read in conjunction with the audited consolidated financial statements and notes included in the 2018 Form 10-K. The results for interim periods are not necessarily indicative of a full fiscal year's results.

**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 1 - SIGNIFICANT ACCOUNTING POLICIES** (continued):

**f. Newly issued and recently adopted Accounting Pronouncements**

In May 2014, the FASB issued ASC 606 which supersedes existing revenue recognition guidance, including industry-specific guidance. Under the new standard, a good or service is transferred to the customer when (or as) the customer obtains control of the good or service, which differs from the risk and rewards approach under current guidance. The guidance provides a five-step analysis of transactions to determine when and how revenue is recognized. Other major provisions include capitalization of certain contract costs, consideration of the time value of money in the transaction price, and allowing estimates of variable consideration to be recognized before contingencies are resolved in certain circumstances. The guidance also requires enhanced disclosures regarding the nature, amount, timing and uncertainty of revenue and cash flows arising from an entity's contracts with customers. The guidance is effective in annual reporting periods beginning after December 15, 2017, including interim reporting periods within that reporting period. The Company has implemented the guidance for its annual period ending on August 31, 2019 and interim periods within such annual period. The Company adopted the standard using the modified retrospective method. See additional information regarding the adoption in note 1c.

In January 2016, the FASB issued guidance on recognition and measurement of financial assets and financial liabilities (ASU 2016-01) that supersedes most current guidance. Changes to the U.S. GAAP model primarily affect the accounting for equity investments, financial liabilities under the fair value option and the presentation and disclosure requirements for financial instruments. In addition, the FASB clarified guidance related to the valuation allowance assessment when recognizing deferred tax assets resulting from unrealized losses on available-for-sale debt securities. The accounting for other financial instruments, such as loans, investments in debt securities, and financial liabilities, is largely unchanged. The classification and measurement guidance under ASU 2016-01 became effective as of September 1, 2018. See additional information regarding the adoption in note 1d.

**NOTE 2 - COMMITMENTS:**

In March 2011, the Subsidiary sold shares of its investee company, Entera, to D.N.A, retaining 117,000 ordinary **a.** shares (after giving effect to a stock split by Entera in July 2018). In consideration for the shares sold to D.N.A, the Company received, among other payments, ordinary shares of D.N.A (see also note 4).



**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 2 - COMMITMENTS** (continued):

As part of this agreement, the Subsidiary entered into a patent transfer agreement (the "Patent Transfer Agreement") according to which the Subsidiary assigned to Entera all of its right, title and interest in and to a certain patent application related to the oral administration of proteins that it has licensed to Entera since August 2010. Under this agreement, the Subsidiary is entitled to receive from Entera royalties of 3% of Entera's net revenues (as defined in the agreement) and a license back of that patent application for use in respect of diabetes and influenza. As of November 30, 2018, Entera had not yet realized any revenues and had not paid any royalties to the Subsidiary. On December 11, 2018, Entera announced that it had entered into a research collaboration and license agreement (the "Amgen License") with Amgen related to research of inflammatory disease and other serious illnesses. As reported by Entera, under the terms of the Amgen License, Entera will receive a modest initial technology access fee from Amgen and will be responsible for preclinical development at Amgen's expense. Entera will be eligible to receive up to \$270,000 in aggregate payments, as well as tiered royalties up to mid-single digits, upon achievement of various clinical and commercial milestones if Amgen decides to move all of these programs forward. Amgen is responsible for clinical development, manufacturing and commercialization of any of the resulting programs. To the extent the Amgen License results in net revenues as defined in the Patent Transfer Agreement, the Subsidiary will be entitled to the aforementioned royalties.

In addition, as part of a consulting agreement with a third party, dated February 15, 2011, the Subsidiary is obliged to pay this third party royalties of 8% of the net royalties received in respect of the patent that was sold to Entera in March 2011.

- b.** On January 3, 2017, the Subsidiary entered into a lease agreement for its office facilities in Israel. The lease agreement is for a period of 60 months commencing October 1, 2016.

The annual lease payment was New Israeli Shekel ("NIS") 119,000 (\$32) from October 2016 through September 2018 and NIS 132,000 (\$36) from October 2018 through September 2021, and is linked to the increase in the Israeli consumer price index ("CPI") (as of November 30, 2018, the future lease payments will be \$101 until the expiration of the lease agreement, based on the exchange rate as of November 30, 2018).

As security for its obligation under this lease agreement, the Company provided a bank guarantee in an amount equal to three monthly lease payments.

c. On March 3, 2016, the Subsidiary entered into an agreement with a vendor for process development and production of its capsules and on November 24, 2016, April 3, 2017 and July 10, 2017 the Subsidiary entered into amendments to such agreement in an amount of up to Swiss Franc (“CHF”) 1,000,000 (\$1,003), CHF 665,000 (\$675) of which was recognized in research and development expenses through November 30, 2018.

d. On May 11, 2016, the Subsidiary entered into a Master Service Agreement with a vendor to retain its services for a pre-clinical toxicology trial for an oral GLP-1 analog capsule for type 2 diabetes patients. As consideration for its services, the Subsidiary will pay the vendor a total amount of \$1,283 during the term of the engagement and based on achievement of certain milestones, of which \$1,275 was recognized in research and development expenses through November 30, 2018.

e. On June 13, 2016, the Subsidiary entered into a four-year service agreement with a third party and on December 19, 2016, this agreement and all of the third party rights and obligations thereunder were assigned to another third party. This agreement is required by the License Agreement as described in note 1 and will support the Company’s research and development. The Subsidiary is obligated to pay the third party a total amount of up to €2,360,000 (\$2,706), of which €1,700,415 (\$1,956) was recognized in research and development expenses through November 30, 2018.

**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 2 - COMMITMENTS** (continued):

On July 24, 2016, the Subsidiary entered into a General Technical Agreement with a vendor for the scale-up process development and production of one of its oral capsule ingredients in the amount of \$4,300 that will be paid **f.** over the term of the engagement and based on the achievement of certain development milestones, \$4,087 of which was recognized in research and development expenses through November 30, 2018. This agreement is part of the requirements of the License Agreement as described in note 1.

On February 21, 2017, the Subsidiary entered into an agreement with a vendor to retain its services for a pre-clinical toxicology trial for an oral insulin capsule. As consideration for its services, the Subsidiary will pay the **g.** vendor a total of up to \$952 during the term of the engagement and based on achievement of certain milestones, of which \$857 was recognized in research and development expenses through November 30, 2018.

On April 8, 2018, the Company entered into a consulting agreement with a third party advisor for a period of one year, pursuant to which such advisor provides investor relations services and is entitled to receive a monthly cash **h.** fee and 10,000 shares of the Company's common stock issued in four equal quarterly installments commencing August 1, 2018. As of November 30, 2018, the Company had issued to such advisor 5,000 shares. The fair value of the shares at the grant date was \$25, which was recognized in general and administrative expenses.

On June 5, 2017, the Subsidiary entered into a clinical research agreement with a vendor, for the conduct of its clamp clinical trial for an oral insulin capsule for type 1 diabetes patients. As consideration for its services, the **i.** Subsidiary will pay the vendor a total amount of \$958 during the term of the engagement and based on achievement of certain milestones, \$578 of which was recognized in research and development expenses through November 30, 2018.

On December 18, 2017, the Subsidiary entered into an agreement with a vendor for the process development and production of one of its oral capsule ingredients in the amount of \$2,905 that will be paid over the term of the **j.** engagement and based on the achievement of certain development milestones, \$1,037 of which was recognized in research and development expenses through November 30, 2018.

On February 14, 2018, the Subsidiary entered into a Clinical Research Organization Services Agreement with a third party, effective as of November 1, 2017, to retain it as a clinical research organization (“CRO”) for the Subsidiary’s three-month dose-ranging clinical trial for its oral insulin capsule for type 2 diabetes patients. As consideration for its services, the Subsidiary will pay the CRO a total amount of \$7,030 during the term of the engagement and based on achievement of certain milestones, \$4,106 of which was recognized in research and development expenses through November 30, 2018.

On May 21, 2018, the Subsidiary entered into a CRO Services Agreement with a third party to retain it as a CRO for the Subsidiary’s food effect clinical trial for its oral insulin capsule. As consideration for its services, the Subsidiary will pay the CRO a total amount of \$1,166 during the term of the engagement and based on achievement of certain milestones, \$758 of which was recognized in research and development expenses through November 30, 2018.

**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 2 - COMMITMENTS** (continued):

m. On July 4, 2018, the Subsidiary entered into an agreement with a vendor to retain its services for a pre-clinical six months toxicology trial for its oral insulin capsule. As consideration for its services, the Subsidiary will pay the vendor a total of up to \$971 during the term of the engagement and based on achievement of certain milestones, of which \$360 was recognized in research and development expenses through November 30, 2018.

n. On July 15, 2018, the Company entered into a consulting agreement with a third party advisor for a period of one year, pursuant to which such advisor provides investor relations services and is entitled to receive a monthly cash fee and shares of the Company's common stock issued in four quarterly installments in an amount equal to \$25 per quarter, pursuant to and in accordance with the terms of the agreement, commencing July 15, 2018. As of November 30, 2018, the Company had issued to such advisor 9,874 shares and the related expense was recognized in general and administrative expenses. The Company terminated this consulting agreement in December 2018.

**o. Grants from the Israel Innovation Authority ("IIA")**

Under the terms of the Company's funding from the IIA, royalties of 3% are payable on sales of products developed from a project so funded, up to a maximum amount equaling 100%-150% of the grants received (dollar linked) with the addition of interest at an annual rate based on LIBOR.

At the time the grants were received, successful development of the related projects was not assured. The total amount that was received through November 30, 2018 was \$2,194.

The royalty expenses which are related to the funded project were recognized in cost of revenues in the quarter ended November 30, 2018 and in prior periods.



**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 3 - FAIR VALUE:**

The Company measures fair value and discloses fair value measurements for financial assets. Fair value is based on the price that would be received to sell an asset in an orderly transaction between market participants at the measurement date. In order to increase consistency and comparability in fair value measurements, the guidance establishes a fair value hierarchy that prioritizes observable and unobservable inputs used to measure fair value into three broad levels, which are described as follows:

- Level 1: Quoted prices (unadjusted) in active markets that are accessible at the measurement date for assets or liabilities. The fair value hierarchy gives the highest priority to Level 1 inputs.
- Level 2: Observable prices that are based on inputs other than quoted prices included within Level 1 that are observable for the asset or liability, either directly or indirectly.
- Level 3: Unobservable inputs are used when little or no market data is available. The fair value hierarchy gives the lowest priority to Level 3 inputs.

As of November 30, 2018, the assets measured at fair value are comprised of equity securities (Level 1). The fair value of held to maturity bonds as presented in note 4 was based on a Level 1 measurement.

As of November 30, 2018, the carrying amounts of cash equivalents, short-term deposits and accounts payable approximate their fair values due to the short-term maturities of these instruments.

As of November 30, 2018, the carrying amounts of long-term deposits approximate their fair values due to the stated interest rates which approximate market rates.

The amounts funded in respect of employee rights are stated at cash surrender value which approximates its fair value.

There were no Level 3 items for the three-month periods ended November 30, 2018 and 2017.



**ORAMED PHARMACEUTICALS INC.**

NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 4 - MARKETABLE SECURITIES:**

The Company's marketable securities include investments in equity securities of D.N.A and Entera, and in held to maturity bonds.

**a. Composition:**

	<b>November 30,</b>	August
	<b>2018</b>	31, 2018
Short-term:		
D.N.A (see b below)	\$ 737	\$666
Entera (see c below)	621	632
Held to maturity bonds (see d below)	3,785	3,294
	\$ 5,143	\$4,592
Long-term:		
Held to maturity bonds (see d below)	\$ 2,290	\$2,785

**b. D.N.A**

The D.N.A ordinary shares are traded on the Tel Aviv Stock Exchange. The fair value of those securities is measured at the quoted prices of the securities on the measurement date.

As of November 30, 2018, the Company owns approximately 6.9% of D.N.A's outstanding ordinary shares.

The cost of the securities as of November 30, 2018 and August 31, 2018 is \$595.

**c. Entera**

Entera ordinary shares have been traded on The Nasdaq Capital Market since June 28, 2018. The Company measures the investment at fair value from such date, since it has a readily determinable fair value (prior to such date the investment was accounted for as a cost method investment (amounting to \$1)).

**ORAMED PHARMACEUTICALS INC.**

NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 4 - MARKETABLE SECURITIES** (continued):

**d. Held to maturity securities**

The amortized cost and estimated fair value of held-to-maturity securities as of November 30, 2018, are as follows:

	November 30, 2018		
	Amortized cost	Gross unrealized losses	Estimated fair value
Short-term:			
Commercial bonds	\$3,739	\$ (26 )	\$ 3,713
Accrued interest	46	-	46
Long-term	2,290	(19 )	2,271
	\$6,075	\$ (45 )	\$ 6,030

As of November 30, 2018, the contractual maturities of debt securities classified as held-to-maturity are as follows: after one year through two years, \$2,290, and the yield to maturity rates vary between 1.65% to 3.20%.

The amortized cost and estimated fair value of held-to-maturity securities as of August 31, 2018, are as follows:

	August 31, 2018		
	Amortized cost	Gross unrealized losses	Estimated fair value
Short-term:			

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Commercial bonds	\$3,259	\$ (17	)	\$ 3,242
Accrued interest	35	-		35
Long-term	2,785	(17	)	2,768
	\$6,079	\$ (34	)	\$ 6,045

As of August 31, 2018, the contractual maturities of debt securities classified as held-to-maturity are as follows: after one year through two years, \$2,785 and the yield to maturity rates vary between 1.45% to 3.13%.

Held to maturity securities which will mature during the 12 months from the balance sheet date are included in short-term marketable securities. Held to maturity securities with maturity dates of more than one year are considered long-term marketable securities.

**ORAMED PHARMACEUTICALS INC.**

**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**

U.S. Dollars in thousands (except share and per share data)

(UNAUDITED)

**NOTE 5 - STOCKHOLDERS' EQUITY:**

On April 2, 2015, the Company entered into an At The Market Issuance Sales Agreement (the "Sales Agreement") with B. Riley FBR, Inc., as successor to FBR Capital Markets & Co. ("FBR"), as amended, pursuant to which the Company may, from time to time and at its option, issue and sell shares of its common stock having an aggregate offering price of up to \$25,000 through FBR as its sales agent, subject to certain terms and conditions. Any shares sold will be sold pursuant to the Company's effective shelf registration statement on Form S-3 including a prospectus dated February 2, 2017, as supplemented by a prospectus supplement dated April 5, 2017. The Company will pay FBR a commission of 3.0% of the gross proceeds of the sale of any shares sold through FBR. Through November 30, 2018, 576,834 shares were sold under the Sales Agreement for aggregate net proceeds of \$5,198. During the three months ended November 30, 2018, the Company did not issue shares under the Sales Agreement.

**NOTE 6 - RELATED PARTIES - TRANSACTIONS:**

On July 1, 2008, the Subsidiary entered into two consulting agreements with KNRY Ltd. ("KNRY"), an Israeli company owned by the Chief Scientific Officer (the "CSO"), whereby the Chief Executive Officer (the "CEO") and the CSO, through KNRY, provide services to the Company (the "Consulting Agreements"). The Consulting Agreements are both terminable by either party upon 140 days prior written notice. The Consulting Agreements, as amended, provide that KNRY will be reimbursed for reasonable expenses incurred in connection with performance of the Consulting Agreements and that the monthly consulting fee paid to the CEO and the CSO is NIS 127,570 (\$34) and NIS 80,454 (\$22), respectively.

In addition to the Consulting Agreements, based on a relocation cost analysis prepared by consulting company ORI - Organizational Resources International Ltd., the Company pays for certain direct costs, related taxes and expenses incurred in connection with the relocation of the CEO to New York. During the three months ended November 30, 2018, such relocation expenses totaled \$131 compared to \$88 for the three months ended November 30, 2017.



## 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 condensed consolidated financial statements and the related notes included elsewhere herein and in our consolidated financial statements, accompanying notes and "Management's Discussion and Analysis of Financial Condition and Results of Operations" contained in our Annual Report (as defined below).*

### Forward-Looking Statements

The statements contained in this Quarterly Report on Form 10-Q that are not historical facts are "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995 and other federal securities laws. Words such as "expects," "anticipates," "intends," "plans," "planned expenditures," "believes," "seeks," "estimates" and expressions or variations of such words are intended to identify forward-looking statements, but are not deemed to represent an all-inclusive means of identifying forward-looking statements as denoted in this Quarterly Report on Form 10-Q. Additionally, statements concerning future matters are forward-looking statements. We remind readers that forward-looking statements are merely predictions and therefore inherently subject to uncertainties and other factors and involve known and unknown risks that could cause the actual results, performance, levels of activity, or our achievements, or industry results, to be materially different from any future results, performance, levels of activity, or our achievements, or industry results, expressed or implied by such forward-looking statements. Such forward-looking statements include, among other statements, statements regarding the following:

the expected development and potential benefits from our products in treating diabetes;

the prospects of entering into additional license agreements, or other partnerships or forms of cooperation with other companies or medical institutions;

future milestones, conditions and royalties under the license agreement with Hefei Tianhui Incubator of Technologies Co., Ltd., or HTIT;

our research and development plans, including pre-clinical and clinical trials plans and the timing of enrollment, obtaining results and conclusion of trials, including without limitation, our expectation that we will initiate two six-month Phase III clinical trials if our Phase IIb three-month dose-ranging clinical trial is successful, and our expectation to file a New Drug Application thereafter;

our belief that our technology has the potential to deliver medications and vaccines orally that today can only be delivered via injection;

the competitive ability of our technology based product efficacy, safety, patient convenience, reliability, value and patent position;

the potential market demand for our products;

our expectation that in the upcoming year our research and development expenses, net, will continue to be our major expenditure;

our expectations regarding our short- and long-term capital requirements;

our outlook for the coming months and future periods, including but not limited to our expectations regarding future revenue and expenses; and

information with respect to any other plans and strategies for our business.



Although forward-looking statements in this Quarterly Report on Form 10-Q reflect the good faith judgment of our management, such statements can only be based on facts and factors currently known by us. Consequently, forward-looking statements are inherently subject to risks and uncertainties and actual results and outcomes may differ materially from the results and outcomes discussed in or anticipated by the forward-looking statements. Factors that could cause or contribute to such differences in results and outcomes include, without limitation, those specifically addressed under the heading “Item 1A. Risk Factors” in our Annual Report on Form 10-K for the fiscal year ended August 31, 2018, or our Annual Report, as filed with the Securities and Exchange Commission, or the SEC, on November 28, 2018, as well as those discussed elsewhere in our Annual Report and this Quarterly Report on Form 10-Q and expressed from time to time in our other filings with the SEC. In addition, historic results of scientific research, clinical and preclinical trials do not guarantee that the conclusions of future research or trials would not suggest different conclusions. Also, historic results referred to in this Quarterly Report on Form 10-Q could be interpreted differently in light of additional research, clinical and preclinical trials results. Readers are urged not to place undue reliance on these forward-looking statements, which speak only as of the date of this Quarterly Report on Form 10-Q. Except as required by law, we undertake no obligation to revise or update any forward-looking statements in order to reflect any event or circumstance that may arise after the date of this Quarterly Report on Form 10-Q. Readers are urged to carefully review and consider the various disclosures made throughout the entirety of this Quarterly Report on Form 10-Q which attempt to advise interested parties of the risks and factors that may affect our business, financial condition, results of operations and prospects.

## **Overview of Operations**

We are a pharmaceutical company currently engaged in the research and development of innovative pharmaceutical solutions, including an oral insulin capsule to be used for the treatment of individuals with diabetes, and the use of orally ingestible capsules or pills for delivery of other polypeptides.

### ***Recent business developments***

#### *Product Candidates*

##### ***Oral Insulin***

In April 2018, we initiated a three-month dose-ranging Phase IIb clinical trial of our proprietary flagship product, an orally ingestible insulin capsule, or ORMD-0801. This placebo controlled, randomized, 90 day treatment clinical trial is being conducted on approximately 285 type 2 diabetic patients in multiple centers throughout the United States pursuant to an Investigational New Drug application, or IND, with the U.S. Food and Drug Administration, or FDA.

The primary endpoints of the trial are to assess the safety and evaluate the effect of ORMD-0801 on HbA1c levels over a 90 day treatment period. Secondary endpoints of the trial include measurements of fasting plasma glucose, or FPG, post-prandial glucose, or PPG levels, during a mixed-meal tolerance test, or MMTT, and weight.

We had a call with the FDA in August 2017 regarding ORMD-0801 after the completion of a Phase IIb clinical trial on 180 diabetic patients, which indicated a statistically significant blood glucose lowering effect of ORMD-0801 versus placebo across several endpoints. During the call, the FDA advised that the regulatory pathway for the submission of ORMD-0801 would be a Biologics License Application, or BLA. The BLA pathway would grant us 12 years of marketing exclusivity for ORMD-0801, from the approval date, and an additional six months of exclusivity may be granted to us if the product also receives approval for use in pediatric patients. The FDA confirmed that the approach to nonclinical toxicology, chemistry manufacturing controls and qualification of excipients would be driven by their published guidance documents.

In June 2018, we initiated a glucose clamp study which will quantify insulin absorption in type 1 diabetic patients treated with ORMD-0801. The glucose clamp is a method for quantifying insulin absorption in order to measure a patient's insulin sensitivity and how well a patient metabolizes glucose. This exploratory, randomized, double-blind glucose clamp study is evaluating exposure-response profiles of type 1 diabetic patients treated with ORMD-0801. Six patients with HbA1c levels of 10% or below, aged 18-50, are enrolled in the study.

In June 2018, we also initiated a food effect trial in the United States for ORMD-0801. This single-blind, five period, randomized, placebo-controlled crossover trial is evaluating the pharmacokinetics and pharmacodynamics of ORMD-0801 taken at different times in relation to meals in healthy volunteers and patients with type 1 diabetes. Up to 48 patients will be enrolled, including up to 24 healthy volunteers and 24 patients with type 1 diabetes.

In September 2018, we initiated a six month toxicology study of our oral insulin capsule, following the FDA's request.

Should our Phase IIb three-month dose-ranging clinical trial successfully meet its primary or secondary endpoints, we anticipate initiating two six-month Phase III clinical trials on both type 1 and type 2 diabetic patients, following which we expect to file a NDA with potential FDA approval by the second half of calendar year 2023.

### ***Oral GLP-1 Analog***

In addition to our flagship product, the ORMD-0801 insulin capsule, we are using our technology for an orally ingestible GLP-1/exenatide capsule, or ORMD-0901. In September 2018, the FDA cleared our IND application for human trials of ORMD-0901. We initiated in the first quarter of calendar year 2019 a Phase I pharmacokinetic trial which will evaluate the safety and the pharmacokinetics of ORMD-0901 compared to placebo. This study is being conducted pursuant to the IND and will be followed by a Phase II trial on type 2 diabetic patients which will be conducted in the United States under an IND.

### ***Other products***

In April 2017, Israel's Ministry of Health approved our commencement of a proof of concept single dose study for our oral leptin drug candidate to evaluate its pharmacokinetic and pharmacodynamics (glucagon reduction) in 10 type 1 adult diabetic patients. The study is projected to be initiated in calendar year 2019 and be completed during calendar year 2019.

In October 2018, we initiated an exploratory clinical study of ORMD-0801 in patients with nonalcoholic steatohepatitis, or NASH. The three-month treatment study, which was approved by Israel's Ministry of Health, will assess the effectiveness of ORMD-0801 in reducing liver fat content, inflammation and fibrosis in patients with NASH. We expect to complete the study during calendar year 2019.

The table below gives an overview of our primary product pipeline (calendar quarters):

	<b>Phase I</b>	<b>Phase II</b>	<b>Phase III</b>	<b>Timeline</b>
<b>ORMD-0801</b>				Q2 '18: Phase IIb 90-day multi-center study initiated (projected completion Q4 '19) Q3 '20: Phase III study projected initiation (projected completion Q3 '22)
<b>oral insulin</b>				Q2 '18: Clamp study initiated (projected completion Q1 '19) Q2 '18: Food effect study initiated (projected completion Q2 '19) Q3 '20: Phase III projected initiation (projected completion Q3 '22)
<b>ORMD-0901</b>				Q1 '19: Pharmacokinetics clinical study initiated (projected completion Q1 '19)
<b>oral GLP-1</b>				Q4 '19: Phase II projected initiation (projected completion Q1 '21)

### *Out-Licensed Technology*

On November 30, 2015, we, our Israeli subsidiary and HTIT entered into a Technology License Agreement, and on December 21, 2015 these parties entered into an Amended and Restated Technology License Agreement that was further amended by the parties on June 3, 2016 and July 24, 2016, or the License Agreement. According to the License Agreement, we granted HTIT an exclusive commercialization license in the territory of the People's Republic of China, Macau and Hong Kong, or the Territory, related to our oral insulin capsule, ORMD-0801, or the Product. Pursuant to the License Agreement, HTIT will conduct, at its own expense, certain pre-commercialization and regulatory activities with respect to our subsidiary's technology and ORMD-0801 capsule, and will pay (i) royalties of 10% on net sales of the related commercialized products to be sold by HTIT in the Territory, or Royalties, and (ii) an aggregate of \$37.5 million, of which \$3 million was payable immediately, \$8 million was paid subject to our entry into certain agreements with certain third parties, and \$26.5 million will be payable upon achievement of certain milestones and conditions. In the event that we will not meet certain conditions, the Royalties rate may be reduced to a minimum of 8%. Following the final expiration of our patents covering the technology in the Territory in 2033, the Royalties rate may be reduced, under certain circumstances, to 5%. The royalty payment obligation shall apply during the period of time beginning upon the first commercial sale of the Product in the Territory, and ending upon the later of (i) the expiration of the last-to-expire licensed patents in the Territory; and (ii) 15 years after the first commercial sale of the Product in the Territory, or the Royalty Term. The License Agreement shall remain in effect until the expiration of the Royalty Term. The License Agreement contains customary termination provisions. Through November 30, 2018, we received aggregate milestone payments of \$17.5 million and an additional milestone payment of \$3 million was received in January 2019.

On November 30, 2015, we also entered into a separate Securities Purchase Agreement with HTIT, or the SPA, pursuant to which, in December 2015, we issued to HTIT 1,155,367 shares of our common stock for total consideration of \$12 million. In connection with the License Agreement and the SPA, we received a non-refundable payment of \$500,000 as a no-shop fee.

## **Results of Operations**

### *Comparison of three month periods ended November 30, 2018 and 2017*

The following table summarizes certain statements of operations data of the Company for the three month periods ended November 30, 2018 and 2017 (in thousands of dollars except share and per share data):

#### **Three months ended**

	<b>November 30,</b>	
	<b>2018</b>	<b>2017</b>
Revenues	\$674	\$611
Cost of revenues	35	-
Research and development expenses	4,347	2,327
General and administrative expenses	932	1,016
Financial income, net	338	201
Net loss for the period	\$4,302	\$2,531
Loss per common share - basic and diluted	\$0.25	\$0.18
Weighted average common shares outstanding	17,448,744	14,239,346

### ***Revenues***

Revenues consist of proceeds related to the License Agreement that are recognized on a cumulative basis when it is probable that a significant reversal in the amount of cumulative revenue recognized will not occur, through the expected product submission date of June 2023 using the input method.

Revenues for the three month period ended November 30, 2018 increased by 10% to \$674,000, from \$611,000 for the three month period ended November 30, 2017. The increase is primarily attributable to the effect of recognizing revenue of the variable consideration on a cumulative basis following the adoption of Accounting Standards Update 2014-09, "Revenue from Contracts with Customers (Topic 606)", or ASC 606.

***Cost of revenues***

Cost of revenues consists of royalties related to the License Agreement that will be paid over the term of the License Agreement in accordance with revenue recognition accounting and the Law for the Encouragement of Industrial Research, Development and Technological Innovation, 1984, as amended, including any regulations or tracks promulgated thereunder.

Cost of revenues for the three month period ended November 30, 2018 increased to \$35,000 compared to no cost of revenues for the three month period ended November 30, 2017. The increase is attributable to the inclusion of additional milestone payments under the License Agreement in the transaction price as part of the implementation of ASC 606.

***Research and development expenses***

Research and development expenses include costs directly attributable to the conduct of research and development programs, including the cost of salaries, employee benefits, costs of materials, supplies, the cost of services provided by outside contractors, including services related to our clinical trials, clinical trial expenses, the full cost of manufacturing drugs for use in research and preclinical development. All costs associated with research and development are expensed as incurred.

Clinical trial costs are a significant component of research and development expenses and include costs associated with third-party contractors. We outsource a substantial portion of our clinical trial activities, utilizing external entities such as contract research organizations, or CROs, independent clinical investigators and other third-party service providers to assist us with the execution of our clinical studies.

Clinical activities which relate principally to clinical sites and other administrative functions to manage our clinical trials are performed primarily by CROs. CROs typically perform most of the start-up activities for our trials, including document preparation, site identification, screening and preparation, pre-study visits, training, and program management.

Clinical trial and pre-clinical trial expenses include regulatory and scientific consultants' compensation and fees, research expenses, purchase of materials, cost of manufacturing of the oral insulin and exenatide capsules, payments for patient recruitment and treatment, as well as salaries and related expenses of research and development staff.

Research and development expenses for the three month period ended November 30, 2018 increased by 87% to \$4,347,000, from \$2,327,000 for the three month period ended November 30, 2017. The increase is primarily due to expenses related to our Phase IIb three-month treatment clinical trial, food effect and clamp clinical trials and is partially offset by a decrease in expenses related to the scale-up process development and production of our oral capsule ingredients. Stock-based compensation costs for the three month period ended November 30, 2018 totaled \$39,000, as compared to \$171,000 during the three month period ended November 30, 2017. The decrease is mainly attributable to the progress in amortization and the forfeiture of awards granted in prior periods.

*Government grants*

In the three month periods ended November 30, 2018 and 2017, we did not recognize any research and development grants. As of November 30, 2018, we incurred liabilities to pay royalties to the Israel Innovation Authority of the Israeli Ministry of Economy & Industry of \$425,000.



***General and administrative expenses***

General and administrative expenses include the salaries and related expenses of our management, consulting costs, legal and professional fees, travel expenses, business development costs, insurance expenses and other general costs.

General and administrative expenses for the three month period ended November 30, 2018 decreased by 8% to \$932,000 from \$1,016,000 for the three month period ended November 30, 2017. The decrease in costs related to general and administrative activities during the three month period ended November 30, 2018 is primarily attributable to a decrease in stock-based compensation costs and is partially offset by an increase in salaries and related expenses. Stock-based compensation costs for the three month period ended November 30, 2018 totaled \$199,000, as compared to \$352,000 during the three month period ended November 30, 2017. The decrease is primarily attributable to the progress in amortization of awards granted to employees and directors during fiscal year 2017 and is partially offset by an increase due to awards granted during fiscal year 2018.

***Financial income, net***

Net financial income increased by 68% from net income of \$201,000 for the three month period ended November 30, 2017 to net income of \$338,000 for the three month period ended November 30, 2018. The increase is primarily attributable to an increase in income from bank deposits and held to maturity bonds as a result of an increase in interest rates, as well as an increase in fair value of the ordinary shares of D.N.A Biomedical Solutions Ltd., or D.N.A, and Entera Bio Ltd., or Entera, which was classified in other comprehensive income in fiscal year 2018, prior to the implementation of Accounting Standards Update 2016-01, "Recognition and Measurement of Financial Assets and Financial Liabilities."

***Other comprehensive income***

No unrealized gains on available for sale securities were recognized for the three month period ended November 30, 2018 as compared to gains of \$326,000 for the three month period ended November 30, 2017. The decrease is due to the implementation of Accounting Standards Update 2016-01, "Recognition and Measurement of Financial Assets and Financial Liabilities," under which changes in fair value of the ordinary shares of D.N.A and Entera that we hold are recognized as financial income.

***Liquidity and capital resources***

From inception through November 30, 2018, we have incurred losses in an aggregate amount of \$71,050,000. During that period we have financed our operations through several private placements of our common stock, as well as public offerings of our common stock, raising a total of \$77,736,000, net of transaction costs. During that period, we also received cash consideration of \$5,877,000 from the exercise of warrants and options. We will seek to obtain additional financing through similar sources in the future, as needed. As of November 30, 2018, we had \$3,861,000 of available cash, \$31,533,000 of short-term and long-term bank deposits and \$7,433,000 of marketable securities.

Management continues to evaluate various financing alternatives for funding future research and development activities and general and administrative expenses through fundraising in the public or private equity markets. Although there is no assurance that we will be successful with those initiatives, management believes that it will be able to secure the necessary financing as a result of future third party investments. Based on our current cash resources and commitments, we believe we will be able to maintain our current planned development activities and the corresponding level of expenditures for at least the next 12 months and beyond.

As of November 30, 2018, our total current assets were \$29,651,000 and our total current liabilities were \$4,164,000. On November 30, 2018, we had a working capital surplus of \$25,487,000 and an accumulated loss of \$71,050,000. As of August 31, 2018, our total current assets were \$31,037,000 and our total current liabilities were \$4,553,000. On August 31, 2018, we had a working capital surplus of \$26,484,000 and an accumulated loss of \$69,223,000. The decrease in working capital from August 31, 2018 to November 30, 2018 was primarily due to the cash used in operating activities.

During the three month period ended November 30, 2018, cash and cash equivalents decreased to \$3,861,000 from the \$4,996,000 reported as of August 31, 2018, which is due to the reasons described below.

Operating activities used cash of \$4,131,000 in the three month period ended November 30, 2018, as compared to \$2,728,000 used in the three month period ended November 30, 2017. Cash used in operating activities in the three month period ended November 30, 2018 primarily consisted of net loss resulting from research and development and general and administrative expenses, as well as changes in contract liabilities due to the License Agreement and is partially offset by changes in accounts payable and accrued expenses, while cash used in operating activities in the three month period ended November 30, 2017 primarily consisted of net loss resulting from research and development and general and administrative expenses, as well as changes in contract liabilities due to the License Agreement and is partially offset by changes in stock-based compensation.

Investing activities provided cash of \$2,995,000 in the three month period ended November 30, 2018, as compared to \$5,146,000 used in the three month period ended November 30, 2017. Cash provided by investing activities in the three month period ended November 30, 2018 consisted primarily of the maturity of short-term deposits and held to maturity securities and is partially offset by the purchase of held to maturity securities, while cash used in investing activities in the three month period ended November 30, 2017 consisted primarily of the purchase of short-term and long-term bank deposits and held to maturity securities.

Financing activities did not provide cash in the three month period ended November 30, 2018, as compared to \$5,160,000 provided in the three month period ended November 30, 2017. Financing activities in the three month period ended November 30, 2017 consisted of aggregate net proceeds of \$4,230,000 from our issuance of 453,919 shares of common stock under an At The Market Issuance Sales Agreement, dated April 2, 2015, or the Sales Agreement, with B. Riley FBR, Inc., as successor to FBR Capital Markets & Co., or FBR, as amended, and proceeds from the exercise of warrants and options. Pursuant to the Sales Agreement, we may, from time to time and at our option, issue and sell shares of our common stock having an aggregate offering price of up to \$25,000,000 through FBR as sales agent, subject to certain terms and conditions. Any shares sold will be sold pursuant to our effective shelf registration statement on Form S-3 including a prospectus dated February 2, 2017, as supplemented by a prospectus supplement dated April 5, 2017. We will pay FBR a commission of 3.0% of the gross proceeds of the sale of any shares sold through FBR.

#### *Off-balance sheet arrangements*

As of November 30, 2018, we had no off-balance sheet arrangements that have had or that we expect would be reasonably likely to have a future material effect on our financial condition, changes in financial condition, revenues or expenses, results of operations, liquidity, capital expenditures or capital resources.

#### *Critical accounting policies and estimates*

Our significant accounting policies are described in the notes to the consolidated financial statements as of August 31, 2018 included in our Annual Report and in the notes to the condensed consolidated financial statements contained in this Quarterly Report on Form 10-Q.

#### **Planned Expenditures**

We invest heavily in research and development, and we expect that in the upcoming years our research and development expenses will continue to be our major operating expense.

### **ITEM 3 - QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK**

There has been no significant change in our exposure to market risk during the quarter ended November 30, 2018. For a discussion of our exposure to market risk, refer to Part II, Item 7A, “Quantitative and Qualitative Disclosures About Market Risk,” contained in our Annual Report.

### **ITEM 4 - CONTROLS AND PROCEDURES**

#### **Disclosure Controls and Procedures**

Our management, including our Chief Executive Officer and Chief Financial Officer, evaluated the effectiveness of our disclosure controls and procedures as of November 30, 2018. Based upon that evaluation, our Chief Executive Officer and Chief Financial Officer concluded that our disclosure controls and procedures are effective.

#### **Changes in Internal Control over Financial Reporting**

There were no changes in our internal control over financial reporting that occurred during the quarter ended November 30, 2018 that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting, except for new controls with regard to the implementation of ASC 606.

## PART II – OTHER INFORMATION

### ITEM 2 - UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS

On November 1, 2018, we issued 2,500 shares of our common stock to Corporate Profile, LLC, or Corporate Profile, in payment of a portion of the consulting fee for investor relations services owed to Corporate Profile pursuant to a Letter Agreement, dated April 8, 2018, between us and Corporate Profile.

On October 15, 2018, we issued 5,694 shares of our common stock to Acorn Management Partners, L.L.C., or Acorn, in payment of a portion of the consulting fee for investor relations services owed to Acorn pursuant to a Consulting Agreement, dated July 15, 2018, between us and Acorn.

We issued these shares pursuant to an exemption from registration contained in Section 4(a)(2) of the Securities Act of 1933, as amended.

### ITEM 6 - EXHIBITS

#### Exhibit Number

- 10.1\* Consulting Agreement, dated December 12, 2018, between Oramed Ltd. and Joshua Hexter.
- 31.1\* Certification of Principal Executive Officer pursuant to Rule 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as amended.
- 31.2\* Certification of Principal Financial Officer pursuant to Rule 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as amended.
- 32.1\*\* Certification of Principal Executive Officer pursuant to 18 U.S.C. Section 1350.
- 32.2\*\* Certification of Principal Financial Officer pursuant to 18 U.S.C. Section 1350.
- 101.1\* The following financial statements from the Company's Quarterly Report on Form 10-Q for the quarter ended November 30, 2018 formatted in XBRL: (i) Condensed Consolidated Balance Sheets, (ii) Condensed Consolidated Statements of Comprehensive Loss, (iii) Condensed Consolidated Statement of Changes in Stockholders' Equity, (iv) Condensed Consolidated Statements of Cash Flows and (v) the Notes to Condensed Consolidated Financial Statements.

\* Filed herewith

\*\*Furnished herewith

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**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.

**ORAMED PHARMACEUTICALS INC.**

Date: January 14, 2019 By: /s/ Nadav Kidron  
Nadav Kidron  
President and Chief Executive Officer

Date: January 14, 2019 By: /s/ Hilla Eisenberg  
Hilla Eisenberg  
Chief Financial Officer  
(Principal Financial and Accounting Officer)