UNITED STATES SECURITIES AND EXCHANGE COMMISSION Washington, DC 20549

FORM 10-Q

[X] QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934 For the quarterly period ended March 31, 1998

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

Commission file number: 0-15006

T CELL SCIENCES, INC. (Exact name of registrant as specified in charter)

Delaware (State of Incorporation)

No. 13-3191702 (I.R.S. Employer Identification No.)

119 Fourth Avenue, Needham, Massachusetts 02194-2725

(Address of principal executive offices) (Zip code)

(617) 433-0771 (Registrant's telephone number, including area code)

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports) and (2) has been subject to such filing requirements for the past 90 days. Yes X No

Class

Outstanding as of May 8, 1998

Common Stock, par value \$.001 28,531,285

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PART I. FINANCIAL INFORMATION

Item 1. Financial Statements

T CELL SCIENCES, INC. CONDENSED CONSOLIDATED BALANCE SHEET March 31, 1998 and December 31, 1997

	March 31, 1998	December 31, 1997
ASSETS		(audited)
Current Assets:	Ф 0 101 000	Ф 6 426 200
Cash and Cash Equivalents Current Portion Restricted Cash	\$ 8,181,900 750,000	\$ 6,436,300 750,000
Accounts Receivable	12,600	22,900
Inventories	7,300	15,000
Prepaid Expenses and Other	187,800	165,400
Total Current Assets	9,139,600	7,389,600
Property and Equipment, Net	346,400	364,500
Restricted Cash	500,000	525,000
Other Noncurrent Assets	1,598,600	1,547,500
Total Assets	\$ 11,584,600	\$ 9,826,600
LIADILITIES AND STOCKHOLDERS LIGHTLY		
LIABILITIES AND STOCKHOLDERS' EQUITY Current Liabilities:		
Accounts Payable	\$ 225,700	\$ 201,200
Accrued Expenses	757,000	1,059,900
Deferred Revenue	500,000	750,000
Short-Term Note Payable	750,000	750,000
Total Current Liabilities	2,232,700	2,761,100
Long-Term Note Payable	750,000	750,000
Stockholders' Equity:		
Common Stock, \$.001 Par Value	28,500	26,500
Additional Paid-in Capital	80,256,800	76,561,400
Less: Common Treasury Shares at Cost	(31,300)	(35,800)
Accumulated Deficit	(71,652,100)	(70,236,600)
Total Stockholders' Equity	8,601,900	6,315,500
Total Liabilities and Stockholders' Equity	\$ 11,584,600	\$ 9,826,600

See accompanying notes to condensed consolidated financial statements

	March 31, 1998	March 31, 1997
OPERATING REVENUE:		
Product Development and Licensing Agreements Product Sales	27,400	\$ 62,100 1,300
Total Operating Revenue	361,000	63,400
OPERATING EXPENSE:		
Cost of Product Sales Research and Development General and Administrative Marketing and Sales	13,100 1,108,800 735,700 18,000	400 1,335,900 774,100 40,800
Total Operating Expenses	1,875,600	2,151,200
Operating Loss	(1,514,600)	(2,087,800)
Non-Operating Income, Net	99,100	152,200
Net Loss	\$ (1,415,500)	\$(1,935,600)
Net Loss Per Common Share	\$ (0.05)	\$ (0.08)
Weighted Average Common Shares Outstanding	26,774,000	24,948,400

See accompanying notes to condensed consolidated financial statements

	March 31, 1998	March 31, 1997
Cash Flows from Operating Activities: Net Loss Adjustments to Reconcile Net Loss to Net Cash Used by Operating Activities:	\$(1,415,500)	\$(1,935,600)
Depreciation and Amortization	86,400	88,400
Write-off of Capitalized Patent Costs Net Change in Current Assets and Current Liabilities	(532,800)	51,100 (288,700)
Net Cash Used by Operating Activities	(1,861,900)	(2,084,800)
Cash Flows from Investing Activities: Acquisition of Property and Equipment Other Noncurrent Assets	(27,800) (91,600)	(47,100) (22,800)
Decrease in Restricted Cash Sale of Investment in Common Stock of Endogen, Inc.	25,000 	1,802,700
Net Cash Provided (Used) by Investing Activities	(94,400)	1,732,800
Cash Flows from Financing Activities: Proceeds from Issuance of Common Stock	3,701,900	2,500
Net Cash Provided by Financing Activities	3,701,900	2,500
Increase (Decrease) in Cash and Cash Equivalents	1,745,600	(349,500)
Cash and Cash Equivalents at Beginning of Period	6,436,300	12,591,800
Cash and Cash Equivalents at End of Period	\$ 8,181,900	\$12,242,300

See accompanying notes to condensed consolidated financial statements

T CELL SCIENCES, INC. Notes to Condensed Consolidated Financial Statements March 31, 1998

(1) Nature of Business

T Cell Sciences, Inc. (the "Company") is a biopharmaceutical company engaged in the discovery and development of innovative drugs using novel applications of immunology to prevent and treat cardiovascular, pulmonary and immune disorders. The Company develops and commercializes products on a proprietary basis and in collaboration with pharmaceutical partners, including Novartis Pharma AG, Astra AB and Yamanouchi Pharmaceutical Co., Ltd.

The condensed consolidated financial statements include the accounts of T Cell Sciences, Inc. and its wholly owned subsidiary, T Cell Diagnostics, Inc. All intercompany transactions have been eliminated.

(2) Interim Financial Statements

The accompanying condensed consolidated financial statements for the three months ended March 31, 1998 and 1997 include the consolidated accounts of the Company, and have been prepared in accordance with generally accepted accounting principles and with the instructions to Form 10-Q and article 10 of Regulation S-X. In the opinion of management, the information contained herein reflects all adjustments, consisting solely of normal recurring adjustments, that are necessary to present fairly the financial positions at March 31, 1998 and December 31, 1997, the results of operations for the quarters ended March 31, 1998 and 1997, and the cash flows for the three months ended March 31, 1998 and 1997. The results of operations for the quarter ended March 31, 1998 are not necessarily indicative of results for any future interim period or for the full year.

Certain information and footnote disclosures normally included in financial statements prepared in accordance with generally accepted accounting principles have been omitted, although the Company believes that the disclosures included are adequate to make the information presented not misleading. The condensed consolidated financial statements and the notes included herein should be read in conjunction with footnotes contained in the Company's Annual Report on Form 10-K for the year ended December 31, 1997.

(3) Issuance of Common Stock

In March 1998, the Company completed a private placement of approximately 2,043,000 shares of common stock to institutional investors at a price of \$1.90 per share. Net proceeds from the common stock issuance totaled approximately \$3,699,900. The Company believes that its current cash and cash equivalents, which includes the private placement proceeds, together with cash flows from existing SBIR grants and collaborations, and interest income on invested funds will be sufficient to meet working capital requirements and fund operations into 1999. The working capital requirements will depend on several factors including, but not limited to, the progress and costs associated with research and development programs, preclinical and clinical studies, and the timing and scope of collaborative arrangements.

(4) Statement of Financial Accounting Standards Nos. 130 and 131

In June 1997, the Financial Accounting Standards Board issued Statement of Accounting Standards Nos. 130, "Reporting Comprehensive Income" ("SFAS 130"), and 131, "Disclosures about Segments of an Enterprise and Related Information" ("SFAS 131") to become effective for interim and annual periods beginning after December 15, 1997. The Company adopted SFAS 130 and SFAS 131 on January 1, 1998. SFAS 130 establishes standards for the reporting of comprehensive income and its components in the consolidated financial statements. To date the Company has not had material adjustments between net income as reported and comprehensive income as defined by SFAS 130. SFAS 131 establishes standards for the reporting of information on operating segments in interim and annual financial statements beginning with the annual financial statements for the year ending December 31, 1998.

Safe Harbor Statement under the Private Securities Litigation Reform Act of 1995: Statements contained in the following, Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations, that are not historical facts are forward-looking statements within the meaning of section 2xx of the Securities Exchange Act of 1934 that are subject to a variety of risks and uncertainties. There are a number of important factors that could cause the actual results to differ materially from those expressed in any forward-looking statements made by the Company. These factors include, but are not limited to: (i) the Company's ability to successfully complete product research and development, including pre-clinical and clinical studies, and commercialization; (ii) the Company's ability to obtain substantial additional funding; (iii) the Company's ability to obtain required governmental approvals; (iv) the Company's ability to attract manufacturing, sales, distribution and marketing partners and other strategic alliances; and (v) the Company's ability to develop and commercialize its products before its competitors.

Item 2. Management's Discussion And Analysis Of Financial Condition And Results
Of Operations

Overview

T Cell Sciences is a biopharmaceutical company engaged in the discovery and development of innovative drugs using novel applications of immunology to prevent and treat cardiovascular, pulmonary and immune disorders. The Company's technology platforms are based on its understanding of the ways in which the body triggers its natural defense mechanisms. Product development efforts focus on three therapeutic programs: (i) developing compounds that inhibit inappropriate complement activation, which is part of the body's immune defense system; (ii) discovery and development of T cell activation inhibitors for the prevention of transplant rejection and treatment of autoimmune diseases; and (iii) development of a therapeutic vaccine for the management of atherosclerosis.

The Company concluded clinical trials in November 1997 for its lead complement inhibitor, TP10, in patients undergoing lung transplantation. Final results released in April 1998 showed that TP10 therapy appears safe and well tolerated and demonstrated significant efficacy. The Company's collaborative partner, Astra AB, completed a Phase I clinical trial for ATM027 in patients with multiple sclerosis. ATM027 is one of the products derived from the Company's T cell antigen receptor (TCAR) program. The Phase I data showed that ATM027 had an effect on the target cells and there have been no serious adverse effects in the study to date. In January the Company was awarded its fourth Small Business Innovation Research (SBIR) grant from the National Institutes of Health. The grant, in the amount of \$96,000, will support the development of a peptide vaccine to prevent or treat atherosclerosis. In March 1998, the Company completed a private placement of approximately 2,043,000 shares of common stock to institutional investors at a price of \$1.90 per share. Net proceeds from the common stock issuance totaled approximately \$3,699,900.

Results of Operations

The Company reported a consolidated net loss of \$1,415,500, or \$.05 per share, for the first quarter ended March 31, 1998, a decrease of \$520,000, or 26.9%, compared with a net loss of \$1,935,600, or \$.08 per share, for the first quarter ended March 31, 1997. The decrease in net loss for the first quarter of 1998 compared to the first quarter of 1997 is due to a increase in operating revenue of \$297,600, or 469.4%, to \$361,000 in 1998 from \$63,400 in 1997 combined with a decrease in operating expense of \$275,600, or 12.8%, to \$1,875,600 in 1998 from \$2,151,200 in 1997. The Company reported a net operating loss of \$1,514,600 for the first quarter of 1998 a decrease of \$573,200, or 27.5%, compared with a net operating loss of \$2,087,800 for the first quarter of 1997. The decrease in net operating loss was partially offset by a decrease in non-operating income of \$53,100 to \$99,100 in 1998 compared to \$152,200 in 1997.

Total operating revenue increased \$297,600 to \$361,000 for the first quarter of 1998 compared to \$63,400 for the first quarter of 1997. The increase in operating revenue for the quarter was primarily due

to the option payment received from Novartis Pharma AG in November 1997 which is being recognized over the option term. Product sales revenue was \$27,400 for the quarter compared to \$1,300 for the same period last year.

Total operating expense decreased \$275,600, or 12.8%, for the first quarter of 1998 compared to \$2,151,200 for the first quarter of 1997. The decrease in operating expense is primarily due to a \$227,100, or 17.0%, decrease in research and development expense to \$1,108,800 in 1998 compared to \$1,335,900 in 1997. The decrease in research and development expense is primarily due to lower costs associated with clinical trials. During the first quarter of 1997, there was an ongoing Phase IIa clinical trial for TP10 in patients with ARDS and a Phase I/II clinical trial for TP10 in patients undergoing lung transplant. The Phase IIa clinical trial was completed in December 1997 and the Phase I/II clinical trial was completed in November 1997 following a six month safety review. There were no ongoing clinical trials during the first quarter of 1998.

Non-operating income decreased \$53,100, or 34.9%, to \$99,100 for the first quarter ended March 31, 1998 compared to non-operating income of \$152,200 for the first quarter ended March 31, 1997. The decrease in non-operating income is primarily the result of a \$71,000, or 41.7% decrease in interest income primarily due to lower cash balances for the first quarter of 1998 compared to the first quarter of 1997. Included in non-operating income for the first quarter of 1997 is a gain recognized from the conversion of the convertible note receivable from Endogen, Inc. and subsequent sale of the stock received.

Liquidity and Capital Resources

The Company ended the first quarter of 1998 with cash and cash equivalents of \$8,181,900 compared to \$6,436,300 at December 31, 1997. Net cash used in operations decreased 10.7% to \$1,861,900 for the first quarter of 1998 compared to \$2,084,800 for the first quarter of 1997. For the first quarter of 1998, net cash used in operations was offset by net proceeds of \$3,699,900 from the private placement of 2,043,000 shares of common stock. For the first quarter of 1997, net cash used in operations was partially offset by \$1,829,000 received from the conversion and subsequent sale of the convertible note receivable from Endogen, Inc.

In March 1998, the Company completed a private placement of approximately 2,043,000 shares of common stock to institutional investors at a price of \$1.90 per share. Net proceeds from the private placement totaled approximately \$3,699,900. The Company believes that its current cash and cash equivalents, which includes the private placement proceeds, together with cash flows from existing SBIR grants and collaborations, and interest income on invested funds will be sufficient to meet working capital requirements and fund operations into 1999. The working capital requirements will depend on several factors including, but not limited to, the progress and costs associated with research and development programs, preclinical and clinical studies, and the timing and scope of collaborative arrangements.

PART II. OTHER INFORMATION

Item 1. Legal Proceedings

No material changes since the Company's annual report of Form 10-K for the year ended December $31,\ 1997.$

Item 2. Changes in Securities

None

Item 3. Defaults Upon Senior Securities

None

Item 4. Submissions of Matters to a Vote of Security Holders

None

Item 5. Other Information

On May 12, 1998, the Company entered into a definitive merger agreement whereby it will acquire Virus Research Institute, Inc. (VRI). Under the terms of the merger agreement, which is subject to shareholder and regulatory approval, the Company will issue 1.55 shares of its common stock and 0.2 warrants for each share of VRI common stock. Each warrant represents the right to purchase one share of the Company's common stock for \$6.00 per share and will expire five years from the closing date. The number of shares of common stock outstanding on May 12, 1998 was 28,531,285 and 9,039,355 for T Cell Sciences and VRI, respectively.

Item 6. Exhibits and Reports on Form 8-K

A. Exhibits

None

B. Reports on Form 8-K

None

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.

T CELL SCIENCES, INC.

BY: /s/ Norman W. Gorin

Norman W. Gorin

Vice President, Finance and Chief Financial Officer

Dated: May 14, 1998

This schedule contains summary financial information extracted from the condensed financial statements of T Cell Sciences, Inc. for the Three Months ended March 31, 1998 and is qualified in its entirety by reference to such financial statements.

US Dollars

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3-M0S
          DEC-31-1998
              JAN-01-1998
               MAR-31-1998
                       1
                        8,181,900
                    18,600
                    (6,000)
                       7,300
             9,139,600
                        3,130,700
              (2,784,300)
              11,584,600
        2,232,700
                               0
                 0
                           0
                         28,500
                    8,573,400
11,584,600
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                361,000
                            13,100
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             (1,415,500)
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                        0
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                    (0.05)
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