UNITED STATES SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 10-Q

(Mark One)	
QUARTERLY REPORT PURSUANT TO SE ACT OF 1934	ECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE
For the quarterly period	ended September 30, 2000
	OR
// TRANSITION REPORT PURSUANT TO SECT EXCHANGE ACT	TION 13 OR 15(d) OF THE
For the transition period	from to
Commission fil	e number 0-22613
AVI BIOPH	IARMA, INC.
(Exact name of registran	t as specified in its charter)
Oregon (State or other jurisdiction of incorporation or organization)	93-0797222 (I.R.S. Employer Identification No.)
One SW Columbia Street, Suite 1105, Portland, Oregon (Address of principal executive offices)	97258 (Zip Code)
Issuer's telephone number, in	cluding area code: 503-227-0554
Check whether the issuer (1) filed all reports required to be filed by Sepreceding 12 months (or for such shorter period that the registrant was requirements for the past 90 days. Yes /x/ No //	
Indicate the number of shares outstanding of each of the issuer's class	sses of common stock, as of the latest practicable date.
Common Stock with \$.0001 par value (Class)	21,447,931 (Outstanding at October 31, 2000)
AVI BIOPI	HARMA, INC.
FOR	M 10-Q
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PART I—FINANCIAL INFORMA	TION

	Statements of Operations—Three and Nine Months Ended September 30, 2000 and 1999 and from July 22, 1980 (Inception) to September 30, 2000	4
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AVI BIOPHARMA, INC.

(A Development Stage Company)

BALANCE SHEETS

	September 30, 2000		December 31, 1999	
Assets				
Current Assets:				
Cash and cash equivalents	\$	29,554,775	\$	8,683,005
Short-term securities—available-for-sale		8,592,886		2,937,500
Other current assets		121,556		31,242
Total Current Assets		38,269,217		11,651,747
Property and Equipment, net of accumulated depreciation and amortization of \$2,637,250 and \$2,518,494		811,975		403,303
Patent Costs, net of accumulated amortization of \$529,268 and \$418,268		903,721		844,731
Other Assets		29,847		29,847
Other Assets		29,047		29,047
Total Assets	\$	40,014,760	\$	12,929,628
	_			
Liabilities and Shareholders' Equity Current Liabilities:				
Accounts payable	\$	1,014,102	\$	727,673
Accrued liabilities	Ψ	302,223	Ψ	312,481
Accided liabilities		302,223		312,401
Total Current Liabilities		1,316,325		1,040,154
Shareholders' Equity:				
Preferred Stock, \$.0001 par value, 2,000,000 shares authorized;				
none issued and outstanding		_		_
Common stock, \$.0001 par value, 50,000,000 shares authorized;				
21,444,132 and 16,236,428 issued and outstanding		2,144		1,624
Additional paid-in capital		105,286,681		62,901,227
Accumulated other comprehensive income (loss)		(9,304,114)		40,500
Deficit accumulated during the development stage		(57,286,276)		(51,053,877)
Total Shareholders' Equity		38,698,435		11,889,474
Total Liabilities and Shareholders' Equity	\$	40,014,760	\$	12,929,628

The accompanying notes are an integral part of these balance sheets

(A Development Stage Company)

STATEMENTS OF OPERATIONS

		Three months ended September 30,		Nine mon Septem	July 22, 1980	
		2000	1999	2000	1999	(Inception) to September 30, 2000
Revenues, from license fees,						
grants and research contracts	\$	122,215 \$	3,558	\$ 1,272,338	\$ 7,783	\$ 2,113,555
Operating expenses:						
Research and development		2,155,538	1,739,728	6,575,953	4,709,856	31,303,586
General and administrative Acquired in-process research		557,911	504,607	1,484,159	1,341,099	10,682,827
and development	_				61,337	19,545,028
		2,713,449	2,244,335	8,060,112	6,112,292	61,531,441
Other Income:						
Interest income, net		339,130	35,696	555,375	162,784	2,034,860
Realized gain on sale of short-term investments		_	_	_	_	96,750
		339,130	35,696	555,375	162,784	2,131,610
Net loss	\$	(2,252,104) \$	(2,205,081)	\$ (6,232,399)	\$ (5,941,725)	\$ (57,286,276)
Net loss per share—basic and diluted	\$	(0.11) \$	(0.17)	\$ (0.35)	\$ (0.45)	
Weighted average number of common shares outstanding for computing basic and diluted earnings per share		20,303,112	13,351,206	17,800,160	13,350,597	

The accompanying notes are an integral part of these balance sheets

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AVI BIOPHARMA, INC.

(A Development Stage Company)

STATEMENTS OF CASH FLOWS

	Nine months end	For the Period July 22, 1980	
	2000	1999	(Inception) to September 30, 2000
Cash flows from operating activities:	 		
Net loss	\$ (6,232,399)	\$ (5,941,725)	\$ (57,286,276)
Adjustments to reconcile net loss to net cash flows used in operating activities:	, ,	,	·
Depreciation and amortization	235,170	222,802	3,288,701
Realized gain on sale of short-term investments—available for sale	<u> </u>	_	(96,750)
Compensation expense on issuance of common stock and partnership units	_	_	251,992
Compensation expense on issuance of options and warrants to purchase common stock or partnership units	_	_	562,353
Conversion of interest accrued to common stock	_	_	7,860
Acquired in-process research and development	_	61,337	19,545,028
(Increase) decrease in:			
Other current assets	(90,314)	478,186	(121,556)
Other assets	_	_	(29,847)
Net increase (decrease) in accounts payable and accrued liabilities	276,171	(508,806)	1,316,325

Net cash used in operating activities		(5,811,372)		(5,688,206)		(32,562,170)
Cash flows from investing activities:						
Proceeds from sale or redemption of						
short-term investments		_		_		247.750
Purchase of property and equipment		(532,842)		(133,686)		(3,514,728)
Patent costs		(169,990)		(197,651)		(1,489,669)
Acquisition costs		_		(61,337)		(2,377,616)
					_	, , , , , ,
Net cash used in investing activities		(702,832)		(392,674)		(7,134,263)
Cash flows from financing activities:						
Proceeds from sale of common stock,						
warrants, and partnership units, net of						
offering costs, and exercise of options		27,385,974		15,000		69,636,645
Buyback of common stock pursuant to						
rescission offering		_		_		(288,795)
Withdrawal of partnership net assets		_		_		(176,642)
Issuance of convertible debt		_		_		80,000
Net cash provided by financing activities		27,385,974		15,000		69,251,208
Increase (decrease) in cash and cash equivalents		20,871,770		(6,065,880)		29,554,775
				,		, ,
Cash and cash equivalents:						
Beginning of period		8,683,005		8,510,020		_
Ford of a color	<u> </u>	00 554 775	<u> </u>	0.444.440	Φ.	00 554 775
End of period	\$	29,554,775	\$	2,444,140	\$	29,554,775
OURRI EMENTAL COURRI II E OF						
SUPPLEMENTAL SCHEDULE OF						
NONCASH INVESTING ACTIVITIES AND FINANCING ACTIVITIES:						
Short-term securities—available-for-sale						
received in connection with the private						
offering	\$	15,000,000	\$	_	\$	17,897,000
Unrealized loss on short-term	Ψ	10,000,000	Ψ		Ψ	17,007,000
securities—available-for-sale	\$	(9,344,614)	\$	_	\$	(9,304,114)
	Ť	(0,0,0 / 1)	Ψ		Ψ.	(5,55.,.11)

The accompanying notes are an integral part of these statements

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AVI BIOPHARMA, Inc.

NOTES TO FINANCIAL STATEMENTS

(Unaudited)

Note 1. Basis of Presentation

The financial information included herein for the three and nine-month periods ended September 30, 2000 and 1999 and the financial information as of September 30, 2000 is unaudited; however, such information reflects all adjustments consisting only of normal recurring adjustments which are, in the opinion of management, necessary for a fair presentation of the financial position, results of operations and cash flows for the interim periods. The financial information as of December 31, 1999 is derived from AVI BioPharma, Inc.'s (the Company's) Form 10-K. The interim financial statements should be read in conjunction with the financial statements and the notes thereto included in the Company's Form 10-K. The results of operations for the interim periods presented are not necessarily indicative of the results to be expected for the full year.

Note 2. Earnings Per Share

Basic EPS is calculated using the weighted average number of common shares outstanding for the period and diluted EPS is computed using the weighted average number of common shares and dilutive common equivalent shares outstanding. Given that the Company is in a loss position, there is no difference between basic EPS and diluted EPS since the common stock equivalents would be antidilutive.

Three Months Ended September 30,	2000	1999
Net loss Weighted average number of shares of common stock and common	\$ (2,252,104)	\$ (2,205,081)
stock equivalents outstanding:		
Weighted average number of common shares outstanding for computing basic earnings per share	20,303,112	13,351,206
Dilutive effect of warrants and stock options after application of the treasury stock method	*	*

Weighted average number of common shares outstanding for computing diluted earnings per share		20,303,112	13,351,206
Net loss per share—basic and diluted		\$ (0.11)	\$ (0.17)
Nine Months Ended September 30,		2000	1999
Net loss		\$ (6,232,399)	\$ (5,941,725)
Weighted average number of shares of common stock and common stock equivalents outstanding:			,
Weighted average number of common shares outstanding for computing basic earnings per share		17,800,160	13,350,597
Dilutive effect of warrants and stock options after application of the treasury stock method		*	*
Weighted average number of common shares outstanding for computing diluted earnings per share		17,800,160	13,350,597
Net loss per share—basic and diluted		\$ (0.35)	\$ (0.45)
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* The following common stock equivalents are excluded from earnings per share calculation as their effect would have been antidilutive:

Three Months Ended September 30,	2000	1999
Warrants and stock options Nine Months Ended September 30,	10,298,111 2000	7,077,082 1999
Warrants and stock options	10,298,111	7,077,082

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Item 2. Management's Discussion and Analysis

Forward-Looking Information

The Financial Statements and Notes thereto should be read in conjunction with the following discussion. The discussion in this Form 10-Q contains certain forward-looking statements that involve risks and uncertainties, including, but not limited to, the results of research and development efforts, the results of pre-clinical and clinical testing, the effect of regulation by FDA and other agencies, the impact of competitive products, product development, commercialization and technological difficulties, and other risks detailed in the Company's Securities and Exchange Commission filings.

Overview

From its inception in July 1980, the Company has devoted its resources primarily to fund its research and development efforts. The Company has been unprofitable since inception and, other than limited interest, license fees and grants, has had no material revenues from the sale of products or other sources, and does not expect material revenues for at least the next 12 months. The Company expects to continue to incur losses for the foreseeable future as it expands its research and development efforts. As of September 30, 2000, the Company's accumulated deficit was \$57,286,276.

Results of Operations

Revenues, from license fees, grants and research contracts, increased to \$122,215 in the third quarter of 2000 from \$3,558 in the third quarter of 1999. Revenues, from license fees, grants and research contracts, increased to \$1,272,338 for the nine months ended September 30, 2000 from \$7,783 for the comparable period of 1999 due primarily to the receipt of a \$1,000,000 fee for expansion of a license for diagnostic applications.

Operating expenses increased to \$2,713,449 in the third quarter of 2000 from \$2,244,335 in the third quarter of 1999 and to \$8,060,112 for the nine months ended September 30, 2000 from \$6,112,292 for the comparable period of 1999 due to increases in research and development staffing and increased expenses associated with outside collaborations and pre-clinical and clinical testing of the Company's technologies. Additionally, increased general and administrative costs were incurred to support the research expansion, and to continue to broaden the Company's investor and public relations efforts. Net interest income increased to \$339,130 in the third quarter of 2000 from \$35,696 in the third quarter of 1999 and to \$555,375 for the nine months ended September 30, 2000 from \$162,784 for the comparable period in 1999 due to earnings on increased cash balances.

Liquidity and Capital Resources

The Company's cash and cash equivalents were \$29,554,775 at September 30, 2000, compared with \$8,683,005 at December 31, 1999. The increase of \$20,871,770 was primarily due to net proceeds of \$19,886,619 from the Company's secondary offering completed August 1, 2000, net proceeds of \$4,744,720 from the alliance with SuperGen, Inc. for shared development and marketing rights for Avicine and \$2,754,635 from the exercise of options and warrants, offset by \$5,811,372 used in operations and \$702,832 used for investing activities which consist primarily of purchases of property and equipment and patent related costs. In addition the Company's short-term securities increased \$5,655,386 to \$8,592,886 at September 30, 2000 due to receiving 347,826 shares of SuperGen, Inc. common stock from the alliance with SuperGen, Inc. for shared development and marketing rights for Avicine, offset by unrealized losses in the value of these securities.

In April 2000, the Company entered into an alliance with SuperGen, Inc. for shared development and marketing rights for Avicine. Under the terms of the agreement, AVI and SuperGen will equally share in future clinical development and FDA registration costs as well as in profits from product sales

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in the United States. At closing the Company received from SuperGen, Inc. \$5,000,000 in cash and 347,826 shares of SuperGen, Inc. common stock in exchange for 1,684,211 shares of AVI common stock and a warrant to purchase 1,665,878 shares of AVI common stock, subject to anti-dilution provisions. Closing of the transaction occurred during the third quarter of 2000.

In July 2000, the Company completed a secondary offering for 3,000,000 shares of common stock at \$7.25 per share. Closing occurred in August 2000. Net proceeds were \$19,886,619. In addition, representatives' warrants to purchase 300,000 shares of AVI common stock were issued to the underwriters' of the secondary offering.

The Company's future expenditures and capital requirements will depend on numerous factors, including without limitation, the progress of its research and development programs, the progress of its pre-clinical and clinical trials, the time and costs involved in obtaining regulatory approvals, the cost of filing, prosecuting, defending and enforcing any patent claims and other intellectual property rights, competing technological and market developments, the ability of the Company to establish collaborative arrangements and the terms of any such arrangements, and the costs associated with commercialization of its products. The Company's cash requirements are expected to continue to increase significantly each year as it expands its activities and operations. There can be no assurance, however, that the Company will ever be able to generate product revenues or achieve or sustain profitability.

The Company expects that its cash requirements over the next twenty-four months will be satisfied by existing cash resources.

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PART II—OTHER INFORMATION

Item 6. Exhibits and Reports on Form 8-K

The exhibit filed as a part of this report is listed below and this list constitutes the exhibit index.

Exhibit No

(a)

27 Financial Data Schedule

(b) Reports on Form 8-K

The Company did not file any Reports on Form 8-K during the guarter ended September 30, 2000.

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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.

Date: November 4, 2000 AVI BIOPHARMA, INC.

By: /s/ DENIS R. BURGER, PH.D.

Denis R. Burger, Ph.D. Chief Executive Officer and Chairman (of the Board of Directors) (Principal Executive Officer)

By:	/s/ MARK M	WERRER
Dy.	/ S/ IVI/AI AI A IVI	. WEDDER

Mark M. Webber Chief Financial Officer (Principal Financial and Accounting Officer)

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QuickLinks

(Mark One)
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SIGNATURES

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