SECURITIES AND EXCHANGE COMMISSION

FORM 10-Q

Quarterly report pursuant to sections 13 or 15(d)

Filing Date: **2001-08-03** | Period of Report: **2001-06-30** SEC Accession No. 0001021408-01-504445

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FILER

HOLLIS EDEN PHARMACEUTICALS INC /DE/

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SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

Form 10-Q

(Mark one)

Quarterly Report Under Section 13 or 15 (d)
X Of the Securities Exchange Act of 1934

For Quarterly Period Ended June 30, 2001

Transition Report Pursuant to Section 13 or 15(d) of the Securities Exchange Act 1934 for the period from $__$ to $__$.

HOLLIS-EDEN PHARMACEUTICALS, INC (Exact name of registrant as specified in its charter)

DELAWARE

(State or other jurisdiction of incorporation)

000-24672 (Commission File No.)

13-3697002

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

9333 Genesee Ave., Suite 200
SAN DIEGO, CALIFORNIA 92121
(Address of principal executive offices and zip code)

Registrant's telephone number, including area code: (858) 587-9333

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days.

YES X N

As of August 3, 2001 there were 11,615,803 shares of registrant's Common Stock, \$.01 par value, outstanding.

HOLLIS-EDEN PHARMACEUTICALS, INC. Form 10-Q FOR THE QUARTER ENDED JUNE 30, 2001

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Part I. Financial Information			
Item I. Financial Statements			
Hollis-Eden Pharmaceuticals, Inc. (A Development Stage Company) Balance Sheets (Unaudited)			
<table></table>			
<caption> All numbers in thousands</caption>	June 30, 2001	Dec. 31, 2000	
<\$>	<c></c>	 <c></c>	
ASSETS: Current assets:			
Cash and cash equivalents		\$ 34,298	
Prepaid expenses Deposits		96 27	
Total current assets	27,716	34,421	
Property and equipment, net of accumulated			
depreciation of \$267 and \$204 Other receivables	468 35	422	
Other receivable from related party	279	256	
Total assets	\$ 28,498 ======	\$ 35,099 =======	
LIABILITIES AND STOCKHOLDERS' EQUITY:			
Current liabilities: Accounts payable and accrued expenses	\$ 3,279	\$ 2,636	
Total liabilities		2,636	
Commitments and contingencies			
Stockholders' equity:			
Preferred stock, \$.01 par value, 10,000 shares			
authorized; no shares outstanding Common stock, \$.01 par value,	_	-	
30,000 shares authorized; 11,616 and			
11,590 shares issued and outstanding	116 80,669	116 80,503	
Deficit accumulated during development stage	(55,566)	(48,156)	
Total stockholders' equity	25 , 219	32,463	
Total liabilities and stockholders' equity	\$ 28,498	\$ 35,099 ======	

			The accompanying notes are an integral part of these fi	nancial stateme	ents.	
	acrar stateme					
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Hollis-Eden Pharmaceuticals, Inc. (A Development Stage Company) Statements of Operations (Unaudited)						
All numbers in thousands, except per share amounts

<TABLE> <CAPTION>

	3 months er 2001	aded June 30, 2000	6 months e 2001	ended June 30, 2000	Inception (Aug.15,1994) to June 30 2001
<\$>	<c></c>	<c></c>	<c></c>	<c></c>	<c></c>
Operating expenses:					
Research and development: R&D operating expenses R&D costs related to common stock, option, & warrant grants	\$ 2,942	\$ 2,617	\$ 5,659	\$ 6,815	\$ 32,249
for collaborations	24	24	48	2,478	5,228
G&A operating expenses	1,242	1,025	2,507	2,101	15,493
stock, option, & warrant grants					9,490
Total operating expenses	4,208	3,666	8,214	11,394	62,460
Other income (expense): Interest income	330	673 -	804	1,316	6,944 (50)
Total other income	330	673	804	1,316	6,894
Net loss	\$ (3,878) ======	\$ (2,993) ======	\$ (7,410) ======	\$ (10,078) ======	\$ (55,566) ======
Net loss per share-basic and diluted	(0.33)	(0 27)	(0.64)	(0.90)	
Weighted average number of common shares outstanding-basic and diluted	11,616	11,233	11,609	11,203	

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

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Hollis-Eden Pharmaceuticals, Inc. (A Development Stage Company) Statements of Cash Flows (Unaudited)

All numbers in thousands

<TABLE> <CAPTION>

		6 months 2001		June 30, 2000		Inception 15, 1994) to June 30 2001
<\$>	<c></c>		<c></c>		<c></c>	
Cash flows from operating activities:						
Net loss Adjustments to reconcile net loss to net cash used in operating activities:	\$	(7,410)	\$	(10,078)	\$	(55,566)
Depreciation		63		52		2.67
Common stock issued for the company 401k/401m plan Common stock issued as consideration for amendments/		95		63		158
termination of agreementsExpense related to common stock issued for the purchase of		-		-		67
technology Common stock and options issued as consideration for license		-		1,848		1,848
fees and services		48		630		1,742
Common stock issued as consideration for In Process R&D Expense related to warrants issued as consideration to		-		-		2,000
consultants Expense related to warrants issued to a director for		-		_		2,140
successful closure of merger		-		_		570
Expense related to stock options issued		-		_		5,140
Deferred compensation expense related to options issued		=		-		1,210
Changes in assets and liabilities:						
Prepaid expenses		(128)		(75)		(224)
Deposits		_		_		(27)
Other receivables Loan receivable from related party		(35) (23)		(13)		(35) (279)

Period from

1,224 (581)	511 - -	3,279 - 7
(6,747)	(7,062)	(37,703)
(109)	(49)	(742)
(109)	(49)	(742)
- - - - - 23 	- - - - 765 765	104 4,000 42,172 371 6,271 12,992
(6,833) 34,298	(6,346) 47,486	27 , 465
\$ 27,465 ======	\$ 41,140 ======	\$ 27,465 =======
	(109)	(581) ————————————————————————————————————

</TABLE>

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

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HOLLIS-EDEN PHARMACEUTICALS, INC.
(A Development Stage Company)

NOTES TO FINANCIAL STATEMENTS

(UNAUDITED)

1. Basis of Presentation

The information at June 30, 2001, and for the three-month and six-month periods ended June 30, 2001 and 2000, is unaudited. In the opinion of management, these financial statements include all adjustments, consisting of normal recurring adjustments, necessary for a fair presentation of the results for the interim periods presented. Interim results are not necessarily indicative of results for a full year. These financial statements should be read in conjunction with the Hollis-Eden Pharmaceuticals, Inc. ("Hollis-Eden" or the "Company") Annual Report on Form 10-K for the year ended December 31, 2000, which was filed with the United States Securities and Exchange Commission on March 30, 2001.

While management believes that the discussion and analysis in this report is adequate for a fair presentation of the information, management recommends that this discussion and analysis be read in conjunction with Management's Discussion and Analysis of Results of Operations and Financial Condition included in the Company's Annual Report on Form 10-K for the year ended December 31, 2000.

2. Technology Acquisition - Commitments and Contingencies

Pursuant to a Technology Assignment Agreement dated January 20, 2000, Patrick Prendergast and Colthurst Limited assigned to Hollis-Eden ownership of all patents, patent applications and current or future improvements of the technology previously licensed to Hollis-Eden under the Colthurst License Agreement dated May 18, 1994 (as amended), including HE2000, Hollis-Eden's lead clinical compound. In exchange for certain covenants made by Mr. Prendergast and Colthurst regarding their future activities (the "Covenants") as well as for the consideration described above, Hollis-Eden agreed to issue to Colthurst 660,000 shares of Common Stock and a warrant to purchase an aggregate of 400,000 shares of Common Stock at \$25 per share, of which only 132,000 shares were to be issued in 2000, with the remaining 528,000 shares to be issued over the next four years conditioned on continued compliance with the Covenants. The shares underlying the warrant were to vest over four years and were likewise conditioned on continued compliance with the Covenants.

In accordance with Emerging Issues Task Force No. 96-18, Accounting for Equity Instruments That Are Issued to Other Than Employees for Acquiring, or in Conjunction with Selling, Goods or Services, certain future events could not be determined at the date of the agreements (January 2000). Accordingly, the shares and warrants were to be accounted for as they vest or were issued. During

2000, we recorded a research and development charge for \$1.9 million representing the fair value of the 132,000 shares issued under the agreement.

Because Mr. Prendergast and Colthurst failed to comply with the Covenants, Hollis-Eden has not issued any additional shares to Colthurst and believes it has no obligation to issue to Colthurst any additional shares. While Hollis-Eden is confident in its analysis, if any dispute should arise in this matter, Hollis-Eden cannot guarantee that, subject to the resolution of any such dispute, it will not be required to issue additional equity to Mr. Prendergast and Colthurst or that it will

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not incur additional accounting charges or other adverse accounting reporting as a result of any such resolution.

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

The forward-looking comments contained in the following discussion involve risks and uncertainties. Our actual results may differ materially from those discussed here. Factors that could cause or contribute to such differences can be found in the following discussion, as well as in the Company's Annual Report on Form 10-K for the year ended December 31, 2000.

General

Hollis-Eden Pharmaceuticals, Inc., a development-stage pharmaceutical company, is presently engaged in the discovery and development of products for the treatment of infectious diseases and immune system disorders, including HIV/AIDS, hepatitis B and C, and malaria.

We are focusing our initial development efforts on a potent series of immune regulating hormones and hormone analogs. We are currently developing three clinical stage compounds with potentially broad therapeutic applications: HE2000, HE2200 and, through our relationship with Aeson Therapeutics, HE2500. Our lead compound, HE2000, is currently in Phase II clinical studies. By altering cytokine production, HE2000 appears from early clinical studies to help reestablish immune system balance in situations such as HIV, where the immune system is dysregulated. In the case of HIV, we believe that by reestablishing this balance the immune system may be able to better control virus levels and potentially delay or prevent the progression to AIDS. In addition, based on the mechanism of action, we believe these compounds will have an attractive safety profile and will avoid issues of resistance that plague many existing antiviral drugs.

We have been unprofitable since our inception and we expect to incur substantial additional operating losses for at least the next few years as we increase expenditures on research and development and begin to allocate significant and increasing resources to clinical testing and other activities. In addition, during the next few years, we may have to meet the substantial new challenge of developing the capability to market products. Accordingly, our activities to date are not as broad in depth or scope as the activities we may undertake in the future, and our historical operations and financial information may not be indicative of our future operating results or financial condition or our ability to operate profitably as a commercial enterprise when and if we succeed in bringing any drug candidates to market.

On March 26, 1997, Hollis-Eden, Inc., a Delaware corporation, was merged with and into us, then known as Initial Acquisition Corp. ("IAC"), a Delaware corporation. Upon consummation of the merger of Hollis-Eden, Inc. with IAC (the "Merger"), Hollis-Eden, Inc. ceased to exist, and IAC changed its name to Hollis-Eden Pharmaceuticals, Inc.

Results of Operations

We have not generated any revenues for the period from August 15, 1994 (inception of Hollis-Eden) through June 30, 2001. We have devoted substantially all of our resources to the payment of licensing fees and research and development expenses plus expenses related to the startup of our business. From inception until June 30, 2001, we have incurred expenses of approximately \$37.5 million in research and development (\$5.2 million are non-cash expenses) and \$25.0 million in general and administrative expenses (\$9.5 million are non-cash expenses), which have been partially offset by \$6.9 million in net interest income resulting in a loss of \$55.6 million for the period.

Research and development expenses were \$3.0 million and \$5.7 million for the three- and six-month periods ended June 30, 2001 and \$2.6 million and \$9.3 million for the same periods in 2000. The research and

development expenses relate primarily to the ongoing development, preclinical testing, and clinical trials for our first drug candidate, HE2000. Included in the six-month period ended June 30, 2000 was \$2.2 million related to collaborations and \$2.5 million in non-cash charges related to the acquisition of technology and in-process R&D. There were no such expenses in the six-month period ended June 30, 2001. The increase in research and development expenses (net of collaborations and technology acquisitions) for the three- and six-month periods ended June 30, 2001, compared to the same period in 2000, was due to increased staffing and clinical trial activities.

General and administrative operating expenses were \$1.2 million and \$2.5 million for the three- and six-month periods ended June 30, 2001, compared to \$1.0 million and \$2.1 million for the same period in 2000. The general and administrative expenses relate to staffing, facilities, supplies, benefits, recruiting, legal, investor relations and travel. The increase in general and administrative operating expenses was mainly due to expenses associated with investor relations and the growth of the Company's operations.

Net interest income was \$330,000 and \$804,000 for the three- and six-month periods ended June 30, 2001, compared to \$673,000 and \$1,316,000 for the same periods in 2000. The decline in interest income is due to lower interest rates and lower average balances of cash and cash equivalents as a result of ongoing operating losses.

Liquidity and Capital Resources

We have financed our operations since inception through the sale of equity. During the year ended December 31, 1995, we received cash proceeds of \$250,000 from the sale of securities. In May 1996, we completed a private placement of shares of Common Stock, from which we received aggregate gross proceeds of \$1.3 million. In March 1997, the Merger of IAC and Hollis-Eden, Inc. provided us with \$6.5 million in cash and other receivables. In May 1998, we completed a private placement of shares and warrants, from which we received gross proceeds of \$20 million. During January 1999, we completed two private placements raising approximately \$25 million. In addition, since inception Hollis-Eden has received a total of \$13 million from the exercise of warrants and stock options.

Our operations to date have consumed substantial capital without generating any revenues, and we will continue to require substantial and increasing amounts of funds to conduct necessary research and development and preclinical and clinical testing of our drug candidates, and to market any drug candidates that receive regulatory approval. We do not expect to generate revenue from operations for the foreseeable future, and our ability to meet our cash obligations as they become due and payable is expected to depend for at least the next several years on our ability to sell securities, borrow funds or some combination thereof. Based upon our current plans, we believe that our existing capital resources, together with interest thereon, will be sufficient to meet our operating expenses and capital requirements well into 2002. However, changes in our research and development plans or other events affecting our operating expenses may result in the expenditure of such cash before that time. We may not be successful in raising necessary funds. Our future capital requirements will depend upon many factors, including progress with preclinical testing and clinical trials, the number and breadth of our programs, the time and costs involved in preparing, filing, prosecuting, maintaining and enforcing patent claims and other proprietary rights, the time and costs involved in obtaining regulatory approvals, competing technological and market developments, and our ability to establish collaborative arrangements, effective commercialization, marketing activities and other arrangements. We expect to continue to incur increasing negative cash flows and net losses for the foreseeable future.

Item 3. Quantitative and Qualitative Disclosures about Market Risk Not applicable.

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PART II Other Information

Item 1. Legal Proceedings

From time to time, we may be involved in litigation relating to claims arising out of our operations in the normal course of business. As of the date of this Quarterly Report on Form 10-Q, we are not engaged in any legal proceedings that are expected, individually or in the aggregate, to have a material adverse effect on our business, financial condition or operating results.

Item 2. Changes in Securities

In May 2001, the Company entered into a financial advisory and investment banking agreement with H.C. Wainwright & Co., Inc. As an initial retainer fee, the Company agreed to issue to H.C. Wainwright & Co. Inc, \$25,000 and a warrant to purchase up to 25,000 shares of Common Stock at an exercise price of \$3.09 per share.

The sale and issuance of securities in the transaction described in the foregoing paragraph was deemed to be exempt from registration under the Securities Act of 1933, as amended, by virtue of Section 4(2) and/or Regulation D promulgated under such Act. The recipient represented their intention to acquire the securities for investment only and not with a view to the distribution thereof. Appropriate legends are affixed to the securities issued in such transaction. All recipients either received adequate information about the Company or had access, through employment or other relationships, to such information.

Item 3. Defaults upon Senior Securities

None

Item 4. Submission of Matters to a Vote of Securities Holders

The Annual Meeting of Stockholders of Hollis-Eden Pharmaceuticals was held on June 29, 2001. At this meeting, the Company solicited the vote of the stockholders on the proposals set forth below and received for each proposal the votes indicated below:

- (1) To elect three Class I directors to hold office until the 2004 Annual Meeting of Stockholders. Elected to serve as Class I directors were Paul Bagley, William H. Tilley and Salvatore J. Zizza. For each elected director the results of voting were: 9,974,219 for, 542,099 withheld, and 0 abstained. The continuing directors are Thomas Charles Merigan, Jr., M.D., Brendan R. McDonnell, Richard B.Hollis and Leonard Makowka M.D., Ph.D., FRCS(C), FACS.
- (2) To approve the Company's 1997 Incentive Stock Option Plan, as amended, to increase the aggregate number of shares of Common Stock authorized for issuance under such plan by 500,000 shares to a total of 3,250,000 shares. The 1997 Incentive Stock Option Plan, as amended, was approved with the following votes: 9,799,259 for, 655,081 against, and 61,978 abstained.
- (3) To approve the Company's Restated Certificate of Incorporation, as amended, to increase the authorized number of shares of Common Stock to 50,000,000. The Company's Restated Certificate of Incorporation, as amended, was approved with the following votes: 10,289,416 for, 202,861 against, and 24,041 abstained.
- (4) To ratify the selection of BDO Seidman, LLP as independent auditors of the Company for its fiscal year ending December 31, 2001. The selection of BDO Seidman, LLP as independent auditors of the Company for its fiscal year ending December 31, 2001 was ratified with the following votes: 10,489,259 for, 8,256 against, and 18,803 abstained.

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Item 5. Other Information

None

Item 6. Exhibits and Reports on Form 8-K

(a) The following exhibits are included as part of this report:

Exhibit Number Description

3.4 Certificate of Amendment to Certificate of Incorporation

(b) Reports on Form 8-K: None

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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 hereunto duly authorized.

Dated: August 3, 2001

By: /s/ Daniel D. Burgess

Daniel D. Burgess Chief Operating Officer/ Chief Financial Officer (Principal Financial Officer)

By: /s/ Robert W. Weber

Robert W. Weber Vice President-Controller/ Chief Accounting Officer (Principal Accounting Officer)

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CERTIFICATE OF AMENDMENT OF CERTIFICATE OF INCORPORATION OF HOLLIS-EDEN PHARMACEUTICALS, INC.

Hollis-Eden Pharmaceuticals, Inc. (the "Corporation"), a corporation organized and existing under and by virtue of the General Corporation Law of the State of Delaware (the "DGCL"), does hereby certify that:

- 1. The name of the Corporation is Hollis-Eden Pharmaceuticals, Inc. The Corporation was originally incorporated under the name Initial Acquisition Corp. The original Certificate of Incorporation for Initial Acquisition Corp. was filed with the Secretary of State of the State of Delaware (the "Secretary of State") on November 18, 1992. The name of the Corporation was changed from Initial Acquisition Corp. to Hollis-Eden Pharmaceuticals, Inc. through a Certificate of Merger filed with the Secretary of State on March 26, 1997.
- 2. Pursuant to a meeting of the Board of Directors of the Corporation in accordance with the provisions of Sections 141 and 242 of the DGCL, the Corporation adopted resolutions setting forth a proposed Amendment to Certificate of Incorporation of the Corporation, declaring said Amendment to Certificate of Incorporation to be advisable and authorizing the officers of the Corporation to present the proposed Amendment to Certificate of Incorporation to the stockholders of the Corporation for their consideration.
- 3. Thereafter, the proposed Amendment to Certificate of Incorporation of the Corporation was approved by the holders of a majority of the outstanding shares of stock of the Corporation entitled to vote thereon at a duly convened annual meeting of stockholders called in accordance with Section 222 of the DGCL.
- 4. Pursuant to Section 242 of the DGCL, this Amendment to Certificate of Incorporation amends the provisions of the Certificate of Incorporation of this Corporation filed with the Secretary of State on March 26, 1997.
- 5. This Certificate of Incorporation was duly adopted in accordance with the provisions of Sections 141 and 242 of the DGCL.
- 6. Paragraph A of Article IV of the Certificate of Incorporation of the Corporation is hereby amended to read in its entirety as follows:
 - "IV(A). This corporation is authorized to issue two classes of stock to be designated, respectively, "Common Stock" and "Preferred Stock." The total number of shares which the corporation is authorized to issue is sixty million (60,000,000) shares. Fifty million (50,000,000) shares shall be Common Stock, each having a par value of one cent (\$.01). Ten million (10,000,000) shares shall

be Preferred Stock, each having a par value of one cent (\$.01)."

In Witness Whereof, Hollis-Eden Pharmaceuticals, Inc. has caused this Certificate of Amendment to be signed by its Vice President and Corporate General Counsel this 30th day of July, 2001.

Hollis-Eden Pharmaceuticals, Inc.

By: /s/ Eric J. Loumeau

Eric J. Loumeau, Esq.

Vice President, Corporate General Counsel