BOVIE MEDICAL CORP Form 10QSB November 16, 2006

U.S. Securities and Exchange Commission Washington D.C. 20549

FORM 10-QSB

(Mark One)

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

For the quarterly period ended September 30, 2006

[] TRANSITION REPORT UNDER SECTION 13 OR 15(d) OF THE EXCHANGE ACT

For the transition period from to Commission file number <u>0-12183</u>

BOVIE MEDICAL CORPORATION

(Exact name of small business issuer as specified in its charter)

Delaware (State or other jurisdiction Of incorporation or organization)

11-2644611 (IRS Employer Identification No.)

734 Walt Whitman Rd., Melville, New York 11747 (Address of principal executive offices)

(631) 421-5452 (Issuer's telephone number)

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

Yes [X] No []

APPLICABLE ONLY TO CORPORATE ISSUERS

Indicate the number of shares outstanding of each of the issuer's class of common stock, as of the latest practicable date: 14,726,938.

BOVIE MEDICAL CORPORATION INDEX TO FORM 10-QSB

FOR THE QUARTER ENDED SEPTEMBER 30, 2006

(This filing was originally mistakenly filed under the header of 10KSB, for the Quarter ended September 30, 2006 instead of Form 10QSB header. This filing is to correct that error.)

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PART I. FINANCIAL INFORMATION

ITEM 1: CONSOLIDATED FINANCIAL STATEMENTS

BOVIE MEDICAL CORPORATION CONSOLIDATED BALANCE SHEET SEPTEMBER 30, 2006 AND DECEMBER 31, 2005

Assets

	(Unaudited) September 30, 2006	(Audited) December 31, 2005
Current assets:		
Cash	\$ 2,309,492	\$ 1,295,266
Trade accounts receivable (net)	3,016,709	2,316,761
Inventories	3,377,987	2,996,832
Prepaid expenses	424,447	335,492
Deferred tax asset	386,200	386,200
Total current assets	9,514,835	7,330,551
Property and equipment, (net)	3,163,142	2,595,641
Other assets:		
Brand name/trademark (net)	1,509,662	1,509,662
Purchased technology (net)	316,084	33,663
License rights (net)	250,000	280,000
Deposits	21,215	21,215
	2,096,961	1,844,540
	\$ 14,774,938	\$ 11,770,732

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

BOVIE MEDICAL CORPORATION CONSOLIDATED BALANCE SHEET SEPTEMBER 30, 2006 AND DECEMBER 31, 2005 (CONTINUED)

Liabilities and Stockholders' Equity

Current liabilities:	(Unaudited) September 30, 2006	(Audited) December 31, 2005
Current naomities:		
Accounts payable Accrued expenses and other liabilities Deferred revenue Notes payable	\$ 921,937 775,909 109,636 203,241	868,212 471,006 141,586 348,328
Total current liabilities	2,010,723	1,829,132
Minority interest	125,000	140,000
Stockholders' equity:		
Preferred stock, par value \$.001 10,000,000 shares authorized 0 issued and outstanding on September 30, 2006 and December 31, 2005		
Common stock par value \$.001; 40,000,000 shares authorized, issued and outstanding 14,704,438 shares and 14,040,728 shares on September 30, 2006 and		
December 31, 2005 respectively Additional paid in capital Accumulated deficit	14,722 21,107,143 (8,482,650)	, , , , , , , , , , , , , , , , , , ,
Accumulated deficit	(8,482,030)	(10,742,349)
Total stockholders' equity	12,639,215	9,801,600
Total liabilities and stockholders' equity	\$14,774,938	\$11,770,732
The accompanying notes are an integral part of the financial statements		

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

BOVIE MEDICAL CORPORATION CONSOLIDATED STATEMENTS OF OPERATIONS FOR THE THREE MONTHS ENDED SEPTEMBER 30, 2006 AND 2005 AND FOR THE NINE MONTHS ENDED SEPTEMBER 30, 2006 AND 2005 (UNAUDITED)

	Three Months Ended September 30,		Nine Months Sep	tember 30,
	2006	2005	2006	2005
Sales	\$ 6,999,054	\$ 5,038,908	\$ 19,751,250	\$ 14,839,711
Cost of sales	4,118,252	2,924,168	11,743,121	9,489,840
Gross Profit	2,880,802	2,114,740	8,008,129	5,349,871
Costs and expenses:				
Research and development	261,923	303,850	631,674	657,939
Professional services	154,264	126,846	400,945	361,495
Salaries and related costs	647,002	447,210	1,847,328	1,469,673
Selling, general and	952,453	1,038,418	2,789,005	2,653,983
administrative				
Development joint venture	34,506	34,890	112,506	88,772
	2,050,148	1,951,214	5,781,458	5,231,862
Gain from operations	830,654	163,526	2,226,671	118,009
Other income (expense):				
Interest (net of expense)	28,872	5,594	50,795	16,797
Net income before minority				
interest and income tax	859,526	169,120	2,277,466	134,806
Minority interest	5,000	2,500	15,000	7,500
Provision for income tax Realized benefit of loss	(292,239)	(66,300)	(774,338)	(54,000)
carryforward	284,672	66,300	741,771	54,000
Net income	\$ 856,959	\$ 171,620	\$ 2,259,899	\$ 142,306
Earnings per share				
Net income:				
Basic	.06	.01	.16	.01
Diluted	.05	.01	.13	.01
Weighted average number of shares outstanding	14,610,828	13,928,162	14,351,324	13,907,579

Weighted average number of shares outstanding adjusted for dilutive securities 17,483,781

15,925,494

16,895,099

16,020,454

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

BOVIE MEDICAL CORPORATION CONSOLIDATED STATEMENTS OF SHAREHOLDERS' EQUITY FOR THE PERIOD ENDED JANUARY 1, 2005 TO SEPTEMBER 30, 2006

	Options Outstanding	Comm Shares	on Value	Paid-in Capital	Deficit	Total
January 1, 2005	3,951,200	13,862,128	\$ 13,881	\$20,391,407	\$(11,148,757)	\$9,256,531
Options granted	475,500					
Options exercised	(178,600)	178,600	178	3 138,683		138,861
Options forfeited	(31,230)					
Income for period					406,208	406,208
December 31, 2005	4,168,870	14,040,728	\$ 14,059	\$20,530,090	\$ (10,742,549)	\$9,801,600
Options exercised	(663,710)	663,710	663	493,362		494,025
Options forfeited	(22,360)					
Stock based compensation				20,390		20,390
Stock options issued to acquire assets				63,301		63,301
Income for period					2,259,899	2,259,899
September 30, 2006	3,482,800	14,704,438	\$ 14,722	2 \$ 21,107,143	\$ (8,482,650)	\$12,639,215
6						

BOVIE MEDICAL CORPORATION CONSOLIDATED STATEMENT OF CASH FLOWS INCREASE (DECREASE) IN CASH AND CASH EQUIVALENTS FOR THE NINE MONTHS ENDED SEPTEMBER 30, 2006 AND 2005 (UNAUDITED)

	2006	2005
Cash flows from operating activities		
Net income	\$ 2,259,899	\$ 142,306
Adjustments to reconcile net income	Ψ 2,237,077	ψ 1+2,500
to net cash provided by (used in) operating		
activities:		
Depreciation and amortization	394,106	344,292
Loss on disposals of fixed assets	6,727	, ,
Stock-based compensation	20,390	
Stock-based expense for Henvil asset purchase	20,886	
Changes in current assets and liabilities:		
Receivables	(699,948)	(331,421)
Inventories and repair parts	(381,155)	(760,468)
Prepaid expenses	1,045	110,265
Other receivable		(55,000)
Accounts payable	53,725	98,051
Customer deposits		78,588
Accrued expense	304,903	111,115
Deferred revenue	(31,950)	(6,610)
Net cash provided (applied) by operating activities	1,948,628	(268,882)
Cash flows from investing activities		
Increase in fixed assets	(929,253)	(781,621)
Increase in deposits	(727,233)	(2,000)
Increase in purchased technology	(264,088)	(2,001)
Net cash used in investing activities	(1,193,341)	(785,622)
Cash flows from financing activities		
Decrease in mortgage payable		
(:	348,328)	(23,748)
Increase in notes payable	132,067	
Decrease in notes payable	(18,825)	
Common shares purchased	494,025	50,363
Net cash provided in financing activities	258,939	26,615
	1,014,226	(1,027,889)

Net increase (decrease) in cash and cash equivalents

Cash and cash equivalents, beginning of period 1,295,266 2,294,746

Cash and cash equivalents, end of period \$2,309,492 \$1,266,857

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

BOVIE MEDICAL CORPORATION CONSOLIDATED STATEMENT OF CASH FLOWS INCREASE (DECREASE) IN CASH AND CASH EQUIVALENTS FOR THE NINE MONTHS ENDED SEPTEMBER 30, 2006 AND 2005

Cash paid during the nine months ended September 30:

2006 2005

Interest paid \$ 15,969 \$ 16,524 Income taxes 25,000 17,692

SUPPLEMENTAL SCHEDULE OF NON-CASH INVESTING AND FINANCING ACTIVITIES:

FOR THE NINE MONTHS ENDED SEPTEMBER 30, 2006 AND 2005:

There were two non-cash investing and financing activities in the first three quarters of the year 2006. The first was \$20,390 for stock based compensation to employees. The second was \$63,301, which was the calculated fair value of stock options given as consideration in the purchase of assets under the Henvil agreement, of which \$20,886 was expensed in the first nine months of 2006.

The fair value of the Henvil agreement options were estimated on the grant date using the binomial lattice option-pricing model with the following assumptions: expected volatility of 25%, expected term of 5 years, risk-free interest rate of 5.0%, and expected dividend yield of 0%. Expected volatility is based on a weighted average of the historical volatility of the Company's stock and peer company volatility. The average expected life was calculated using the simplified method under SAB 107. The risk-free rate is based on the rate of U.S. Treasury zero-coupon issues.

During the first nine months of 2006 we purchased patent pending rights and an exclusive license for technology. The patent and technology rights were valued at \$306,503 of which the full amount had been paid as of September 30, 2006.

NOTE 1. INTERIM FINANCIAL INFORMATION

The accompanying unaudited condensed consolidated financial statements have been prepared in accordance with accounting principles generally accepted in the United States of America (U.S.) for interim financial information and with the instructions to Form 10-QSB and Article 10 of Regulation S-X. Accordingly, they do not include all of the information necessary for a fair presentation of results of operations, financial position, and cash flows in conformity with accounting principles generally accepted in the U.S. In the opinion of management, the condensed consolidated financial statements reflect all adjustments (consisting of normal recurring adjustments) considered necessary for a fair presentation of the results of Bovie Medical Corporation and its subsidiaries for the periods presented. Operating results for interim periods are not necessarily indicative of results that may be expected for the fiscal year as a whole. The preparation of the financial statements in conformity with accounting principles generally accepted in the U.S. requires management to make estimates and assumptions that affect the reported amounts of assets, liabilities, revenues, expenses, and the related disclosures at the date of the financial statements and during the reporting period. Actual results could materially differ from these estimates. For further information, refer to the consolidated financial statements and notes thereto included in the Company's Annual Report on Form 10-KSB for the year ended December 31, 2005. Certain prior year amounts may have been reclassified to conform with the presentation used in 2006.

NOTE 2. INVENTORIES

Inventories are stated at the lower of cost or market. Cost is determined principally on the average cost method. Inventories at September 30, 2006 and December 31, 2005 were as follows:

	September 30, 2006	December 31, 2005
Raw materials	\$1,307,296	\$ 1,139,730
Work in process	1,406,072	1,267,991
Finished goods	664,619	589,111
Total	\$ 3,377,987	\$ 2,996,832

NOTE 3. INTANGIBLE ASSETS

At September 30, 2006 and December 31, 2005 intangible assets consisted of the following:

	September 30, 2006	December 31, 2005	
Goodwill acquired: Trade name (life indefinite)	\$ 1,509,662	\$ 1,509,662	
Other intangibles:			
License rights (20 yr life)	400,000	400,000	
Less: Accumulated amortization	(150,000)	(120,000)	
Net carrying amount	250,000	280,000	

Purchased technology (5 yr \$ 587,267 \$ 280,764

life)

Less: Accumulated amortization (271,183) (247,101)

Net carrying amount \$

316,084 \$ 33,663

NOTE 4. NEW ACCOUNTING PRONOUNCEMENTS

FASB Interpretation No. 46R, Consolidation of Variable Interest Entities - An Interpretation of ARB51

The FASB finalized FIN 46R in December 2003. FIN 46R expands the scope of ARB51 and various EITFs and can require consolidation of legal structures, called *Variable Interest Entities (VIEs)*. Companies with investments in *Special Purpose Entities (SPEs)* were required to implement FIN 46R in 2003; however, companies with VIEs were permitted to implement in the first quarter of 2004. While we do not have SPEs, we do have a VIE that we have determined will qualify for consolidation. Our joint venture with Jump Agentur Fur Electrotechnik GMBH ("the Joint Venture", "JAG") qualifies as a VIE. We have consolidated this VIE for the period ended September 30, 2006 and for the year ended December 31, 2005. The most significant impact to our financial statements is to include the net intangible assets of JAG, totaling \$250,000 for the period ended September 30, 2006, and minority interest of \$125,000 as of September 30, 2006 to our balance sheets. The impacts on our consolidated statements of net income or cash flows are not material.

In November 2004, the FASB issued SFAS No. 151, "Inventory Costs, an amendment of Accounting Research Bulletin No. 43, Chapter 4," which adopts wording from the International Accounting Standards Board's (IASB) IAS 2 "Inventories" in an effort to improve the comparability of cross-border financial reporting. The FASB and IASB both believe the standards have the same intent; however, an amendment to the wording was adopted to avoid inconsistent application. The new standard indicates that abnormal freight, handling costs, and wasted materials (spoilage) are required to be treated as current period charges rather than as a portion of inventory cost. Additionally, the standard clarifies that fixed production overhead should be allocated based on the normal capacity of a production facility. The Statement is effective beginning in fiscal year 2007. Adoption is not expected to have a material impact on our consolidated earnings, financial position or cash flows.

In December 2004, the FASB issued FSP FAS 109-1, "Application of FASB Statement No. 109, Accounting for Income Taxes, to the Tax Deduction on Qualified Production Activities Provided by the American Jobs Creation Act of 2004." The FSP clarifies that the manufacturer's deduction provided for under the American Jobs Creation Act of 2004 (the Act) should be accounted for as a special deduction in accordance with SFAS No. 109, "Accounting for Income Taxes," and not as a tax rate reduction. The Qualified Production Activities Deduction will not impact our consolidated earnings, financial position or cash flows for fiscal year 2006 because the deduction is not available to us. We are currently evaluating the effect that this deduction will have in subsequent years.

BOVIE MEDICAL CORPORATION NOTES TO FINANCIAL STATEMENTS

NOTE 4. NEW ACCOUNTING PRONOUNCEMENTS (Continued)

Accounting for Stock-Based Compensation

Effective January 1, 2006, the Company adopted Statement No. 123R, Share-Based Payment ("SFAS 123R"), which requires companies to measure and recognize compensation expense for all share-based payment awards made to employees and directors based on estimated fair values. SFAS 123R is being applied on the modified prospective basis. Prior to the adoption of SFAS 123R, the Company accounted for its stock-based compensation plans under the recognition and measurement principles of Accounting Principles Board (APB) Opinion No. 25, Accounting for Stock Issued to Employees, as provided by SFAS 123, "Accounting for Stock Based Compensation" ("SFAS 123") and accordingly, recognized no compensation expense related to the stock-based plans as stock options granted to employees and directors were equal to the fair market value of the underlying stock at the date of grant. In March 2005, the SEC issued Staff Accounting Bulletin No. 107 ("SAB 107") relating to SFAS 123R. The Company has applied the provisions of SAB 107 in its adoption of SFAS 123R.

Under the modified prospective approach, SFAS 123R applies to new awards and to awards that were outstanding on January 1, 2006 that are subsequently modified, repurchased or cancelled. Under the modified prospective approach, compensation cost recognized includes compensation cost for all share-based payments granted prior to, but not yet vested on, January 1, 2006, based on the grant-date fair value estimated in accordance with the provisions of SFAS 123, and compensation cost for all share-based payments granted subsequent to January 1, 2006, based on the grant-date fair value estimated in accordance with the provisions of SFAS 123R. Prior periods were not restated to reflect the impact of adopting the new standard. During the first, second, and third quarters for 2006, the Company recorded \$0, \$11,305 and \$9,085, respectively, in non-cash charges for the implementation of SFAS 123R. As of September 30, 2006, there was approximately \$76,030 of total unrecognized compensation costs related to unvested options. That cost is expected to be recognized over a weighted average period of 3.5 years.

The weighted average grant date fair value of options granted during the nine months ended September 30, 2006 were estimated on the grant date using the binomial lattice option-pricing model with the following assumptions: expected volatility of 25%, expected term of 5 years, risk-free interest rate of 5.0%, and expected dividend yield of 0%. Expected volatility is based on a weighted average of the historical volatility of the Company's stock and peer company volatility. The average expected life was calculated using the simplified method under SAB 107. The risk-free rate is based on the rate of U.S. Treasury zero-coupon issues with a remaining term equal to the expected life of option grants. The Company uses historical data to estimate pre-vesting forfeiture rates.

Allocation of non-cash stock based compensation expense for the nine months ended September 30, 2006:

September 30, 2006

Cost of Sales \$ 1,101

Research and Development	6,725
Salaries and related costs	12,564

Total \$20,390

NOTE 5. SHAREHOLDERS' EQUITY

During the nine-month period ending September 30, 2006, we issued 663,710 common shares in exchange for employee and non-employee exercised options. The issuance of the common stock resulted in an increase in capital of \$494,025.

NOTE 6. EARNINGS PER SHARE

We compute basic earnings per share ("basic EPS") by dividing net income by the weighted average number of common shares outstanding for the reporting period. Diluted earnings per share ("diluted EPS") gives effect to all dilutive potential shares outstanding resulting from employee stock options during the period. The following table sets forth the computation of basic and diluted earnings per share for the nine month periods ended September 30, 2006 and 2005.

Nine months ended September 30				
	2006	2005		
Net income	\$ 2,260	\$ 142		
Basic-weighted average shares outstanding	14,351	13,908		
Effect of dilutive potential securities	2,544	2,112		
Diluted - weighted average shares outstanding	16,895	16,020		
Basic EPS	\$.16	\$.01		
Diluted EPS	\$.13	\$.01		

All above figures are in thousands except basic and diluted earnings per share which are not. The shares used in the calculation of diluted EPS exclude options to purchase shares where the exercise price was greater than the average market price of common shares for the period. Such shares aggregated approximately zero and .5 million in the nine months ended September 30, 2006 and 2005, respectively.

NOTE 7. Endoscopic Modular Instruments

In January, 2006, pursuant to agreement to acquire technology from Henvil Corp. LTD and Steve Livneh, its principal, we acquired patent pending technology for new endoscopic disposable and reusable modular instruments ("the Product") from Henvil Corp. LTD. ('Henvil"). The innovative modular forceps are ergonomically designed to provide surgeons' added comfort and improved safety while reducing per-procedure costs. The modular forceps offer an innovative and simpler assembly process for laparoscopic procedures and is a modular design for the arthroscopy market. Commercial prototypes have been developed and based on current projections we expect to commence marketing during fiscal 2006. The estimated annual worldwide market size for instruments of these categories is estimated to exceed \$200 million.

The agreement requires us to purchase certain equipment and machinery (\$400,000.00 value) and to retain Henvil to develop the technology and complete the design of the arthroscopic and laparoscopic instruments. In addition we must also purchase tools and molds expected to cost approximately \$120,000. Henvil and Steve Livneh have also been hired as consultants for the development of the technology. The consultants are to be paid \$30,000 per month for twelve months from the day of execution of the agreement. In addition, commencing with the year following the first sale or commercial delivery of the Product, Bovie shall pay to Henvil's principal, Steve Livneh, an initial minimum

royalty of the greater of \$35,000 per year or 3% of adjusted gross revenues received from the sale and marketing of the instruments. Thereafter, a royalty equal to 2.5% or adjusted gross sales for the life of the patents issuable for the technology.

Mr. Livneh also shall receive 50,000 5-year restricted stock options to purchase Bovie common stock for each category of instrumentation (a total of 100,000 stock options) exercisable at the closing price of Bovie common stock on the American Stock Exchange on the date of the contract. The respective options shall vest independently and immediately subject to Bovie receiving FDA clearance for marketing each of the products. We have received FDA 510k clearance to market the modular laparoscopic instruments.

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

Executive Level Overview

We are a medical device company engaged in the manufacturing and marketing of electrosurgical devices. Our medical products include a wide range of devices including electrosurgical generators and accessories, cauteries, medical lighting, nerve locators and other products.

We divide our operations into three reportable business segments: Electrosurgical products, battery operated cauteries and other products. The electrosurgical segment manufactures and markets electrosurgical generators, electrodes, electrosurgical pencils and various ancillary disposable products. These products are used in surgery for the cutting and coagulation of tissue. Battery operated cauteries are used for precise hemostasis (to stop bleeding) in ophthalmology and in other fields. Our other revenues are derived from nerve locators, disposable and reusable penlights, medical lighting, license fees, development fees and other miscellaneous income. Domestic sales accounted for 88% of total revenues for the nine months ended September 30, 2006 as compared to 84% in the first nine months of 2005. Most of the Company's products are marketed through medical distributors which distribute to a market of more than 6,000 hospitals and to doctors and other health-care facilities.

International sales were slightly lower in the first nine months of 2006 and accounted for 12% of total revenues for that period, as compared to 16% for the period ended September 30, 2005. The Company's products are sold in more than 150 countries through local dealers. Local dealer support is coordinated by sales and marketing personnel at the St. Petersburg, Florida facility. We have no branch offices other than the Florida facility. We sell our products to distributors that distribute them in the following countries: Argentina, Australia, Belgium, Brazil, Canada, Chile, Denmark, Finland, France, Germany, Greece, India, Italy, Japan, Korea, Mexico, The Netherlands, New Zealand, Norway, Poland, Portugal, Singapore, South Africa, Spain, Sweden, Switzerland, Taiwan, United Kingdom, China, the CIS (former Soviet Union), Cyprus, Indonesia, Ireland, Latin America, Malaysia, the Philippines, Thailand, Turkey, and Vietnam. Our business is generally not seasonal in nature.

Outlook for 2006

The Company's outlook for the fiscal year ending December 31, 2006 continues positive, as we expect revenues to be at a record level which will translate into substantially improved net earnings when compared to fiscal year 2005. Our optimism is based on our sales during the first ten months of fiscal year 2006 and on current budget estimates. The outlook excludes any sales which may be generated by Bovie's new products, some of which we expect to begin marketing though not necessarily impacting sales during the current year. As part of our growth, we continue experiencing higher costs in selling, payroll, professional fees, and administrative costs. Growth, to a significant degree, is being fueled by higher OEM sales and generally increased sales in most product areas. The markets for our products are highly competitive; however, we believe that our competitive advantage is rooted in our ability to offer high quality products that meet changing demand. In addition, we offer the flexibility, quality of products and responsiveness that a smaller company can offer.

Fiscal year 2006 featured new product introductions either by Bovie alone, and/or in collaborations with other larger medical companies. The shifting away from being highly dependent on OEM contracts and toward designing, developing and marketing our own brand of products, is accelerating and we anticipate that this will result in improved margins in the future. We have received 510k clearances to market the suture removal device, the ICON GI and the modular laparoscopic instruments.

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS(CONTINUED)

Result of Operations (to be read in conjunction with the profit and loss statement)

The table below outlines the components of the consolidated statements of earnings as a percentage of net sales and the year-to-year percentage change in dollar amounts:

Analysis of Quarter Ended September 30, 2006/2005

	3 rd Quarter		Percentage Change in Dollar Amounts Nine m		onths	Percentage Change in Dollar amounts
	2006	2005		2006	2005	
	%	%	%	%	%	%
Sales	100.0	100.0	39.0	100.0	100.0	33.0
Cost of sales	59.0	58.0	41.0	60.0	64.0	24.0
Gross profit	41.0	42.0	36.0	40.0	36.0	50.0
Other costs:						
R & D	4.0	6.0	(14.0)	3.0	4.0	(4.0)
Professional fees	2.0	2.0	22.0	2.0	2.0	11.0
Salaries	9.0	9.0	45.0	9.0	10.0	26.0
SGA	13.0	21.0	(8.0)	14.0	18.0	5.0
Equity in loss of	1.0	1.0	(1.0)	1.0	1.0	27.0
Unconsolidated affiliate						
Total other costs	29.0	39.0	5.0	29.0	35.0	11.0
Gain/ from operations	12.0	3.0	408.0	11.0	1.0	1,787.0
Other income	0.0	0.0	416.0	0.0	0.0	202.0
Net Income before Minority Interest and Income Tax	12.0	3.0	408.0	11.0	1.0	1,589.0
Minority Interest	0.0	0.0	100.0	0.0	0.0	100.0
Income tax expense	(4.0)	(1.0)	341.0	(4.0)	(0.0)	1,334.0
Income tax benefit	4.0	1.0	329.0	4.0	0.0	1,274.0

Net earnings 12.0 3.0 399.0 11.0 1.0 1,488.0

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS(CONTINUED)

The table below sets forth domestic/international and product line sales information for the third quarter of 2006 and 2005.

Net Sales (in thousands)		F	Percentage change	Increase/
	2006	2005	2006/2005	(Decrease)
Domestic/international sales:				
Domestic	\$ 6,160	\$ 4,138	49.0	2,022
International	839	901	(7.0)	(-62)
Total net sales	\$ 6,999	\$ 5,039	39.0	1,960
Product line sales:				
Electrosurgical	\$ 5,041	\$ 3,388	49.0	1,653
Cauteries	1,465	1,429	3.0	36
Other	493	222	17.0	72
Total net sales	\$ 6,999	\$ 5,039	39.0	1,960

The table below sets forth domestic/international and product line sales information for the nine months of 2006 and 2005.

Net Sales (in thousands)	2006	2005	Percentage change 2006/2005	Increase/ (Decrease)	
Domestic/international sales:				(= :::::)	
Domestic	\$ 17,278	\$ 12,330	40.0	4,948	
International	2,473	2,510	(1.0)	(37)	
Total net sales	\$ 19,751	\$ 14,840	33.0	4,911	
Product line sales:					
Electrosurgical	\$ 13,435	\$ 8,613	56.0	4,822	
Cauteries	4,343	4,031	8.0	312	
Other	1,973	2,196	(10.0)	(223)	
Total net sales	\$ 19,751	\$ 14,840	33.0	4,911	

Nine months ended September 30, 2006 compared to nine months ended September 30, 2005

The results of operations for the nine months ended September 30, 2006 show increased sales and increased profitability, as compared to the first nine months of 2005. Sales were \$19.8 million for the nine months ended September 30, 2006 as compared to \$14.8 million for the same period of 2005, an increase of 33% or \$5.0 million. Sales of electrosurgical products increased by 56% or \$4.8 million compared to the same period of 2005 while sales of cauteries increased by 8% from \$4.0 million to \$4.3 million. Other sales decreased by 10% from \$2.2 million to \$2.0 million. Over the last nine months we have instituted price increases on cauteries and other products of 3%. No sales of one particular electrosurgical product dominated the number of units sold.

Arthrex sales of generators and accessories increased \$2.3 million or 112% from \$2.05 million in the first nine months of 2005 to \$4.4 million in the first nine months of 2006.

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS(CONTINUED)

Domestic sales were \$17.3 million for first nine months of 2006, representing an increase of 40% from the same period last year. International sales were \$2.5 million for the first nine months of 2006, representing a decrease of 1% over the same period 2005.

Cost of sales represented 60% of sales in the first nine months of 2006 as compared to 64% of sales in the first nine months of 2005, a total of \$11.7 million and \$9.5 million, respectively, an increase of \$2.2 million. The reason for the decrease in cost of sales percentage was due to a decrease of 5.7% in indirect costs as a percentage of sales and a decrease in labor cost as a percentage of sales of 2.72%. As our sales have increased our indirect costs and labor costs as a dollar amount have not increased in the same manner and have remained relatively constant. Material cost as a percentage of sales increased slightly from 32% for the first nine months of 2005 to 33% for the first nine months of 2006.

Research, development and engineering expenses were 3.0% and 4.0% of sales for the first nine months of 2006 and 2005, respectively. These expenses, as a dollar amount, decreased 4% in 2006 to \$631,674 a decrease over the corresponding period of 2005 of \$26,265. New products under development are the modular forceps instruments, suture removal device, plasma technology, GI device and various improvements to our line of electrosurgical generators.

Professional fees increased from \$361,495 in the first nine months of 2005 to \$400,945 in the first nine months of 2006, an increase of \$39,449 or 11%. The company had increased legal costs in patent research and filings for some of the new products under development for first nine months ended September 30, 2006 compared to the previous year's first nine months.

Salaries and related costs increased in the first nine months of 2006 by 26% to \$1.8 million as compared to the first nine months of 2005 at \$1.5 million. The increase was mainly attributable to additional employees and annual salary increases needed to foster the growth of the company in various areas.

Selling, general and administrative expenses increased by 5.0% for the first nine months of 2006 as compared to the first nine months of 2005, to a total of \$2.8 million.

Total costs and expenses went from \$5,231,862 for the nine months ended September 30, 2005 to \$5,781458 for the same period in 2006, an increase of \$549,596 or 11%.

Net interest earned increased by \$33,998 during the first nine months of 2006 when compared to the first nine months of 2005 as a result of our higher cash balances being invested and yielding higher interest rates.

The effective income tax rate was 36% in the first nine months of 2006 and the first nine months of 2005. There was also a tax loss carryover benefit of 35.6 % for the first nine months of 2006 and 36% for first nine months of 2005. The difference between the income tax and the tax loss carryover benefit for the first nine months of 2006 is \$32,567, an estimated amount for the AMT (alternative minimum tax).

Diluted net earnings increased \$.12 to \$.13 per share or \$2,259,899 in the first nine months of 2006 as compared to a net income of \$142,306 or \$.01 per share in the first nine months of 2005. The increase in earnings of the first nine

months of 2006 over the first nine months of 2005 was mostly attributable to an increase in electrosurgical sales while labor costs and indirect costs of sales as a dollar amount remained constant.

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS(CONTINUED)

Results of Operations- Three months ended September 30, 2006 and September 30, 2005

Sales for the three month period ended September 30, 2006 were \$7.0 million as compared to \$5.0 million for the same period in 2005, an increase of \$2.0 million or 39%. The increase was mainly attributed to increased sales of electrosurgical products.

Cost of goods sold increased from \$2.9 million to \$4.1 million an increase of \$1.2 million or 41% for the three month period ended September 30, 2006 as compared to the same period in 2005.

Gross profit increased from \$2.1 million to \$2.9 million an increase of \$.8 million or 36%. Gross profit percentage decreased from 42% in 2005 to 41% in 2006. The reason for the decrease was due to a slight increase in materials cost, as a percentage of sales, allocated to the product mix sold during the quarter. Factory overhead costs and labor costs, as a dollar amount, remained the same.

Research and development decreased by \$41,927 or 14% from \$303,850 to \$261,923 for the quarters ended September 30, 2005 and September 30, 2006, respectively. The decrease is due to reduced costs for new products under development as they approach completion (i.e. modular forceps instruments, suture removal device, and GI device).

Professional fees increased by \$27,418 or 22% from \$126,846 to \$154,264 for the quarters ended September 30, 2005 to September 30, 2006, respectively. This increase is mainly attributed to increased legal costs in patent research and filings for some of the new products under development.

Salaries and related costs increased from \$447,210 to \$647,002 for the quarters ended September 30, 2005 to September 30, 2006, respectively, an increase of \$199,792 or 45%. This increase was mainly attributable to adding employees and salary increases needed to foster the growth of the company.

Selling, general and administrative expenses decreased by \$85,965 or 8% from \$1,038,418 to \$952,453 for the quarters ended September 30, 2005 to September 30, 2006, respectively. The largest areas of increased costs were for commissions and liability insurance.

Total costs and expenses went from \$1,951,214 for the three months ended September 30, 2005 to \$2,050,148 for the same period in 2006, an increase of \$98,934 or 5%.

Net interest income increased from \$5,594 to \$28,872 in income for the quarter ended September 30, 2006 as compared to quarter ended September 30, 2005. The increase is a direct result from the investment of our higher cash balances yielding higher interest rates.

Net income for the three months ended September 30, 2006 was \$856,959 or \$.05 diluted earnings per share as compared to net income of \$171,620 for the same quarter in 2005, an increase of \$685,339 or 399%. The main reason for the increase in net income of the three months ended September 30, 2006 over the same period for 2005 was mostly attributable to an increase in electrosurgical sales coupled with labor and indirect costs, as a dollar amount of

cost of goods sold, which remained relatively unchanged.

Marketing and Sales

We sell our products through distributors both overseas and in U.S. markets. New distributors are contacted through responses to our advertising in domestic and international medical journals and domestic or international trade shows.

We have agreements with various sales representatives to develop markets for our new products and to maintain customer relations. Our current representatives receive an average commission of approximately 4% of sales in

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS(CONTINUED)

their market areas. In the first nine months of 2006 and 2005, commissions paid were \$448,956 and \$328,444 respectively, an increase of 37%.

An adequate supply of raw materials is available from both domestic and international suppliers. The relationship between us and our suppliers is generally limited to individual purchase order agreements, supplemented by contractual arrangements with key vendors to ensure availability of certain products. We have developed multiple sources of supply where possible.

In order to provide additional working capital, we have secured a \$1.5 million credit facility with Bank of America. This facility is payable on demand. For the period ended September 30, 2006, we had zero funds drawn down on this credit facility.

Our ten largest customers accounted for approximately 72% of net revenues for the first nine months of 2006 as compared to 64% in the same period of 2005. For both periods ended September 30, 2006 and 2005, our ten largest trade receivables accounted for approximately 78% and 60% of outstanding receivables, respectively. In the first nine months of 2006 and 2005 one customer accounted for 22% and 14% of total sales, respectively.

Product Development

Most of the Company's products and product improvements have been developed internally. Funds for this development have come from internal cash flow and the issuance of common stock upon the exercise of stock options. The Company maintains close working relationships with physicians and medical personnel in hospitals and universities who assist in product research and development. New and improved products play a critical role in the Company's sales growth. The Company continues to place emphasis on the development of proprietary products and product improvements to complement and expand its existing product lines. The Company has a centralized research and development focus, with its one manufacturing location responsible for new product development and product improvements. Our research, development and engineering units at the manufacturing location maintain relationships with distribution locations and customers in order to provide an understanding of changes in the market and product needs. During 2005 and into 2006 we invested in the J Plasma Technology, the Suture Removal Technology, the Gastrointestinal "GI" device and undertook development of Cardio and Urological Electrosurgical devices for a contractual partner. The suture removal device, the GI device, modular laparoscopic instruments and the Bovie Button are being marketed although no significant sales are anticipated in 2006. The ongoing cost for this development will be paid from operating cash flows.

In the next year we do not contemplate any material purchase or acquisition of assets which our ordinary cash flow and or credit line would not be able to sustain.

We believe that Bovie has the financial resources needed to meet business requirements in the foreseeable future, including capital expenditures needed for the expansion of our manufacturing site, working capital requirements, and product development programs, subject to Bovie maintaining compliance with our credit facility.

Reliance on Collaborative, Manufacturing and Selling Arrangements

We are dependent on certain contractual OEM customers for product development wherein we are to provide the manufacturing of the product developed. However, the customers have no legal obligation to purchase the developed products. Should the collaborative customer fail to give us purchase orders for the product after development, our future business and value of related assets could be negatively affected. Furthermore, no assurance can be given that a collaborative customer may give sufficient high priority to our products. In addition, disagreements or disputes may arise between Bovie and its contractual customers which could adversely affect production of our products. We also have similar informal collaborative arrangements with two foreign suppliers except that we request the development of certain items and components and we purchase them from the foreign supplier pursuant to purchase orders. Our purchase orders are never for more than one year and are supported by customer purchase orders from our customers.

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS(CONTINUED)

Liquidity and Capital Resources

Our working capital at September 30, 2006 increased \$2.0 million to \$7.5 million from \$5.5 million at December 31, 2005. The increase in working capital was primarily a result of cash provided from operating activities. Accounts payable and other accrued liabilities together increased \$.2 million in the first nine months of 2006 as a result of the growth in the business. Accounts receivable day sales outstanding were 47.3 days and 45.8 days at September 30, 2006 and September 30, 2005 respectively.

We generated cash from operations of \$1.9 million for the nine months ended September 30, 2006 compared with applying cash to operations of \$.27 million in the same period of 2005. The increase in cash from operations for the period ended September 30, 2006 compared to the prior year is primarily due to the increase in sales volume.

In the first nine months ended September 30, 2006 we applied \$.9 million for the purchase of fixed assets. Total borrowing decreased by \$258,939, which was the net amount of paying off our first mortgage coupled with proceeds received from the exercise of stock options and the incurred debt from short term financing of insurance premiums.

We had \$2.3 million in cash and cash equivalents at September 30, 2006. We also had outstanding borrowings totaling \$.2 million at that date which all consisted of current note payable debt at September 30, 2006. We believe our cash on hand, as well as anticipated cash flows from operations, will be sufficient to fund future operating capital requirements, future manufacturing facility construction and other capital expenditures and future acquisitions to supplement our current product offerings. Should additional funds be required, we have \$1.5 million of additional borrowing capacity available under our existing credit facility.

The Company's future contractual obligations for agreements with initial terms greater than one year, including agreements to purchase materials in the normal course of business, are summarized as follows (in thousands):

As o	of September		Payment Pe	eriod	
30,					
	2006	2007	2008	2009	2010
Current debt	137	66	-0-	-0-	-0-
Operating leases	36	135	115	-0-	-0-
Unconditional purchase	1,009	3,026	-0-	-0-	-0-
obligations					

The Company's additional borrowing capacity, along with the expected expiration period of the commitments, is summarized as follows (in millions):

				Amount of Commitment			
	Total			Expiration Per Period			
	Amount Committed			Less t	han	In excess of	
			1 year		1 year		
Secured revolving credit agreement and other lines							
of credit	\$		1.5	\$	1.5	-0-	

As of September 30, 2006 the total amount is available.

Our future results of operations and the other forward-looking statements contained herein, particularly the statements regarding growth in the medical products industry, capital spending, research and development, and marketing and general and administrative expenses, involve a number of risks and uncertainties. In addition to the factors discussed above, there are other factors that could cause actual results to differ materially, such as business conditions and the general economies; competitive factors including rival manufacturers' availability of components at reasonable prices; risk of nonpayment of accounts receivable; risks associated with foreign operations; and litigation involving intellectual property and consumer issues.

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS(CONTINUED)

We believe that we have the product mix, facilities, personnel, competitive edge, operating cash flows and financial resources for business success in the immediate (1 year) future and distant future (after 1 year), but future revenues, costs, margins, product mix and profits are all subject to the influence of a number of factors, as discussed above.

Critical Accounting Estimates

We have adopted various accounting policies to prepare the consolidated financial statements in accordance with accounting principles generally accepted (GAAP) in the United States of America (U.S.).

The preparation of the consolidated financial statements, in conformity with U.S. GAAP, requires us to make estimates and assumptions that affect the amounts reported in the consolidated financial statements and accompanying notes. Our estimates and assumptions, including those related to bad debts, inventories, intangible assets, property, plant and equipment, minority investment, legal proceedings, research and development, warranty obligations, product liability, sales returns and discounts, and income taxes are updated as appropriate, which in most cases is at least quarterly. We base our estimates on historical experience, actuarial valuations, or various assumptions that are believed to be reasonable under the circumstances and the results form the basis for making judgments about the reported values of assets, liabilities, revenues and expenses. Actual results may materially differ from these estimates.

Estimates are considered to be critical if they meet both of the following criteria: (1) the estimate requires assumptions about material matters that are uncertain at the time the accounting estimates are made, and (2) other materially different estimates could have been reasonably made or material changes in the estimates are reasonably likely to occur from period to period. Our critical accounting estimates include the following:

Allowance for doubtful accounts

We maintain an allowance for doubtful accounts for estimated losses in the collection of accounts receivable. We make estimates regarding the future ability of our customers to make required payments based on historical credit experience and expected future trends. If actual customer financial conditions are less favorable than projected by management, additional accounts receivable write-offs may be necessary, which could unfavorably affect future operating results.

Inventory Reserves

We maintain reserves for excess and obsolete inventory resulting from the potential inability to sell our products at prices in excess of current carrying costs. The markets in which we operate are highly competitive, with new products and surgical procedures introduced on an ongoing basis. Such marketplace changes may cause our products to become obsolete. We make estimates regarding the future recoverability of the costs of these products and record a provision for excess and obsolete inventories based on historical experience, and expected future trends. If actual product life cycles, product demand or acceptance of new product introductions are less favorable than projected by management, additional inventory write-downs may be required, which could unfavorably affect future operating results.

Insurance Deductible Reserves

We maintain reserves for fifty percent of our maximum insurance deductible exposure related to our product liability insurance coverage. This reserve amount is estimated using a historical average for the Company.

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS(CONTINUED)

Income Taxes

We operate in multiple tax jurisdictions both inside and outside the United States. Accordingly, management must determine the appropriate allocation of income to each of these jurisdictions. Tax audits associated with the allocation of this income and other complex issues may require an extended period of time to resolve and may result in income tax adjustments if changes to the income allocation are required between jurisdictions with different tax rates. Because tax adjustments in certain jurisdictions can be significant, we record accruals representing our best estimate of the probable resolution of these matters. To the extent additional information becomes available, such accruals are adjusted to reflect the revised estimated probable outcome.

Other Matters

We distribute our products throughout the world. As a result, our financial results could be significantly affected by factors such as changes in foreign currency exchange rates or weak economic conditions in foreign markets. Our operating results are primarily exposed to changes in exchange rates among the United States dollar and European currencies, in particular the Euro and the British pound. When the United States dollar weakens against foreign currencies, the dollar value of sales denominated in foreign currencies increases. When the United States dollar strengthens, the opposite situation occurs. We manufacture our products in the United States, China and Bulgaria and incur the costs to manufacture in US dollars. This worldwide deployment of factories serves to partially mitigate the impact of the high costs of manufacturing in the US.

ITEM 3. CONTROLS AND PROCEDURES

(a) Evaluation of disclosure controls and procedures

An evaluation of the effectiveness of the design and operation of the Company's disclosure controls and procedures [as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)] as of September 30, 2006 was carried out under the supervision and with the participation of the Company's management, including the President and Chief Executive Officer and the Chief Financial Officer ("the Certifying Officers"). Based on that evaluation, the Certifying Officers concluded that the Company's disclosure controls and procedures are effective.

Disclosure controls and procedures are designed to ensure that information required to be disclosed in our reports filed or submitted under the Securities Exchange Act is recorded, processed, summarized and reported within the time periods specified in the SEC's rules and forms. Disclosure controls and procedures include, without limitation, controls and procedures designed to ensure that information required to be disclosed in our reports filed or submitted under the Securities Exchange Act is accumulated and communicated to management, including our President and Chief Financial Officer, as appropriate, to allow timely decisions and timely reporting regarding required disclosure.

(b) Changes in internal controls

There was no change to the Company's internal control over financial reporting during the quarter ended September 30, 2006 that materially affected, or is reasonably likely to materially affect, the Company's internal control over financial reporting.

PART II. OTHER INFORMATION

ITEM 1. LEGAL PROCEEDINGS

There were no legal proceedings during the quarterly period ended September 30, 2006 pending that could have a material effect on our financial position.

ITEM 2. UNREGISTERED SALES OF EQUITY SECURITIES AND OF PROCEEDS

None

ITEM 3. DEFAULTS ON SENIOR SECURITIES HOLDERS

None

ITEM 4. SUBMISSION OF MATTERS TO VOTE BY SECURITY HOLDERS

None

ITEM 5. OTHER INFORMATION

The Company filed a Form 510-K application, which has since been approved, with the Food and Drug Administration (FDA) for its "In-a-Flash" Suture Removal Device which is designed to remove sutures with a tension free cut. This device is to be utilized in various human and animal medical procedures.

The Company has received 510-K approval to market its ICON GI and modular laparoscopic instruments.

ITEM 6. EXHIBITS

- 31.1 Certifications of Andrew Makrides, President and Chief Executive Officer of Registrant pursuant to Rule 13a-14 adopted under the Securities Exchange Act of 1934, as amended, and Section 302 of the Sarbanes-Oxley Act of 2002.
- Certifications of Gary D. Pickett, Chief Financial Officer of Registrant pursuant to Rule 13a-14 adopted under the Securities Exchange Act of 1934, as amended, and Section 302 of the Sarbanes-Oxley act of 2002.
- 32.1 Certification pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
- 32.2 Certification pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.

SIGNATURES:

In accordance with the requirements of the Exchange Act, the registrant caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

Bovie Medical Corporation. (Registrant)

Date: November 7, 2006

/s/Andrew Makrides Chief Executive Officer - Andrew Makrides

/s/Gary D. Pickett Chief Financial Officer- Gary D. Pickett