
**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549**

FORM 10-Q

(Mark One)

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

For the quarterly period ended September 30, 2010

or

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

For the transition period from _____ to _____

Commission file number: 0-16159

LECTEC CORPORATION

(Exact Name of Registrant as Specified in Its Charter)

Minnesota

(State or Other Jurisdiction of
Incorporation or Organization)

41-1301878

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

1407 South Kings Highway, Texarkana, TX

(Address of Principal Executive Offices)

75501

(Zip Code)

(903)-832-0993

(Registrant's telephone number, including area code)

Not Applicable

(Former Name, Former Address and Former Fiscal Year, if Changed Since Last Report)

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

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Web site, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit and post such files). YES NO

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, or a smaller reporting company. See the definitions of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act.

Large accelerated filer Accelerated filer Non-Accelerated filer Smaller reporting company
(Do not check if a smaller reporting company)

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). YES NO

As of November 15, 2010 the registrant had 4,305,026 shares of common stock outstanding.

LECTEC CORPORATION
REPORT ON FORM 10-Q FOR THE QUARTERLY PERIOD ENDED SEPTEMBER 30, 2010
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Forward-Looking Statements

From time to time, in reports filed with the Securities and Exchange Commission (including this Form 10-Q), in press releases, and in other communications to shareholders or the investment community, we may provide forward-looking statements concerning possible or anticipated future results of operations or business developments which are typically preceded by the words “believes,” “wants,” “expects,” “anticipates,” “intends,” “will,” “may,” “should,” or similar expressions. Such forward-looking statements are subject to risks and uncertainties which could cause results or developments to differ materially from those indicated in the forward-looking statements. Such risks and uncertainties include, but are not limited to, our dependence on royalty payments from Novartis Consumer Health, Inc., which is selling an adult vapor patch licensed by us, our dependence on key personnel and Board of Director members, the success or failure of any attempt by us to protect or enforce our patents and territories of coverage, the outcome of pending patent infringement litigation against Chattem, Inc., and Prince of Peace Enterprises, Inc., the issuance of new accounting pronouncements, the availability of opportunities for license, sale or strategic partner agreements related to patents that the Company holds, limitations on market expansion opportunities, and other risks and uncertainties as described in “Risk Factors” included in Item 1A as filed in our Form 10-K for the year ended December 31, 2009.

PART 1. FINANCIAL INFORMATION**ITEM 1. CONDENSED FINANCIAL STATEMENTS AND NOTES TO CONDENSED FINANCIAL STATEMENTS****LECTEC CORPORATION
CONDENSED BALANCE SHEETS**

| | September 30 2010 (Unaudited) | December 31, 2009 |
|--|-------------------------------------|----------------------------|
| ASSETS | | |
| CURRENT ASSETS: | | |
| Cash and cash equivalents | \$ 9,283,697 | \$15,766,107 |
| Royalty receivable | 26,026 | 31,525 |
| Prepaid expenses and other | 634,476 | 975,423 |
| Deferred tax asset | <u>409,000</u> | <u>—</u> |
| Total current assets | <u>10,353,199</u> | <u>16,773,055</u> |
| FIXED ASSETS: | | |
| Office equipment | 9,847 | 8,590 |
| Accumulated depreciation | <u>(5,404)</u> | <u>(3,021)</u> |
| | <u>4,443</u> | <u>5,569</u> |
| OTHER ASSETS: | | |
| Patent costs | <u>47,178</u> | <u>29,811</u> |
| TOTAL ASSETS | <u>\$10,404,820</u> | <u>\$16,808,435</u> |
| LIABILITIES AND SHAREHOLDERS' EQUITY | | |
| CURRENT LIABILITIES: | | |
| Accounts payable | \$ 89,089 | \$ 84,659 |
| Accrued expenses | 51,973 | 322,854 |
| Dividend payable | — | 4,298,350 |
| Income tax payable | — | 993,403 |
| Deferred tax liability | <u>—</u> | <u>48,000</u> |
| Total current liabilities | <u>141,062</u> | <u>5,747,266</u> |
| COMMITMENTS AND CONTINGENCIES | | |
| SHAREHOLDERS' EQUITY: | | |
| Common stock, \$.01 par value; 15,000,000 shares authorized; 4,305,026 and 4,290,026 shares issued and outstanding at September 30, 2010 and December 31, 2009, respectively | 43,050 | 42,900 |
| Additional contributed capital | 11,161,935 | 11,018,269 |
| Accumulated deficit | <u>(941,227)</u> | <u>—</u> |
| | <u>10,263,758</u> | <u>11,061,169</u> |
| TOTAL LIABILITIES AND SHAREHOLDERS' EQUITY | <u>\$10,404,820</u> | <u>\$16,808,435</u> |

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

LECTEC CORPORATION
CONDENSED STATEMENTS OF OPERATIONS
(Unaudited)

| | Three Months Ended September 30, | | Nine Months Ended September 30, | |
|--|-------------------------------------|---------------------|------------------------------------|---------------------|
| | 2010 | 2009 | 2010 | 2009 |
| REVENUE: | | | | |
| Royalty and licensing fees | \$ 26,373 | \$ 24,215 | \$ 49,156 | \$ 76,633 |
| Infringement income | — | — | — | 600,000 |
| TOTAL REVENUE | <u>26,373</u> | <u>24,215</u> | <u>49,156</u> | <u>676,633</u> |
| OPERATING EXPENSES | <u>639,878</u> | <u>155,221</u> | <u>1,376,230</u> | <u>803,809</u> |
| Loss from operations | (613,505) | (131,006) | (1,327,074) | (127,176) |
| INTEREST INCOME | <u>4,836</u> | <u>32</u> | <u>13,089</u> | <u>732</u> |
| LOSS BEFORE INCOME TAXES | (608,669) | (130,974) | (1,313,985) | (126,444) |
| INCOME TAX BENEFIT | <u>143,758</u> | <u>—</u> | <u>372,758</u> | <u>—</u> |
| NET LOSS | <u>\$ (464,911)</u> | <u>\$ (130,974)</u> | <u>\$ (941,227)</u> | <u>\$ (126,444)</u> |
| WEIGHTED AVERAGE COMMON SHARES OUTSTANDING: | | | | |
| Basic and diluted | <u>4,305,026</u> | <u>4,290,026</u> | <u>4,303,927</u> | <u>4,290,026</u> |
| LOSS PER COMMON SHARE: | | | | |
| Basic and diluted | <u>\$ (0.11)</u> | <u>\$ (0.03)</u> | <u>\$ (0.22)</u> | <u>\$ (0.03)</u> |

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

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LECTEC CORPORATION
CONDENSED STATEMENTS OF CASH FLOWS
(Unaudited)

| | Nine Months Ended September 30, | |
|---|---------------------------------|-------------------|
| | 2010 | 2009 |
| Cash flows from operating activities: | | |
| Net loss | \$ (941,227) | \$ (126,444) |
| Adjustments to reconcile net loss to net cash used in operating activities: | | |
| Depreciation expense | 2,383 | 1,659 |
| Amortization of patent costs | 14,274 | 16,189 |
| Compensation expense related to issuance of stock options | 104,816 | — |
| Deferred income tax | (457,000) | — |
| Changes in operating assets and liabilities: | | |
| Infringement and royalty receivable | 5,499 | (91,351) |
| Prepaid expenses and other | 340,947 | 43,137 |
| Income tax payable | (993,403) | — |
| Accounts payable | 4,430 | 20,845 |
| Accrued expenses | (270,881) | 37,324 |
| Net cash used in operating activities | <u>(2,190,162)</u> | <u>(98,641)</u> |
| Cash flows from investing activities: | | |
| Purchase of office equipment | (1,257) | — |
| Investment in patents | (31,641) | (6,725) |
| Net cash used in investing activities | <u>(32,898)</u> | <u>(6,725)</u> |
| Cash flows from financing activities: | | |
| Payments of dividend | (4,298,350) | — |
| Stock option exercised | 39,000 | — |
| Net cash used in financing activities | <u>(4,259,350)</u> | <u>—</u> |
| Net decrease in cash and cash equivalents | (6,482,410) | (105,366) |
| Cash and cash equivalents — beginning of period | <u>15,766,107</u> | <u>332,848</u> |
| Cash and cash equivalents — end of period | <u>\$ 9,283,697</u> | <u>\$ 227,482</u> |

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

LECTEC CORPORATION
Notes to Condensed Financial Statements
September 30, 2010 and 2009

(Unaudited)

(1) Basis of Presentation

The accompanying condensed financial statements include the accounts of LecTec Corporation (the “Company” or “we,” “us” or “our”) as of September 30, 2010 and December 31, 2009 and for the three and nine month periods ended September 30, 2010 and 2009. Our condensed financial statements have been prepared in accordance with accounting principles generally accepted in the United States of America and should be read in conjunction with our Annual Report on Form 10-K for the year ended December 31, 2009. The interim condensed financial statements are unaudited and in the opinion of management, reflect all adjustments necessary for a fair presentation of results for the periods presented. Results for interim periods are not necessarily indicative of results for the full year.

(2) Business/Premises Summary and Critical Accounting Policies

Business Summary. We are an intellectual property (“IP”) licensing and holding company with approximately \$9,300,000 in cash as of September 30, 2010, plus a cash trust reserve of approximately \$604,000 to fund future patent litigation efforts. We hold multiple domestic and international patents based on our original hydrogel patch technology and have also filed for a provisional patent for a hand sanitizer patch. Our hydrogel patch technology allows for a number of potential applications, including our previously marketed TheraPatch® products, while our anti-microbial hand sanitizer patch is a consumer product which kills targeted infectious organisms and is intended to be dry, thereby rendering the patch harmless in the event that it is licked, chewed or exposed to the eye. An initial prototype of our hand sanitizer patch has been developed. Although we are conducting limited research and development on our hand sanitizer patch, we intend to engage a strategic partner to complete its development and the pursuit of manufacturing and marketing/co-marketing partners for products from LecTec’s IP is also ongoing. We have a licensing agreement (“Novartis Agreement”) with Novartis Consumer Health, Inc. (“Novartis”), which pays royalties to us from time to time, within the terms of the Novartis Agreement, based upon a percentage of Novartis’ net sales of licensed products. We take legal action as necessary to protect our IP and are currently involved in two patent infringement actions. Finally, we are pursuing a merger/acquisition strategy with the intent to leverage our cash assets and improve shareholder value and liquidity.

Corporate Office and Premises Summary. We have two leased facilities as of September 30, 2010.

In July, 2008, the Company moved its corporate headquarter facilities (principal executive office) from Edina, Minnesota to Texarkana, Texas. In connection with this relocation, we entered into a Lease Agreement with Lockaway Storage, Inc. (the “Lessor”) pursuant to which we agreed to lease approximately 1,200 square feet of space located at 1407 South Kings Highway, Texarkana, Texas 7550 (the “Texas Lease”). In March 2010, we renewed the Texas Lease until March 1, 2011 at a monthly rate of \$750 per month and we must also pay our pro rata share of the costs and expenses incurred by the Lessor to operate the common areas of the office and warehouse complex. The Texas Lease contains customary representations, warranties and covenants on the part of the Company and the landlord.

In January, 2009, we entered into a lease amendment (the “Lease Amendment”) amending our lease, dated May 23, 2003, between us and SMD Lincoln Investments (the “Minnesota Lease”), regarding our previous headquarters located at 5610 Lincoln Drive, Edina, Minnesota (the “Leased Premises”). The Lease Amendment will continue to renew for successive one-month periods until such lease is terminated by the landlord upon 30 days written notice to us or by us upon 90 days written notice to the landlord. We use the space for liquidating saleable assets, storage of current accounting records, and managing an orderly wind down of operations at this facility. We maintain approximately 3,300 square feet of space at this facility. The Company pays \$2,400 per month for use of this space. Notice has been provided by the Company to the landlord of its intention to vacate this facility by December 31, 2010.

In July, 2008, we opened an office in India, Level 2, Connaught Place, Bund Garden Road, Pune (India), 411001, to explore research, development and manufacturing opportunities for our advanced skin interface technologies and products. Having completed an evaluation of our IP portfolio, we terminated this lease effective July 31, 2010.

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Critical Accounting Policies

The Company's most critical accounting policies include:

Revenue Recognition. Royalty and licensing fees are recognized when earned under the terms of the Novartis Agreement, based upon sales information of licensed products provided by Novartis, and when collection is reasonably assured. Infringement income is recognized when settlement agreements have been signed and collection is reasonably assured.

Patent Costs. The carrying value of patent costs is reviewed periodically or when factors indicating impairment are present. Any impairment loss is measured as the amount by which the carrying value of the assets exceeds the fair value of the assets. We believe no such impairment currently exists.

Royalty Receivable. We grant credit to our only customer, Novartis, in the normal course of business and under the terms contained in the Novartis Agreement. Novartis pays royalty income to us pursuant to the terms of the Novartis Agreement. At September 30, 2010, we had an outstanding royalty receivable with Novartis of \$26,026, which was collected in October, 2010.

Use of Estimates. In preparing financial statements in conformity with accounting principles generally accepted in the United States of America, management is required to make estimates and assumptions that affect certain reported amounts of assets and liabilities, the disclosure of contingent assets and liabilities at the date of the financial statements, and the reported amounts of revenues and expenses during the reporting period. Actual results could differ from those estimates.

Share-Based Compensation. We account for share-based compensation in accordance with Financial Accounting Standards Board ("FASB") Accounting Standards Codifications ("ASC") Topic 718, *Compensation—Stock Compensation*, which requires that compensation cost relating to share-based payment transactions (including the cost of all employee stock options) be recognized in the financial statements. That cost is measured based on the estimated fair value of the equity or liability instruments issued.

Off-Balance Sheet Arrangements. We do not have any "off-balance sheet arrangements" (as such term is defined in Item 303 of Regulation S-K) that are reasonably likely to have a current or future effect on our financial condition, changes in financial condition, revenues or expenses, operating results, liquidity, capital expenditures or capital resources.

Reclassification. The December 31, 2009 balance sheet has been reclassified to allocate \$1.6 million of dividends to additional contributed capital from accumulated deficit. This reclassification had no impact on total shareholders' equity, net income or net cash flows.

Recent Accounting Pronouncements:

In October 2009, FASB issued an update to the accounting and reporting guidance for multiple-deliverable revenue arrangements. The new accounting guidance removes the separation criterion that objective and reliable evidence of the fair value of the undelivered item must exist for the delivered items to be considered a separate unit or separate units of accounting. This FASB-issued update requires an entity to determine the selling price of qualifying deliverables based on a hierarchy of evidence. In considering the hierarchy of evidence, the entity must first determine the selling prices by using vendor-specific objective evidence ("VSOE"), if it exists; otherwise, third-party evidence ("TPE") of selling price must be used. If neither VSOE nor TPE of selling price exists for a deliverable, an entity must use its best estimate of the selling price for that deliverable in allocating consideration among deliverables in an arrangement. This update is effective for arrangements entered into in the fiscal years beginning on or after June 15, 2010, unless the vendor elects early application. We are evaluating the potential impact, if any, of the adoption of this update on our financial position and results of operations.

In January 2010, FASB issued an update to the existing disclosure requirements related to fair value measurements which requires entities to make new disclosures about recurring or nonrecurring fair value measurements including significant transfers into and out of Level 1 and Level 2 fair value measurements and information on purchases, sales, issuances, and settlements on a gross basis in the reconciliation of Level 3 fair value

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measurements. This update is effective for annual and interim periods beginning after December 15, 2009, except for Level 3 reconciliation disclosures which are effective for annual periods beginning after December 15, 2010. The adoption of the required portion of this standard did not have a material impact on our financial position or results of operations.

In April 2010, FASB issued new accounting guidance to provide clarification on the classification of a share-based payment award as either equity or a liability. Under ASC 718, *Compensation-Stock Compensation*, a share-based payment award that contains a condition that is not a market, performance, or service condition is required to be classified as a liability. The amendments clarify that a share-based payment award with an exercise price denominated in the currency of a market in which a substantial portion of the entity's equity securities trades should not be considered to contain a condition that is not a market, performance, or service condition. Therefore, such an award should not be classified as a liability if it otherwise qualifies as equity. The amendments are effective for fiscal years, and interim periods within those fiscal years, beginning on or after December 15, 2010. Earlier application is permitted. Adoption of this standard is not expected to have a material impact on our financial position or results of operations.

In May 2010, FASB issued new guidance on the use of the milestone method of recognizing revenue for research and development arrangements under which consideration to be received by the vendor is contingent upon the achievement of certain milestones. The update provides guidance on the criteria that should be met for determining whether the milestone method of revenue recognition is appropriate. A vendor can recognize consideration in its entirety as revenue in the period in which the milestone is achieved only if the milestone meets all criteria to be considered substantive. Additional disclosures describing the consideration arrangement and the entity's accounting policy for recognition of such milestone payments are also required. The new guidance is effective for fiscal years, and interim periods within such fiscal years, beginning on or after June 15, 2010, with early adoption permitted. The guidance may be applied prospectively to milestones achieved during the period of adoption or retrospectively for all prior periods. Adoption of this standard is not expected to have a material impact on our financial position or results of operations.

(3) Loss per Common Share

Basic loss per common share is computed by dividing net loss by the weighted average number of common shares outstanding. Diluted loss per common share is computed by dividing net loss by the weighted average number of common shares outstanding and common share equivalents related to stock options and warrants when dilutive. Common stock options to purchase 374,000 shares of common stock with a weighted average exercise price of \$3.85 were outstanding for the three and nine month periods ended September 30, 2010, respectively. Because we had a loss from operations during the three and nine months ended September 30, 2010 and 2009, those shares were excluded from the loss per share computations because they were antidilutive.

(4) Income Taxes

Deferred income taxes are provided for temporary differences between the financial reporting and tax basis of assets and liabilities. Deferred taxes are reduced by a valuation allowance when, in the opinion of management, it is more likely than not that some portion or all of the deferred tax assets will not be realized. Deferred tax assets and liabilities are adjusted for the effects of changes in tax laws and rates on the date of the enactment.

In evaluating the ultimate realization of deferred income tax assets, management considers whether it is more likely than not that the deferred income tax assets will be realized. Management establishes a valuation allowance if it is more likely than not that all or a portion of the deferred income tax assets will not be utilized. The ultimate realization of deferred income tax assets is dependent on the generation of future taxable income, which must occur prior to the expiration of the net operating loss carryforwards.

(5) Novartis Supply and License Agreement

In July, 2004, we entered into a supply and licensing agreement with Novartis, effective January 1, 2004. By December 31, 2004, the supply portion of the Novartis Agreement was completed and we no longer manufactured any product. Under the Novartis Agreement, we granted Novartis an exclusive license (the "License") to all of our intellectual property to the extent that it is used or is useful in the production of the vapor patches that Novartis is selling under the Novartis Agreement. The License will continue in effect for the duration of the patents' life permitted under applicable law. Upon the expiration of the patents included in the licensed intellectual property,

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Novartis will have a non-revocable, perpetual, fully paid-up license to the intellectual property used or useful in the production of vapor patches for the pediatric and the adult cough/cold market. Novartis is required by the Novartis Agreement to pay royalties to us at an agreed upon percentage based on net sales of vapor patches by Novartis for each year the License is in effect.

In June 2006, Novartis issued a nationwide recall of all of its Triaminic® vapor patch products. In a press release issued by Novartis pertaining to the recall, Novartis explained that the recall was “due to the serious adverse health effects that could result if the product is ingested by a child removing the patch and chewing on it.” At the same time that Novartis announced this voluntary recall, the U.S. Food and Drug Administration (“FDA”) issued a release warning consumers “not to use the Triaminic Vapor Patch due to reports of serious adverse events associated with accidental ingestion by children.” According to news reports, the recall resulted from an adverse event experienced by a child who suffered a seizure after chewing on a Triaminic Vapor Patch. Novartis confirmed to us that the patch involved in this incident was not manufactured by us. As a result of this recall, we were proactive in assisting Novartis to resolve the FDA issues surrounding the product recall in order to restore our royalty income stream. We have met with Novartis representatives to discuss how to prevent an incident where a child or pet chews or ingests a patch.

In July 2007, Novartis began shipping a new adult vapor patch product in the United States, Canada and Mexico for the cough, cold, and flu season. Novartis has not announced whether it will re-introduce a vapor patch for the pediatric market.

During the three and nine months ended September 30, 2010, we recorded revenue of \$26,373 and \$49,156, respectively, compared to revenue recorded for the three and nine months ended September 30, 2009 of \$24,215 and \$76,633, respectively, for royalties covered under the Novartis Agreement. We do not know the marketing intent of Novartis for our vapor patch product. Year to year royalties paid to us have declined and although 3rd quarter 2010 revenue is slightly higher than 3rd quarter 2009, we expect royalties received during the current year to continue on the per annum downward trend.

(6) Discontinued Operations

We ceased manufacturing operations of topical patches and sold all of our manufacturing assets related to the production of patches to our only remaining customer, Novartis, as of December 31, 2004. We had a liability for discontinued operations that consisted of a reserve for sales returns and credits of \$130,000 related to sales prior to the discontinuance of operations. This reserve was written off against operating expenses during December 2009.

(7) Patents and Trademarks

Our policy is to protect our proprietary position by securing U.S. and foreign patents that cover the technology, inventions and improvements important to our business. We have 17 U.S. and 43 international patents related to our patch technology. We have three U.S. patent pending applications, two international patent pending applications, and two foreign applications through the Patent Cooperation Treaty (“PCT”). The issued U.S. patents most pertinent to our major products have a remaining legal duration ranging from one to 13 years.

In 2008, we converted our two new provisional patents to PCT international applications. These applications include: (1) adding an aversive agent to our licensed patch or other patches to prevent ingestion by children or pets; and (2) a hand sanitizer patch that will kill targeted infectious organisms. Moreover, we filed an additional provisional patent application in 2010 to further expand the scope of our hand sanitizer intellectual property.

Issued patents can later be held invalid by the patent office issuing the patent or by a court. We cannot be certain that our patents will not be challenged, invalidated or circumvented or that the rights granted under our patents will provide a competitive advantage.

We use both patents and trade secrets to protect our proprietary property and information. To the extent we rely on confidential information to maintain our competitive position, there can be no assurance that other parties will not independently develop the same or similar information.

On July 25, 2008, we filed a complaint for patent infringement against five companies, alleging that those companies had infringed upon two of our patents relating to our medicated patch technology. We settled with three of

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the parties during fiscal 2009 and are actively pursuing litigation against the remaining defendants. Currently, the Company has not filed for any additional patents. See PART II, ITEM 1 of this Form 10-Q for additional information.

(8) Stock Option

On September 22, 2010, our shareholders adopted and approved the LecTec Corporation 2010 Stock Incentive Plan (the "Stock Incentive Plan"). The purpose of the Stock Incentive Plan is to promote the interests of LecTec and our shareholders by aiding us in attracting and retaining employees, officers, consultants, advisors and non-employee directors who we expect will contribute to our success and to enable these individuals to participate in our long-term success and growth by giving them a proprietary interest in LecTec. The Stock Incentive Plan is intended to replace our existing LecTec Corporation 2001 Stock Incentive Plan, which is scheduled to expire by its terms on July 1, 2011. The aggregate number of shares of our common stock that may be issued under all stock-based awards made under the Stock Incentive Plan will be 450,000.

On June 1, 2010, our CEO, as partial compensation for his employment, was granted a Non-Qualified Stock Option to purchase 125,000 shares of our common stock at \$3.50 per share. The option has a 10 year term and vests as to 25,000 shares every 90 days, with the first vesting date August 30, 2010, until fully vested. Notwithstanding the vesting schedule, the entire option vests immediately if we have a change in control and the CEO's employment is terminated by us within 15 months following the change in control for any reason other than the CEO's death, by us for cause or by the CEO other than for good reason. The fair value of the option is the estimated present value at the grant date using the Black-Scholes-Merton option pricing model with the following assumptions: risk free interest rate 3.29%, expected life 10 years, expected volatility 195.84% and expected dividend rate 0.00%. We recorded share-based compensation of \$78,660 and \$104,816 during the three and nine month periods ended September 30, 2010. We did not record any share-based compensation during the three and nine months ended September 30, 2009. At September 30, 2010, there was approximately \$288,244 of total unrecognized compensation cost related to nonvested share-based compensation arrangements. The cost is expected to be recognized over 0.9 years.

(9) Advisory Service Agreement

We have an Advisory Service Agreement with our previous Chairman of the Board, Chief Executive Officer and Chief Financial Officer. The scope of the Advisory Service Agreement is to evaluate our opportunities in Asia and support our efforts regarding the development of our IP portfolio and the protection thereof. Compensation pursuant to this agreement is \$11,000 per month, in addition to reimbursements for normal out of pocket expenses including a monthly stipend of \$500 per month for phone expenses, and may be terminated on 30 days notice by either party.

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

OVERVIEW

We are focused on four major areas: (1) engage a strategic partner to complete development of the hand sanitizer patch and pursue manufacturing and marketing/co-marketing arrangements; (2) further monetization, if possible, of our IP portfolio, including our previously marketed TheraPatch® products, but excluding our hand sanitizer patch, through licensing, selling or engaging strategic partners for all or a portion of our hydrogel IP; (3) support our current litigation strategy; and (4) pursue merger/acquisition opportunities.

Hand Sanitizer Patch. Due to the growing worldwide concern regarding the spread of germs through hand contact, we filed patents for, and screened, identified and tested technologies suitable for, an anti-microbial hand sanitizer patch. This activity has led to the development of a prototype that is ready to begin efficacy and other testing to determine its market viability. We will continue limited research and development efforts to continue this progress and will pursue engaging a strategic partner to complete the development of our hand sanitizer patch and pursue manufacturing and marketing/co-marketing arrangements. Because the hand sanitizer patch is a consumer product, we believe that engaging an established strategic partner is the best go-to-market strategy because we will be able to leverage any such partner's competencies regarding the development and manufacturing of products, customer requirements and marketing and distribution strategies. We expect that Asian entities will provide the greatest opportunities because the Asian topical patch market represents the most significant portion of the worldwide market. If we are not able to engage an acceptable strategic partner, we will evaluate increasing the amount of resources designated for developing our hand sanitizer patch in light of progress made in our other strategic initiatives.

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IP Portfolio, Excluding Hand Sanitizer Patch. We completed an evaluation of our IP portfolio, which included conducting both a current analysis of our portfolio and referring to our 2007 extensive market research and intellectual property report. Based on this evaluation, we believe that the best strategy to derive further value, if any, from our IP portfolio, other than our hand sanitizer patch, is to pursue licensing of this IP, and to engage strategic partners to help us further develop, if necessary, manufacture and/or market this IP, including reintroduction of TheraPatch® products or sell all or a portion of this IP. At this time, we do not intend to conduct any further research and development with respect to our hydrogel IP. We will begin to identify those parties that we believe may have interest in this IP and approach them. If we are not able to identify suitable alternatives regarding the licensing, sale or strategic partnering of our hydrogel IP, we will reevaluate our position with respect to the foregoing in light of progress made in our other strategic initiatives.

Litigation. In April 2007, we were granted a re-examination certificate that expanded our prior claims related to a patent that we hold. During 2008, we retained a legal firm on a contingency fee basis to assist us in enforcing our rights related to potential patent infringement claims by the Company. As a result, we sued five parties and, during 2009, we settled our claims with three of such parties. We remain diligent in pursuing our patent infringement claims against the remaining two parties. See PART II, ITEM 1 of this Form 10-Q for additional information.

Merger/Acquisition Opportunities. We believe that our cash balance and public company status provide the potential for merger/acquisition opportunities in addition to our work surrounding our IP portfolio, which we continue to explore. In evaluating any such opportunities, primary consideration will be given to companies generating revenue and addressing sizable markets which we believe may attract significant investment interest. Any transaction under consideration must also be expected to provide increased liquidity for our shareholders. Our current intention is not to seek multiple investments, but to focus our efforts on identifying a single transaction in which to apply our cash balance and public company status. Although opportunities related to our current business areas will be of greatest interest, we will evaluate situations in other areas in which we have the capability to make an appropriate and informed review.

Our strategy described above will remain fluid as we pursue each area of such strategy. Although we believe that our strategy will result in increased value for our shareholders, there can be no assurance that our strategy, or any component thereof, will be successful.

COMPARISON OF THE THREE AND NINE MONTHS ENDED SEPTEMBER 30, 2010 AND 2009

Results of Operations

We recorded royalty income of \$26,373 and \$24,215 for the three months ended September 30, 2010 and 2009, respectively, and \$49,156 and \$76,633 for the nine months ended September 30, 2010 and 2009, respectively. This reflects an increase of \$2,158 for the three months ended September 30, 2010 over 2009, but a decrease of \$27,477 for the nine months ended September 30, 2010 over 2009. Although Novartis royalties for our 3rd quarter 2010 were higher than for our 3rd quarter 2009, the decrease in such revenue year to date is a result of the previous trend of lower sales by Novartis of its patch products using our licensed IP. The royalty income recorded during the three and nine month periods ended September 30, 2010 and 2009 was based on information provided by Novartis.

Infringement income was \$600,000 for the nine months ended September 30, 2009. The lack of infringement income in 2010 is a result of no litigation being settled or concluded during the first three quarters of 2010.

Our operating expenses increased \$484,657 to \$639,878 for the three months ended September 30, 2010, from operating expenses of \$155,221 for the comparable period in 2009. Such increase was the result of the following operating expenses: (1) litigation expenses of \$275,708, funded from our litigation escrow account, which are direct out of pocket costs (other than attorney fees for which we have a contingent fee arrangement), resulting from preparation for our January 2011 trial; (2) expenses of approximately \$64,500 related to our management and director transition, including costs of the Annual Meeting of Shareholders; and (3) a non-cash compensation expense of \$78,660 for options granted to our current CEO. Our operating expenses increased \$572,421 to \$1,376,230 for the nine months ended September 30, 2010, from operating expenses of \$803,809 for the comparable period in 2009. The increase in operating expenses for the nine month period resulted primarily from increases in salaries, non-cash compensation, management and director transition, consulting, litigation expenses and travel expenses. Actions to reduce our operating expenses have begun to be realized, and will continue to be reflected over the next two quarters, in a number of areas including compensation, rent, consulting fees and professional services. However, operating expenses relating to litigation and M&A activity will continue to fluctuate based upon activity in a particular quarter.

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We recorded a net loss of \$(464,911), or \$(0.11) per basic and diluted share, for the three months ended September 30, 2010, compared to a net loss of \$(130,974), or \$(0.03) per basic and diluted share, for the same period in 2009 and a net loss of \$(941,227), or \$(0.22) per basic and diluted share, for the nine months ended September 30, 2010, compared to a net loss of \$(126,444), or \$(0.03) per basic and diluted share, for the same period in 2009. The increase in net loss for the three and nine month periods ended September 30, 2010 from the comparable periods in 2009 is due to the increase in operating expenses combined with the decrease in royalty income from Novartis for the nine months ended 2010 and lack of infringement income. Interest income of \$4,836 and \$13,089 for the three and nine month periods ended September 30, 2010, respectively, were higher than comparable periods due to greater cash assets being available. Interest rates continue to be at historical lows and we continue to weigh risks against returns, taking a conservative approach to our cash investment strategy. Certain action has been, and will continue to be, taken to try to increase interest returns, however, we do not believe there will be a significant positive change in our interest returns for the foreseeable future. However, we have been able to receive slighter higher rates on part of our cash assets in accounts that are FDIC insured.

Income Taxes

The income tax benefit for the three months and nine months ended September 30, 2010 was \$143,758 and \$372,758, respectively, as compared to no benefit in corresponding periods in 2009, as realization of net deferred taxes was not reasonably assured in the 2009 periods. The benefit for the three and nine months ended September 30, 2010 was principally the result of the federal tax benefit of the operating loss for such period.

Effect of Inflation

Inflation has not had a significant impact on our operations or cash flow.

Liquidity and Capital Resources

Our cash and cash equivalents decreased \$6,482,410 for the nine month period ended September 30, 2010, to \$9,283,697 from cash and cash equivalents of \$15,766,107 at December 31, 2009. The decrease in cash and cash equivalents resulted primarily from cash dividend payments of \$4,298,350 and from our current operating expenses. Our litigation escrow account and other prepaid expenses decreased \$340,947 for the nine month period ended September 30, 2010 to \$634,476 as a result of expenses for preparation of our January 2011 trial.

We had no material commitments for capital expenditures at September 30, 2010 or 2009.

We had working capital of \$10,212,137 and a current ratio of 73.39 at September 30, 2010 compared to working capital of \$11,025,789 and a current ratio of 2.92 at December 31, 2009. The decrease in working capital and increase in the current ratio at September 30, 2010, compared to December 31, 2009, was primarily due to our payment of a cash dividend of \$4,298,350, our payment of income taxes of approximately \$1.1 million, coupled with our net loss of \$(941,227).

Shareholders' equity decreased \$797,411 to \$10,263,758 at September 30, 2010 from \$11,061,169 at December 31, 2009, due to the net loss we incurred during the nine months ended September 30, 2010, offset with stock options exercised and the grant of a stock option.

We entered into a contingency fee agreement with Rader, Fishman & Grauer PLLC, who is our legal counsel in the pending patent infringement litigation. See Part II, Item 1 of this Form 10-Q for additional information concerning this litigation. Under this agreement, the Rader firm will receive a percentage of any recovery in the litigation or other proceeds resulting from a settlement of the litigation as its primary compensation for representing us in this matter. We are also obligated (i) to reimburse the Rader firm for its out-of-pocket expenses in connection with the litigation through an up-front advance of \$50,000 and monthly advances of \$10,000, and (ii) to engage and pay for expert services needed in the litigation, provided that our obligation to advance such funds and pay such expert expenses will be suspended if our cash levels fall below certain thresholds. Thereafter, if our cash levels exceed such thresholds, or there is a recovery in or other proceeds from the litigation, then the Rader firm will be reimbursed for any expenses it has covered while such advances and payments were suspended. To date we have expended an aggregate of approximately \$8.3 million under the agreement.

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We currently estimate that we will receive less than \$75,000 per year in royalty income based upon historical royalty income and cash receipt activity from Novartis, coupled with the downward trending of this income. Royalty income is uncertain because it is subject to factors that we cannot control. There can be no assurance that the anticipated revenue stream or the anticipated expenses will be as planned, or that we will be successful in negotiating new licensing opportunities with Novartis or other companies or having success in our litigation due to the uncertainties and risks described in “Risk Factors” in Item 1A filed on Form 10-K for the period ending December 31, 2009.

CRITICAL ACCOUNTING POLICIES

Management believes that the Company has not adopted any critical accounting policies which, if changed, would result in a material change in financial estimates, financial condition, results of operation or cash flows for the three months ended September 30, 2010 and 2009. The critical accounting policies appear in Note 2 of the Notes to Condensed Financial Statements in this Form 10-Q.

ITEM 3. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISKS

Not Applicable.

ITEM 4T. CONTROLS AND PROCEDURES

Evaluation of Disclosure Controls and Procedures

We maintain “disclosure controls and procedures” as defined in Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934, (the “Exchange Act”), that are designed to ensure that information required to be disclosed by us in reports we file or submit under the Exchange Act is recorded, processed, summarized and reported within the time periods specified in the SEC’s rules and forms, and that such information is accumulated and communicated to our management, including our principal executive officer and principal financial officer, and Board of Directors, as appropriate, to allow timely decisions regarding required disclosure. In designing and evaluating our disclosure controls and procedures, management recognizes that disclosure controls and procedures, no matter how well conceived and operated, can provide only reasonable assurance of achieving the desired objectives, and we necessarily are required to apply our judgment in evaluating the cost-benefit relationship of possible disclosure controls and procedures.

Our management, including our principal executive officer and principal financial officer, evaluated the effectiveness of the design and operation of our disclosure controls and procedures as of September 30, 2010 and concluded that our disclosure controls and procedures were effective.

Changes in Internal Controls Over Financial Reporting

During the quarter ended September 30, 2010, there were no changes in our internal control over financial reporting (as defined in Rule 13a-15(f) and 15d-15(f) under the Exchange Act) that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

PART II –OTHER INFORMATION

ITEM 1. LEGAL PROCEEDINGS

On July 25, 2008, we filed a complaint for patent infringement (the “Complaint”) against five companies, including Chattem, Inc. (Ticker: CHTT), Endo Pharmaceuticals, Inc. (Ticker: ENDP), Johnson & Johnson Consumer Company, Inc. (Ticker: JNJ), The Mentholatum Company, Inc. (Division of Rohto Pharmaceuticals, Ticker RPHCF.PK), and Prince of Peace Enterprises, Inc. (Private Company) (collectively, the “Defendants”) in the U.S. District Court for the Eastern District of Texas. The Complaint alleges, among other things, that the Defendants have infringed two of our patents (the “Patents–In–Suit”), which relate to our medicated patch technology. We are seeking to enjoin the Defendants from infringing the Patents–In–Suit and to recover monetary damages related to such infringement, as well as interest and litigation costs.

In October 2008, all five of the Defendants filed answers (the “Answers”) in response to the Complaint denying our claims therein, and asserting certain affirmative defenses and counterclaims against us, including assertions that the Patents–In–Suit are invalid and unenforceable, and claims for attorneys’ fees and costs. On October 20, 2008, we filed our replies to the Answers, denying such counterclaims and affirmative defenses, including the claims that the Patents–In–Suit are invalid and unenforceable.

On December 3, 2008, our counsel in the litigation, Rader, Fishman & Grauer PLLC (the “Counsel”), participated in a scheduling conference in this case. As a result of that conference, the Court scheduled a Markman hearing for May 6, 2010 and a final pretrial conference for January 3, 2011 to address any final matters before the start of trial on January 4, 2011. Based on the schedule established by the Court, it is clear that pursuing our claims in this litigation through trial will be a lengthy process.

In February 2009, Counsel filed with the Court a motion to preliminarily enjoin the five defendants from infringing the Patents-In Suit pending the trial.

On May 29, 2009, we entered into a Settlement Agreement and Mutual Release (the “Mentholatum Settlement Agreement”) with The Mentholatum Company (“Mentholatum”) to settle our claims against Mentholatum that Mentholatum infringed the Patents–In–Suit. Pursuant to the Mentholatum Settlement Agreement, Mentholatum paid us an aggregate of \$600,000 in \$100,000 monthly installments from May through October 2009. In addition, under the Mentholatum Settlement Agreement (a) we agreed to dismiss the litigation against Mentholatum with prejudice, (b) the parties agreed to mutual general releases of all claims other than their prospective obligations under the Mentholatum Settlement Agreement and claims arising after the date of the Mentholatum Settlement Agreement, (c) we agreed not to sue Mentholatum or Rohto Pharmaceutical Co., Ltd., the parent company of Mentholatum, for any infringement of the Patents–In–Suit, any patent that claims priority, directly or indirectly, from the Patents–In–Suit, or any foreign counterparts of the Patents–In–Suit, and (d) we agreed not to transfer any such patents unless the transferee agrees to be bound by the covenant not to sue. Mentholatum and Rohto agreed not to challenge the validity or enforceability of such patents. The proceeds received from this settlement were reduced by the amounts due to the Rader firm per our contingent fee arrangement, out of pocket expenses, and other costs incurred related to depositions of parties in the infringement lawsuits, travel expenses, and other related costs. We received approximately \$300,000 in net cash proceeds from the Mentholatum settlement.

On November 11, 2009, we entered into a Settlement and License Agreement (the “Endo Settlement Agreement”) with Endo Pharmaceuticals Inc. (“Endo”). Pursuant to the Endo Settlement Agreement, Endo agreed to pay us a one–time license fee of \$23,000,000 and we granted to Endo an exclusive license to the Patents–In–Suit for use in the field of prescription pain medicines and treatment. In addition, under the Endo Settlement Agreement: (a) the parties agreed to the dismissal of the litigation with prejudice and without costs; (b) we agreed to release all claims against Endo that were asserted by or could have been asserted by us against Endo in the litigation or that relate to, arise from or are in any manner connected to the Patents–In–Suit; (c) Endo agreed to release all claims against us that were asserted by or could have been asserted by Endo against us in the litigation; (d) we agreed not to sue Endo for any infringement of any U.S. or foreign patents or patent applications owned or controlled by us as of November 11, 2009, any continuation, continuation–in–part or divisional of any such patent, any U.S. patent resulting from the reissue or reexamination of any such patents and any U.S. or foreign patent or patent application claiming common priority with any of such patents; and (e) we agreed not to transfer either of the Patents–In–Suit or any other such patent unless the transferee agrees in writing to the terms and conditions of the Endo Settlement Agreement. We received approximately \$16,000,000 in net cash proceeds from this settlement in December 2009. From these proceeds, we replenished the trust

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fund we have with the Rader Firm with \$1,000,000 dollars to fund ongoing patent litigation. The trust fund balance at September 30, 2010 was \$603,690 compared to a balance of \$931,954 at December 31, 2009. If funds are not completely expended, then the remaining cash balance in the trust fund will revert to us.

On December 18, 2009, we entered into a Settlement Agreement and Mutual Release (the “JJCC Settlement Agreement”) with Johnson & Johnson Consumer Companies, Inc. (“JJCC”) to settle our claims against JJCC that JJCC infringed our Patents–In–Suit. Pursuant to the JJCC Settlement Agreement, JJCC paid us a one–time sum of \$1,200,000 and we granted to JJCC a fully paid–up, world–wide, non–exclusive and irrevocable license to (a) the Patents–In–Suit, (b) any patent that claims priority, directly or indirectly, from the Patents–In–Suit (the “Family Patents”), including, without limitation, U.S. Patent Nos. 6,096,333, 6,096,334 and 6,361,790, (c) any foreign counterparts of the Patents–In–Suit or any of the Family Patents to make, have made, sell, offer for sale, use, import, export or otherwise dispose of any apparatus, method, product, component, service, product by process or any device associated with JJCC or its subsidiaries, affiliates or other controlled entities, for the past, present and future until the expiration of the last patent described above and (d) any patents that we own or currently have an interest in to make, have made, sell, offer for sale, use, import, export or otherwise dispose of any non–prescription, non–occlusive medicated hydrogel patch products that are used to alleviate pain associated with JJCC (collectively, the License Grant”); provided, however, that the License Grant under clauses (a), (b) and (c) above excludes over–the–counter vapor patches which emit vapors that provide cough and cold relief when inhaled, and prescription, non–occlusive, medicated hydrogel patch products that are used to alleviate pain. The proceeds received from this settlement were reduced by the amounts due to the Rader firm per our contingent fee arrangement, out of pocket expenses, and other costs incurred related to depositions of parties in the infringement lawsuits, travel expenses, and other related costs. After these expenses we received net cash proceeds of approximately \$720,000.

On May 6, 2010 a Markman hearing occurred in Texarkana, Texas and the US District Court for the Eastern District of Texas issued Orders concerning it on May 20, 2010. The first Order was based on our motion to strike an exhibit from Chattem, Inc.’s Opposition Brief, and our motion to strike was granted by the Court. The second Order issued by the court denied Defendant’s motion request for leave to file for summary judgment as to non–infringement, but granted the request for leave to file for summary judgment as to invalidity of patents. The Court also issued its Markman ruling interpreting the terms “cured” and “non–occlusive” contained in our patents.

We engaged in voluntary mediation with Chattem, Inc. in July 2010. A Report of Mediation by the Hon. Harlan A. Martin was filed stating that the parties were unable to reach settlement. On September 28, 2010 the United States District Court for the Eastern District of Texas issued an Order regarding Prince of Peace’s and Chattem’s (“Defendants”) requests to file motions for summary judgment: (1) of invalidity due to the on–sale bar of 35 U.S.C. § 102(b); and (2) regarding Defendants’ defenses of equitable estoppels and laches and our motions: (3) on, and to preclude testimony related to, Defendants’ 35 U.S.C. § 102(b) defense based on the Aqua–Patch; and (4) on infringement by Defendants. The Order granted Defendants’ the right to file a summary judgment motion regarding on–sale bar, but denied them the opportunity regarding the equitable defenses of estoppel and laches. With regard to the equitable issues, the Court stated that the custom in patent cases is to hold a separate bench proceeding after the jury trial on such issues. The Order granted us the right to file summary judgment motions on infringement and to preclude Defendants Aqua–Patch defense.

We are diligent in pursuing our patent infringement lawsuit against the remaining two defendants, Chattem, Inc. and Prince of Peace Enterprises, Inc. We will oppose Defendants’ expected Motion for Summary Judgment and our Counsel is preparing for trial, which is scheduled to begin in January 2011.

We are unable to determine, based on current information available, whether we will be successful in our legal pursuits against the remaining two defendants. We give no assurance as to the outcome of the ongoing lawsuit or whether our Patents–In–Suit and claims asserted in the related patents could be deemed invalid by a court of law.

ITEM 1A. RISK FACTORS

Item 1A (“Risk Factors”) of our most recently filed Form 10-K sets forth information relating to important risks and uncertainties that could materially have an adverse effect on our business, financial condition, or operating results. There have been no material changes to the risk factors described in our most recently filed Form 10-K; however, those risk factors continue to be relevant to an understanding of our business, financial condition, and operating results, etc. Accordingly, potential and current investors should review and consider these risk factors in making any investment decision with respect to our securities. An investment in our securities continues to have a high degree of risk.

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ITEM 2. UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS

None.

ITEM 3. DEFAULTS UPON SENIOR SECURITIES

None.

ITEM 4. (REMOVED AND RESERVED)

This item was removed and reserved pursuant to SEC Release No. 33-9089A issued on February 23, 2010.

ITEM 5. OTHER INFORMATION

None.

ITEM 6 — EXHIBITS

| <u>Exhibit No.</u> | <u>Description</u> |
|--------------------|--|
| 3.01 | Articles of Incorporation of LecTec Corporation, as amended (Incorporated herein by reference to the Company's Form S-1 Registration Statement (file number 33-9774C) filed on October 31, 1986 and amended on December 12, 1986). |
| 3.02 | Bylaws of LecTec Corporation (Incorporated herein by reference to the Company's Form S-1 Registration Statement (file number 33-9774C) filed on October 31, 1986 and amended on December 12, 1986). |
| 10.01 | LecTec Corporation 2010 Stock Incentive Plan (incorporated by reference to Appendix A to the Company's Definitive Proxy Statement on Schedule 14A filed on August 20, 2010). |
| 10.02 | Form of Non-Qualified Stock Option Agreement for grants to Directors under 2010 Stock Incentive Plan, filed herewith. |
| 31.01 | Certification of Principal Executive Officer Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002, filed herewith. |
| 31.02 | Certification of Principle Financial Officer Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002, filed herewith. |
| 32.01 | Certification Pursuant to 18 U.S.C. §1350, as Adopted Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, filed herewith. |

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned thereunto duly authorized.

LECTEC CORPORATION

Date: November 15, 2010

By: /s/ Greg G. Freitag

Greg G. Freitag

Chief Executive Officer, Chief Financial Officer, & Director

(Principal Financial Officer and Principal Executive Officer)

EXHIBIT INDEX

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**LECTEC CORPORATION
NON-INCENTIVE STOCK OPTION AGREEMENT**

This **Non-Incentive Stock Option Agreement**, made as of this ____ day of ____, 20__ (the "*Effective Date*"), by and between LecTec Corporation, a Minnesota corporation (the "*Company*"), and _____ ("*Optionee*").

WHEREAS, the Company wishes to grant this stock option to Optionee pursuant to the LecTec Corporation 2010 Stock Incentive Plan (the "*Plan*").

NOW, THEREFORE, in consideration of the premises and of the mutual covenants herein contained, the parties hereto hereby agree as follows:

1. Grant of Option. The Company hereby grants to Optionee the right and option (the "*Option*") to purchase all or any part of an aggregate of ____ shares (the "*Shares*") of the common stock, par value \$0.01 per share (the "*Common Stock*"), of the Company at the price of \$ ____ per Share on the terms and conditions set forth herein. It is understood and agreed that such price is not less than 100% of the fair market value of each such Share on the date of this Agreement. The Option is not intended to qualify as an incentive stock option within the meaning of Section 422A of the Internal Revenue Code of 1986, as amended (the "*Code*").

2. Duration and Exerciseability. The Option may not be exercised by Optionee except as set forth herein, and the Option shall in all events terminate seven (7) years from the date hereof (the "*Termination Date*"). Subject to the other terms and conditions set forth herein, the Option shall vest and may be exercised by Optionee in cumulative installments as follows:

| On or after each of the following dates | Percentage of Shares as to which the Option is exercisable |
|--|---|
| | |
| | |
| | |
| | |

During the lifetime of Optionee, the Option shall be exercisable only by Optionee. The Option shall not be assignable or transferable by Optionee, other than by will or the laws of descent and distribution. The vesting of the Option is subject to acceleration under the circumstances described in Section 4.

3. Exercise of Option After Death or Departure from Board of Directors.

(a) In the event Optionee ceases to serve as a member of the Board of Directors of the Company or its subsidiaries, if any, for any reason other than Optionee's gross and willful



misconduct, death or disability, voluntary resignation or Requested Resignation, Optionee shall continue to have the right to exercise this Option at any time within the term of this Option to the extent of the full number of Shares Optionee was entitled to purchase under this Option on the date of such termination.

(b) In the event Optionee ceases to serve as a member of the Board of Directors of the Company or its subsidiaries, if any, by reason of Optionee's gross and willful misconduct during the course of services, which shall include, but not be limited to, the wrongful appropriation of funds of the Company or the commission of a gross misdemeanor or felony, this Option shall be terminated as of the date of the misconduct.

(c) In the event Optionee dies while serving as a member of the Board of Directors of the Company or its subsidiaries, if any, or within three months after termination of services for any reason other than gross and willful misconduct, or becomes disabled (within the meaning of Section 22(e)(3) of the Code) while serving as a member of the Board of Directors of the Company or its subsidiaries, if any, and Optionee has not fully exercised this Option, this Option may be exercised at any time within twelve months after Optionee's death or disability by Optionee, the personal representatives, administrators or, if applicable, guardian of Optionee or by any person or persons to whom this Option is transferred by will or the applicable laws of descent and distribution to the extent of the full number of Shares Optionee was entitled to purchase under this Option on the date of death, disability or termination of services, if earlier.

(d) In the event Optionee ceases to serve as a member of the Board of Directors of the Company or its subsidiaries, as a result of voluntary resignation by the Optionee, the Optionee shall continue to have the right to exercise this Option at any time within the original seven year term of this Option to the extent of the full number of Shares Optionee was vested and entitled to purchase under this Option on the date of termination.

(e) In the event Optionee ceases to serve as a member of the Board of Directors of the Company, or its subsidiaries, as a result of resignation at the request of the Company and in connection with an acquisition, merger or substantial investment in a new business venture by the company (a "Requested Resignation"), this Option shall vest and become fully exercisable as to the total amount of 20,000 Shares as of the date of such Requested Resignation and shall continue to be exercisable during its original seven year term.

(f) Notwithstanding the above, in no case may this Option be exercised to any extent by anyone after the Termination Date.

4. Change in Control.

(a) In the event that a "Change in Control" (as hereinafter defined) occurs, all outstanding Options, whether or not vested, shall be subject to the agreement pursuant to which such Change in Control is consummated. Such agreement shall provide for one or more of the following:

- (i) the continuation of such outstanding Options by the Company (if the Company is the surviving corporation);
- (ii) the assumption of such outstanding Options by the surviving corporation or its parent in a manner that complies with Section 424(a) of the Code; or
- (iii) the substitution by the surviving corporation or its parent of new options for such outstanding Options in a manner that complies with Section 424(a) of the Code.

(b) A “*Change in Control*” of the Company shall be deemed to have occurred if:

(i) any “*person*” (as such term is used in Sections 13(d) and 14(d)(2) of the Securities Exchange Act of 1934, as amended (the “*Exchange Act*”)) shall, together with his, her or its “*Affiliates*” and “*Associates*” (as such terms are defined in Rule 12b-2 promulgated under the Exchange Act), become the “*Beneficial Owner*” (as such term is defined in Rule 13d-3 promulgated under the Exchange Act), directly or indirectly, of securities of the Company representing 50% or more of the combined voting power of the Company’s then outstanding securities (any such person being hereinafter referred to as an “*Acquiring Person*”);

(ii) the “*Continuing Directors*” (as hereinafter defined) shall cease to constitute a majority of the Company’s Board of Directors;

(iii) there should occur (A) any consolidation or merger involving the Company and the Company shall not be the continuing or surviving corporation or the shares of the Company’s capital stock shall be converted into cash, securities or other property; *provided, however,* that this subclause (A) shall not apply to a merger or consolidation in which (i) the Company is the surviving corporation and (ii) the stockholders of the Company immediately prior to the transaction have the same proportionate ownership of the capital stock of the surviving corporation immediately after the transaction; (B) any sale, lease, exchange or other transfer (in one transaction or a series of related transactions) of all or substantially all of the assets of the Company; or (C) any liquidation or dissolution of the Company; or

(iv) the majority of the Continuing Directors determine, in their sole and absolute discretion, that there has been a Change in Control.

(c) “*Continuing Director*” shall mean any person who is a member of the Board of Directors of the Company, while such person is a member of the Board of Directors, who is not an Acquiring Person, an Affiliate or Associate of an Acquiring Person or a representative of an Acquiring Person or of any such Affiliate or Associate and who (i) was a member of the Company’s Board of Directors on the date of grant of the Option or (ii) subsequently became a member of the Board of Directors, upon the nomination or recommendation, or with the approval of, a majority of the Continuing Directors.

5. Manner of Exercise.

(a) The Option may only be exercised by Optionee or other proper party within the option period by delivering written notice of exercise to the Company at its principal executive office. The notice shall state the number of Shares as to which the Option is being exercised and shall be accompanied by payment in full of the option price for all of the Shares designated in the notice.

(b) Payment of the exercise price shall be made by certified or bank cashier's check payable to the Company, or by tender of shares of the Company's Common Stock, which, unless the Committee provides its consent, must have been, previously owned by Optionee, having a fair market value on the date of exercise equal to the exercise price of the Option, or a combination of cash and shares equal to such exercise price.

6. **Adjustments.** In the event that any dividend or other distribution (whether in the form of cash, Common Stock, other securities or other property), recapitalization, stock split, reverse stock split, reorganization, merger, consolidation, split—up, spin—off, combination, repurchase or exchange of Common Stock or other securities of the Company, issuance of warrants or other rights to purchase Common Stock or other securities of the Company or other similar corporate transaction or event affects the Common Stock such that an adjustment is necessary pursuant to Section 4(c) of the Plan in order to prevent dilution or enlargement of the benefits or potential benefits intended to be made available under the Plan, and all or any portion of the Option shall then be unexercised and not yet expired, then appropriate adjustments in the outstanding Option shall be made as determined by the Committee in accordance with the provisions of Section 4(c) of the Plan in order to prevent dilution or enlargement of Option rights.

7. Miscellaneous.

(a) *Plan Provisions Control.* In the event that any provision of this Agreement conflicts with or is inconsistent in any respect with the terms of the Plan, the terms of the Plan shall control.

(b) *No Rights of Shareholders.* Neither Optionee, Optionee's legal representative nor a permissible assignee of this Option shall have any of the rights and privileges of a shareholder of the Company with respect to the Shares, unless and until such Shares have been issued in the name of Optionee, Optionee's legal representative or permissible assignee, as applicable.

(c) *No Right to Continuance of Services.* This Agreement shall not confer on Optionee any right with respect to the continuance of any relationship with the Company or any subsidiary of the Company, nor will it interfere in any way with the right of the Company to terminate such relationship at any time.

(d) *Governing Law.* The validity, construction and effect of the Plan and this Agreement, and any rules and regulations relating to the Plan and this Agreement, shall be determined in accordance with the laws of the State of Minnesota.

(e) *Severability*. If any provision of this Agreement is or becomes or is deemed to be invalid, illegal or unenforceable in any jurisdiction or would disqualify this Agreement under any law deemed applicable by the Committee (as defined in the Plan), such provision shall be construed or deemed amended to conform to applicable laws, or if it cannot be so construed or deemed amended without, in the determination of the Committee, materially altering the purpose or intent of the Plan or this Agreement, such provision shall be stricken as to such jurisdiction or this Agreement, and the remainder of this Agreement shall remain in full force and effect.

(f) *No Trust or Fund Created*. Neither the Plan nor this Agreement shall create or be construed to create a trust or separate fund of any kind or a fiduciary relationship between the Company or any affiliate and Optionee or any other person.

(g) *Headings*. Headings are given to the sections and subsections of this Agreement solely as a convenience to facilitate reference. Such headings shall not be deemed in any way material or relevant to the construction or interpretation of this Agreement or any provision thereof.

(h) *Conditions Precedent to Issuance of Shares*. Shares shall not be issued pursuant to the exercise of the Option unless such exercise and the issuance and delivery of the applicable Shares pursuant thereto shall comply with all relevant provisions of law, including, without limitation, the Securities Act of 1933, as amended, the Securities Exchange Act of 1934, as amended, the rules and regulations promulgated thereunder, the requirements of the NASDAQ Global Market or any other applicable stock exchange and the Minnesota Business Corporation Act. As a condition to the exercise of the purchase price relating to the Option, the Company may require that the person exercising or paying the purchase price represent and warrant that the Shares are being purchased only for investment and without any present intention to sell or distribute such Shares if, in the opinion of counsel for the Company, such a representation and warranty is required by law.

(i) *Withholding*. In order to provide the Company with the opportunity to claim the benefit of any income tax deduction which may be available to it upon the exercise of the Option and in order to comply with all applicable federal or state income tax laws or regulations, the Company may take such action as it deems appropriate to assure that, if necessary, all applicable federal or state payroll, withholding, income or other taxes are withheld or collected from Optionee.

(j) *Consultation With Professional Tax and Investment Advisors*. Optionee acknowledges that the grant, exercise, vesting or any payment with respect to this Option, and the sale or other taxable disposition of the Shares acquired pursuant to the exercise thereof, may have tax consequences pursuant to the Code or under local, state or international tax laws. Optionee further acknowledges that such Optionee is relying solely and exclusively on Optionee's own professional tax and investment advisors with respect to any and all such matters (and is not relying, in any manner, on the Company or any of its employees or representatives). Finally, Optionee understands and agrees that any and all tax consequences resulting from this Option and its grant, exercise, vesting or any payment with respect thereto, and the sale or other taxable disposition of the Shares acquired pursuant to the Plan, is solely and exclusively the

responsibility of Optionee without any expectation or understanding that the Company or any of its employees or representatives will pay or reimburse such holder for such taxes or other items.

IN WITNESS WHEREOF, the parties hereto have caused this Agreement to be executed on the Effective Date.

LECTEC CORPORATION

By: _____
Name:
Its:

OPTIONEE

[Optionee]

**CERTIFICATION OF PRINCIPAL EXECUTIVE OFFICER PURSUANT TO SECTION 302 OF THE
SARBANES-OXLEY ACT OF 2002**

I, Greg G. Freitag, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of LecTec Corporation;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officers and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
 - b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principals;
 - c) Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
 - d) Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
5. The registrant's other certifying officers and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: November 15, 2010

/s/ Greg G. Freitag

Greg G. Freitag
Chief Executive Officer

**CERTIFICATION OF PRINCIPAL FINANCIAL OFFICER PURSUANT TO SECTION 302 OF THE
SARBANES-OXLEY ACT OF 2002**

I, Greg G. Freitag, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of LecTec Corporation;

2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;

4. The registrant's other certifying officers and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:

- a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
- b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principals;
- c) Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
- d) Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

5. The registrant's other certifying officers and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):

- a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
- b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: November 15, 2010

/s/ Greg G. Freitag

Greg G. Freitag
Chief Financial Officer

**CERTIFICATION PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002
(SUBSECTIONS (A) AND (B) OF SECTION 1350, CHAPTER 63 OF TITLE 18, UNITED STATES
CODE)**

In connection with the Quarterly Report of LecTec Corporation (the "Company") on Form 10-Q for the quarter ended September 30, 2010 as filed with the Securities and Exchange Commission (the "Report"), I, Greg G. Freitag, Chief Executive Officer and Chief Financial Officer of the Company, certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, to the best of my knowledge that:

1. The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and
2. The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

/s/ Greg G. Freitag

Greg G. Freitag
Chief Executive Officer
(Principal Executive and Financial Officer)
November 15, 2010