SECURITIES AND EXCHANGE COMMISSION WASHINGTON, D. C. 20549

FORM 10-K

ANNUAL REPORT PURSUANT TO SECTION 13 OR 15(d) OF
THE SECURITIES EXCHANGE ACT OF 1934
FOR THE FISCAL YEAR ENDED OCTOBER 31, 1998 COMMISSION FILE NO. 1-8597

THE COOPER COMPANIES, INC. (Exact name of registrant as specified in its charter) $% \left(\frac{1}{2}\right) =\frac{1}{2}\left(\frac{1}{2}\right) +\frac{1}{2}\left(\frac{1}$

Delaware (State or other jurisdiction of incorporation) 94-2657368 (I.R.S. Employer Identification No.)

6140 Stoneridge Mall Road, Suite 590 Pleasanton, California (Address of principal executive offices) 94588 (Zip Code)

925-460-3600 (Registrant's telephone number, including area code)

SECURITIES REGISTERED PURSUANT TO SECTION 12(b) OF THE ACT:

Title of each class

Name of each exchange on which registered

Common Stock, \$.10 Par Value, and associated Rights

New York Stock Exchange Pacific Exchange

SECURITIES REGISTERED PURSUANT TO SECTION 12(g) OF THE ACT: None

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, and (2) has been subject to such filing requirements for the past 90 days. Yes [X] No $[\]$

Indicate by check mark if disclosure of delinquent filers pursuant to Item 405 of Regulation S-K is not contained herein, and will not be contained, to the best of registrant's knowledge, in definitive proxy or information statements incorporated by reference in Part III of this Form 10-K or any amendment to this Form 10-K. [X]

Aggregate market value of the voting stock held by non-affiliates of the registrant as of December 31, 1998: Common Stock, \$.10 Par Value -- \$297,773,840.

Number of shares outstanding of the registrant's common stock, as of December 31, 1998: 14,913,957.

DOCUMENTS INCORPORATED BY REFERENCE:

Document

Part of Form 10-K

Portions of the Annual Report to Stockholders for the fiscal year ended October 31, 1998 Portions of the Proxy Statement for the Annual Meeting of Stockholders to be held March 18, 1999

Parts I and II

Part III

ITEM 1. BUSINESS.

INTRODUCTION

The Cooper Companies, Inc. ("Cooper" or the "Company"), through its major subsidiaries, develops, manufactures and markets healthcare products, including hard and soft daily, flexible and extended wear contact lenses, and diagnostic products and surgical instruments and related products. In October 1998, the Company's Management and Board of Directors declared its Hospital Group of America, Inc. ("HGA") business a discontinued operation.

FORWARD-LOOKING STATEMENTS

Statements in this report that are not based on historical fact may be "forward-looking statements" as defined by the Private Securities Litigation Reform Act of 1995. They include words like "may," "will," "expect," "estimate," "anticipate," "continue" or similar terms and reflect Cooper's current analysis of existing trends. Actual results could differ materially from those indicated due to: major changes in business conditions and the economy, loss of key senior management, major disruptions in the operations of Cooper's manufacturing facilities, new competitors or technologies, significant disruptions caused by the failure of third parties to address the Year 2000 issue or by unforeseen delays in completing Cooper's Year 2000 compliance program, acquisition integration costs, foreign currency exchange exposure including the potential impact of the Euro, investments in research and development and other start-up projects, dilution to earnings per share from acquisitions or issuing stock, regulatory issues, significant environmental clean-up costs above those already accrued, litigation costs, costs of business divestitures, in the Company's SEC reports, including this section entitled "Business" in this Form 10-K and the related portions of the Company's 1998 Annual Report to Stockholders (the "1998 Annual Report") incorporated by reference herein, which 1998 Annual Report is included as Exhibit 13 to this Form 10-K.

GENERAL DESCRIPTION AND DEVELOPMENT OF BUSINESSES

The information required for this item is contained under the caption "Letter to Shareholders" in the 1998 Annual Report, which information is incorporated herein by reference.

RESEARCH AND DEVELOPMENT

Company-sponsored research and development expenditures during the fiscal year ended October 31, were \$1.9 million in 1998, \$1.7 million in 1997 and \$1.2 million in 1996. During fiscal 1998, CooperVision spent about 49% and CooperSurgical spent about 51% of the total. Cooper did not conduct any customer-sponsored research and development programs.

Cooper employs 21 people in its research and development and manufacturing engineering departments. Outside specialists in lens design formulation science, polymer chemistry, microbiology and biochemistry support product development and clinical research for CooperVision products. CooperSurgical conducts research and development in-house and also employs outside surgical specialists, including members of its surgical advisory board.

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GOVERNMENT REGULATION

The U.S. Food and Drug Administration ("FDA"), other federal agencies and foreign ministries of health regulate the development, testing, production and marketing of the Company's products. The Federal Food, Drug and Cosmetic Act and other statutes and regulations govern the testing, manufacturing, labeling, storage, advertising and promotion of such products. If applicable regulations are not followed, companies are subject to fines, product recall or seizure, suspension of production and criminal prosecution.

Cooper develops and markets medical devices under different levels of FDA regulation depending upon the classification of the device. Class III devices, such as flexible and extended wear contact lenses, require extensive premarket testing and approval, while Class I and II devices require substantially lower levels of regulation.

Before a new contact lens can be sold commercially, CooperVision ("CVI") must complete these steps: (1) compile data on its chemistry and toxicology, (2) determine its microbiological profile and (3) define the proposed manufacturing process. This data must be submitted to the FDA to support an application for an Investigational Device Exemption. Once this is granted, clinical trials can begin. These are subject to review and approval by an Institutional Review Board and, where a lens is determined to have a significant risk, the FDA. After the clinical trials are completed, a Premarket Approval Application must be submitted and approved by the FDA.

In connection with some of its new surgical products, CooperSurgical ("CSI") can submit an expedited procedure known as a 510(k) application for premarket notification to the FDA. Any product that can demonstrate that it is substantially equivalent to another device marketed before May 28, 1976 can use this procedure. If the new product is not substantially equivalent to a preexisting device or if the FDA rejected a claim of substantial equivalence, FDA approval to market would require extensive preclinical and clinical testing. This would increase the cost and would delay product marketing substantially.

FDA and state regulations also require the Company to adhere to applicable "good manufacturing practices" ("GMP"). They require detailed quality assurance and record keeping and periodic unscheduled regulatory inspections. The Company believes it is in substantial compliance with GMP regulations.

Health authorities in foreign countries regulate Cooper's human device clinical trials and medical device sales. The regulations vary widely from country to country. Even if the FDA has approved a product, the regulatory agencies in each country must approve new products before they are marketed.

These regulatory procedures require considerable resources and usually result in a substantial time lag between new product development and marketing. Cooper cannot assure that all necessary approvals will be obtained, or obtained in a timely manner. If the Company does not maintain compliance with regulatory standards or if problems occur after marketing, product approval may be withdrawn.

ISO 9000 CERTIFICATION AND CE MARK APPROVAL

In addition to the FDA regulatory requirements, the Company also maintains ISO 9000 certification and CE Mark approvals for all lens products. These quality programs and approvals are required by the European Medical Device Directive and must be maintained for all products intended to be sold in the European market. In order to maintain these prestigious quality benchmarks, the Company is subjected to rigorous biannual reassessment audits of their quality systems and procedures by globally recognized notified bodies and agencies.

RAW MATERIALS

In general, CVI's raw materials consist of various polymers and packaging materials. There are alternative supply sources of all of these materials. Raw materials used by CSI or its suppliers are generally available from more than one source. However, because some products require specialized manufacturing procedures, CSI could experience inventory shortages if it needed an alternative manufacturer on short notice.

MARKETING AND DISTRIBUTION

In the United States, Canada and certain European countries, CVI markets its products through its field sales representatives, who call on ophthalmologists, optometrists, opticians and optical chains. In the United States, field sales representatives also call on distributors. In certain other European counties, CVI uses distributors and has given them the exclusive right to market our products.

CSI's products are marketed worldwide by a network of field sales representatives and distributors. In the United States, CSI also uses telemarketing, direct mail, advertising in professional journals and a direct mail catalog.

PATENTS, TRADEMARKS AND LICENSING AGREEMENTS

Cooper owns or licenses a variety of domestic and foreign patents which, in total, are material to its businesses. The names of certain of Cooper's products are protected by trademark registrations in the United States Patent and Trademark Office and, in some cases, also in foreign trademark offices. Applications are pending for additional trademark registrations. These trademarks are valuable as they contribute to the brand identity of Cooper's products. Cooper aggressively enforces and defends its patents and other proprietary technology.

DEPENDENCE ON CUSTOMERS

Cooper's business does not materially depend on any one customer or any one affiliated group of customers.

GOVERNMENT CONTRACTS

Cooper's business is not materially subject to profit renegotiation or termination of contracts or subcontracts at the election of the United States government.

COMPETITION

Each of Cooper's businesses operates in a highly competitive environment. Competition in the healthcare industry revolves around the search for technological and therapeutic innovations in the prevention, diagnosis and treatment of illness or disease. Cooper competes primarily on the basis of product quality, program differentiation, technological benefit, service and reliability.

Many companies develop and manufacture contact lenses. CVI competes primarily on its product quality, service and reputation among medical professionals and by participating in specialty niche markets.

It sponsors clinical studies to generate medical information to improve its lenses. Major competitors have greater financial resources and larger research and development and sales forces than CVI. Many of these competitors offer a greater range of contact lenses and a variety of other eyecare products, including lens care products and ophthalmic pharmaceuticals, which may give them a competitive advantage in marketing their lenses to high volume contract accounts.

In the surgical segment, competitive factors include technological and scientific advances, product quality, price and effective communication of product information to physicians and hospitals. CSI believes that it benefits, in part, from the technological advantages of certain of its products and from developing new medical procedures that can create new markets for equipment and instruments. CSI competes by focusing on distinct niche markets and by supplying these with high quality equipment, instruments and disposable products. For certain procedures, medical practitioners can obtain all of the equipment, instruments and disposable products from CSI. As CSI develops products for new medical procedures, it offers to train medical professionals to perform them. CSI competes with a number of manufacturers in each of its niche markets, including larger manufacturers with greater financial and personnel resources who sell a substantially larger number of product lines.

BACKLOG

Backlog is not a material factor in Cooper's businesses.

SEASONALITY

CVI's contact lens sales in the first fiscal quarter are generally lower than subsequent quarters as fewer patients visit practitioners during the holiday season.

COMPLIANCE WITH ENVIRONMENTAL LAWS

Federal, state and local provisions that regulate the discharge of materials into the environment, or relate to the protecting of the environment, do not currently materially effect Cooper's capital expenditures, earnings or competitive position. See "Environmental" in Note 11 of Notes to Consolidated Financial Statements of the Company included in the 1998 Annual Report, regarding certain anticipated remediation costs, which information is incorporated herein by reference.

WORKING CAPITAL

Cooper's businesses have not required any material working capital arrangements in the past five years.

FINANCIAL INFORMATION ABOUT BUSINESS SEGMENTS, GEOGRAPHIC AREAS, FOREIGN OPERATIONS AND EXPORT SALES

The information required for this item is included in Note 12 "Business Segment Information" of Notes to Consolidated Financial Statements of the Company included in the 1998 Annual Report, which information is incorporated herein by reference.

EMPLOYEES

On October 31, 1998, Cooper and its continuing operations employed approximately 1,900 persons. The Company believes that its relations with its employees are good.

ITEM 2. PROPERTIES.

The following are the principal facilities of Cooper's continuing operations as of October 31, 1998:

	LOCATION	OPERATIONS	APPROXIMATE FLOOR AREA (SQ. FT.)	OWNED OR LEASED	LEASE EXPIRATION
United S	States				
	Pleasanton, CA Irvine, CA	Executive Offices Executive Offices, CVI Offices, Distribution	13,700	Leased	Sept. 2000
	Huntington Beach, CA	and Customer Service CVI Manufacturing &	15,400	Leased	Jan. 2000
	,	Technical Offices CVI Administrative	16,500	Leased	March 2002
	Fairport, NY Scottsville, NY	Offices & Marketing CVI Manufacturing,	20,100	Leased	April 2002
	Shelton, CT	Distribution and Warehouse Facilities CSI Manufacturing, Research and Development, Marketing,	49,500	Owned	N/A
		Distribution and Warehouse Facilities	35,000	Leased	Dec. 2002
Canada	Markham, Ont.	CVI Offices, Manufacturing Distribution and Warehouse Facilities	21,000	Leased	Feb. 2000
United I	Kingdom Hamble, Hampshire, England	Aspect Manufacturing, Research and Development, Marketing and Admin.			
	Faraham Hampahira	Offices Distribution and Customer	93,800	Owned	N/A
	Fareham, Hampshire, England	Service	30,800	Leased	Jan. 2018
	Fareham, Hampshire, England	Manufacturing and Warehouse	27,100	Leased	June 2018

The Company believes its properties are suitable and adequate for its businesses. $% \begin{center} \end{center} \begin{center} \end{center}$

ITEM 3. LEGAL PROCEEDINGS.

The information required for this item is contained under the caption "Pending Litigation - GT Labs" in Note 11 of Notes to Consolidated Financial Statements of the Company included in the 1998 Annual Report, which information is incorporated herein by reference.

ITEM 4. SUBMISSION OF MATTERS TO A VOTE OF SECURITY HOLDERS.

PART II

ITEM 5. MARKET FOR REGISTRANT'S COMMON EQUITY AND RELATED STOCKHOLDER MATTERS.

The information required for this item is contained under the caption "Common Stock Price Range" in the 1998 Annual Report, which information is incorporated herein by reference.

ITEM 6. SELECTED FINANCIAL DATA.

The information required for this item is contained under the caption "Five Year Financial Highlights" in the 1998 Annual Report, which information is incorporated herein by reference.

ITEM 7. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS.

The information required for this item is contained under the caption "Management's Discussion and Analysis of Financial Condition and Results of Operations" in the 1998 Annual Report, which information is incorporated herein by reference.

ITEM 7A. QUANTITATIVE AND QUALITATIVE DISCLOSURE ABOUT MARKET RISK

The Company is primarily exposed to market risks that relate to changes in interest rates, foreign currency fluctuations and in the market value of its long-term debt obligations. The Company seeks to minimize its exposure to the impact of changing interest rates and foreign currency fluctuations by entering into interest rate swaps and foreign currency forward exchange contracts. The Company generally does not enter into derivative financial instrument transactions for speculative purposes. Additional information for this item is contained under the caption "Derivatives" in Note 1 "Summary of Significant Accounting Policies" and in Note 7 "Financial Instruments" in the 1998 Annual Report, which information is incorporated herein by reference.

LONG-TERM DEBT

The following table sets forth as of October 31, 1998, the Company's long-term debt obligations, principal cash flows by scheduled maturity, weighted average interest rates and estimated fair market value.

	Expected Maturity Date - Fiscal Year							
	1999	2000	2001	2002	2003	There- after	Total	Fair Value
(\$ in Millions) Long-term Debt								
Fixed interest rate(\$US) Average interest rate	\$ 4.0 8.00%	\$ - 8.00%	\$ - 8.00%	\$ - 8.00%	\$ 23.6 8.00%	\$ - -	\$ 27.6	\$ 27.6
Variable interest rate (\$US) Average interest rate	\$ 0.2 6.40%	\$ 0.2 6.51%	\$ 0.3 6.64%	\$ 22.1 6.74%	\$ 17.7 5.25%	\$ 1.6 4.99%	\$ 42.1	\$ 42.1

INTEREST RATE EXPOSURES

The Company enters into interest rate swap agreements to reduce the impact of changes in interest rates on its variable rate long-term debt obligations. The Company currently has two interest rate swap agreements on a total of \$20.5 million of its outstanding variable rate debt obligations. These instruments have the effect of converting variable rate instruments to fixed rate instruments. The interest rate swap agreements assure that the Company will pay 6.19% and 4.88% on the aforementioned \$20.5 million long-term debt obligations for the periods ending November 2002 (principal amount \$17.5 million) and January 2012 (principal amount \$3 million), respectively. The table below sets forth the notional amount and weighted average interest rates of each of the Company's interest rate swaps by maturity. The receive rate is based on October 31, 1998 rates, and projected based on the consumer price index. Notional amounts are used to calculate the contractual payments to be made under the contracts.

Notional	Amounts	Maturing	in	Fiscal	Year

	1999	2000	2001	2002	2003	There- after	Total	Fair Value
(\$ in Millions) Interest rate swaps Variable to fixed (\$US) Average pay rate	\$ - 6.19%	\$ - 6.19%	\$ - 6.19%	\$ 17.5 6.19%	\$ - -	\$ - -	\$ 17.5 6.19%	\$ 18.2
Average receive rate Variable to fixed (\$US) Average pay rate Average receive rate	5.93% \$ 0.2 4.88% 4.27%	6.03% \$ 0.2 4.88% 4.34%	6.14% \$ 0.3 4.88% 4.42%	6.24% \$ 0.3 4.88% 4.49%	- \$ 0.3 4.88% 4.57%	- \$ 1.7 4.88% 4.99%	6.03% \$ 3.0 4.88% 4.75%	\$ 3.2

FOREIGN CURRENCY EXPOSURES

The Company uses forward exchange contracts to minimize the effect of foreign currency fluctuations on its long-term debt obligations denominated in Great Britain Pounds ("GBP"), which debt was incurred to fund a portion of the Company's acquisition of Aspect Vision Care Ltd. (see caption "Aspect Acquisition" in Note 2 "Acquisitions" in the 1998 Annual Report, which information is incorporated herein by reference). The following table provides information on the Company's foreign currency forward exchange contracts. The information is provided in U.S. Dollar equivalent amounts, as presented in the Company's financial statements. The table presents the notional amounts at the contract exchange rates and the weighted average contractual foreign currency exchange rates by expected maturity dates.

Notional	Amounts	Maturing	in	Fiscal	Year

	nocional randards matering in risolar roa.							
	1999	2000	2001	2002	2003	There- after	Total	Fair Value
Foreign Contracts to Buy G.B.P.: Notional amount (in millions) Average contractual exchange rate	\$ 3.5 \$ 1.62	\$ 1.8 \$ 1.61	\$ 8.5 \$ 1.63	\$ 3.5 \$ 1.62	\$ 27.3 \$ 1.62	\$ -	\$ 44.6 \$ 1.62	\$ 43.8

ITEM 8. FINANCIAL STATEMENTS AND SUPPLEMENTARY DATA.

The information required for this item is included under the captions "Consolidated Balance Sheets," "Consolidated Statements of Income," "Consolidated Statements of Cash Flows," "Notes to Consolidated Financial Statements," "Independent Auditors' Report" and "Two Year Quarterly Financial Data" in the 1998 Annual Report, which information is incorporated herein by reference.

ITEM 9. CHANGES IN AND DISAGREEMENTS WITH ACCOUNTANTS ON ACCOUNTING AND FINANCIAL DISCLOSURE.

Not applicable.

PART III

ITEM 10. DIRECTORS AND EXECUTIVE OFFICERS OF THE REGISTRANT.

The information contained under the heading "Election of Directors" and "Executive Officers of the Company" in the Company's Proxy Statement for the Annual Meeting of Stockholders scheduled to be held on March 18, 1999 (the "1999 Proxy Statement") is incorporated herein by reference with respect to each of the Company's directors and the executive officers who are not also directors of the Company.

ITEM 11. EXECUTIVE COMPENSATION.

The information contained under the subheadings "Executive Compensation" and "Compensation of Directors" of the "Election of Directors" section of the 1999 Proxy Statement is incorporated herein by reference with respect to the Company's chief executive officer, the four other most highly compensated executive officers of the Company and the Company's directors.

ITEM 12. SECURITY OWNERSHIP OF CERTAIN BENEFICIAL OWNERS AND MANAGEMENT.

The information contained under the subheadings "Securities Held by Management" and "Principal Security Holders" of the "Election of Directors" section of the 1999 Proxy Statement is incorporated herein by reference with respect to certain beneficial owners, the directors and management.

ITEM 13. CERTAIN RELATIONSHIPS AND RELATED TRANSACTIONS.

The information required for this item is contained in the heading "Aspect Acquisition" in Note 2 "Acquisitions" in the 1998 Annual Report, which information is incorporated herein by reference.

PART IV

ITEM 14. EXHIBITS, FINANCIAL STATEMENT SCHEDULES, AND REPORTS ON FORM 8-K.

- (a) Documents filed as part of this report:
 - 1. Description of the Business.

The general description of the business and its general development are contained under the caption "Letter to Shareholders" in the 1998 Annual Report, which information is incorporated herein by reference.

- 2. Accountants' Consent and Report on Schedule.
- 3. Financial Statement Schedule of the Company.

SCHEDULE

NUMBER DESCRIPTION

Schedule II Valuation and Qualifying Accounts

All other schedules for which provision is made in the applicable accounting regulations of the Securities and Exchange Commission are not required under the related instructions or are not applicable and, therefore, have been omitted.

ACCOUNTANTS' CONSENT AND REPORT ON SCHEDULE

The Board of Directors
THE COOPER COMPANIES, INC.

The audits of the consolidated financial statements of The Cooper Companies, Inc. and subsidiaries referred to in our report dated December 10, 1998, which is incorporated herein by reference, included the related financial statement schedule for each of the years in the three-year period ended October 31, 1998 as listed in Item 14 of the Annual Report on Form 10-K. This financial statement schedule is the responsibility of the Company's management. Our responsibility is to express an opinion on this financial statement schedule based on our audits. In our opinion, such financial statement schedule, when considered in relation to the basic consolidated financial statements taken as a whole, presents fairly in all material respects the information set forth therein.

We consent to incorporation by reference in the Registration Statement Nos. 33-50016, 33-11298, 333-22417, 333-25051 and 333-27639 on Form S-3 and Registration Statement Nos. 333-10997, 33-27938, 33-36325, 33-36326 and 333-58839 on Form S-8 of The Cooper Companies, Inc. of our reports dated December 10, 1998, relating to the consolidated balance sheets of The Cooper Companies, Inc. and subsidiaries as of October 31, 1998 and 1997 and the related consolidated statements of income and cash flows for each of the years in the three-year period ended October 31, 1998, and related schedule, which reports appear in or are incorporated by reference in the October 31, 1998 Annual Report on Form 10-K of The Cooper Companies, Inc.

KPMG LLP

San Francisco, California January 25, 1999

SCHEDULE II

THE COOPER COMPANIES, INC. AND SUBSIDIARIES VALUATION AND QUALIFYING ACCOUNTS THREE YEARS ENDED OCTOBER 31, 1998

	BALANCE AT BEGINNING OF YEAR	ADDITIONS CHARGED TO COSTS AND EXPENSES	DEDUCTIONS/ RECOVERIES/ OTHER(1)	BALANCE AT END OF YEAR
		(IN THOU	ISANDS)	
		(11/11/100	ionivo)	
Allowance for doubtful accounts:				
Year ended October 31, 1998	.\$ 721	\$ 283	\$ 83	\$ 1,087
	======	=======	=======	=======
Year ended October 31, 1997	.\$ 716	\$ 155	\$ (150)	\$ 721
	=======	=======	=======	=======
Year ended October 31, 1996	.\$ 548	\$ 11	\$ 157	\$ 716
	=======	=======	=======	=======

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All prior periods have been restated to exclude Hospital Group of America as it was declared a discontinued operation.

⁽¹⁾ Principally uncollectible accounts written off, net of accounts recovered that were previously written off.

EXHIBIT NUMBER

PAGE

3.1	-	Restated Certificate of Incorporation, as partially amended, incorporated by reference to Exhibit 4(a) to the Company's Registration Statement on Form S-3 (No. 33-17330) and Exhibits 19(a) and 19(c) to the Company's Quarterly Report on Form 10-Q for the Fiscal Quarter ended April 30, 1988
3.2	-	Certificate of Amendment of Restated Certificate of Incorporation dated September 21, 1995 incorporated by reference to Exhibit 3.2 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1995
3.3		Company's Report on Form 8-A dated January 18, 1994
4.1	-	Certificate of Elimination of Series A Junior Participating Preferred Stock of The Cooper Companies, Inc. filed with the Delaware Secretary of State on October 30, 1997, incorporated by reference to Exhibit 4.1 on Form 10-K for fiscal year ended October 31, 1997
4.2	-	Rights Agreement, dated as of October 29, 1997, between the Company and American Stock Transfer & Trust Company, incorporated by reference to Exhibit 4.0 to the Company's Current Report on Form 8-K dated October 29, 1997
4.3	-	Amendment No. 1 to Rights Agreement dated September 25, 1998, incorporated by reference to Exhibit 99.1 of the Company's Current Report on Form 8-K dated September 25, 1998
4.4	-	Certificate of Designations of Series A Junior Participating Preferred Stock of The Cooper Companies, Inc., incorporated by reference to Exhibit 4.0 of the Company's Current Report on Form 8-K dated October 29, 1997
10.1	-	1998 Long-Term Incentive Plan, incorporated by reference to Exhibit A of the Company's Proxy Statement for its 1998 Annual Meeting of Shareholders held on April 2, 1998
10.2	-	Amendment No. 1 to 1998 Long-Term Incentive Plan of The Cooper Companies, Inc. dated April 2, 1998, incorporated by reference to Exhibit 4.7 to the Company's post-effective Amendment No. 1 to Form S-8 Registration Statement filed on January 20, 1999.
10.3	-	Severance Agreement entered into as of June 10, 1991, by and between CooperVision, Inc. and A. Thomas Bender, incorporated by reference to Exhibit 10.26 to Amendment No. 1 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1992
10.4	-	Letter dated March 25, 1994, to A. Thomas Bender from the Chairman of the Compensation Committee of the Company's Board of Directors, incorporated by reference to Exhibit 10.4 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1994
10.5	-	Severance Agreement entered into as of April 26, 1990, by and between Nicholas J. Pichotta and the Company incorporated by reference to Exhibit 10.8 to the Company's Annual Report on Form 10-K for fiscal year ended October 31, 1995
10.6	-	Letter Agreement dated November 1, 1992, by and between Nicholas J. Pichotta and the Company incorporated by reference to Exhibit 10.9 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1995
10.7	-	Severance Agreement entered into as of August 21, 1989, by and between Robert S. Weiss and the Company, incorporated by reference to Exhibit 10.28 to Amendment No. 1 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1992
10.8	-	1996 Long-Term Incentive Plan for Non-Employee Directors of The Cooper Companies, Inc., incorporated by reference to the Company's Proxy Statement for its 1996 Annual Meeting of Stockholders
10.9	-	Amendment No. 1 to 1996 Long-Term Incentive Plan for Non-Employee Directors of The Cooper Companies, Inc., dated October 10, 1996, incorporated by reference to Exhibit 10.14 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1996

10.10	-	Amendment N	0. 2	to 1	996 L	ong-Term	Incentiv	e Pl	an for	Non-	Emplo	yee	Dire	ctors	of
		The Cooper	Compa	nies	, Inc	., dated	October	29,	1997,	incor	porat	ed b	y re	ference	е
		to Exhibit :	10.15	to '	the	Company's	s Annual	Repo	rt on	Form	10-K	for	the	fiscal	
		vear ended	Octob	er 3	1. 19	97									

- Operations, the Consolidated Financial Statements and the Notes thereto, and the
- Financial Data Schedule, incorporated by reference to Exhibit 27 on the Company's Current Report on Form 8-K dated December 14, 1998.....
- * The information called for in this exhibit is contained in Note 4, "Earnings Per Share," in the 1998 Annual Report, which information is incorporated herein by reference.
- (b) REPORTS ON FORM 8-K.

August 26, 1998 -- Item 5. Other Events.
September 03, 1998 -- Item 5. Other Events.
September 25, 1998 -- Item 5. Other Events.
October 02, 1998 -- Item 5. Other Events.
October 21, 1998 -- Item 5. Other Events.
October 26, 1998 -- Item 5. Other Events.

SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) 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, on January 27, 1999.

THE COOPER COMPANIES, INC.

By: /s/ A. THOMAS BENDER

A. THOMAS BENDER

PRESIDENT, CHIEF EXECUTIVE

OFFICER AND DIRECTOR

Pursuant to the requirements of the Securities Exchange Act of 1934, this report has been signed below by the following persons on behalf of the registrant and in the capacities indicated on the dates set forth opposite their respective names.

SIGNATURE	CAPACITY	DATE 		
/s/ ALLAN E. RUBENSTEIN (ALLAN E. RUBENSTEIN)	Chairman of the Board of Directors	January 27, 1999		
/s/ A. THOMAS BENDER (A. THOMAS BENDER)	President, Chief Executive Officer and Director	January 27, 1999		
/s/ ROBERT S. WEISS (ROBERT S. WEISS)	Executive Vice President, Treasurer, Chief Financial Officer and Director	January 27, 1999		
/s/ STEPHEN C. WHITEFORD(STEPHEN C. WHITEFORD)	Vice President and Corporate Controller	January 27, 1999		
/s/ MICHAEL H. KALKSTEIN (MICHAEL H. KALKSTEIN)	Director	January 27, 1999		
/s/ MOSES MARX(MOSES MARX)	Director	January 27, 1999		
/s/ DONALD PRESS (DONALD PRESS)	Director	January 27, 1999		
/s/ STEVEN ROSENBERG	Director	January 27, 1999		
(STEVEN ROSENBERG) /s/ STANLEY ZINBERG (STANLEY ZINBERG)	Director	January 27, 1999		

LOCATION OF EXHIBIT IN SEQUENTIAL NUMBER SYSTEM

EXHIBIT NUMBER DESCRIPTION OF DOCUMENT

Restated Certificate of Incorporation, as partially amended, incorporated by 3.1 reference to Exhibit 4(a) to the Company's Registration Statement on Form S-3 (No. 33-17330) and Exhibits 19(a) and 19(c) to the Company's Quarterly Report 3.2 3.3 4.1 The Cooper Companies, Inc. filed with the Delaware Secretary of State on October 30, 1997, incorporated by reference to Exhibit 4.1 on Form 10-K for fiscal year ended October 31, 1997..... Rights Agreement, dated as of October 29, 1997, between the Company and American Stock Transfer & Trust Company, incorporated by reference to Exhibit 4.2 4.0 to the Company's Current Report on Form 8-K dated October 29, 1997..... Amendment No. 1 to Rights Agreement dated September 26, 1998, incorporated by 4.3 reference to Exhibit 99.1 of the Company's Current Report on Form 8-K dated September 25, 1998..... 4.4 Certificate of Designations of Series A Junior Participating Preferred Stock of The Cooper Companies, Inc., incorporated by reference to Exhibit 4.0 of the Company's Current Report on Form 8-K dated October 29, 1997...... 10.1 -1998 Long-Term Incentive Plan, incorporated by reference to Exhibit A of the Company's Proxy Statement for its 1998 Annual Meeting of Shareholders held on 10.2 dated April 2, 1998, incorporated by reference to Exhibit 4.7 to the Company's post-effective Amendment No. 1 to Form S-8 Registration Statement filed on January 20, 1999..... 10.3 CooperVision, Inc. and A. Thomas Bender, incorporated by reference to Exhibit 10.26 to Amendment No. 1 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1992...... 10.4 Letter dated March 25, 1994, to A. Thomas Bender from the Chairman of the

- 10.4 Letter dated March 25, 1994, to A. Thomas Bender from the Chairman of the Compensation Committee of the Company's Board of Directors, incorporated by reference to Exhibit 10.4 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1994.......
 10.5 Severance Agreement entered into as of April 26, 1990, by and between Nicholas
- Severance Agreement entered into as of April 26, 1990, by and between Nicholas
 Pichotta and the Company incorporated by reference to Exhibit 10.8 to the
 Company's Annual Report on Form 10-K for fiscal year ended October 31, 1995....
- 10.6 Letter Agreement dated November 1, 1992, by and between Nicholas J. Pichotta and the Company incorporated by reference to Exhibit 10.9 to the Company's
- Annual Report on Form 10-K for the fiscal year ended October 31, 1995..........

 10.7 Severance Agreement entered into as of August 21, 1989, by and between Robert S. Weiss and the Company, incorporated by reference to Exhibit 10.28 to Amendment No. 1 to the Company's Annual Report on Form 10-K for the fiscal year ended October 31, 1992.......

LOCATION OF EXHIBIT IN SEQUENTIAL NUMBER SYSTEM

EXHIBIT NUMBER DESCRIPTION OF DOCUMENT

10.8	-	1996 Long-Term Incentive Plan for Non-Employee Directors of The Cooper
		Companies, Inc., incorporated by reference to the Company's Proxy Statement
		for its 1996 Annual Meeting of Stockholders

- 10.9 Amendment No. 1 to 1996 Long-Term Incentive Plan for Non-Employee Directors of The Cooper Companies, Inc., dated October 10, 1996, incorporated by reference to Exhibit 10.14 to the Company's Annual Report on Form 10-K for the fiscal year ended October 21, 1996

- * The information called for in this exhibit is contained in Note 4, "Earnings Per Share," in the 1998 Annual Report, which information is incorporated herein by reference.
- (b) REPORTS ON FORM 8-K.

August 26, 1998 -- Item 5. Other Events. September 03, 1998 -- Item 5. Other Events. September 25, 1998 -- Item 5. Other Events. October 02, 1998 -- Item 5. Other Events. October 21, 1998 -- Item 5. Other Events. October 26, 1998 -- Item 5. Other Events.

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STATEMENTS OF DIFFERENCES

The	trademark symbol shall be expressed as	'TM'
The	registered symbol shall be expressed as	'r'
The	British pound sterling sign shall be expressed as	'L'

1998 ANNUAL REPORT

NARROWING OUR FOCUS . EXPANDING OUR REACH

THE COOPER COMPANIES, INC.

THE COOPER COMPANIES, INC.

IS A RAPIDLY GROWING SPECIALTY HEALTHCARE COMPANY. ITS BUSINESS UNITS SERVE ATTRACTIVE NICHE MARKETS IN THE MEDICAL DEVICE MARKET WITH HIGH QUALITY PRODUCTS AND SERVICES. COOPERVISION MARKETS A BROAD RANGE OF CONTACT LENSES IN NORTH AMERICA AND EUROPE. COOPERSURGICAL MARKETS DIAGNOSTIC PRODUCTS, SURGICAL INSTRUMENTS AND ACCESSORIES TO THE WOMEN'S HEALTHCARE MARKET. IN OCTOBER 1998, HOSPITAL GROUP OF AMERICA, COOPER'S PSYCHIATRIC SERVICES BUSINESS, WAS DECLARED A DISCONTINUED OPERATION.

FORWARD-LOOKING STATEMENTS

Statements in this report that are not based on historical fact may be "forward-looking statements" as defined by the Private Securities Litigation Reform Act of 1995. They include words like "may," "will," "expect," "estimate," "anticipate," "continue" or similar terms and reflect Cooper's current analysis of existing trends. Actual results could differ materially from those indicated due to: major changes in business conditions and the economy, loss of key senior Management, major disruptions in the operations of Cooper's manufacturing facilities, new competitors or technologies, significant disruptions caused by the failure of third parties to address the Year 2000 issue or by unforeseen delays in completing Cooper's Year 2000 compliance program, acquisition integration costs, foreign currency exchange exposure including the potential impact of the Euro, investments in research and development and other start-up projects, dilution to earnings per share from acquisitions or issuing stock, regulatory issues, significant environmental clean-up costs above those already accrued, litigation costs, costs of business divestitures, significant delay or failure to complete the sale of Hospital Group of America (HGA), and items listed in the Company's SEC reports, including its Annual Report on Form 10-K for the year ended October 31, 1998.

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FINANCIAL HIGHLIGHTS

Restated to reflect Hospital Group of America as a Discontinued Operation

Fiscal years ended October 31,

			,		,	
(In millions except per share data)		1998	% change vs. 1997		1997	% change vs. 1996
REVENUE						
CooperVision	\$	119.2	86%	\$	64.0	31%
CooperSurgical	\$	28.0	13%	\$	24.8	44%
Total	\$	147.2	66%	\$	88.8	34%
OPERATING INCOME						
CooperVisiion	\$	34.6	50%	\$	23.1	21%
CooperSurgical	\$	2.1	(14%)	\$	2.5	49%
Corporate Expenses	\$	(7.0)	21%		(5.8)	(11%)
Total	\$	29.7	50%	\$	19.8	39%
As a Percent of Revenue		20%			22%	
EARNINGS						
Net Income	\$	39.8	27%	\$	31.4	89%
As a Percent of Revenue	•	27%		•	35%	
From Continuing Operations	\$	57.8	32%	\$		180%
As a Percent of Revenue	•	39%		•	49%	
DILUTED EARNINGS PER SHARE						
Continuing Operations	\$	3.79	14%	\$	3.33	152%
Income Before Items Below:	\$	1.51	17%	\$	1.29	36%
Tax Benefit	\$	2.28	12%	\$	2.04	451%
Extraordinary Items	Ψ		n/m	\$	(0.11)	
Discontinued Operations	\$	(1.18)		\$	(1.05)	n/m
Net Income	\$	2.61	9%	\$	2.39	70%
OTHER FINANCIAL INFORMATION						
Depreciation and Amortization	\$	6.4	130%	\$	2.8	29%
Cash Flow from Operating Activities	э \$	11.5	(2%)	э \$	2.0 11.7	239%
Cash and Cash Equivalents	\$	7.3	(60%)	\$	18.2	167%
Working Capital	\$	69.4	(3%)	\$	71.5	121%
Total Assets	\$	296.0	74%	\$	170.6	103%
Total Liabilities	\$	150.8	155%	\$	59.1	(14%)
Stockholders' Equity	\$	145.2	30%	\$	111.5	628%
Average Shares Used for EPS calculation	~	15.3	16%	+	13.1	11%
		20.0				

TO OUR SHAREHOLDERS:

THERE HAVE BEEN IMPORTANT CHANGES AT YOUR COMPANY SINCE WE LAST WROTE TO YOU.

When Cooper completes the previously announced divestiture of Hospital Group of America (HGA), our mental health services business, we will be a pure medical device company, competing in two attractive markets, vision care and women's healthcare.

To update the divestiture plan: we have completed the sale of MeadowWood Hospital to Focus Healthcare, LLC, for approximately \$5 million in cash and trade receivables. In addition, we have signed a letter of intent with Universal Health Services, Inc., for the other three HGA properties. Universal will pay Cooper up to \$27 million in cash when the transaction closes plus up to \$3 million if certain contingent events occur. The closing remains, at this writing, subject to completion of a definitive agreement, ongoing due diligence and certain contingencies.

HGA is now a discontinued operation and all financial data in this report have been restated to reflect only the two medical device businesses, CooperVision and CooperSurgical.

COOPERVISION, OUR CONTACT LENS UNIT, AND COOPERSURGICAL, OUR WOMEN'S HEALTHCARE BUSINESS, ALSO MADE IMPORTANT TRANSITIONS THIS YEAR.

CooperVision (CVI) expanded its geographic reach by integrating Aspect Vision Care, Ltd. of Hampshire, England, acquired in December 1997. Aspect markets contact lenses in the United Kingdom and other European countries and supplies CooperVision with products for the United States market. With Aspect, CVI has significantly expanded its share of the spherical lens market-lenses that correct near- and farsightedness and not astigmatism--which is the largest segment of the contact lens market. At the same time, CVI continues to pursue its goal to become the worldwide leader in high-margin specialty contact lenses, especially toric lenses to correct astigmatism.

On October 27, 1998, The New York Stock Exchange hosted a meeting of Cooper's Board of Directors and Management. At the meeting, the president of the NYSE, William Johnston, presented Cooper's CEO Tom Bender with this replica of the famous NYSE bull.

CooperSurgical (CSI), our women's healthcare business, changed its new product development strategy. Future growth will come not only from acquisition but also from internal new product development. Adding to its base of products acquired since 1990, CSI introduced three new technologically advanced products, while continuing business development activities to consolidate the in-office women's healthcare market through acquisition.

In 1998, CVI sales grew 86% to \$119.2 million--including revenue of \$35 million from Aspect Vision for the eleven months it has been part of CVI. Operating income at CVI reached \$34.6 million, a 50% improvement over fiscal 1997. CSI sales grew 13% to \$28 million with operating income at \$2.1 million, down 14% from last year due primarily to new product introduction expenses and manufacturing scale-up delays.

Together, sales of Cooper's two medical device businesses reached \$147.2 million, up 66% from last year with operating income of \$36.7 million, before headquarters expense, up 44%.

Cooper's diluted earnings per share from continuing operations, before tax credits and charges for discontinued operations, were \$1.51 in 1998 compared with \$1.29 in 1997, up 17%. When tax credits and discontinued operations' charges are included, earnings per share were \$2.61 in 1998 versus \$2.39 in 1997 (see the "Diluted Earnings Per Share" section of the Financial Highlights table).

The tax benefits recorded in 1998 reflect the remaining tax savings that Cooper expects from its existing NOLs (net operating loss carry forwards). Generally Accepted Accounting Principles require that companies with NOLs must account for their remaining tax benefits when they determine that they expect sustainable profit going forward. Cooper has made this determination and in the future, will record expenses for income taxes using, initially, a tax rate of approximately 40%. From a cash standpoint, however, it will pay only state and foreign taxes for as long as the NOLs last. For federal tax purposes, the NOLs, now about \$180 million, will continue to shelter Cooper's federal tax liability. As you can see in the "Diluted Earnings Per Share" section of the Financial Highlight table, this net tax benefit was \$2.28 in 1998 and \$2.04 per share in

For 1999, we expect revenue and earnings from our combined medical device businesses to grow in the range of 20% to 25%. In fact, with HGA now a discontinued operation, Cooper's total 1999 revenue and earnings growth and its operating margin should each be above what we would have expected with HGA continuing as an operating unit.

FACTS AND FIGURES

ABOUT COOPER'S MEDICAL DEVICE BUSINESSES

With Cooper's imminent exit from the mental health services business, CooperVision and CooperSurgical form a medical device company serving two attractive healthcare segments, contact lenses and the women's healthcare market. A summary of the financial performance of these two businesses and their total as a medical device business follows.

REVENUE UP 66% FROM 1997 30% COMPOUND GROWTH FROM 1994 (In millions)

[GRAPH]

OPERATING INCOME(1) UP 43% FROM 1997 35% COMPOUND GROWTH FROM 1994 (In millions)

[GRAPH]

(1) Before Headquarters expense

NEW PRODUCTS AND ACQUISITIONS

COOPERVISION NEW PRODUCTS AND ACQUISITIONS 1994-1998 Recent acquisitions and new product introductions make CooperVision a challenging competitor in every segment of the contact lens market

Product Market Segment

Preference Three-month planned replacement spherical lens

Preference Toric Three-month planned replacement toric lens

CooperFlex One-month planned replacement spherical lens

Aspec Vision product line Disposable-planned replacement spherical lens

Hydrasoft Toric Options Quarterly custom toric lens

Frequency 55 Toric Disposable-planned replacement toric lens

Frequency 55 Sphere Disposable-planned replacement spherical lens

Natural Touch Conventional opaque lens

Alliance Toric Disposable-planned replacement toric lens

COOPERSURGICAL NEW PRODUCTS AND ACQUISITIONS SINCE 1990 During the past several years, CooperSurgical acquired new products and businesses and built critical mass. In the future, CSI will build on this platform through strategic acquisitions and new specialized women's healthcare products.

PRODUCT COMPANY DESCRIPTION

Frigitronics Colposcopes; cryosurgery equipment

Euro-Med, Inc. Biopsy instruments; instrument cleaning systems;

gynecology instruments

RUMI Uterine manipulator with disposable tip

Unimar, Inc. Disposable endometrial biopsy device; disposable

uterine manipulator; disposable cervical PAP smear

device, disposable infertility device

Marlow Surgical Technologies Disposable intrauterine catheter; laproscopic

instrument with disposable tip; disposable balloon cannula;

micro laproscopic instruments

Hyskon solution Diagnostic and surgical aid

FemExam Test Card System Diagnostic tests for vaginal infections

Cerveillance System Digital colposcopy instrumentation and software packages

Cooper Surgical Infrared

Coagulator Non-traumatic, nonsurgical instrument to treat

genital lesions

COOPERVISION

CooperVision (CVI) markets a broad range of contact lenses in North America and Europe.

The worldwide market for contact lenses grew 11% in dollars during 1998, according to our estimates. As it has for the past several years, the specialty lens segment--toric lenses for astigmatism, multifocal lenses for presbyopia and opaque lenses that can modify the eye's natural color--continued to grow significantly faster than the spherical lens market. In the United States, we estimate that during the first three quarters of the calendar year, the market for all toric lenses grew 16% in value, while the sphere market was flat. Improved toric technology and the continuing popularity of more frequently replaced toric lens regimens continue to drive demand.

Manufacturers' revenue in the disposable-planned replacement toric market, the fastest growing segment of the U.S. toric market, grew 42% in the first nine months of calendar 1998, according to our latest market research. Sales of CVI's disposable-planned replacement toric lenses in the U.S. grew 77% in fiscal 1998, led by its well regarded Preference Toric brand, which we believe is the fastest growing product today in the worldwide contact lens market.

Outside North America, the specialty market is not yet well-developed (see "How CooperVision Views the Contact Lens Market"), and we plan major educational efforts with practitioners in those markets to generate acceptance of CVI's toric products. In Europe, Aspect introduced CVI's toric products late in fiscal 1998 and expects to expand the product line in 1999.

During 1998, CVI's "divide and conquer" strategy took shape. We plan to offer products that can profitably meet most contact lens wearers' needs, segmenting the market by type of vision correction required, distribution channel, price point, geography, replacement cycle or lens material.

To implement this strategy in the U.S. toric market this year, we aimed at the lower priced segment of the market and introduced Frequency 55 Toric, a planned replacement lens for two-week or monthly use. Frequency 55 Toric attacks both the toric market's leading product and the new toric lenses recently launched by competitors, all positioned at the lower price point.

With its wider range of lens parameters compared with other low priced torics, Frequency 55 Toric is an attractive alternative to the competitive lenses. Moreover, even with Frequency 55 entering the lower priced segment, we expect continued share growth with Preference Toric. Many practitioners have learned that a few extra dollars for Preference Toric, with 15,500 lens parameters that can fit patients more efficiently, is a good investment that maximizes the value of their

SEGMENT	ESTIMATED 1998 MARKET (\$'s millions)
Soft Toric Lenses Conventional Lenses (lenses replaced annually) Disposable-Planned Replacement (various replacement schedules) Custom Lenses (for special prescriptions) Total Toric Market	\$ 65 \$ 104 \$ 39 \$ 208
Soft Spherical Lenses Conventional Lenses Disposable-Planned Replacement Total Spherical Lens Market	\$ 200 \$ 730 \$ 930
Rigid Gas Permeable Lenses	\$ 90
Total North American Soft Contact Lens Market Estimate	\$1,228

(1) CooperVision Estimates

time. (For more about this, see the "Prices, parameters and manufacturing technology" in "How CooperVision Views the Contact Lens Market").

We also introduced Hydrasoft Toric Options in 1998, a quarterly custom planned replacement lens program for another niche: astigmatic patients who have complex vision correction requirements.

In the U.S. spherical lens market, we describe our strategy as "comb and brush". Comb sales representatives can easily add brushes to their line because they call on customers who use both. We think the same way about selling spherical lenses to customers who already fit our toric lenses. Spherical lenses are the largest segment of the worldwide contact lens market, and the Aspect acquisition gives us the state-of-the-art technology we need to compete effectively in both North America and Europe with our recently introduced Frequency 55 Sphere.

While competitors have the lion's share of the North American disposable-planned replacement spherical lens market, we hope to capitalize on our strong relationships with our toric customers and the clinical benefits of a comfortable lens edge design, to help us sell Frequency 55 Sphere. In 1998, we valued the North American market at about \$930 million. We expect to capture a relatively small but meaningful share with Frequency 55 Sphere: around 5%, or a cumulative total of about \$40 million in sales, over the next several years.

After acquiring Aspect, CVI invested \$2.7 million to consolidate Aspect's facilities and integrate the worldwide manufacturing of spherical lenses in one location in Hampshire.

Before the rationalization of our manufacturing facilities could be completed, U.S. demand for Frequency 55 Sphere exceeded Aspect's capacity. Rather than lose the long-term annuity value to competitors, we decided to accept higher costs and meet the demand, including the additional costs to hire and train a larger work force, which we paid at premium rates. Manufacturing yields also declined because of the lower level of experience of the new workers.

Shifting sphere manufacturing from the Scottsville plant to the Hampshire site in the U.K. relieved a toric lens capacity constraint that developed while we built inventory for the introduction of Frequency 55 Toric. However, during this transition, the new toric manufacturing lines were not yet at full efficiency, and we incurred temporarily higher costs.

The manufacturing transition coupled with demand above our capacity in both plants, caused CVI's gross margins to fall from 65% after the first three quarters of 1998 to 60% in the fourth quarter. Both manufacturing facilities have since lowered their costs through increased automation. Lens output per employee has improved, and shift premiums are being eliminated. We expect gross margins to return to previous levels in the 1999 fiscal year.

In addition to our manufacturing investment, our 1998 capital spending also included construction of an expanded, up-to-date distribution center to ship lenses manufactured in Scottsville and Hampshire. In 1999, we estimate that we will spend about 50% less on our capital expenses than we did in 1998, so our cash flow should improve.

With market data still showing rapid growth in the toric lens segment and a continued shift from conventional to planned replacement toric lenses, it's important that CVI's business keep pace. At fiscal year end, CVI's toric lens sales accounted for 38% of its worldwide business growing 38% over 1997. In the U.S., independent market research indicates that CVI is gaining share in every market segment in which it competes. Worldwide, we expect most of the future market growth from the disposable-planned replacement segment. CVI's disposable-planned replacement toric and spherical lenses now comprise more than 75% of its business.

HOW COOPERVISION VIEWS THE CONTACT LENS MARKET

First, toric lenses.

CVI is concentrating its efforts in an attractive segment. In the U.S., toric dollar volume is currently growing 16%, while the spherical lens market is declining. Torics remain a proprietary segment with good margins available to suppliers, unlike the price sensitive spherical lens segment. Astigmatic patients have complicated vision requirements that make lens performance very important. Our market research indicates that the practitioner's toric lens purchasing decision is driven about 80% by how well the lens performs and 20% by price. With spherical lenses, it's just the opposite: price drives the decision.

CVI estimates that the toric contact lens market will grow more than 20% to about \$370 million worldwide in calendar 1999--about 12% of the total market--and reach about \$1 billion in 2003--or about 20% of the market. The table to the right shows how markets around the world will share this estimated potential.

PROJECTED GEOGRAPHIC GROWTH OF TORIC CONTACT LENSES(1)

Market (millions of \$'s)	1998	% of 1998 Total	2003	% of 2003 Total	CAGR2
North America	\$208	74%	\$462	50%	17%
Japan	15	5%	124	13%	53%
Europe	48	17%	163	17%	28%
Rest of the world	12	4%	184	20%	73%
Total toric market	\$283	100%	\$933	100%	27%

- (1) CooperVision estimates
- (2) Compound annual growth rate

To understand these markets, we need to look at the characteristics of contact lens wear in each of them.

North America is the most highly developed toric market. Total contact lens penetration among those requiring vision correction is about 20%--roughly 33 million wearers--the highest in the world. Among these, 45% of myopic (nearsighted) patients have astigmatism, but only about half of them have an astigmatic disability that requires a toric lens to achieve clear vision. In 1998, these percentages translated to a North American toric lens potential of about 8.3 million people. Currently, we estimate that about 3.8 million people wear them. Clearly, there's still room to grow.

Two forces drive North American market growth: first, improvements in contact lens technology allow more precise fits and more patients can now wear torics successfully; second the switch from conventional lenses that are replaced once a year to lenses replaced more frequently--daily, weekly, twice a month, monthly or quarterly--means that more lenses will be consumed. Contact lenses generate a continuing stream of income for suppliers while a patient remains in the market. As long as patients continue to move to toric lenses or move from conventional torics to disposable--planned replacement lenses, the market will continue to grow.

Japan is the second largest contact lens market in the world. We estimate that the penetration of contact lenses there is about 15% of the population requiring vision correction. The incidence of both myopia and astigmatism in the Asian population is the highest in the world, and about 75% of the myopic population also requires an astigmatic correction. This would translate into a potential Japanese toric market of about 6.1 million people, including a large number of patients currently wearing hard lenses. Until recently, Japan was predominantly a hard lens market, as concerns about chemicals from contact lens care solutions accumulating in soft lenses prevented their wide spread acceptance. Until the health authorities favorably reviewed the safety data, soft lenses grew slowly. Now many brands of soft lenses are available in various replacement cycles, but the acceptance of toric contact lenses has been slow. As Japan was a hard lens market for so long, practitioners tend to use them for their astigmatic patients as well, and it is difficult to change this practice.

Surprisingly, in economically well-developed Western Europe, revenue estimates for the toric market are relatively low, and even lower in lesser-developed Eastern Europe. Only about 8% of the Western Europeans who require vision correction wear contact lenses and only 5% of Eastern Europeans. The reasons behind this are primarily cultural and economic and vary from country to country. In Western Europe, spectacles and sunglasses with fashion frames have been preferred. In Germany, correcting astigmatism with hard lenses is still a popular technique. Moreover, Europeans in general have less astigmatism than North Americans do--about 35% of nearsighted patients have the condition, according to prevalence studies.

TORIC LENS PARAMETERS AND PRICE SEGMENTS

	Number of Parameters	
Competitors	Available	Comments
LOWER PRICE SEGMENT		
Cooper Toric	7,776	offers flexible wearing schedule
Cooper's Frequency 55 Toric	5,832	on the market and ready to meet emerging competition
Brand C	2,952	the current market leader
New Product B	1,300	introduced in late 1998
Brand W	750	introduced in 1997
HIGHER PRICE SEGMENT		
Cooper's Preference Toric Cooper's Hydrasoft Toric	15,500	FIPS manufacturing
Hydrasoft Toric Options	potential	custom lathed products
	13 million	for complex cases
Brand S	potential	a proprietary process with
	13 million	higher manufacturing costs and prices than Preference Toric

Before the market acceptance of our toric lenses in Japan and Europe reaches that of North America, we will have to educate practitioners there about the clinical and financial benefits of these products.

Prices, parameters and manufacturing technology. Understanding these three variables is the key to learning why CVI believes it has a sustainable competitive advantage in the toric market.

We believe that the U.S. toric lens market has two distinct and barely overlapping price segments. In the lower priced segment, practitioners remain willing to trade off performance in favor of a price advantage.

They can choose among several brands of lower priced toric lenses that offer as few as 750 to as many as nearly 3,000 different fitting parameters--combinations of 1.) "Power" that corrects for near and far defects, 2.) "Cylinder" that corrects for the astigmatism itself and 3.) "Axis," the exact position on the toric lens that will match the location of astigmatic defect on the cornea. CVI's recent entry in this lower priced segment--Frequency 55 Toric--has excellent prospects because it will offer over 5,800 parameters when all lenses are available in mid-1999.

In the higher priced segment, fitters view lens performance as more important than price. This segment is growing at the expense of the lower price segment, as these fitters begin to recognize the clinical and financial benefits of these lenses. CVI's Preference Toric, the market leader in this segment, offers 15,500 parameters. Practitioners can fit more Preference Toric patients correctly in less time than when using lenses with fewer choices.

CVI's manufacturing technology allows it to compete profitably in the higher priced segment. Historically, contact lens manufacturers have used either labor intensive lathing, which can generate a large number of parameters, or volume oriented lens molding where costs are low but only a limited number of parameters can be produced profitably. CVI's patented

manufacturing process called FIPS (finished inside polymerization system) combines the benefits of both processes--it molds the inside surface and lathes the front surface--yielding the highest number of mass produced toric parameters currently available at reasonable cost.

Next, spherical lenses.

In 1999, we estimate that the worldwide sphere market will grow about 10% over 1998 to nearly \$2.6 billion. Most of the growth, we think, will come from markets outside North America, led by Japan. In North America we estimate 1999 growth in spherical lenses will remain flat in dollars at about \$930 million. Longer term, we see the total sphere market growth compounding at about 11%, reaching about \$4 billion in 2003. With the Aspect acquisition, we now have significant volume in the market for spherical lenses. In Europe and North America together, our 1998 revenue from spheres totaled about \$73 million.

With Rohto Pharmaceuticals, Ltd. as a marketing partner in Japan, CVI will participate in the second largest spherical lens market in the world when the health authorities have favorably reviewed our products. We're expecting Rohto to begin selling spherical lenses in Japan during 1999. CooperVision will manufacture the lenses and sell them to Rohto to market in Japan and throughout the Far East. While the revenue from these sales will reflect our OEM manufacturing status, our operating margins should reach their usual levels, as we will not incur many of our normal marketing expenses.

Finally, a thought about laser surgery and contact lenses. We think that Laser Vision Correction (LVC) and contact lenses are complementary, not competitive. About 30 million people in North America wear contact lenses, and we estimate that about 20 million former lens wearers have dropped out of the market. They left because they no longer wanted the inconvenience of regularly caring for their lenses or because they no longer needed the social benefits that lenses can provide. Many left the market because toric technology had not advanced to where it is today and their vision was never crisp. We believe that most of the 200,000 or so patients who underwent LVC in 1998, came from this large pool of former lens wearers. These patients still want the benefits of contact lenses, but not the drawbacks, or they have occupational requirements that prevent their use. Even if all 200,000 LVC patients had given up contacts for the laser treatment, the negative impact on the 30-million wearer contact lens market would be minimal. In some cases, there's even a positive effect. When patients learn about the recent advances in contact lenses, especially those that correct astigmatism, and think about the potential negatives of LVC, some want to try lenses again.

What's ahead?

We expect growth in the North American toric market to continue at its current rate for the next several years. Improved toric technology and the continuing popularity of more frequently replaced toric lens regimens will help push the market ahead. We're also looking for the demand for soft toric lenses outside of North America to pick up as practitioners begin to appreciate the benefits of these products.

As for potential downsides, the North American spherical lens market was disappointing in 1998, and although we have a relatively small stake in it, we still need to determine the magnitude and duration of this market decline. While we do not see the toric market currently declining--it's up 16% this year--we must watch these trends closely and develop contingency plans to protect the growth of our operating income.

Note:

THE MARKET DATA IN THIS REPORT IS FROM CVI'S "CONTACT LENS MARKET ESTIMATES 1998-2003". IF YOU'D LIKE A COPY, PLEASE CONTACT COOPER'S INVESTOR RELATIONS DEPARTMENT.

COOPERSURGICAL

COOPERSURGICAL (CSI) MARKETS DIAGNOSTIC PRODUCTS, SURGICAL INSTRUMENTS AND ACCESSORIES TO THE WOMEN'S HEALTHCARE MARKET.

The case for opportunity in the women's healthcare market continues to be compelling. Like many attractive markets, it's driven primarily by favorable demographics, and in this case, it gets an added boost from U.S. national politics. Women's life expectancy has increased by 30 years over the last century. In 1990, women comprised 59% of Americans over the age of 65 and 72% of the over 85 age group. As women live longer and join the work force in increasing numbers, industry, government and medical practitioners are recognizing their special healthcare requirements.

Seven years ago, the National Institutes of Health, spurred in part by the Congressional Caucus for Women's Issues, began a major undertaking called the Women's Health Initiative. This is a \$625 million study lasting 16 years that is designed to increase the understanding of women's health during the last third of their lives. In October 1998, Congress approved legislation to extend these and several other women's health programs at the Centers for Disease Control and Prevention. The bill also authorizes research on cardiovascular disease in women at the National Heart, Lung, and Blood Institute.

The women's healthcare market is large and growing with three major segments--pharmaceuticals, capital equipment for hospitals and large clinics and in-office treatment. Each year, the 34,000 gynecologists in the United States record approximately 60 million office visits, assist in 4.6 million births and perform over two million surgical procedures. They treat conditions such as vaginitis, excessive menstrual bleeding, cancer and its precursors, non-malignant fibroid tumors and endometritis (an inflammation of the uterine lining). With the recent emphasis on preventive care, many managed care organizations now routinely reimburse common screening services such as PAP smears, osteoporosis evaluations and mammography. The cost pressures of managed care continue to move procedures from the hospital to the physician's office, and many women--some estimates are as high as 55% of women--now use their general practitioner.

CSI primarily targets the in-office practice where physicians screen, diagnose and treat the most commonly occurring gynecological conditions such as vaginitis. CSI also provides products for hospitals and clinics (including products for minimally invasive procedures) and reproductive medicine. In each of these, our strategy is to identify the most frequently performed procedures and surround physicians with the products used to do them.

Historically, CSI has attempted to consolidate the in-office women's healthcare products market through acquisition. Since 1990, it has acquired nine companies or product lines, and its 1998 revenue of \$28 million came primarily from these acquisitions. Typically, CSI consolidates an

TESTING FOR VAGINAL INFECTIONS

"No longer will physicians have to rely on costly laboratory tests or access to a microscope to diagnose bacterial vaginosis. The FemExam pH and Amines TestCard improves office testing by saving time and eliminating the need for in-office equipment."

Dr. Mark Newman,

an investigator for the U.S. Centers for Disease Control and Prevention and a maternal medicine specialist investigator.

VAGINAL INFECTIONS: HOW PREVALENT ARE THEY?

Based on data from the Centers for Disease Control and Prevention and the National Disease and Therapeutic Index.

Infection	Number Reported in 1996	Estimated Actual Annual Incidence (Number of Cases)
BACTERIAL		
Gonorrhea Chlamydia Syphilis Chancroid	325,883 490,080 52,995 386	
VIRAL		
Human papillomavirus virus	180,000 office visits	up to 1 million
Herpes simplex	210,000 office visits	up to 500,000
OTHER		
Trichomoniasis	250,000 office visits	up to 3 million
Other vaginitis (largely bacterial vaginosis vulvovaginal candidiasis, i.e. yeast)	3.5 million office visits	N/A

SOURCE: NYIRJESY, M.D., PAUL, "MANAGING CHRONIC VAGINITIS" OBG MANAGEMENT, MAY 1998. THE TABLE WAS ADAPTED FROM A PRESENTATION BY JOHN M. DOUGLAS, M.D., DIRECTOR OF STD CONTROL, DENVER PUBLIC HEALTH DEPARTMENT, "CARING FOR WOMEN WITH VAGINAL INFECTIONS," A NATIONAL SATELLITE VIDEOCONFERENCE PRESENTED BY THE NATIONAL NETWORK OF STD/HIV PREVENTION TRAINING CENTERS, MARCH 12,1998.

The diagnostic technology on the FemExam pH and Amines TestCard is con-tained on a disposable card that is about the size of a credit card (shown half-size on the left). It is designed to replace the costly, subjective and inconvenient testing practices currently used. It provides accurate, definitive indications of elevated pH (pH greater than or equal to 4.7) and detects the presence of amines, which together, along with other symptoms, indicate that bacteria are present. Industry estimates of the potential for vaginitis testing approach 125 million tests annually.

Today, physicians diagnose bacterial vaginosis using a combination of pH measurement, subjective vaginal fluid amine evaluation, microscopy and symptom evaluation. The FemExam pH and Amines Test Card does not require a culture, microscopic evaluation or capital equipment and provides almost immediate results. Collecting vaginal fluid with a cotton swab and wiping it across the reagent portion of the card performs the tests. Clinical studies show that the FemExam pH and Amines TestCard, when used in conjunction with clinical impressions can accurately detect bacterial vaginosis. It is not only equivalent to the standard measurement criteria currently used, but it also can accurately predict when vaginosis is not present 97% of the time.

Eventually, CSI plans to introduce a series of test cards for other vaginal infections. The primary target markets are Trichomoniasis and yeast infection. The adjacent table indicates the large market potential for the diagnosis of vaginal infections.

NOTE:

"OPTIMAL DIAGNOSIS OF VAGINITIS," A SUPPLEMENT TO THE NOVEMBER 1998 EDITION OF OBG MANAGEMENT IS AVAILABLE ON REQUEST FROM COOPER'S INVESTOR RELATION DEPARTMENT. IT REVIEWS CURRENT TRENDS IN THE DIAGNOSIS OF VAGINITIS.

acquired company's operations, including manufacturing, into its Connecticut facility within sixty days. This generates significant economies of scale that eventually boost margins. These acquisitions gave CSI the financial "critical mass" to allow it to introduce its own proprietary new products.

One of these is the innovative digital colposcopy system, Cerveillance Scope, introduced in May 1998. Using Cerveillance, physicians can examine the cervix and then document, store and recall digital images of their findings. Cerveillance sales to date are meeting expectations.

Another promising product introduced this year is the CooperSurgical InfraRed Coagulator, a device that creates infrared energy for contact coagulation of condylomas (genital lesions). Infrared coagulation is a simple, safe, rapid and exact technique that can be used on an outpatient basis without special training for physicians.

CSI also introduced the FemExam pH and Amines TestCard in 1998. It is the first in a planned series of point-of-care diagnostic products in its FemExam Test Card System. This product can improve the quality of the information that practitioners now have to diagnose vaginitis, the most common gynecological condition presenting in the physicians' office, and bacterial vaginosis (BV), the most prevalent form of vaginitis. When not treated, BV has been associated with amniotic fluid infections, premature rupture of the amniotic sac, pre-term and low birth weight infants, endometritis and post-surgical complications.

While early acceptance of this first FemExam card has been below our expectations due to yet unanswered questions about its medical economic benefit, we continue to believe that the FemExam System is a breakthrough technology. The slower than expected early sales do not change our long-term view of its potential.

The FemExam pH and Amines TestCard is more objective than current testing practices and can save time, thus improving diagnostic accuracy and practice economics. We have begun formal cost benefit studies and programs to raise the third-party reimbursement level for the product. We have lowered our revenue expectations for 1999, but this slower ramp-up will not impact operating income, as 1999's planned marketing expenses have also been lowered.

With CSI's new products and selective acquisitions, our objective is to double revenue to about \$60 million in the next three years and then to grow it at twenty percent per year in the next five-year period.

COOPER GOING FORWARD

WE STRONGLY BELIEVE THAT OUR STRATEGIES FOR OUR VISION CARE AND WOMEN'S HEALTHCARE BUSINESSES CAN ALLOW US TO CAPTURE AN EXPANDING SHARE OF THESE TWO ATTRACTIVE MEDICAL DEVICE MARKETS OVER THE COMING YEARS.

Our success, as always, depends on the hard work of our employees and our continuing commitment to shareholder value. We look forward to the challenges and opportunities of 1999.

We also want to thank the many loyal Cooper shareholders that remained confident in our future throughout the dramatic decline in the market for small capitalization stocks during 1998. We are confident that Cooper's outlook for 1999 and beyond will reward your patience.

January 27, 1999

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FIVE YEAR FINANCIAL HIGHLIGHTS

FIVE YEAR FINANCIAL HIGHLIGHTS

CONSOLIDATED OPERATIONS

(In thousands, except per share amounts)	1998	Years 1997	s Ended Octobe 1996	r 31, 1995	1994
Net operating revenue	\$ 147,192	\$ 88,769	\$ 66,118	\$ 55,296	\$ 51,034
Gross profit	\$ 91,428	\$ 61,444		\$ 37,747	\$ 33,128
Income from continuing operations before income taxes (Benefit of) provision for income taxes	\$ 23,087 (34,723)	\$ 16,936	\$ 11,167 (4,438)	\$ 6,121 43	\$ (9,536) (4,600)
Income (loss) from continuing operations before extraordinary item Discontinued operations, net of taxes:	57,810	43,671	15,605	6,078	(4,936)
Income before extraordinary item Loss from disposal Extraordinary item	4,336 (22,300) 	4,719 (18,000) (469)		(5,963) 	239
	(17,964)	(13,750)	998	(5,963)	239
Income (loss) before extraordinary item Extraordinary item, net		29,921 1,461		115 	(4,697)
Net income (loss) Less, preferred stock dividends	39,846 	31,382		115 	(4,697) 89
Net income (loss) applicable to common stock	\$ 39,846	\$ 31,382	\$ 16,603		\$ (4,786)
Diluted earnings (loss) per share: Continuing operations Discontinued operations Extraordinary item, net	\$ 3.79 (1.18)	\$ 3.33 (1.05) 0.11	\$ 1.32 0.09	\$ 0.52 (0.51)	\$ (0.49) 0.02
Earnings (loss) per share	\$ 2.61	\$ 2.39	\$ 1.41	\$ 0.01	\$ (0.47)
Average number of shares used to compute diluted earnings per share	15,269	13,120	11,794	11,667	10,193
Memo diluted earnings per share data: Income from continuing operations before income taxes	\$ 1.51	\$ 1.29	\$ 0.95	\$ 0.52	\$ (0.94)
CONSOLIDATED FINANCIAL POSITION					
			October 31,		
(In thousands)	1998	1997	1996	1995	1994
Current assets* Property, plant and equipment, net Intangible assets, net	\$ 116,077 34,234 84,308	\$ 100,574 7,634 32,274	\$ 58,712 4,650 16,864	\$ 52,185 3,974 9,901	\$ 58,927 3,591 9,669
Other assets	61,422	30,142	4,004	1,417	937
Total assets	\$ 296,041	\$ 170,624	\$ 84,230	\$ 67,477	\$ 73,124
Current liabilities** Long-term debt Other long-term liabilities	\$ 46,701 78,677 25,410	\$ 29,118 9,125 20,848	\$ 26,318 37,912 4,670	\$ 27,321 34,268 7,637	\$ 32,391 34,815 9,572
Total liabilities Stockholders' equity (deficit)	150,788 145,253	59,091 111,533	68,900 15,330	69,226 (1,749)	76,778 (3,654)
Total liabilities and stockholders' equity	\$ 296,041	\$ 170,624	\$ 84,230	\$ 67,477	\$ 73,124

All prior periods have been restated to present Hospital Group of America as a discontinued operation (see Note 3)

*Includes net assets of discontinued operations.

**Includes current installments of long-term debt

TWO YEAR QUARTERLY FINANCIAL DATA

	First Quarter	Second Quarter	Third Quarter	Fourth Quarter
Net sales	\$ 29,384	\$ 37,450	\$ 39,709	\$ 40,649
Gross profit	\$ 18,107	\$ 24,423	\$ 24,836	\$ 24,062
Income from continuing operations before income taxes senefit of income taxes**				\$ 3,891 (32,859)
Income from continuing operations Discontinued operations, net of taxes: Income before extraordinary item	5,343 650	7,378 1,105	8,339 1,835	36,750 746
Loss from disposal				(22,300)
Loss from discontinued operations			1,835	
Net income	\$ 5,993 	\$ 8,483	\$ 10,174	\$ 15,196
Diluted earnings per share*: Continuing operations Discontinued operations	\$ 0.35 0.04	\$ 0.48 0.07	\$ 0.54 0.12	\$ 2.45 (1.44)
Net income	\$ 0.39	\$ 0.55	\$ 0.66	\$ 1.01
Number of shares used to compute diluted earnings per share	15,354	15,443	15,342	14,978
Memo diluted earnings per share data: Income from continuing operations before income taxes	\$ 0.32	\$ 0.45	\$ 0.48	\$ 0.26
1997 -				
- 1997 - Net sales	\$ 17,027	\$ 20,630	\$ 24,951	\$ 26,161
			\$ 24,951 \$ 16,674	
Net sales Gross profit Income from continuing operations before income taxes		\$ 14,526	\$ 16,674	\$ 18,248 \$ 5,934
Net sales Gross profit Income from continuing operations before income taxes Genefit of income taxes** Income from continuing operations before extraordinary item	\$ 11,996 \$ \$ 2,630	\$ 14,526 \$ 3,714	\$ 16,674 \$ 4,658	\$ 18,248 \$ 5,934
Net sales Gross profit Income from continuing operations before income taxes Benefit of income taxes** Income from continuing operations before extraordinary item Discontinued operations, net of taxes: Income before extraordinary item	\$ 11,996 \$ 2,630 (422) 3,052	\$ 14,526 \$ 3,714 (455) 4,169 1,204	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465	\$ 18,248 \$ 5,934 (24,795 30,729
Net sales Gross profit Uncome from continuing operations before income taxes Benefit of income taxes** Uncome from continuing operations before extraordinary item Discontinued operations, net of taxes: Income before extraordinary item Loss from disposal	\$ 11,996 \$ 2,630 (422) 3,052 258	\$ 14,526 \$ 3,714 (455) 4,169 1,204	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465	\$ 18,248 \$ 5,934 (24,795 30,729 1,792 (18,000 (469
Net sales Gross profit Encome from continuing operations before income taxes Genefit of income taxes** Encome from continuing operations before extraordinary item Discontinued operations, net of taxes: Income before extraordinary item Loss from disposal Extraordinary item	\$ 11,996 \$ 2,630 (422) 3,052 258	\$ 14,526 \$ 3,714 (455) 4,169 1,204	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465	\$ 18,248 \$ 5,934 (24,795 30,729 1,792 (18,000 (469
Net sales Gross profit Income from continuing operations before income taxes Genefit of income taxes** Income from continuing operations before extraordinary item Discontinued operations, net of taxes: Income before extraordinary item Loss from disposal Extraordinary item Loss from discontinued operations Income before extraordinary item	\$ 11,996 \$ 2,630 (422) 3,052 258 	\$ 14,526 \$ 3,714 (455) 4,169 1,204 	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465 1,465 7,186	\$ 18,248 \$ 5,934 (24,795 30,729 1,792 (18,000 (469 (16,677 14,052 1,461
Het sales Gross profit Gross	\$ 11,996 \$ 2,630 (422) 3,052 258 258 3,310	\$ 14,526 \$ 3,714 (455) 4,169 1,204 1,204 5,373	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465 1,465 7,186	\$ 18,248 \$ 5,934 (24,795 30,729 1,792 (18,000 (469 (16,677 14,052 1,461
Net sales Gross profit Income from continuing operations before income taxes Genefit of income taxes** Income from continuing operations before extraordinary item Discontinued operations, net of taxes: Income before extraordinary item Loss from disposal Extraordinary item Loss from discontinued operations Income before extraordinary item Extraordinary item Extraordinary item, net Net income	\$ 11,996 \$ 2,630 (422) 3,052 258 	\$ 14,526 \$ 3,714 (455) 4,169 1,204 	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465 	\$ 18,248 \$ 5,934 (24,795 30,729 1,792 (18,000 (469 (16,677 14,052 1,461 \$ 15,513 \$ 2.02 (1.10 0.10
Net sales Gross profit Encome from continuing operations before income taxes Genefit of income taxes** Encome from continuing operations before extraordinary item Discontinued operations, net of taxes: Income before extraordinary item Loss from disposal Extraordinary item Loss from discontinued operations Encome before extraordinary item Extraordinary item, net Net income Diluted earnings per share*: Continuing operations Discontinued operations	\$ 11,996 	\$ 14,526 \$ 3,714 (455) 4,169 1,204 1,204 5,373 \$ 5,373 \$ 0.34 0.10	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465 1,465 7,186 \$ 7,186 \$ 0.44 0.11	\$ 18,248 5,934 (24,795 30,729 1,792 (18,000 (469 (16,677 14,052 1,461 \$ 15,513 \$ 2.02 (1.10 0.10
Net sales Gross profit Income from continuing operations before income taxes Benefit of income taxes** Income from continuing operations before extraordinary item Discontinued operations, net of taxes: Income before extraordinary item Loss from disposal Extraordinary item Loss from discontinued operations Income before extraordinary item Extraordinary item, net Net income Diluted earnings per share*: Continuing operations Discontinued operations Extraordinary item	\$ 11,996 \$ 2,630 (422) 3,052 258 	\$ 14,526 \$ 3,714 (455) 4,169 1,204 	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465 1,465 7,186 \$ 7,186 \$ 0.44 0.11 \$ 0.55	\$ 18,248 \$ 5,934 (24,795) 30,729 1,792 (18,000 (469) (16,677) 14,052 1,461 \$ 15,513 \$ 2.02 (1.10 0.10
Net sales Gross profit Income from continuing operations before income taxes Benefit of income taxes** Income from continuing operations before extraordinary item Discontinued operations, net of taxes: Income before extraordinary item Loss from disposal Extraordinary item Loss from discontinued operations Income before extraordinary item Extraordinary item, net Net income Diluted earnings per share*: Continuing operations Discontinued operations Extraordinary item Net income Number of shares used to compute diluted	\$ 11,996 	\$ 14,526 \$ 3,714 (455) 4,169 1,204 	\$ 16,674 \$ 4,658 (1,063) 5,721 1,465 1,465 7,186 \$ 7,186 \$ 0.44 0.11	\$ 18,248 \$ 5,934 (24,795) 30,729 1,792 (18,000 (469) (16,677) 14,052 1,461 \$ 15,513 \$ 2.02 (1.10 0.10 \$ 1.02

All prior periods have been restated to present Hospital Group of America as a discontinued operation (see Note 3) $\,$

^{*}The sum of earning per share for the four quarters is different from the full year amount as a result of computing the quarterly and full year amounts on the weighted average number of common shares outstanding in the respective periods.

^{**}Includes a tax benefit of \$33.3 million and \$25 million for the reduction of the valuation allowance against the deferred tax assets in the fourth quarters

of fiscal 1998 and 1997, respectively.

MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

Note numbers refer to the "Notes to Consolidated Financial Statements" of the Company beginning on page 32 of this report.

RESULTS OF OPERATIONS

Comparison of each of the fiscal years in the three year period ended October 31.1998.

NET SALES

Consolidated net sales grew 66% in 1998 and 34% in 1997.

Net sales of the Company's CooperVision ("CVI") and CooperSurgical ("CSI") business units have shown consistent growth over the three-year period:

(In thousands)	1998 v	s. 1997	1997 vs	. 1996
BUSINESS UNIT				
CVI	\$55,197	86%	\$15,121	31%
CSI	\$ 3,220	13%	\$ 7,536	44%

1998 vs. 1997

Net sales of CVI products increased primarily due to the acquisition of Aspect Vision Care Limited ("Aspect") (see Note 2) and sales growth achieved in planned replacement contact lenses in North America. The acquisition of Aspect accounted for 63% of the sales growth and represented approximately 29% of CVI's 1998 sales. In North America, sales of disposable-planned replacement toric lenses grew by approximately 74%, and sales of disposable-planned replacement spherical lenses grew approximately 79%. Sales of toric lenses to correct astigmatism, CVI's leading product group, grew 38% for the year and accounted for 38% of CVI's sales. In March 1997, the Company acquired Natural Touch, a line of opaque, cosmetic contact lenses that contributed \$5.4 million to 1998 sales. These increases were partially offset by anticipated declines in sales of mature product lines.

In February 1998, CVI introduced the Frequency 55 disposable-planned replacement spherical lens in the United States. The worldwide market for disposable-planned replacement spherical lenses represents about 60% of the total worldwide contact lens market.

In May 1998, CVI introduced two new toric products: Hydrasoft Toric Options, a custom planned replacement toric lens for astigmatic patients with complex corrections, and Frequency 55 Toric, a planned replacement lens designed for two-week or monthly replacement, positioned in the low-priced segment of the disposable-planned replacement toric market.

The Company believes that CVI is well-positioned to compete successfully in the contact lens market, particularly with its Preference and Frequency 55 line of planned replacement lenses and its line of custom toric lenses.

At CSI, net sales increased by 13% principally due to sales of Marlow Surgical Technologies, Inc. ("Marlow") products, acquired in April 1997 and Hyskon, a hysteroscopy fluid used by gynecologists in certain surgical procedures, acquired in December 1997.

CSI introduced three new product lines at the 1998 meeting of the American College of Obstetricians and $\mathsf{Gynecologists}$:

The Cerveillance Scope, an instrument that uses digital imaging and proprietary software to provide enhanced visualization and documentation in examinations of the cervix.

The CooperSurgical Infrared Coagulator, an instrument to perform a nonsurgical, noninvasive procedure to treat genital lesions in the physician's office. This technique coagulates tissue without carbonization, providing the surgeon with a smoke free environment that reduces the possibility of contamination.

The FemExam pH and Amines TestCard, the first in a series of patented diagnostic tests in the FemExam TestCard System that CSI recently licensed. These tests are used, primarily in the physician's office, to rapidly and economically screen and diagnose common vaginal infections such as bacterial vaginosis, yeast and trichomonasis.

1997 VS. 1996

CVI's net sales grew 31% due primarily to increased sales of toric lenses, CVI's leading product group, which grew by 40% and sales of the Preference spherical product lines, which grew 22%. Also, the addition of two new products, Natural Touch, a line of opaque, cosmetic lenses acquired in March 1997, and Encore, a line of planned replacement lenses, increased net sales 6%.

Net sales of CSI increased 44%. Women's healthcare products grew approximately 56%, primarily because of sales of Marlow products and Unimar products, acquired in April 1996. The increased sales of these products were partially offset by expected reductions in non-strategic or non-gynecologic product sales.

COST OF SALES/GROSS PROFIT

Gross profit as a percentage of net sales ("margin") was as follows:

MARGIN	1998	1997	1996
CVI	64%	76%	77%
CSI	55%	52%	51%
Consolidated	62%	69%	70%

CVI's margin declined in 1998 vs. 1997 due to the acquisition of Aspect, whose products have lower margins, and increased sales of lower margin Natural Touch products. Also, in the fourth quarter of 1998, CVI incurred estimated costs of \$1.7 million for rationalizing contact lens manufacturing, filling backorders and new product start-up inefficiencies. Margins on CVI's toric and other specialty lines of contact lenses have maintained their strong levels. Despite the anticipated margin decrease and the additional fourth quarter costs, CVI's gross profit grew by 57% over 1997, fueled by rapid sales growth.

CVI's margin declined in 1997 compared with 1996 due primarily to a write-off of approximately \$300,000 of inventory related to an unsuccessful attempt to enter the over-the-counter ophthalmic pharmaceutical market in Canada and increased sales of lower margin Natural Touch products.

CSI's margins have improved over the three-year period, reflecting the successful implementation of programs to more efficiently manufacture Unimar and Marlow products. In the absence of a material acquisition of lower margin products, Management expects that new and future proprietary products, after initial start-up phase, will command higher margins and that CSI's margins will continue to improve.

SELLING, GENERAL AND ADMINISTRATIVE EXPENSE ("SGA")

(In thousands)	1998	1997	1996
CVI CSI Corporate/Other	\$38,530 10,686 7,010	\$23,756 8,813 5,768	\$17,281 6,243 6,193
	\$56,226	\$38,337	\$29,717

Consolidated SGA increased by 29% in 1997 and 47% in 1998. Over the same periods, consolidated revenue grew 34% and 66%, respectively, resulting in consistent improvement in the ratio of SGA to sales from 45% of sales in 1996 to 43% in 1997 and 38% in 1998.

SGA at CVI increased by 62% in 1998 and 37% in 1997. The increase in 1998 was primarily due to the Aspect acquisition. Also, in the fourth quarter, CVI incurred an estimated \$1 million in SGA related to product launches, some of which experienced delays. The increase in 1997 resulted primarily from investments in selling, promotion and distribution costs required by the 31% increase in net sales, and an accrual for a potential environmental cleanup at one of its locations (see Note 11). As a percentage of its sales, CVI's SGA was 32% in 1998, 37% in 1997 and 35% in 1996.

The 1998 and 1997 SGA increases at CSI were due primarily to the acquisition of Marlow in 1997 and Unimar in 1996 (see Note 2).

The 1998 increase in Corporate/Other SGA was caused primarily by additional legal costs incurred to settle certain litigations and higher headquarters operating costs due to expanded responsibilities for international operations. The decrease in 1997 vs. 1996 SGA was primarily due to ongoing savings realized from reduced insurance costs and a 1995 restructuring.

RESEARCH AND DEVELOPMENT EXPENSE

Research and development expense was \$1.9 million or 1% of net sales in 1998, \$1.7 million or 2% in 1997 and \$1.2 million or 2% in 1996.

The current level of research and development spending is expected to remain stable as a percentage of sales, as the $\,$

THE COOPER COMPANIES, INC. AND SUBSIDIARIES

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Company is focusing on acquiring products that will be marketable immediately or in the short-term, rather than on funding longer-term, higher-risk research and development projects.

AMORTIZATION OF INTANGIBLES

Amortization of intangibles was \$3.6 million in 1998, \$1.6 million in 1997 and \$1 million in 1996. The increase in each year reflects the effect of acquisition activity during the three-year period (see Note 2).

INCOME FROM OPERATIONS

As a result of the activities discussed above, income from operations improved by \$15.4 million in 1998 vs. 1996. Income from operations by business unit and Corporate/Other was:

(In thousands)	Yea 1998	ers Ended October 31, 1997	1996
CVI CSI Corporate/other	\$ 34,574 2,136 (7,010)	\$ 23,101 2,476 (5,774)	\$ 19,065 1,667 (6,462)
	\$ 29,700	\$ 19,803	\$ 14,270
Percent growth	50%	39%	

INVESTMENT INCOME, NET

Investment income, net includes interest income of \$311,000, \$344,000 and \$207,000 in 1998, 1997 and 1996, respectively. The decrease in interest income in 1998 reflected the expenditure of cash to partially fund the Aspect acquisition. Interest income increased in 1997 because of higher investment balances primarily from cash received from the Company's follow-on offering, net of certain debt repayments.

SETTLEMENT OF DISPUTES, NET

In 1998, the Company recorded a charge to income of \$1.3 million (\$1.1 million in the fourth quarter) to settle the dispute between the Company and GT Laboratories (see Note 11) and certain other smaller matters. In 1997, the \$104,000 credit resulted from the reversal of an accrual no longer required.

OTHER INCOME (LOSS), NET

The change in other income (loss), net from a loss of \$141,000 in 1997 to income of \$561,000 in 1998, primarily related to foreign exchange transactions. In 1997, the weakness of the Canadian dollar against the U.S. dollar resulted in a foreign exchange loss of \$142,000. In 1998, the Company had a net foreign exchange gain of \$591,000, including a gain of \$850,000 as a result of the U.S. dollar amount of Pound Sterling denominated liabilities on the Company's books being reduced by the weakening in the Pound Sterling exchange rate prior to such liability being hedged. In 1996, the change in the foreign exchange rate was not a material factor.

INTEREST EXPENSE

Interest expense was \$6.3 million in 1998, \$3.2 million in 1997 and \$3.4 million in 1996. The increase in interest expense in 1998 vs. 1997 reflects debt used to finance a portion of the Aspect acquisition (see Note 2). The decrease in interest expense in 1997 vs. 1996 resulted from the redemption of the Company's 10 5/8% Convertible Subordinated Reset Debentures in April 1997 and 10% Senior Subordinated Secured Notes in September 1997.

INCOME TAXES

Details of the Company's income tax benefit for each year in the three-year period ended October 31, 1998 are set forth in Note 5. The 1998 provision for federal and state taxes of \$933,000 and foreign provision of \$131,000 and the utilization of foreign deferred tax asset of \$168,000 was offset by the recognition of an income tax benefit of \$36 million from reducing the valuation allowance against net U.S. deferred tax assets, based on Management's belief that the Company's future results will enable it to utilize this asset. The 1997 provision for federal and state taxes of \$545,000 was offset by a reversal of \$215,000 of tax accruals no longer required and the recognition of an income tax benefit of \$27.1 million from reducing the valuation allowance against net deferred tax assets. The 1996 provision for federal and state taxes of \$325,000 was offset by a reversal of \$615,000 of tax accruals no longer required and the recognition of an income tax benefit of \$4.1 million from reducing the valuation allowance against net deferred tax assets.

INCOME FROM DISCONTINUED OPERATIONS

Income from discontinued operations is income derived from the Company's Hospital Group of America, Inc. ("HGA") business unit, which was declared a discontinued operation by the Board of Directors in October 1998 (see Note 3). The reported income of \$4.3 million, \$4.7 million and \$1 million for fiscal years ended 1998, 1997 and 1996, respectively, is net of income tax expense (benefit) of \$130,000, \$129,000 and (\$50,000), respectively.

LOSS FROM DISPOSAL OF DISCONTINUED OPERATIONS

In 1998, the Company wrote down the net assets of HGA by \$22.3 million to the estimated fair market value of its net assets in anticipation of the sale of the business. HGA is accounted for as a discontinued operation (see Note 3).

In 1997, \$18 million was charged to discontinued operations which related to a settlement made in 1993 with Medical Engineering Corporation (see Note 11).

EXTRAORDINARY ITEM, NET

Continuing Operations:

In 1997, the Company recorded a net extraordinary gain of \$1.5 million on the early extinguishment of a portion of its long-term debt.

Discontinued Operations:

The \$500,000 charge in 1997 reflected early extinguishment of debt recorded by HGA.

CAPITAL RESOURCES & LIQUIDITY

The Company grew significantly in 1998, primarily due to the Aspect acquisition. Aspect provides distribution channels for CVI products in European markets and an additional range of products for CVI to sell in North America. Aspect also enabled CVI to enter the biweekly and monthly lens replacement market in the U.S.--the largest segment of the U.S. contact lens market. The integration of Aspect and the expansion of manufacturing capacity at both Aspect and CVI's United States plants resulted in an unusually high level of capital expenditures in 1998. The Company's investment in inventory for new products was also unusually high in 1998. Management expects investments in both of these to decrease in 1999.

Cash provided by operating activities totaled \$11.4 million in 1998 and \$11.7 million in 1997. Pre-launch inventory buildup of \$6.9 million and the Aspect acquisition significantly reduced 1998 operating cash flow. Aspect made over \$3 million of one-time payments shortly after the acquisition, and made a tax payment of approximately \$3 million in the second quarter for its tax liability on pre-acquisition operations.

Cash used by investing activities was \$59.3 million in 1998 vs. \$17.5 million in 1997. Primary uses of cash for investing activities included payments of approximately \$21.6 million for the acquisition of Aspect (see Note 2); the Hyskon product line purchase for \$2.3 million; the purchase, for \$10 million, of a 10% equity position in Litmus Concepts Inc. and an exclusive license to distribute Litmus' FemExam TestCard System in the U.S. and Canada. It also included capital expenditures of nearly \$20 million, which included approximately \$9.5 million to increase CVI's manufacturing capacity for disposable-planned replacement lenses. The principal uses of cash in 1997 included capital expenditures of \$7.7 million, \$3 million for the acquisition of the Natural Touch line of opaque contact lenses, \$4.1 million for the acquisition of Marlow and an investment of \$2.2 million in escrow funds restricted to expansion of CooperVision's Scottsville, New York, facility.

In 1998, the Company obtained \$37.3 million in cash flow from financing activities to fund the Aspect acquisition and the other major items discussed under investing cash flows. The financing activities primarily related to a \$21.8 million draw from the KeyBank line of credit, the Midland Bank loan of \$17.4 million, a net increase in capitalized leases of \$8 million and Aspect obtaining \$4.2 million of additional debt. The Company also repaid the \$4.2 million Unimar Promissory Note and \$1 million of the Wesley-Jessen Promissory Note. Financing activities included the purchase of treasury stock, which was authorized by the Board of Directors in September 1998. At October 31, 1998 the Company had repurchased 486,000 shares of the Company's stock at a cost of approximately \$8 million.

For the fiscal year ended October 31, 1998, the Company failed to meet one of the financial covenants in its KeyBank credit facility and received a waiver. KeyBank has amended the credit agreement by reducing the ratio required by such covenant, and the Company anticipates that it will remain in compliance in the future (see Note 6).

RISK MANAGEMENT

The Company is exposed to risks caused by changes in foreign exchange, principally Pounds Sterling denominated debt. The Company has hedged most of this risk by entering into contracts to buy Sterling forward. The Company is also exposed to risks associated with changes in interest rates, as the interest rate on certain of its debt varies with the London Interbank Offered Rate. The Company has protected itself against this risk by entering into agreements to swap most of its variable rate debt for fixed rate debt (see Note 7).

OUTLOOK

Management believes that cash flow from operations will fund ongoing operations. Financings may be required to fund further plant expansion in Europe, additional purchases of the Company's common stock and other acquisitions, if completed. At October 31, 1998, the Company had \$3.6 million available under the KeyBank line of credit. Management anticipates that additional financing would be available when and if required.

YEAR 2000

The Year 2000 ("Y2K") problem exists today because programmers who developed computer systems and applications over the past few decades used two digit date codes to designate the year. This creates a problem in the year 2000 in that many systems will recognize "00" as the year 1900 not the year 2000. Those systems that are not fixed may abort or produce erroneous data once the year 2000 arrives.

The Company has completed an in-depth review of the financial and operational systems at each of its business units and is implementing a Y2K compliance program, which is expected to be substantially completed by mid-1999. It is taking all reasonable steps to confirm that all of its critical business systems, software and equipment that consider and process date-related information will continue to function properly after December 31, 1999.

The Company initiated compliance programs in 1995 to modify its proprietary software, and many of the required changes have been completed. Software that was recently purchased requires minimal modification and the Company will ensure that any new software purchased will be Y2K compliant. It intends to use both internal and external resources to reprogram, or replace, and test its software for Y2K modifications. The Company has identified a person at each of its major operations who is responsible for Y2K compliance and has also appointed a Y2K Compliance Officer for the corporation. The Compliance Officer reports to the Board of Directors on the status of the Company's programs.

The Company has experienced significant growth in the past three years and is planning to begin implementation of a broad-based Enterprise Resource Planning ("ERP") system throughout its major CVI operations in the U.S. and the U.K. in 1999. The new ERP system will be Y2K compliant. In any event, as part of its contingency plan, CVI will assure that all of its existing systems are Y2K compliant prior to the conversion to the new system. In addition, at Aspect Vision, its recently acquired contact lens business in the U.K., plans are in place to build additional contact lens inventory prior to the millennium to assure there is no disruption in the flow of products to its customers.

The Company has or is currently ensuring Y2K compliance of all business systems and does not anticipate Y2K problems with these. It also has or is currently communicating with vendors to determine their Y2K compliance and is not aware of third party Y2K issues that could materially affect its operations.

The Company will continue to devote adequate resources to address its Y2K issues. However, it cannot assure that its systems and products do not contain undetected Y2K problems, that it will not experience operating difficulties as a result of Y2K issues or that its new systems will be implemented in time to avoid the probability of Y2K problems. Further, it cannot assure that the Company's assessment of suppliers and vendors will be accurate.

The Company has developed contingency plans to identify and mitigate potential Y2K problems and assess their impact on its operations. These plans will be designed to protect its assets, continue safe operations and allow any interrupted operations to resume in a timely fashion. The Company has developed contingency plans to respond to equipment failures, emergencies and business interruptions. However, contingency planning for Y2K issues is complicated by possible multiple and simultaneous incidents, which could significantly delay efforts to respond and resume normal business. Such incidents may be outside of the Company's control, for example if third parties, providing goods and/or services critical to the Company's operations, do not successfully address their own material Y2K problems.

Based on its Y2K assessment, the Company anticipates that the cost of upgrading or replacing its programs, systems and equipment will not materially affect its financial position. The total expenditures, including capital, required to be Y2K compliant are currently estimated at \$500,000. Approximately \$74,000 has been expended to date.

INFLATION AND CHANGING PRICES

Inflation has had little effect on the Company's operations in the last three years.

IMPACT OF STATEMENTS OF FINANCIAL ACCOUNTING STANDARDS ISSUED BUT NOT ADOPTED See Footnote 1 on page 32 of this report.

MANAGEMENT'S STATEMENT

The financial statements and other financial information in this report are Management's responsibility. They were prepared according to generally accepted accounting principles and, accordingly, include amounts based on Management's informed estimates and judgments. Other financial information in this report is consistent with that in the financial statements.

The Company's accounting systems include controls to reasonably assure that assets are safeguarded and financial statements conform to generally accepted accounting principles. These systems are supplemented by selecting and training qualified personnel and by an organizational structure that provides for appropriate separation of duties.

The Board of Directors, through its Audit and Finance Committee of three outside directors is responsible to determine that Management fulfills its responsibilities regarding preparation of financial statements and maintenance of financial control over operations. The Audit and Finance Committee recommends to the Board of Directors appointment of the Company's independent certified public accountants subject to ratification by the stockholders. It meets regularly with Management and the independent accountants. The independent accountants have access to the Audit and Finance Committee without Management present, to discuss auditing and financial reporting.

KPMG LLP has been the Company's independent certified public accountants since 1980 when the Company incorporated. KPMG provides an objective, independent review of the fairness of reported operating results and financial position. Their report appears on page 27.

/s/ A. THOMAS BENDER

A. Thomas Bender
President
and Chief Executive Officer

/s/ ROBERT S. WEISS

Robert S. Weiss Executive Vice President Treasurer and Chief Financial Officer

INDEPENDENT AUDITORS' REPORT

INDEPENDENT AUDITORS' REPORT

THE BOARD OF DIRECTORS AND STOCKHOLDERS THE COOPER COMPANIES, INC.:

We have audited the accompanying consolidated balance sheets of The Cooper Companies, Inc. and subsidiaries as of October 31, 1998 and 1997 and the related consolidated statements of income and cash flows for each of the years in the three-year period ended October 31, 1998. These consolidated financial statements are the responsibility of the Company's Management. Our responsibility is to express an opinion on these consolidated financial statements based on our audits.

We conducted our audits in accordance with generally accepted auditing standards. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. An audit includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements. An audit also includes assessing the accounting principles used and significant estimates made by Management, as well as evaluating the overall financial statement presentation. We believe that our audits provide a reasonable basis for our opinion.

In our opinion, the consolidated financial statements referred to above present fairly, in all material respects, the financial position of The Cooper Companies, Inc. and subsidiaries as of October 31, 1998 and 1997, and the results of their operations and their cash flows for each of the years in the three-year period ended October 31, 1998, in conformity with generally accepted accounting principles.

/s/ KPMG LLP

San Francisco, California December 10, 1998

CONSOLIDATED STATEMENTS OF INCOME

CONSOLIDATED STATEMENTS OF INCOME

	`	Years Ended Octob	per 31,
(In thousands, except per share figures)			
Net sales Cost of sales	\$ 147,192 55,764	\$ 88,769 27,325	\$ 66,118 19,911
		27,325	
Gross profit Selling, general and administrative expense	91,428	61,444	46,207
Research and development expense	1.944	1.739	1.176
Amortization of intangibles	3,558	1,565	1,044
Income from operations	29,700	61,444 38,337 1,739 1,565	14,270
Investment income, net	329	344	238
Settlement of disputes, net	1,250	(104)	
Other income (loss), net	561	(141)	80
Interest expense	6,253	344 (104) (141) 3,174	3,421
Income from continuing operations before			
income taxes	23,087	16,936	11,167
Add, benefit of income taxes	34,723	16,936 26,735	4,438
Income from continuing operations before extraordinary item Discontinued operations, net of taxes:	57,810	43,671	
Income before extraordinary item	4.336	4.719	998
Loss from disposal	(22,300)	(18,000)	
Extraordinary item		4,719 (18,000) (469)	
	(17,964)	(13,750)	998
Income before extraordinary item	39,846	29,921	16,603
Extraordinary item, net		29,921 1,461	
Net income	\$ 39,846	\$ 31,382	\$ 16,603
Basic earnings per share: Continuing operations before extraordinary item	\$ 3.90	\$ 3.42	\$ 1.34
Discontinued operations	(1.21)	(1.07)	0.09
Extraordinary item, net	`	\$ 3.42 (1.07) 0.11	
Earnings per share	\$ 2.69	\$ 2.46	\$ 1.43
Diluted earnings per share:			
Continuing operations before extraordinary item	\$ 3.79	\$ 3.33	\$ 1.32
Discontinued operations	(1.18)	(1.05)	0.09
Extraordinary item, net		\$ 3.33 (1.05) 0.11	
Earnings per share	\$ 2.61	\$ 2.39	\$ 1.41
Number of shares used to compute earnings per share: Basic		12,759	
Diluted	1F 260	13,120	11 704
Diluted	15,∠69	13,1∠U	11,794

See accompanying notes to consolidated financial statements.

CONSOLIDATED BALANCE SHEETS

	OC:	October 31,		
(In thousands)	1998	1997		
ASSETS				
Cash and cash equivalents Accounts receivable, less allowances of \$1,087 in 1998 and \$721 in 1997 Inventories	\$ 7,333 24,426 30,349	\$ 18,249 13,150 14,921		
Deferred tax asset Net assets of discontinued operations Prepaid expenses and other current assets	15,057 29,206 9,706	5,031 46,842 2,381		
Total current assets		100,574		
Property, plant and equipment at cost Less accumulated depreciation and amortization	45,079 10,845	16,804 9,170		
	34,234	7,634		
Goodwill and other intangibles, net	84,308 52,754	32,274		
Deferred tax asset Other assets	8,668	3,960		
		\$ 170,624		
LIABILITIES AND STOCKHOLDERS' EQUITY	\$ 4,612	\$		
Current installments of long-term debt Accounts payable	6,958 8,393	438 6,561		
Employee compensation and benefits Other accrued liabilities	5,087 12,664	4,155 8,830		
Accrued income taxes	8,987			
Total current liabilities	46,701	29,118		
Long-term debt Other noncurrent liabilities	78,677 25,410	9,125 20,848		
Total liabilities	150,788	59,091		
Commitments and Contingencies (see Note 11) Stockholders' equity Preferred stock, 10 cents par value, shares authorized:				
1,000: zero shares issued or outstanding Common stock, 10 cents par value, shares authorized: 20,000: issued: 14,912 and 14,798 at October 31, 1998				
and 1997, respectively Additional paid-in capital	1,491 251,167	1,480 249,213		
Other equity	(829)	(731)		
Accumulated deficit Less:	(98,583)	(138, 429)		
Treasury stock at cost, 486 shares at October 31, 1998	(7,993)			
		444 500		
Stockholders' equity	145,253	111,533		

See accompanying notes to consolidated financial statements.

CONSOLIDATED STATEMENTS OF CASH FLOWS

CONSOLIDATED STATEMENTS OF CASH FLOWS

	Years Ended October 31,			
(In thousands)		1997		
Cash flows from operating activities:				
Net income	\$ 39.846	\$ 31,382	\$ 16,603	
Adjustments to reconcile net income to net cash	Ψ 00/0.0	\$ 02,002	Ψ 20,000	
provided by operating activities:				
Deferred income taxes	(35.787)	(27,065)	(4.148)	
Depreciation expense		2,922		
Provision for doubtful accounts		2,336		
Amortization expense		1,345		
Loss from disposal of discontinued operations		18,000		
Extraordinary item		(992)		
Change in operating assets and liabilities excluding		, ,		
effects from acquisitions:				
Receivables	(3,910)	(7,521)	(4,998)	
Inventories	(6,933)	(3,855)	(445)	
Other assets	(952)	(356)	266	
Accounts payable		2,916		
Accrued liabilities	(5,949)	(4,021)	(4,488)	
Income taxes payable	(5,104)	(423)	(459)	
Other long-term liabilities	(3,973)	(3,044)	(4,287)	
Other	471	107	46	
Cash provided by operating activities	11,368	11,731	3,457	
Cash flows from investing activities:				
Purchases of assets and businesses	(34,298)	(7,145)	(4,080)	
Purchases of property, plant and equipment	(19,573)	(7,145) (7,735)	(3, 182)	
Investment in escrow fund		(2,216)		
Investment in options, net	(5,419)			
0ther		(357)	756	
Cash used by investing activities		(17,453)		

See accompanying notes to consolidated financial statements.

CONSOLIDATED STATEMENTS OF CASH FLOWS -- CONCLUDED

CONSOLIDATED STATEMENTS OF CASH FLOWS -- CONCLUDED

(In thousands)	1998		
		1997	1996
each flows from financing activities.			
Cash flows from financing activities: Proceeds from long-term line of credit	\$ 36,500	\$	\$
Repayment of long-term line of credit		ψ 	
Principal payments on long-term obligations	(7,603)		
Proceeds from long-term borrowings	29 682	 3,000	1,320
Net borrowings under short-term agreements	1,011		
Purchase of Treasury Stock	(7,993)		
Net payments of other notes payable and	(.,000)		
current long-term debt		(112)	(1,808
Net proceeds from follow-on offering		50.388	
Early retirement of debt		50,388 (35,740)	
Repayment of line of credit, net			(1.025
Other	430	(402)	(1,025 192
o choi		(402)	
Cash provided (used) by financing activities		17,134	
Effect of exchange rate changes on cash and			
cash equivalents	(321)		
Net increase (decrease) in cash and cash equivalents	(10.916)	11.412	(4.370
Cash and cash equivalents at beginning of year	18.249	6.837	11.207
		11,412 6,837	
Cash and cash equivalents at end of year	\$ 7,333 	\$ 18,249	\$ 6,837
Supplemental disclosures of cash flow information:			
Cash paid for:	A 4 500	A 4 700	4 4 000
Interest (net of amounts capitalized)	\$ 4,536	\$ 4,783	\$ 4,880
Income taxes	\$ 5,846	\$ 742	\$ 119
	1998	1997	1996
Supplemental disclosure of noncash investing			
and financing activities:			
Acquisitions (see Note 2):	¢ 02 406	¢ 10 E7/	6 0 664
Fair value of assets acquired	⊅ 93,400	\$ 18,574	\$ 9,661
.ess:		(45)	(404
Cash acquired	(24.000)	(45)	(404
Cash paid	(34, 298)	(7,145)	(4,080
Company stock issued	(1,492)	(4,662)	
Notes issued	(28,009)	(45) (7,145) (4,662) (4,500)	(4,000
_iabilities assumed and acquisition costs accrued	\$ 29 607	\$ 2,222	\$ 1 177

See accompanying notes to consolidated financial statements.

NOTE 1. SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

General

The Cooper Companies, Inc. (the "Company"), through its major subsidiaries, develops, manufactures and markets healthcare products, including hard and soft daily, flexible and extended wear contact lenses, and diagnostic products and surgical instruments and related products. In December 1997, the Company purchased an English contact lens company (see Note 2). Intercompany transactions and accounts are eliminated in consolidation. In October 1998, the Company's Management and Board of Directors declared its Hospital Group of America, Inc. ("HGA") business a discontinued operation, and prior years' financial statements have been restated to reflect this (see Note 3).

Foreign Currency Translation

Assets and liabilities of the Company's operations located outside the United States are translated at prevailing year-end rates of exchange. Related income and expense accounts are translated at weighted average rates for each year. Gains and losses resulting from the translation of financial statements in foreign currencies into U.S. dollars are recorded in the equity section of the consolidated balance sheet. Gains and losses resulting from the impact of changes in exchange rates on transactions denominated in currencies other than each reporting locations' functional currency are included in the determination of net income or loss for each period. Net foreign exchange income (loss) included in the consolidated statements of income for each of the years ended October 31, 1998, 1997 and 1996 was \$591,000, (\$142,000) and (\$13,000), respectively.

Derivatives

The Company uses derivatives to reduce market risk from changes in foreign exchange and interest rates. The Company generally does not use derivative financial instruments for trading or speculative purposes. It believes that each of the counterparties with whom it enters into forward exchange contracts and interest rate swap agreements is financially sound and that the credit risk of these contracts is low. The Company continually monitors its underlying market risk exposures and believes that it can modify or adapt itshedging strategies when needed (see Note 7).

Estimates in the Preparation of Financial Statements

The preparation of financial statements in conformity with generally accepted accounting principles requires Management to make estimates and assumptions that affect the reported amounts of assets and liabilities and the disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenue and expenses during each of the reporting periods. Actual results could differ from those estimates.

Revenue Recognition

The Company recognizes revenue net of appropriate provisions for returns when risk of ownership has transferred to the buyer. Management believes that there are no significant concentrations of credit risk in trade receivables.

Cash and Cash Equivalents

Cash and cash equivalents include commercial paper and other short-term income producing securities with maturity dates of three months or less. These investments are readily convertible to cash and are carried at cost that approximates market value.

INVENTORIES, AT THE LOWER OF AVERAGE COST OR MARKET

	October 31,		
(In thousands)	1998	1997	
Raw materials Work-in-process Finished goods	\$ 4,886 2,779 22,684	\$ 2,748 1,277 10,896	
	\$30,349	\$14,921	

PROPERTY, PLANT AND EQUIPMENT, AT COST

	October 3	1,
(In thousands)	1998	1997

Buildings and improvements	10,662	4,385
Machinery and equipment	32,909	12,364
	\$45,079 	\$16,804

THE COOPER COMPANIES, INC. AND SUBSIDIARIES

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Depreciation is computed using the straight-line method in amounts sufficient to write off depreciable assets over their estimated useful lives. Leasehold improvements are amortized over the shorter of estimated useful life or the period of the related lease. Buildings are depreciated over 35 to 40 years. Machinery and equipment is depreciated over 3 to 15 years, and software is depreciated over 3 years.

Expenditures for maintenance and repairs are expensed; major replacements, renewals and betterments are capitalized. The cost and accumulated depreciation of depreciable assets retired or otherwise disposed of are eliminated from the asset and accumulated depreciation accounts, and any gains or losses are reflected in operations for the period.

Amortization of Intangibles

Amortization is provided for on all intangible assets (primarily goodwill of \$68.2 million, which represents the excess of purchase price over fair value of net assets acquired) on a straight-line basis over periods of up to 40 years. Accumulated amortization at October 31, 1998 and 1997 was \$8.6 million and \$5.1 million, respectively. The Company assesses the recoverability of goodwill and other long-lived assets by determining whether the amortization of the related balance over its remaining life can be recovered through reasonably expected undiscounted future cash flows. Management evaluates the amortization periods of intangibles to determine whether later events and circumstances warrant revised estimates of useful lives. To date, no such adjustments have been required.

Stock-Based Compensation

The Company adopted Statement of Financial Accounting Standards ("SFAS") 123, Accounting for Stock-Based Compensation in 1997. This statement establishes financial accounting and reporting standards for stock-based compensation, including employee stock option plans. As allowed by SFAS 123, the Company continues to measure compensation expense under the provisions of Accounting Principles Board ("APB") No. 25, Accounting For Stock Issued to Employees, and related interpretations (see Note 9).

Statements of Financial Accounting Standards

Issued But Not Adopted

In June 1998, the Financial Accounting Standards Board ("FASB") issued Statement of Financial Accounting Standards ("SFAS") 133 "Accounting for Derivative Instruments and Hedging Activities," effective beginning with the first quarter of fiscal years beginning after June 15, 1999. SFAS 133 establishes accounting and reporting standards for derivative instruments, and hedging activities. In accordance with SFAS 133, an entity is required to recognize all derivatives as either assets or liabilities on its balance sheet and measure those instruments at fair value. SFAS 133 requires that changes in the derivative's fair value be recognized currently in earnings unless specific hedge accounting criteria are met, in which case gains and losses on the hedging instrument can offset related results on the hedged item in the income statement. The Company will adopt SFAS 133 in the first quarter of fiscal 2000. Under current generally accepted accounting principles, the Company may avail itself of hedge accounting for its forward exchange contracts. Hedge accounting allows the Company to offset the amounts due to and from the counterparty to the contract on its consolidated balance sheet. Under FAS 133, forward exchange contracts will not qualify for hedge accounting and, accordingly, the Company would then be required to include both the receivable and the payable on its consolidated balance sheet, possibly increasing its assets and liabilities materially.

In February 1998, the FASB issued SFAS 132 "Employers' Disclosures About Pensions and Other Postretirement Benefits." SFAS 132 is effective for fiscal years beginning after December 15, 1997. Restatement of disclosures for earlier periods presented is generally required. The Statement revises disclosures about pension and other postretirement benefit plans but does not alter the measurement or recognition of those plans. The Company will adopt SFAS 132 as required in the footnotes to its fiscal 1999 financial statements. Because SFAS 132 is a disclosure only Statement, Management believes adoption will have no impact on the Company's financial position, results of operations or cash flows.

In June 1997, the FASB issued SFAS 131 "Disclosures About Segments of an Enterprise and Related Information" which will be effective for financial statements for periods beginning after December 15, 1997, and establishes standards for disclosures about segments of an enterprise. The Company will make the required disclosures under SFAS 131 beginning with its consolidated financial statements for the year ending October 31, 1999, including the restatement of prior years. As SFAS 131 is a disclosure document only, Management believes adoption will not have any impact on the Company's earnings or cash flows.

In June 1997, the FASB issued SFAS 130 "Reporting Comprehensive Income" which will be effective for financial statements for fiscal years beginning after December 15, 1997. It establishes standards for reporting and displaying comprehensive income and its components in a full set of general purpose financial statements. The Company will report comprehensive income as required beginning with its interim financial statements for its first quarter of fiscal 1999. Upon adoption, reclassification of comparative financial statements for prior periods to reflect application of the provisions of SFAS 130 is required. The Company does not expect that the adoption of this statement will have any impact on its financial position or results of operations.

NOTE 2. ACQUISITIONS

Litmus Acquisition

In February 1998, the Company purchased, for approximately \$10 million in cash, a 10% equity position in Litmus Concepts Inc. and received an exclusive license to distribute Litmus' FemExam TestCard System of diagnostic tests in the U.S. and Canada. Of the \$10 million purchase price, \$5 million has been allocated to the equity investment and \$5 million to the exclusive license. The Company is accounting for its investment in Litmus on the cost basis and is amortizing the license over 17 years. The Company agreed to annual minimum purchases. This commitment ends when the Company purchases 20 million units of the products or on the sixth anniversary of the agreement, whichever occurs first. Under the terms of the agreement, if the Company does not meet the required minimum purchases, Litmus' sole remedy is its ability to cancel the license exclusivity.

Aspect Acquisition

In December 1997, the Company, through Aspect Vision Holdings, Limited ("Holdings"), acquired Aspect Vision Care Ltd. ("Aspect"), a privately-held manufacturer of high quality contact lenses sold primarily in the United Kingdom and other European countries. Aspect is an English company with the Pound Sterling as its functional currency. Holdings' functional currency is the U.S. dollar. Aspect and Holdings are included in CooperVision, Inc. ("CVI's") results from the date of acquisition.

The Company paid approximately \$51 million at closing (\$20 million in cash, 38,000 shares of the Company's common stock with a value of \$1.5 million and \$28 million in 8% five-year notes to the selling shareholders) and will pay an additional amount after approximately three years based Aspect's performance over that period. The minimum amount of the additional payment of 'L'5 million (approximately \$8 million at acquisition) has been discounted at a rate of 8% and will accrete over approximately three years. The \$20 million cash payment made at acquisition was partially financed under the Company's \$50 million line of credit (see "Midland Bank" Note 6) and cash then on hand. The acquisition has been accounted for as a purchase. Based on an independent valuation report, the excess of purchase price over net assets acquired has been recorded at \$44.9 million and is being amortized over 40 years, and other intangibles of \$3.5 million are being amortized over periods from 10 to 30 years.

Following the acquisition, certain of the selling shareholders became employees of the Company. As of October 31, 1998 approximately \$27.6 million of the five-year notes, and the minimum contingent payments, owed by the Company in connection with the acquisition are payable to these employees or members of their immediate family. None of these employees is an officer of the Company. For the year ended October 31, 1998 the Company's consolidated income statement included \$2 million of interest expense paid or payable to these individuals and \$2.3 million of royalty expense paid or payable to them.

In connection with the Aspect acquisition, the Company agreed to make quarterly royalty payments from 5% to 7 1/2% on sales of certain Aspect-manufactured products, with a minimum royalty for five years of 'L'1 million a year. The balance of royalty payable under the agreement was \$656,000 at October 31, 1998.

The following unaudited pro forma statements present consolidated condensed results of operations for the years ended October 31, 1998, and 1997, as if Aspect had been acquired at the beginning of each period. The unaudited pro forma information is not indicative of either the results of operations that would have occurred if Aspect had been purchased during the periods presented or of future results of the combined operations.

	Years Ended October 31,		
(In thousands, except per share figures)	1998 Pro Forma	1997 Pro Forma	
Net operating revenue	\$150,493	\$126,637	
Net income	\$ 40,114	\$ 31,278	
Shares outstanding for: Basic EPS	14,845	12,797	
Diluted EPS	15,286	13,158	
EPS: Basic	\$ 2.70	\$ 2.44	
Diluted	\$ 2.62	\$ 2.38	

Natural Touch'r' Acquisition

In March 1997, the Company acquired the United States rights to Natural Touch, a line of opaque, cosmetic contact lenses, from Wesley-Jessen Corporation ("W-J") for \$7.5 million (\$3 million in cash and a \$4.5 million promissory note, \$4 million of which was repaid) plus an ongoing royalty ranging from 3% to 8% per annum on sales of Natural Touch products other than those supplied by W-J. The Company recorded intangible assets of \$8 million for the patents, trademarks and distribution rights, which are being amortized over 7 to 15 years.

A subsidiary of W-J currently manufactures and supplies the Company with Natural Touch products. A divestiture order issued by the Federal Trade Commission (the "FTC") in connection with the Natural Touch acquisition requires the Company to either develop its own manufacturing capabilities or to find a suitable third-party manufacturer. The FTC could require the Company to divest the Natural Touch line if it has not either developed manufacturing capabilities that meet United States Food and Drug Administration ("FDA") approval or found a suitable third-party manufacturer meeting FDA approval within 42 months from the acquisition date.

Marlow Acquisition

In April 1997, the Company acquired Marlow Surgical Technologies, Inc. ("Marlow"), a women's healthcare products company, for approximately \$3.2 million in cash, liquidation of \$900,000 of Marlow debt and 144,800 shares of the Company's common stock valued at \$2.9 million at closing. As part of the acquisition, the Company agreed to issue an additional \$500,000 of its common stock (valued as of the closing) on the third anniversary of the closing, subject to reduction by the amount of any obligations of the seller to indemnify the Company in connection with the acquisition. Also, the Company has guaranteed that the total value of the shares of its common stock issued or to be issued in the acquisition (valued at \$3.4 million in total at closing) will appreciate by \$1.3 million by the third anniversary of the acquisition. This guarantee has been included in the purchase price, with a corresponding credit to additional paid-in capital. The acquisition has been accounted for as a purchase, with \$8.4 million of goodwill, which is being amortized over 20 years.

Unimar Acquisition

In April 1996, the Company acquired Unimar, Inc., a leading provider of specialized disposable medical devices for women's healthcare, for \$8 million in cash and notes. Goodwill has been recorded at \$7.8 million, which is being amortized over 20 years. As part of the acquisition, the Company granted a warrant to purchase 83,333 shares of the Company's common stock for \$11.375 per share. The warrant is valued at \$231,000. The exercise period of the warrant is from April 11, 1999 to June 10, 1999. The number of shares and the exercise price per share are subject to adjustment as provided in the warrant.

NOTE 3. DISCONTINUED OPERATIONS

In October 1998, the Company's Board of Directors declared the Company's HGA business unit a discontinued operation. Prior periods' financial statements have been restated to show HGA's net earnings as income (loss) from discontinued operations net of tax expense (benefit) of \$130,000, \$129,000 and (\$50,000) for 1998, 1997 and 1996, respectively, and its net assets have been restated to net assets of discontinued operations.

HGA's patient revenues were \$55.5 million, \$52.7 million and \$43 million for fiscal years ended October 31, 1998, 1997 and 1996, respectively. Net assets of discontinued operations at October 31, 1998 consisted primarily of patient receivables, net property, plant and equipment, net of accounts payable and accrued liabilities, including a \$22.3 million reserve for the estimate of the divestiture loss.

In the fourth quarter of 1998, the Company recorded a charge reflecting its estimate of the ultimate loss on this divestiture of \$22.3 million, or \$1.49 per diluted share. The Company has signed two agreements to sell the assets of HGA in two separate transactions. In January 1999, the Company sold the MeadowWood hospital property of its HGA business unit to Focus Healthcare, LLC. The Company will net approximately \$5 million in cash and trade receivables. The closing is subject to customary closing conditions. Under the second transaction, a nonbinding letter of intent, Universal Health Services, Inc. ("UHS") will pay the Company \$27 million in cash for three of its facilities when the transaction closes plus up to \$3 million if certain contingent events occur. The nonbinding letter of intent is subject to execution of a definitive agreement and satisfaction of closing conditions, including regulatory approval. The Company expects that the transaction with UHS will be completed within the first half of fiscal 1999.

NOTE 4. EARNINGS PER SHARE

The Company adopted SFAS 128, "Earnings Per Share" in the first quarter of 1998. This statement requires that earnings per share ("EPS") be determined using the weighted average number of common shares outstanding for Basic EPS, and then outstanding dilutive stock warrants and stock options are added to determine Diluted EPS. All prior period EPS amounts have been restated in accordance with SFAS 128.

	Years Ended October 31,			
In thousands, except per share figures)	1998	1997	1996	
Income from continuing operations				
before extraordinary item	\$ 57,810	\$ 43,671 (13,750)	\$ 15,605	
Discontinued operations, net of income taxes	(17,964)	(13,750)	998	
Income before extraordinary item	39,846	29,921	16,603	
Extraordinary item, net of income taxes		1,461		
Net income		\$ 31,382		
NET THEOME		Ψ 31,302		
ASIC:				
Weighted average common shares	14,828	12,759	11,646	
Basic earnings per common share: Continuing operations before extraordinary item	\$ 3.00	\$ 3.42	\$ 1.3	
Discontinued operations		(1.07)	0.0	
Extraordinary item		0.11		
Basic earnings per share:		\$ 2.46		
ILUTED:				
Weighted average common shares	14,828		11,646	
DD:				
Dilutive warrants	56	62	28	
Dilutive options	385	299	120	
Effect of dilutive securities	111	361	1/19	
Diluted weighted average common shares		13,120		
Diluted earnings per share:	Φ 0.70	Φ 2.22	.	
Continuing operations before extraordinary item Discontinued operations	\$ 3.79	\$ 3.33	\$ 1.33 0.09	
Extraordinary item	(1.18) 	(1.05) 0.11		
•				
Diluted earnings per share:	\$ 2.61	\$ 2.39	\$ 1.4	

In the years ended October 31, 1998, 1997 and 1996, options to purchase 571,250 shares at \$36.00 - \$62.21 per share, 340,000 shares at \$26.00 - \$35.09 per share and 150,000 shares at \$14.50 per share of common stock, respectively, were excluded from the computation of diluted earnings per share because they were antidilutive.

NOTE 5. INCOME TAXES

The income tax provision (benefit) related to income from all operations in the consolidated statements of income consists of:

	Years	Ended October 3	1,
(In thousands)	1998	1997	1996
From continuing operations From discontinued operations	\$(34,723) 130	\$(26,735) 129	\$(4,438) (50)
	\$(34,593)	\$(26,606)	\$(4,488)

	,	Years Ended October	31,
(In thousands)	1998	1997	1996
Current			
Federal State	\$ 462 471	\$ 309 21	\$ 196 (486)
Outside the United States	131		<u>-</u> - '
	1,064	330	(290)
Deferred	(25, 055)	(27, 005)	(4.140)
Federal Outside the United States	(35,955) 168	(27,065) 	(4,148)
	(35,787)	(27,065)	(4,148)
	\$(34,723)	\$(26,735)	\$ (4,438)

A reconciliation of the provision for (benefit of) income taxes attributable to income from continuing operations and the amount computed by applying the federal income tax rate to income from continuing operations before income taxes follows:

	Yea	rs Ended October	31,
(In thousands)	1998	1997	1996
Computed expected provision for taxes from continuing operations	\$ 8,080	\$ 7,407	\$ 4,119
Increase (decrease) in taxes resulting from: Income outside the United States subject to different	Ψ 5/555	Ψ .,	Ψ ., 220
tax rates	431	193	132
Amortization of intangibles	477	394	256
State taxes, net of federal income tax benefit	306	229	70
Reversal of prior years' estimated state tax liabilities			
no longer required		(215)	(615)
Utilization of net operating loss carryforwards	(10,359)	(7,102)	(4,406)
Change in beginning-of-year valuation allowance	(35, 787)	(27,065)	(4,148)
Other, net	2,129	(576)	154
Actual benefit of income taxes	\$(34,723)	\$(26,735)	\$ (4,438)

The tax effects of temporary differences that give rise to significant portions of the deferred tax assets and deferred tax liabilities are as follows:

		ber 31,
(In thousands)	1998	1997
Deferred tax assets:		
Accounts receivable, principally due to allowances for doubtful accounts	\$ 1,492	\$ 1,216
Inventories, principally due to obsolescence reserves	1,215	988
Accrued liabilities, principally due to litigation settlements and reserves,	,	
and compensation accruals	9,327	8,906
Net operating loss carryforwards	64,355	72,579
Capital loss carryforwards		2,523
Tax credit carryforwards	3,715	3,123
Other	1,225	907
Total gross deferred tax assets	91 220	90,242
Less valuation allowance		(52,517)
Less valuation allowance	(1,013)	(52,511)
Deferred tax assets	74,256	37,725
DEFERRED TAX LIABILITIES:		
Plant and equipment, principally due to purchase accounting requirements	(6,445)	(6,512)
Net deferred tax assets	\$ 67,811	\$ 31,213
not do on our tan doore		

The net decrease in the total valuation allowance for the years ended October 31, 1998, 1997 and 1996 was \$45.4 million, \$27.6 million and \$8.5 million, respectively. In 1998, 1997 and 1996, the Company recognized an income tax benefit of \$35.8 million, \$27.1 million and \$4.1 million, respectively, (\$33.3 million, \$25 million and \$4 million in the fourth quarters of fiscal 1998, 1997 and 1996, respectively) from reducing the valuation allowance based primarily on the continued improvement in the Company's operating results and future prospects. The recognition of the net deferred tax assets is based upon the expected utilization of net operating loss carryforwards that the Company believes will more likely than not be realized.

Subsequently recognized tax benefits relating to the valuation allowance as of October 31, 1998 will be allocated to:

(In	thousands)
Consolidated statements of income Goodwill and other intangible assets Additional paid-in capital for stock options	\$3,209 3,555 309
	\$7,073

At October 31, 1998 the Company had net operating loss and tax credit carryforwards for federal tax purposes expiring as follows:

Year of Expiration	Net Operating Losses	Tax Credits
(In thousands)		
1998 1999 2000 2001 2002 2003 2004 2005 2006 2007 2008 2009 2010 Indefinite	\$ 147 20,045 27,326 1,378 22,241 11,006 22,265 22,058 49,535 6,553 1,318	\$ 847 1,132 202 29 330 1,175
THUGH THITTE 1	\$183,872	\$3,715

NOTE 6. LONG-TERM DEBT

	Octob	er 31,
(In thousands)	1998	1997
Aspect promissory notes due December 2, 2002 (see Note 2)	\$27,563	\$
KeyBank line of credit	21,800	
Midland Bank Debt	17,444	
Aspect Vision bank loans	6,754	
County of Monroe Industrial Development Agency ("COMIDA") Bond	2,880	2,975
Capitalized leases, interest rates from 8% to 13% maturing 1998 to 2003	8,620	916
Wesley-Jessen Corporation ("W-J") promissory note	574	1,517
Unimar 12% promissory note		4,155
	85,635	9,563
Less current installments	6,958	438
	\$78,677	\$ 9,125

THE COOPER COMPANIES, INC. AND SUBSIDIARIES

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Annual maturities of capitalized leases and other long-term debt for each of the five years subsequent to October 31, 1998:

	(In thousands)						
	Capitalized Leases	Other Long- Term Debt					
1999 2000 2001 2002 2003	\$1,800 \$1,521 \$1,253 \$1,068 \$1,145	\$ 5,158 \$ 1,465 \$ 2,304 \$23,387 \$43,592					

KeyBank Line of Credit

The Company completed a \$50 million senior secured revolving credit facility with KeyBank National Association ("KeyBank") in September 1997. The facility matures September 11, 2002, with interest rates ranging from 0.5% to 2.0% over the London Interbank Offered Rates (LIBOR) depending on certain financial ratios. The interest rate may be floating or fixed at the Company's option. On October 31, 1998, the effective rates ranged from 6.5% to 6.7%. The Company pays an annual commitment fee of 0.375% on the unused portion of the revolving credit facility. Interest is paid monthly.

 $\mbox{KeyBank}$ syndicated a portion of the facility to other lenders and acts as agent for the lenders.

Terms include a first security interest in all assets of the Company. During the term of the facility, the Company may borrow, repay and re-borrow up to the \$50 million subject to voluntary reductions. The Company has used the KeyBank line of credit to guarantee other foreign borrowings by issuing \$24.6 million of letters of credit against the line of credit, which reduced its unused portion. At October 31, 1998, the Company had \$3.6 million available. This line of credit is guaranteed by the subsidiaries of the Company.

Mandatory prepayments will be required to repay outstanding amounts and permanently reduce the total commitment amount available under certain circumstances when the Company obtains additional debt or equity.

The KeyBank Line of Credit contains various covenants, including maintenance of certain ratios and transaction limitations requiring approval of the lenders. Certain prepayments are subject to penalties. One of the covenants required the Company to achieve the following ratios of earnings before interest, taxes, depreciation and amortization (as defined) to interest expense, capital expenditures and certain other fixed charges:

Ratio
1.3:1
1.3:1
1.3:1 2.0:1

The Company achieved this covenant for all periods except the 12 months ended October 31, 1998. The Company received a waiver for this period from KeyBank. In addition, KeyBank has amended the Credit Agreement by reducing the required ratio and the method of calculating it. The Company anticipates that it will be able to achieve the amended covenant going forward.

Midland Bank

The Aspect acquisition was partially funded by a 'L'10.5 million loan from Midland Bank plc, due November 27, 2002. In March 1998, the Company converted the denomination of the loan to U.S. dollars and entered into an interest rate swap to fix the interest rate at 6.19% per annum (see Note 7). The Midland loan is secured by a letter of credit in its favor from KeyBank National Association. Interest on the Midland loan is 20 basis points (0.2%) over Sterling LIBOR, adjusted monthly, and the Company pays an annual letter of credit fee of 1% of the balance to KeyBank.

Aspect Bank Loans

The balance of the loans at October 31, 1998, was \$6.8 million and is secured by certain assets of Aspect and a \$4.2 million letter of credit in favor of National Westminster Bank ("NWB") from KeyBank National Association. Loan maturity dates range from February 1, 2000, to September 1, 2006. The interest rate on 'L'2.5 million borrowed March 30, 1998 is 0.2625% above Sterling LIBOR. Sterling LIBOR was 7.58% for the period of the loan. The interest rate on other NWB loans is 1.5% above the Base Rate. The Base Rate ranged between 7.25% and 7.5% for the reporting period. Proceeds were used to repay a loan of 'L'827,000 (\$1.4 million), included in acquired debt, and to fund capital expenditures.

Capitalized Leases

The balance of capitalized leases at October 31, 1998, was \$8.6 million. The leases primarily relate to purchases of manufacturing equipment, both in the U.S. and in the United Kingdom. The increase in the amount of capitalized leases for the period was due to the expansion of the Company's manufacturing capacity.

COMIDA Bond

The COMIDA bond is a \$3 million Industrial Revenue Bond ("IRB") to finance the cost of plant expansion, building improvements and the purchase of equipment related to CVI's Scottsville, New York, facility. The interest rate has been fixed at 4.88%, per a Rate Swap Transaction (see Note 7). Principal is repaid quarterly, from July 1997 to October 2012. At October 31, 1998, unutilized proceeds of \$400,000 from the IRB, which must be used for the project described, are carried in other assets. The IRB is secured by substantially all of CVI's rights to the facility.

KeyBank issued a letter of credit to support certain obligations under the COMIDA bond. CVI is obligated to repay KeyBank for draws under and expenses incurred in connection with the letter of credit, pursuant to the terms of a reimbursement agreement, which is guaranteed by the Company. The agreement contains customary provisions and covenants, including the maintenance of certain ratios and levels of net worth. CVI and COMIDA have granted a mortgage lien on the building and real estate located in Scottsville and a first lien security interest on the equipment purchased under the bond proceeds to KeyBank to secure payment under the reimbursement agreement.

W-J

The W-J promissory note was issued for \$4.5 million, due March 17, 2001, in connection with the acquisition of the Natural Touch product line. To date, the Company has repaid \$4 million of the principal plus associated interest. Interest on the W-J promissory note is payable semi-annually and accrues at 12% annually: 8% per year is payable in cash and 4% per year is payable in kind.

Unimar Promissory Note

In April 1996, Cooper Healthcare Group, Inc. (a subsidiary of the Company) acquired Unimar, Inc. and issued a Promissory Note for \$4 million due April 11, 1999. In December 1997, the note plus associated interest was paid in full. Interest on the note was paid annually and accrued at 12% annually: 8% per year was paid in cash and 4% per year was paid in kind.

NOTE 7. FINANCIAL INSTRUMENTS

The fair values of the Company's financial instruments, including cash and cash equivalents, trade receivables, lines of credit, accounts payable and accrued liabilities, approximated their carrying values as of October 31, 1998 and 1997 because of the short maturity of these instruments.

The fair value of the Company's other long-term debt approximated the carrying value at October 31, 1998 and 1997, as the debt was refinanced or obtained within the respective fiscal years.

DERIVATIVES FOREIGN EXCHANGE INSTRUMENTS

The Company enters into forward exchange contracts to hedge the currency exposure of liabilities and firm commitments denominated in foreign currencies. Gains and losses on hedged commitments are deferred and recognized in the Company's results of operations in the same period as the gain or loss from the underlying transactions. As of October 31, 1998, the Company had outstanding forward exchange contracts of \$44.6 million to purchase 27.4 million British Pounds Sterling from November 1998 through December 2002. The fair value of the forward exchange contracts was obtained through KeyBank's Foreign Exchange department. The fair value indicated that termination of the forward exchange contracts at October 31, 1998 would have resulted in a \$740,000 gain.

INTEREST RATE AND OTHER DERIVATIVE INSTRUMENTS

The Company has entered into interest rate swap agreements to reduce the potential negative impact of increases in interest rates on its outstanding variable-rate debt under the Midland Bank Loan and the Industrial Revenue Bond. The Company recognizes in its results of operations over the life of the contract, as interest expense, the amortization of contract premiums incurred from buying interest rate swaps. Net payments or receipts resulting from these agreements are recorded as adjustments to interest expense. The effect of interest rate instruments on the Company's results of operations in fiscal year ended October 31, 1998 was not significant. As of October 31, 1998, the Company had two interest rate swap agreements with notional amounts totaling \$20.5 million. The \$17.5 million interest rate swap matures on November 27, 2002. The \$3 million interest rate swap matures on January 1, 2012. The fair value of the swap agreements was obtained through KeyBank's Derivative department. The fair value indicated that termination of the swap agreements at October 31, 1998 would have resulted in a \$914,000 loss.

In the fourth quarter of fiscal 1998, the Company simultaneously purchased and sold call options in the Semiconductor Index which expired in December 1998. The index options were purchased with temporary surplus funds of approximately \$5.4 million for trading purposes. Before the end of fiscal 1998, the Company traded substantially all of the purchase option position and a small portion of the sell option position and entered into a similar purchase option position and a similar sell option position having the same December 1998 expiration date. As of October 31, 1998, the investments in the purchased and sold call option contracts are netted because the terms of the index option contracts provide for a right of offset. The net investment as of October 31, 1998 in the amount of \$5.4 million is recorded at fair market value as represented by the net cash proceeds realized when the option contracts expired in December 1998 and is included in other current assets. The transaction did not result in any material gain or loss on the Company's financial statements.

NOTE 8. STOCKHOLDERS' EQUITY

	Common Shares	Common Stock	Paid-in Capital	Accumulated Deficit	Treasury Stock
Palance at October 21, 1005	11 576	\$ 1,158	¢ 192 940	¢(106 414)	\$
Balance at October 31, 1995 Exercise of stock options	11,576 22	\$ 1,158 2	\$ 183,840 117	\$(186,414)	ъ
Exercise of stock options Exercise of warrants and warrant valuation Restricted stock amortization and	66	6	297	=======================================	
share issuance	7	1	46		
Net income				16,603	
Balance at October 31, 1996	11,671	1,167	184,300	(169,811)	
Exercise of stock options	36	4	260		
Exercise of warrants Restricted stock amortization and	27	3	147		
share issuance	3		483		
Stock issued for acquisition (see Note 2) Stock issued for 10 5/8% debenture	145	14	4,648		
redemption	616	62	9,217		
Follow-on offering	2,300	230	50, 158		
Net income	,		·	31,382	
Balance at October 31, 1997	14,798	1,480	249,213	(138, 429)	
Exercise of stock options	75	7	419		
Treasury stock purchased Restricted stock amortization and					(7,993)
share issuance	1		47		
Stock issued for acquisition (see Note 2)	38	4	1,488		
Net income			,	39,846	
Balance at October 31, 1998	14,912	\$ 1,491	\$ 251,167	\$ (98,583)	\$ (7,993)

Treasury Stock

In September 1998, the Company's Board of Directors authorized the purchase of up to one million shares of the Company's common stock ("Treasury Stock"). By October 31, 1998, the Company had repurchased 486,000 shares of Treasury Stock at a cost of \$8 million.

Common Stock Offering

In 1997, in an underwritten follow-on stock offering, the Company sold 2.3 million shares of its common stock at \$23.50 per share. The proceeds from the offering of \$50.4 million, net of underwriters discount and transaction costs of \$3.7 million, were primarily used to repay outstanding debt.

Stockholders' Rights Plan

Under the Company's stockholder rights plan, each outstanding share of the Company's common stock carries one preferred share purchase right (a "Right"). The Rights will become exercisable only under certain circumstances involving acquisition of beneficial ownership of 20% or more of the Company's common stock by a person or group (an "Acquiring Person") without the prior consent of the Company's Board of Directors. If a person or group becomes an Acquiring Person, each Right would then entitle the holder (other than an Acquiring Person) to purchase, for the then purchase price of the Right (currently \$145, subject to adjustment), shares of the Company's common stock, or shares of common stock of any person into which the Company is thereafter merged or to which 50% or more of its assets or earning power is sold, with a market value of twice the purchase price. The Rights will expire in October 2007 unless earlier exercised or redeemed. The Board of Directors may redeem the Rights for \$.01 per Right prior to any person or group becoming an Acquiring Person.

Other Equity

At October 31, 1998, 1997 and 1996, the Company's cumulative foreign currency translation adjustments and deferred compensation reported in other equity were (\$829,000), (\$731,000) and (\$326,000), respectively.

NOTE 9. EMPLOYEE STOCK PLANS

At October 31, 1998 the Company has two stock-based compensation plans:

1998 Long-Term Incentive Plan ("1998 LTIP")

The 1998 LTIP is a vehicle for the Company to attract, retain and motivate its key employees and consultants, who are directly linked to the profitability of the Company and to increasing stockholder value. The 1998 LTIP was approved by the Company's stockholders in April 1998.

The 1998 LTIP authorized either a committee consisting of three or more individuals not eligible to participate in the 1998 LTIP or the Company's Board of Directors to grant to eligible individuals during a five-year period, stock options, stock appreciation rights, restricted stock, deferred stock, stock purchase rights, phantom stock units and long-term performance awards for up to 1,000,000 shares of common stock, subject to adjustment for future stock splits, stock dividends, expirations, forfeitures and similar events. Options generally vest based on the Company's stock price, however, in some cases, both stock price and time are used as criteria. As of October 31, 1998, 460,000 shares remained available under the 1998 LTIP for future grants. No restricted shares have been granted under the 1998 LTIP. Under a predecessor plan, the 1988 LTIP which expired, a total of 2,060,635 shares of restricted stock and stock options were awarded.

1996 Long-Term Incentive Plan for Non-Employee Directors ("1996 NEDRSP")

In March 1996, the Company's stockholders approved reducing the annual cash stipend paid to Non-Employee Directors and to award grants of restricted stock and options awarded annually at the start of each fiscal year. Specifically, each Non-Employee Director will be awarded the right to purchase restricted stock worth \$7,500 for \$0.10 per share (or \$9,375 in the case of the Chairman of the Board who is a Non-Employee Director) by January 15 of the year following the date of the grant. Grants of restricted stock not exercised by then will expire. The restrictions on the restricted stock will lapse when the stock reaches certain target values or by the fifth anniversary of the date of grants. In addition, each Non-Employee Director was granted an option to purchase shares of the Company's common stock in fiscal 1998 and 1997 and will be granted 5,000 shares in each subsequent fiscal year (or, in the case of the Chairman of the Board who is a Non-Employee Director, 6,250 shares in each subsequent fiscal year) through fiscal 2000. 215,000 shares of the Company's authorized but unissued common stock have been reserved for this. As of October 31, 1998, 117,240 shares remained available under the 1996 NEDRSP for future grants. Restricted shares of 1,312, 3,501 and 7,393 were granted under the 1996 NEDRSP in fiscal 1998, 1997 and 1996, respectively, and there were no restricted shares with restrictions in place outstanding October 31, 1998.

Common stock activity under the	se plans:								
				Years Ended Octo	ber 31,	,			
	1998			1997		1996			
	Options	A\ Exe	ighted verage ercise Price	Options	A	eighted Average Kercise Price	Options		eighted Average xercise Price
Outstanding at beginning of year Granted Exercised Forfeited	929,564 806,250 (75,017)	\$	19.39 38.16 5.68	459,662 514,165 (36,454) (7,809)	\$	8.90 27.69 7.25 5.78	328,841 192,361 (21,755) (39,785)	\$	5.77 12.77 5.59 3.55
Outstanding at end of year	1,660,797	\$	29.13	929,564	\$	19.39	459,662 	\$	8.90
Options exercisable at year end Weighted-avg. fair value of	605,797	\$	19.99	449,564	\$	9.71	244,164	\$	6.15
options granted during the year		\$	8.57		\$	12.32		\$	5.30

The options outstanding at October 31, 1998 for the stock option plans are:

	Options Outstanding			Options Exercisable	
Exercise Prices	Number Outstanding at 10/31/98	Weighted Average Remaining Contractual Life	Weighted Average Exercise Price	Number Outstanding at 10/31/98	Weighted Average Exercise Price
\$ 1.68 \$ 3.18-6.88 \$ 7.68-16.00 \$20.00-26.00 \$30.69-36.91 \$38.38-43.20 \$51.84 \$62.21	5,490 125,445 280,279 395,833 415,750 285,000 85,000 68,000	0.54 6.77 7.46 8.89 9.13 9.34 9.91	\$ 1.68 6.20 13.10 23.17 34.98 40.55 51.84 62.21	5,490 125,445 230,279 13,333 231,250	\$ 1.68 6.20 12.46 21.00 35.34
\$ 1.68-62.21	1,660,797	8.69	\$ 29.13	605,797	\$ 19.99

The excess of market value over \$.10 per share of 1988 LTIP, 1990 NEDRSP and 1996 NEDRSP restricted shares on respective dates of grant is initially recorded as unamortized restricted stock award compensation, a separate component of stockholders' equity, and charged to operations as earned. Restricted shares and other stock compensation charged against income from operations for the years ended October 31, 1998, 1997 and 1996 was \$260,000, \$107,000 and \$46,000, respectively.

Pro Forma Information

As permitted by FASB 123, the Company applies APB Opinion No.25 and related interpretations in accounting for its plans for stock issued to employees. Accordingly, no compensation cost has been recognized for its stock option plans. Had compensation cost for the Company's stock-based compensation plans been determined under the fair value method included in SFAS 123, the Company's net income and earnings per share would have been reduced to the pro forma amounts indicated below:

(In thousands, except per sh	are amounts)	1998	1997	1996
Net Income	As reported	\$39,846	\$31,382	\$16,603
	Pro forma	\$34,512	\$29,704	\$16,487
Basic earnings per share	As reported	\$ 2.69	\$ 2.46	\$ 1.43
	Pro forma	\$ 2.33	\$ 2.33	\$ 1.42
Diluted earnings per share	As reported	\$ 2.61	\$ 2.39	\$ 1.41
	Pro forma	\$ 2.28	\$ 2.29	\$ 1.40

The above pro forma amounts include compensation expense for options granted since November 1, 1995, and may not be representative of that to be expected in future years.

The fair value of each option grant is estimated on the date of grant using the Black-Scholes option-pricing model with the following weighted average assumptions used for grants in 1998, 1997 and 1996: zero dividend yield; expected volatility of 48 percent; expected option lives of 3.5 years and risk-free interest rates of 4.8%, 6.5% and 5.9%, respectively.

NOTE 10. EMPLOYEE BENEFITS

The Company's Retirement Income Plan The Company's Retirement Income Plan (the "Plan") covers substantially all full-time United States employees of CVI and the Company's corporate headquarters. The Company's contributions are designed to fund normal cost on a current basis and to fund over 30 years the estimated prior service cost of benefit improvements (15 years for annual gains and losses). The unit credit actuarial cost method is used to determine the annual cost. The Company pays the entire cost of the Plan and funds such costs as they accrue. Virtually all of the assets of the Plan are comprised of participation in equity and fixed income funds.

Net periodic pension cost of the Plan was:

		Years Ended October 3	31,
(In thousands)	1998	1997	1996
Service cost Interest cost Actual return	\$ 398 664	\$ 236 622	\$ 256 598
on assets Net amortization and deferral	(199) (571)	(1,446) 786	(1,047) 488
Net periodic pension cost	\$ 292	\$ 198	\$ 295

The actuarial present value of benefit obligations and funded status for the Plan was:

		oer 31,
(In thousands)	1998	1997
Vested benefit obligation	\$ 9,352	\$ 8,120
Non-vested benefit obligation	67	18
Accumulated benefit		
obligation	9,419	8,138
Projected compensation	4 040	040
increases	1,046	819
Projected benefit obligation	10,465	8,957
Fair value of plan assets	8,824	9,012
Projected benefit obligation in		
excess of (less than) assets	1,641	(55)
Add (Deduct): Unrecognized net gain	(401)	1,076
Prior service cost remaining to	(401)	1,070
be amortized, including		
unrecognized net asset	(336)	(358)
Pension liability recognized	\$ 904	\$ 663

Assumptions used in developing the projected benefit obligation were:

 Augus	st 31,
1998	1997

Discount rate on plan liabilities	7.0%	7.5%
Long-range rate of return		
on plan assets	9.0%	9.0%
Salary increase rate	4.0%	4.0%

THE COOPER COMPANIES, INC. AND SUBSIDIARIES

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The Company's 401(k) Savings Plan

The Company's 401(k) Savings Plan provides for the deferral of compensation as described in the Internal Revenue Code and is available to substantially all full-time United States employees of the Company. Employees who participate in the 401(k) Plan may elect to have from 1% to 16% (2% to 10%, prior to October 1, 1996) of their pre-tax salary or wages (but not more than \$5,000 for employees whose salary is more than \$66,000 annually for the calendar year ended December 31, 1996) deferred and contributed to the trust established under the Plan. The Company's contribution on account of participating employees, net of forfeiture credits, was \$396,000, \$218,000 and \$102,000 for the years ended October 31, 1998, 1997 and 1996, respectively.

The Company's Incentive Payment Plan

The Company's Incentive Payment Plan is available to officers and other key executives. Participants may, in certain years, receive bonuses based on performance. Total payments earned for the years ended October 31, 1998, 1997 and 1996, were approximately \$851,000, \$1.8 million and \$1.8 million, respectively. The 1997 and 1996 payments included payments made to HGA executives of \$414,000 and \$326,000 respectively.

NOTE 11. COMMITMENTS, CONTINGENCIES AND PENDING LITIGATION

Total minimum annual rental obligations (net of sublease revenue of approximately \$470,000 per year through March 2000) under noncancelable operating leases (substantially all real property or equipment) in force at October 31, 1998 are payable in subsequent years as follows:

	(In thousands)
1999	\$ 3,971
2000	3,463
2001	2,828
2002	2,356
2003	2,030
2004 and th	ereafter 10,507
	\$25,155

Aggregate rental expense for both cancelable and noncancelable contracts amounted to \$3.2 million, \$3 million and \$2.5 million in 1998, 1997 and 1996, respectively.

An agreement was reached in September 1993 with Medical Engineering Corporation ("MEC"), a subsidiary of Bristol-Myers Squibb Company, which limited the Company's contingent liabilities associated with breast implant litigation involving a former division of the Company (the "MEC Agreement"). The remaining liability recorded for payments to be made to MEC under the MEC Agreement are due on:

December 31,	Other Accrued Liabilities	Other Noncurrent Liabilities
(In thousands)		
1998	\$2,500	\$
1999	·	3,000
2000		3,500
2001		4,000
2002		4,500
2003		3,000
	\$2,500	\$18,000

Payments to MEC of \$18 million beginning December 31, 1999 are contingent upon the Company's earning net income before taxes in each fiscal year beginning with fiscal 1999. They were recorded in the Company's financial statements in the fourth quarter of fiscal 1997 as loss from sale of discontinued operations. They are reflected on the balance sheet in "Other noncurrent liabilities" as Management concluded that the payments would most likely be required. These payments are limited to the lesser of 50% of the Company's net income before

taxes in each fiscal year on a noncumulative basis, or the amounts shown above.

Environmental

In 1997, environmental consultants engaged by the Company identified a contained area of groundwater contamination consisting of industrial solvents including trichloroethane ("TCA") at one of CVI's sites. In the opinion of counsel, the solvents were released into the ground prior to the Company acquiring the business at that site, and the area containingthese chemicals is limited. The New York Department of

Environmental Conservation ("DEC") informed the Company on July 21, 1998 that the site was eligible for the New York Voluntary Cleanup Program. Recently, the Company and DEC reached agreement on a required order and scope of work that involves additional investigation. This further investigation will ultimately result in a state-approved remediation. The Company has accrued approximately \$500,000 for that purpose. In the opinion of Management, the cost of remediation will not be material when considering amounts previously accrued.

Pending Litigation--GT Labs

On October 1, 1992, GT Laboratories, Inc. filed a complaint against the Company in the United States District Court for the Northern District of Illinois alleging that the Company had breached a supply agreement by failing to purchase the requisite number of contact lens blanks used in the manufacture of rigid gas permeable contact lenses. The Company denied that it had breached the contract and asserted its right to terminate the agreement. In the interest of avoiding further litigation costs, the parties have agreed to resolve their dispute by way of settlement. On December 22, 1998, the parties entered into a Settlement Agreement and Release whereby the Company agreed to pay GT Laboratories \$1.3 million, \$1.1 million of which was accrued in the fourth quarter, in return for the plaintiff's release of all claims against the Company. In January 1999, the litigation was dismissed with prejudice.

NOTE 12.

BUSINESS SEGMENT INFORMATION

The Company's operations are attributable to two business segments:

CVI, which develops, manufactures and markets a range of contact lenses, and

CooperSurgical, Inc. ("CSI"), which develops, manufactures and distributes diagnostic products and surgical equipment, instruments and disposables, primarily for obstetrics and women's healthcare.

Total net sales include sales to customers as reported in the Company's consolidated statements of income and sales between geographic areas which are priced at terms that allow for a reasonable profit for the seller. Operating income (loss) is total net sales less cost of sales, research and development expenses, selling, general and administrative expenses and amortization of intangible assets. Corporate operating loss is principally corporate headquarters expense. Investment income, net, settlement of disputes, net, other income (expense), net, and interest expense were not allocated to individual husinesses

Identifiable assets are those assets used in continuing operations exclusive of cash and cash equivalents, which are included as corporate assets.

998	CVI	 CSI	Elin	porate & ninations	Cons	olidate
et revenue from non-affiliates	\$ 119,210				\$	47,192
perating income (loss)	\$ 34,574	 				29,700
nvestment income, net		 			-	329
ettlement of disputes, net						(1,250)
ther income (expense), net						561
nterest expense						(6,253)
ncome from continuing operations before income taxes					\$	23,087
dentifiable assets	\$ 143,888	\$ 41,887	\$	110,266	\$	296,041
epreciation expense	\$ 2,307	\$ 484	\$	81		2,872
mortization expense	\$ 2,090	\$ 1,468	\$		\$	3,558
apital expenditures	\$ 16,941	 		45	\$	17,732
997		 				
et revenue from non-affiliates	\$ 64,007	\$ 24,762	\$		\$	88,769
perating income (loss)	\$ 23,101					19,803
nvestment income, net		 			-	344
ther income (expense), net						(37
nterest expense						(3,174
ncome from continuing operations before income taxes						16,936
dentifiable assets	\$ 43,380	\$		97,701		170,624
epreciation expense	\$ 803	\$ 349	\$		\$	
mortization expense	\$ 674	\$ 891	\$		\$	1,565
apital expenditures	\$ 3,551					4,132
996		 				
et revenue from non-affiliates	\$ 48,892	7,226	\$			66,118
perating income (loss)	\$ 19,065	\$				14,270
nvestment income, net		 			-	238
her income (expense), net						80
terest expense						(3,421
ncome from continuing operations before income taxes					\$	11,167
dentifiable assets	\$ 23,756	\$ 18,089	\$	2,385	\$	84,230
epreciation expense	\$ 800	\$ 236	\$	82	\$	1,118
mortization expense	\$ 314	\$ 461	\$	269	\$	1,044
apital expenditures	\$ 1,293				\$	

Information by geographical area for each of the years in the three-year period ended October 31, 1998 follows: Eliminations 1998 United States Europe Canada & Corporate Consolidated Sales to unaffiliated customers \$ 147,192 Sales between geographic areas (9,261) Net sales **\$ 105,584 \$ 40,810 \$ 10,059 \$ (9,261) \$ 147,192** \$ 29,700 Operating income (loss) \$ 34,134 \$ 2,081 \$ 495 \$ (7,010) \$ 78,042 \$ 2,638 Identifiable assets \$ 105,095 \$ 110,266 \$ 296,041 1997 Sales to unaffiliated customers \$ 79,620 \$ -- \$ 9,149 \$ \$ 88,769 Sales between geographic areas 3,866 (3,866)-----. _____ \$ 88,769 Net sales \$ 83,486 \$ \$ (3,866) \$ 9,149 Operating income (loss) \$ 25,981 \$ \$ (404) \$ (5,774) 19,803 Identifiable assets \$ 3,014 \$ 97,701 \$ 170,624 \$ 69,909 \$ --1996 Sales to unaffiliated customers \$ 57,886 \$ -- \$ 8,232 \$ -- \$ 66,118 --(5,676) Sales between geographic areas Net sales \$ 63,562 \$ -- \$ 8,232 \$ (5,676) \$ 66,118 Operating income (loss) \$ 21,127 \$ \$ (395) \$ (6,462) \$ 14,270 Identifiable assets \$ 39,021 \$ -- \$ 2,824 \$ 42,385 \$ 84,230

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Corporate Information
BOARD OF DIRECTORS:
ALLAN E. RUBENSTEIN, M.D.
President WorldCare Imaging Services, Inc. Chairman
A. THOMAS BENDER
President and Chief Executive Officer
MICHAEL H. KALKSTEIN
Partner Graham & James
MOSES MARX
General Partner United Equities
DONALD PRESS
Executive Vice President Broadway Management Co., Inc.
STEVEN ROSENBERG
Vice President and Chief Financial Officer, Cooper Life Sciences, Inc.
ROBERT S. WEISS
Executive Vice President,
Treasurer and Chief Financial Officer
STANLEY ZINBERG, M.D.
Director of Practice Activities American College of Obstetricians and
Gynecologists
OFFICERS:
A. THOMAS BENDER
President and Chief Executive Officer and President CooperVision, Inc.
ROBERT S. WEISS
Executive Vice President,
Treasurer and Chief Financial Officer
B. NORRIS BATTIN
Vice President Investor Relations and Communications
GREGORY A. FRYLING
Vice President Corporate Development
Vice President of Legal Affairs, Secretary and Chief Administrative Officer
NICHOLAS J. PICHOTTA
President CooperSurgical, Inc.
STEPHEN C. WHITEFORD
Vice President and Corporate Controller
COMMITTEES OF THE BOARD:
MANAGEMENT COMMITTEE
ALLAN E. RUBENSTEIN, M.D. (Chairman)
Donald Press
AUDIT AND FINANCE COMMITTEE
STEVEN ROSENBERG (Chairman)
MICHAEL H. KALKSTEIN, STANLEY ZINBERG, M.D.
COMPENSATION COMMITTEE
MICHAEL H. KALKSTEIN
(Chairman) Donald Press
ÀLLAN E. RUBENSTEIN, M.D.
NOMINATING COMMITTEE
ALLAN E. RUBENSTEIN, M.D. (Chairman)
MOSES MARX
A. THOMAS BENDER, STANLEY ZINBERG, M.D.
CORPORATE OFFICES:
THE COOPER COMPANIES, INC.
10 Faraday, Irvine, CA 92618-1850
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6140 Stoneridge Mall Rd., Suite 590
Pleasanton, CA 94588
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COOPERVISION, INC.
10 Faraday, Irvine, CA 92618-1850
Voice: (949) 597-4700, Fax: (949) 597-0662
COOPERSURGICAL, Inc.
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HOSPITAL GROUP OF AMERICA, INC. 6140 Stoneridge Mall Rd., Suite 590 Pleasanton, CA 94588

15 Forest Parkway, Shelton, CT 06484 Voice: (203) 929-6321, Fax: (203) 925-0135 Voice: (925) 460-3600, Fax: (925) 460-3648

Note: HGA was declared a discontinued operation in October 1998. Its headquarters will close at the end of January, 1999.

COMMON STOCK PRICE RANGE:

Years Ended October 31, 1997 1998 Ouarter Ended Hiah Low High Low January 31 50 34 11/16 18 3/4 14 April 30 22 1/2 51 11/16 16 5/8 34 July 31 40 5/8 30 3/8 30 18 October 31 31 13/16 14 41 1/8

At December 31, 1998 there were 2,150 shareholders of record compared with 2,613 on December 31, 1997. No dividends were paid on the Company's common stock in 1998 or 1997, and the Company does not currently anticipate paying cash dividends in the future.

INVESTOR RELATIONS CONTACT:

B. NORRIS BATTIN

10 Faraday, Irvine, CA 92618-1850 Voice: (949) 597-4700, Fax: (949) 597-0662 email: nbattin@usa.net

INVESTOR INFORMATION:

Corporate information, including the current share price, recent news releases and the Company's annual report on Securities and Exchange Commission Form 10-K without exhibits, is available free of charge through the Company's interactive stockholder communication system. Call 1-800-334-1986, seven days a week, 24 hours a day. Visit The Cooper Companies, Inc. on the Worldwide Web at www.coopercos.com.

ANNUAL MEETING

The annual meeting of stockholders of The Cooper Companies, Inc. will be held on March 18, 1999 at the Marriott East Side Hotel, New York, NY at 10:00 A.M.

TRANSFER AGENT

American Stock Transfer & Trust Company 40 Wall Street, New York, NY 10005

CERTIFIED PUBLIC ACCOUNTANTS

KPMG LLP

STOCK EXCHANGE LISTINGS The New York Stock Exchange The Pacific Exchange Ticker Symbol "COO"

TRADEMARKS

The following trademarks italicized in this report are owned by, licensed to or distributed by The Cooper Companies, Inc., its subsidiaries or affiliates: Alliance Toric, Cerveillance, CooperFlex'TM', CooperSurgical Infrared Coagulator, Cooper Toric, Encore, FemExam TestCard System'TM', FemExam pH and Amines Test Card, Frequency 55, Hydrasoft, Hyskon, Natural Touch'TM', Preference, Rumi and Unimar.

[LOGO]

THE COOPER COMPANIES, INC. 10 Faraday, Irvine, California 92618-1850 Voice: (949) 597-1700, Fax (949) 597-0662, www.coopercos.com

[LOGO]

THE COOPER COMPANIES, INC.
NARROWING OUR FOCUS -- EXPANDING OUR REACH -- 1998 ANNUAL REPORT

Exhibit 21

SUBSIDIARIES OF THE COOPER COMPANIES, INC. A DELAWARE CORPORATION

NAME	JURISDICTION OF INCORPORATION
The Cooper Healthcare Group, Inc. Unimar, Inc. CooperVision Pharmaceuticals, Inc. The Cooper Real Estate Group, Inc. CooperSurgical Acquisition Corp. CooperVision, Inc. CooperVision, Inc. Marlow Surgical Acquisition (dormant)	Delaware California Delaware Delaware Delaware New York Canada Delaware
CooperVision GB Finance, Inc. (dormant) CooperVision GB Services, Inc. (dormant) Hospital Group of America, Inc. HGA Management Services, Inc. Hospital Group of Delaware, Inc. Hospital Group of Illinois, Inc. Hospital Group of Louisiana, Inc. Residential Centers of Indiana, Inc. Hospital Group of New Jersey, Inc. Hampton Learning Center, Inc.	Delaware Delaware Delaware Delaware Delaware Illinois Louisiana Delaware New Jersey
HGNJ, Inc. Arlington Center for Recovery, L.L.C. MeadowWood Health Services, L.L.C. CooperSurgical, Inc. CooperSurgical, Inc. HBH Medizintechnik GmbH Aspect Vision Holdings, Limited Aspect Vision Care Limited Contact Lens Technologies Limited Aspect Specialty Limited New Focus HealthCare Limited Aspect Vision Italia s.r.l. Focus Solutions Limited Averlan Company Limited Aspect Contact Lenses Limited	New Jersey Illinois Delaware Delaware Canada Germany England-Wales England-Wales England-Wales England-Wales Ingland-Wales England-Wales England-Wales Italy England-Wales England-Wales England-Wales England-Wales

NOTE: Except for CooperSurgical and its 52% owned subsidiary, HBH Medizintechnik GmbH, and Aspect Vision Italia s.r.l., each subsidiary is wholly-owned either by The Cooper Companies, Inc. or by the wholly-owned subsidiary under which it is indented in the list above. In the case of CooperSurgical, Inc., 99.8% of the company is owned by The Cooper Companies, Inc. and the remaining .2% is owned by members of CooperSurgical's Medical Advisory Board. In the case of Aspect Vision Italia s.r.l., 75.713% is owned by The Cooper Companies, Inc. and 24.287% is owned by Giacomo Grassi.