SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 10-Q

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

For the quarterly period ended JULY 31, 1996

OR

/ / TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934
For the transition period from _____ to _____

Commission file number 0-17085

TECHNICLONE INTERNATIONAL CORPORATION
(Exact name of Registrant as specified in its charter)

CALIFORNIA	95-3698422
(State or other jurisdiction of incorporation or organization)	(I.R.S. Employer Identification Number)
14282 FRANKLIN AVENUE, TUSTIN, CALIFORNIA	92780
(Address of principal executive offices)	(Zip Code)

(714) 838-0500
(Registrant's telephone number, including area code)

NOT APPLICABLE
(Former name, former address and former fiscal year, if changed, since last report)

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

APPLICABLE ONLY TO ISSUERS INVOLVED IN BANKRUPTCY PROCEEDINGS DURING THE PRECEDING FIVE YEARS

Indicate by check mark whether the registrant has filed all documents and reports required to be filed by Section 12, 13 or 15(d) of the Securities Exchange Act of 1934 subsequent to the distribution of securities under a plan confirmed by a court. YES / / NO / /

APPLICABLE ONLY TO CORPORATE ISSUERS:

Indicate the number of shares outstanding of each of the issuer's classes of common stock, as of the latest practicable date.

21,042,409 shares of Common Stock
as of August 15, 1996

ITEM 1 - FINANCIAL STATEMENTS

The following financial statements required to be provided by this Item 1 and Rule 10.01 of Regulation S-X are filed herewith, at the respective pages indicated on this Quarterly Report, Form 10-Q:

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

This Quarterly Report on Form 10-Q includes certain forward-looking statements, the realization of which may be impacted by certain important factors discussed in "Additional Factors that May Affect Future Results".

The financial statements set forth in this Form 10-Q have been prepared on a going concern basis, which contemplates the realization of assets and the satisfaction of liabilities in the normal course of business. As shown in the financial statements set forth in this Form 10-Q, the Company experienced losses in fiscal 1995 and 1994 and has an accumulated deficit at July 31, 1996. Management has restructured certain of its license agreements to provide it with greater control over the development and clinical trials of its antibodies. If the Company is able to achieve certain goals in relation to these antibodies, it will receive certain additional financing pursuant to the terms of an existing license agreement. Historically, the Company has relied on third party and investor funds to fund its operations and clinical trials, and management expects that additional funds will be required in the future. There can be no assurances that this funding will be received. If the Company does not receive additional funding, it will be forced to scale back operations and it could have a material adverse effect on the Company. The Company's continuation as a going concern is dependent on its ability to generate sufficient cash flow to meet its obligations on a timely basis, to obtain additional financing as may be required and, ultimately to attain successful operations. During the year ended April 30, 1996, the Company received significant funding through the issuance of preferred stock and a foreign distribution agreement which has resulted in cash balances of $6,986,750 as of July 31, 1996. Management believes that the cash and cash equivalents and short-term investments of $6,986,750 as of July 31, 1996 are sufficient to support the Company's estimated operations and other cash needs through at least July 31, 1997.

RESULTS OF OPERATIONS

The Company's net loss of $948,264 for the quarter ended July 31, 1996 represents an increase of $433,162 in comparison to the net loss of $515,102 for the prior year quarter ended July 31, 1995. This increase in the net loss for the quarter ended July 31, 1996 is primarily attributable to a $519,453 increase in total costs and expenses partially offset by a $86,291 increase in total revenues.

Revenues for the quarter ended July 31, 1996 increased $86,291, compared to the same prior year period ended July 31, 1995. This increase resulted from a $86,291 increase in interest income in comparison to the same prior year period ended July 31, 1995. Interest and other income increased during the current year as the level of idle cash funds available for investment has increased in comparison to the prior year. Management expects interest and other income to increase during the remainder of the current year in comparison to the prior year due to the increase in cash and short term investments from the closing of the issuance of the Class B Convertible Preferred Stock in December 1995 and from a distribution agreement consummated in February 1996. Management expects revenues from the sales and licensing of antibodies to increase slightly during the remainder of the fiscal year ending April 30, 1997 as the Company ships its LYM-1 antibody for use in the Phase III clinical trials.

The Company's total costs and expenses increased $519,453 during the quarter ended July 31, 1996, in comparison to the same prior year period ended July 31, 1995. This increase resulted from a $314,066 increase in research and development expenses, a $186,831 increase in general and administrative expenses, and an $18,556 increase in interest expense in comparison to the prior year quarter ended July 31, 1995. Research and development expenses increased $314,066 for the quarter ended July 31, 1996 in comparison to the same prior year period. The increase in research and development expenses resulted from the Company's activities during the current year in preparing for and starting the Phase III clinical trials of the LYM-1 antibody and from the Company's activities in preparing for Phase I clinical trials of the TNT antibody technologies which are expected to commence in early 1997. The Company expects significant research and development expenses in the future as clinical trial activities continue.

General and administrative expenses increased $186,831 for the quarter ended July 31, 1996 in comparison to the same period of the prior year. This increase in current year expenses has resulted primarily from the addition of administration personnel and from expanded public relations activities. Interest expense increased $18,556 during the quarter ended July 31, 1996 in comparison to the same period of the prior year due to a higher level of interest bearing debt outstanding during the current year as a result of a $1,020,000 real estate mortgage loan originated in April 1996 in connection with the Company's purchase of its existing facility. The Company believes that general and administrative costs will increase during the remainder of the current fiscal year due to the continued addition of personnel, the expansion of public relations activities, and the legal and printing costs associated with the Company's September 27, 1996 shareholders' meeting.

The Company has begun Phase III testing in multi-center clinical trials of the LYM-1 antibody in late stage non-Hodgkins lymphoma patients. The clinical trials are being sponsored by Alpha Therapeutic Corporation, a wholly owned subsidiary of Green Cross of Japan. The clinical trials are being held at participating medical centers including M.D. Anderson, The Cleveland Clinic, Cornell University (N.Y.C.), George Washington University and University of Cincinnati. Following the completion of the clinical trials the Company expects to file an application with the FDA to market LYM-1 in the United States.

On February 5, 1996,the Company entered into an agreement with Cambridge Antibody Technology, Ltd. ("CAT") to develop and market a new class of products for cancer therapy and diagnosis. The Agreement provides that the Company and CAT will develop a monoclonal antibody based upon CAT's patented technology for producing fully human monoclonal antibodies and the Company's Tumor Necrosis Technologies ("TNT"). The Agreement provides that equity in the joint venture and costs associated with the development of the product would be shared equally between the Company and CAT. The Company would retain exclusive world-wide manufacturing rights. It is anticipated that the joint venture would conduct clinical trials of TNT in both the United States and Europe.

On February 29, 1996 the Company entered into a Distribution Agreement with Biotechnology Development, Ltd. ("BTD"), a limited partnership controlled by a member of the Board of Directors of the Company and a major shareholder of the Company. Pursuant to the Distribution Agreement, BTD acquired the marketing rights for the LYM-1 antibody technology for certain European countries and other geographic areas not covered by the Company's existing license agreement with Alpha Therapeutic Corporation. BTD paid the Company $3,000,000 for these marketing rights. Under the terms of the Distribution Agreement, the Company retains all manufacturing rights to LYM-1 and will supply LYM-1 to BTD at preset prices. Additionally, the Company has the option under an Option Agreement to repurchase the marketing rights to LYM-1 for a thirty month period. The repurchase price, if repurchase is elected by the Company at its sole discretion, includes a combination of cash, stock options and royalty payments to be made to BTD, the amount of which depends on when the repurchase option is elected by the Company.

LIQUIDITY AND CAPITAL RESOURCES

At July 31, 1996, the Company had $6,986,750 in cash and short term investments and working capital of $6,388,541 compared to $8,078,201 in cash and short term investments and working capital of $7,460,514 at April 30, 1996.

CAPITAL COMMITMENTS

During the remainder of the year ending April 30, 1997 the Company expects to acquire significant additional assets including an additional building and additional building improvements, furniture, fixtures and equipment to expand operations.

As of July 31, 1996, the Company had commitments to spend approximately $600,000 on building improvements, furniture, and fixtures in connection with the construction of office facilities in the unimproved portion of the existing building which was purchased in April 1996. Also, as of July 31, 1996, the Company had commitments to spend approximately $200,000 on additional laboratory and production equipment to expand antibody production capabilities.

On September 11, 1996, the Company entered into a Purchase Agreement for Real Property and Escrow Instructions dated as of September 11, 1996 with TR KOLL TUSTIN TECH CORP. ("Koll"). Under the terms of the agreement, the Company agreed to purchase from Koll land and a building located at 14272 Franklin Avenue, Tustin, California 92780 for the purchase price of $1,524,663. This 24,201 square foot building is adjacent to the Company's existing building which was purchased in April 1996. The building to be purchased is partially occupied by tenants under the terms of leasing arrangements and the balance of the building, consisting of approximately 5,000 square feet, will be occupied by the Company over the next six months to support expanded in-house research and development operations. The Company retains the option to terminate the tenant leases at the end of a two year period if needed to expand its antibody production capabilities. The Company expects to make a down payment of approximately $500,000 towards the building purchase

and to finance the balance under the terms of a mortgage loan currently being negotiated. The closing of the transaction is expected to take place on or before October 15, 1996.

ADDITIONAL FACTORS THAT MAY AFFECT FUTURE RESULTS

FUTURE OPERATING RESULTS. Future operating results may be impacted by a number of factors that could cause actual results to differ materially from those stated herein. These factors include worldwide economic and political conditions, industry specific factors, the Company's ability to maintain access to external financing sources and its financial liquidity, the Company's ability to timely develop and produce commercially viable products at competitive prices, the availability and cost of components of those products, and the Company's ability to manage expense levels.

NEED FOR ADDITIONAL CAPITAL. At July 31, 1996, the Company had $6,986,750 in cash and short term investments which management believes is sufficient to support the Company's estimated operations and other cash needs through at least July 31, 1997. As of July 31, 1996, the Company had significant commitments for expenditures for building improvements, equipment, furniture and fixtures. The Company has continued to experience negative cash flows since its inception and expects the negative cash flow to continue for the foreseeable future. The Company expects that the monthly negative cash flow will increase as a result of increased activities with the Phase III clinical trials for LYM-1 and the significantly increased research and development with the Company's other products, including Tumor Necrosis Therapy ("TNT"). As a result of the increased expenditure of funds, the Company believes that it will be necessary for the Company to raise additional capital to sustain research and development and provide for future clinical trials. The Company must raise additional equity funds in order to continue its operations until it is able to generate sufficient additional revenue from the sale and licensing of its products. There can be no assurance that the Company will be successful in raising such funds on terms acceptable to it or at all, or that sufficient additional capital will be raised to research and develop the Company's additional products. The Company is discussing the possibility of raising additional funds with several investment banking firms, but as of July 31, 1996, the Company had not entered into any firm commitments for additional funds. If the initial results from the Phase III clinical trials of LYM-1 are poor, the results may have a material adverse effect upon the Company's ability to raise additional capital, which would affect the Company's ability to continue a full-scale research and development effort for its antibody technologies. The Company's future success is highly dependent upon its continued access to sources of financing which it believes are necessary for the continued growth of the Company. In the event the Company is unable to maintain access to its existing financing sources, or obtain other sources of financing there would be a material adverse effect on the Company's business, financial position and results of operations.

COMPETITION. The biotechnology industry is intensely competitive and changing rapidly. Substantially all of the Company's existing competitors have larger technical staffs, more established and larger research budgets and significantly greater financial resources than the Company. There can be no assurance that these competitors will not be able to expend resources to develop their products prior to the Company's product being granted approval for marketing by the U.S. Food and Drug Administration. There can be no assurance that the Company will be able to compete successfully or that competition will not have a material adverse effect on the Company's results of operation.

TECHNOLOGY. The Company's future success will depend significantly upon its ability to develop and test workable products which the Company will seek FDA approval to market to certain defined groups. A significant risk remains as to the technological, performance and commercial success of the Company's technology and products. The products currently under development by the Company will require significant additional laboratory and clinical testing and investment over the

foreseeable future. The significant research, development, and testing activities, together with resultant increases in associated expenses, are expected to result in operating losses for the foreseeable future. Although the Company is optimistic that it will be able to successfully complete development of one or more of its products, there can be no assurance that the Company's research and development activities will be successfully completed; that any proposed products will prove to be effective in clinical trials; that the Company will be able to obtain all necessary governmental clearances and approvals to market its products; that such proposed products will prove to be commercially viable or successfully marketed; or that the Company will ever achieve significant revenues or profitable operations. In addition, the Company may encounter unanticipated problems, including development, manufacturing, distribution and marketing difficulties. The failure to adequately address such difficulties could have a material adverse effect on the Company's prospects.

REGULATION. The Company's products are subject to extensive government regulation in the United States by federal, state and local agencies including the Food and Drug Administration. The process of obtaining and maintaining FDA and other required regulatory approvals for the Company's products is lengthy, expensive and uncertain. There can be no assurance that the Company can obtain FDA or other regulatory approval for the marketing of its products or that changes in existing regulations or the adoption of new regulations will not occur which will adversely affect the Company.

EARTHQUAKE RISKS. The Company's corporate headquarters facility, at which the majority of its research and development activities are conducted, is located near major earthquake faults which have experienced earthquakes in the past. The Company does not carry earthquake insurance on its facility due to its prohibitive cost. In the event of a major earthquake or other disaster affecting the Company's facilities, the operations and operating results of the Company could be adversely affected.

STOCK PRICE FLUCTUATIONS AND LIMITED TRADING VOLUME. The Company's participation in the highly competitive biotechnology industry often results in significant volatility in the Company's common stock price. Also, at times there is a limited trading volume in the Company's stock. This volatility in the stock price and limited trading volume are significant risks investors should consider.

FORWARD LOOKING STATEMENTS. This Quarterly Report on Form 10-Q contains certain forward-looking statements that are based on current expectations. In light of the important factors that can materially affect results, including those set forth above and elsewhere in this Form 10-Q, the inclusion of forward-looking information herein should not be regarded as a representation by the Company or any other person that the objectives or plans of the Company will be achieved. The Company may encounter competitive, technological, financial and business challenges making it more difficult than expected to continue to develop, market and manufacture its products; competitive conditions within the industry may change adversely; upon development of the Company's products, demand for the Company's products may weaken; the market may not accept the Company's products; the Company may be unable to retain existing key management personnel; the Company's forecasts may not accurately anticipate market demand; and there may be other material adverse changes in the Company's operations or business. Certain important factors affecting the forward looking statements made herein include, but are not limited to (i) accurately forecasting capital expenditures, and (ii) obtaining new sources of external financing prior to the expiration of existing support arrangements or capital. Assumptions relating to budgeting, marketing, product development and other management decisions are subjective in many respects and thus susceptible to interpretations and periodic revisions based on actual experience and business developments, the impact of which may cause the Company to alter its capital expenditure or other budgets, which may in turn affect the Company's financial position and results of operations.

PART II

Item 1. Legal Proceedings. None.

Item 2. Changes in Securities. None.

Item 3. Defaults Upon Senior Securities. None.

Item 4. Submission of Matters to a Vote of Security Holders. None.

Item 5. Other Information. None.

Item 6. Exhibits and Report on Form 8-K.

(a) Exhibits:

Exhibit Number	Description
27	Financial Data Schedule

(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.

TECHNICLONE INTERNATIONAL CORPORATION

By: /ss/ Lon H. Stone

By: /ss/ William V. Moding

Dated: September 13, 1996

TECHNICLONE INTERNATIONAL CORPORATION

BALANCE SHEETS

	April 30, 1996	July 31, 1996 (Unaudited)
ASSETS		
CURRENT ASSETS:		
Cash and cash equivalents	$ 4,179,313	$ 6,986,750
Short-term investments	3,898,888	--
Accounts receivable, net	95,146	19,578
Inventories, net	93,921	180,880
Prepaid expenses and other current assets	17,294	9,194
Total current assets	8,284,562	7,196,402
PROPERTY:		
Land	525,255	525,255
Building and improvements	1,298,416	1,391,472
Laboratory equipment	1,139,663	1,218,433
Furniture and fixtures	78,155	95,759
Total	3,041,489	3,230,919
Less accumulated depreciation and amortization	(722,436)	(784,395)
Property--net	2,319,053	2,446,524
OTHER ASSETS:		
Patents, net	166,585	164,933
Other	5,557	5,557
Total other assets	172,142	170,490
TOTAL	$10,775,757	$ 9,813,416

See accompanying notes to financial statements.

TECHNICLONE INTERNATIONAL CORPORATION

BALANCE SHEETS

	April 30, 1996	July 31, 1996
		(Unaudited)
LIABILITIES AND STOCKHOLDERS' EQUITY:		
CURRENT LIABILITIES:		
Accounts payable	$ 230,144	$ 212,996
Accrued legal and accounting fees (primarily to a related party)	99,495	89,256
Accrued payroll and related costs	88,791	66,423
Accrued license termination fee	100,000	100,000
Accrued royalties	61,667	81,667
Accrued interest	--	8,303
Reserve for contract losses	173,563	173,563
Current portion of long-term debt	32,968	33,758
Other current liabilities	37,420	41,895
Total current liabilities	824,048	807,861
LONG TERM DEBT - MORTGAGE LOAN	987,032	981,142
COMMITMENTS		
STOCKHOLDERS' EQUITY:		
Preferred Stock--$1.00 par value (authorized, 100,000 shares; Class B Convertible Preferred Stock, outstanding, 6,800 shares at April 30, 1996 and 4,150 shares at July 31, 1996) (liquidation preference of $4,393,315 at July 31, 1996)	6,800	4,150
Common Stock--no par value (authorized, 30,000,000 shares; outstanding, 20,048,014 shares at April 30, 1996 and 20,955,762 shares at July 31, 1996)	21,133,968	23,448,608
Additional paid-in capital	6,061,171	3,757,181
Accumulated deficit	(17,760,680)	(18,708,944)
Total	9,441,259	8,500,995
Less notes receivable from sale of common stock	(476,582)	(476,582)
Net stockholders' equity	8,964,677	8,024,413
TOTAL	$ 10,775,757	$ 9,813,416

See accompanying notes to financial statements.

TECHNICLONE INTERNATIONAL CORPORATION

STATEMENTS OF OPERATIONS

	THREE MONTHS ENDED	
	JULY 31, 1995	JULY 31, 1996
	(Unaudited)	(Unaudited)
REVENUES:		
Net sales	$ --	$ --
Licensing fees	--	
Interest income	11	86,302
Total revenues	11	86,302
COSTS AND EXPENSES:		
Cost of sales	--	--
Research and development	264,056	578,122
General and administrative:		
Unrelated entities	213,352	376,215
Affiliates	31,287	55,255
Interest	6,418	24,974
Total costs and expenses	515,113	1,034,566
NET LOSS	$ (515,102)	$ (948,264)
WEIGHTED AVERAGE SHARES OUTSTANDING	16,930,811	20,686,817
LOSS PER COMMON SHARE	$ (.03)	$ (.05)

See accompanying notes to financial statements.

TECHNICLONE INTERNATIONAL CORPORATION

STATEMENT OF STOCKHOLDERS' EQUITY

	PREFERRED STOCK		COMMON STOCK		ADDITIONAL PAID-IN	ACCUMU-LATED	NOTES RECEIVABLE FROM SALE OF	
	SHARES	AMOUNT	SHARES	AMOUNT	CAPITAL	DEFICIT	STOCK	TOTAL
BALANCE AT April 30, 1996	6,800	$ 6,800	20,048,014	$21,133,968	$ 6,061,171	$(17,760,680)	$(476,582)	$8,964,677
Common stock issued upon exercise of stock options (unaudited)			8,000	8,000				8,000
Common Stock issued upon conversion of Class B Convertible Preferred Stock (unaudited)	(2,650)	(2,650)	899,748	2,306,640	(2,303,990)			--
Net loss (unaudited)						(948,264)		(948,264)
BALANCE AT JULY 31,1996 (unaudited)	4,150	$ 4,150	20,955,762	$23,448,608	$ 3,757,181	$(18,708,944)	$(476,582)	$ 8,024,413

See accompanying notes to financial statements.

TECHNICLONE INTERNATIONAL CORPORATION

STATEMENTS OF CASH FLOWS

	THREE MONTHS ENDED JULY 31, 1995 (Unaudited)	JULY 31, 1996 (Unaudited)
CASH FLOWS FROM OPERATING ACTIVITIES:		
Net loss	$(515,102)	$ (948,264)
Adjustments to reconcile net loss to net cash used by operating activities:		
Depreciation and amortization	47,514	69,786
Common stock issued for services	57,100	--
Changes in operating assets and liabilities:		
Decrease in accounts receivable	2,378	75,568
(Increase) in inventory	(2,475)	(86,959)
Decrease in prepaid expenses	--	8,100
Increase (Decrease) in accounts payable	82,515	(17,148)
Increase in accrued and other current liabilities	95,769	171
Net cash used by operating activities	(232,301)	(898,746)
CASH FLOWS FROM INVESTING ACTIVITIES:		
Proceeds from sale of short-term investments	--	3,898,888
Property acquisitions	(18,434)	(189,430)
Patent costs capitalized	(35,275)	(6,175)
Net cash (used) provided by investing activities	(53,709)	3,703,283

[Continued on next page]

See accompanying notes to financial statements.

TECHNICLONE INTERNATIONAL CORPORATION

STATEMENTS OF CASH FLOWS

[Continued from previous page]

	THREE MONTHS ENDED	
	JULY 31, 1995	JULY 31, 1996
	(Unaudited)	(Unaudited)
CASH FLOWS FROM FINANCING ACTIVITIES:		
Principal payments on short- and long-term borrowings	$ --	$ (5,100)
Proceeds from sale of common stock	298,150	8,000
Net cash provided by financing activities	298,150	2,900
INCREASE IN CASH	12,140	2,807,437
CASH AT BEGINNING OF PERIOD	35,642	4,179,313
CASH AT END OF PERIOD	$ 47,782	$6,986,750
SUPPLEMENTAL INFORMATION:		
Interest paid	$ 1,248	$ 16,671
Income taxes paid	$ 800	$ 800

See accompanying notes to financial statements.

TECHNICLONE INTERNATIONAL CORPORATION

NOTES TO FINANCIAL STATEMENTS

(1) The accompanying unaudited financial statements contain all adjustments (consisting of only normal recurring adjustments) which, in the opinion of management, are necessary to present fairly the financial position of the Company at July 31, 1996, and the results of its operations and its cash flows for the three month periods ended July 31, 1996 and 1995. Certain information and footnote disclosures normally included in the financial statements have been condensed or omitted pursuant to rules and regulations of the Securities and Exchange Commission although the Company believes that the disclosures in the financial statements are adequate to make the information presented not misleading. The financial statements included herein should be read in conjunction with the financial statements of the Company, included in the Company's Annual Report on Form 10-K for the year ended April 30, 1996, filed with the Securities and Exchange Commission on July 26, 1996.

(2) Going Concern -- The accompanying financial statements have been prepared on a going concern basis, which contemplates the realization of assets and the satisfaction of liabilities in the normal course of business. As shown in the financial statements, the Company experienced losses in fiscal 1995 and 1994 and has an accumulated deficit at July 31, 1996. Management has restructured certain of its license agreements to provide it with greater control over the development and clinical trials of its antibodies. If the Company is able to achieve certain goals in relation to these antibodies, it will receive certain additional financing pursuant to the terms of an existing license agreement. Historically, the Company has relied on third party and investor funds to fund its operations and clinical trials, and management expects that additional funds will be required in the future. There can be no assurances that this funding will be received. If the Company does not receive additional funding, it will be forced to scale back operations and it could have a material adverse effect on the Company. The Company's continuation as a going concern is dependent on its ability to generate sufficient cash flow to meet its obligations on a timely basis, to obtain additional financing as may be required and, ultimately to attain successful operations. During the year ended April 30, 1996, the Company received significant funding through the issuance of preferred stock and a foreign distribution agreement which has resulted in cash balances of $6,986,750 as of July 31, 1996. Management believes that the cash and cash equivalents and short-term investments of $6,986,750 as of July 31, 1996 are sufficient to support the Company's estimated operations and other cash needs through at least July 31, 1997.

(3) Results of operations for the interim periods covered by this Report may not necessarily be indicative of results of operations for the full fiscal year.

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THIS SCHEDULE CONTAINS SUMMARY FINANCIAL INFORMATION EXTRACTED FROM FORM 10-K
FOR THE PERIOD ENDED 4/30/96 AND FORM 10-Q FOR THE PERIOD ENDED 07/31/96.

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TECHNICLONE INTERNATIONAL CORPORATION
1,000
U.S. DOLLARS

3-MOS	
APR-30-1997	
MAY-01-1996	
JUL-31-1996	
1,000	
	6,987
	0
	195
	175
	181
7,196	
	3,231
	784
	9,813
808	
	0
0	
	4
	23,449
	(15,429)
9,813	
	0
	86
	0
	1,035
	0
	0
	25
	(948)
	0
(948)	
	0
	0
	0
	(948)
	(.05)
	(.05)