

Hamilton Thorne Ltd.
Management Discussion and Analysis
March 31, 2010

The following discussion and analysis of the operations, results, and financial position of Hamilton Thorne Ltd, (the "Company") for the quarter ended March 31, 2010 should be read in conjunction with the Company's March 31, 2010 unaudited consolidated financial statements and the related notes thereto. Such financial statements have been prepared in accordance with Canadian generally accepted accounting principles. The effective date of this report is May 19, 2010. All financial figures are in United States dollars (US) unless otherwise indicated.

Forward-Looking Statements

Certain statements in this management discussion and analysis ("MD&A") may constitute "forward-looking" statements which involve known and unknown risks, uncertainties and other factors which may cause the actual results, performance or achievements of the Company and its subsidiary, or the industry in which they operate, to be materially different from any future results, performance or achievements expressed or implied by such forward-looking statements. When used in this report, the words "estimate", "believe", "anticipate", "intend", "expect", "plan", "may", "should", "will", the negative thereof or other variations thereon or comparable terminology are intended to identify forward-looking statements. Such forward-looking statements reflect the current expectations of the management of the Company with respect to future events based on currently available information and are subject to risks and uncertainties that could cause actual results, performance or achievements to differ materially from those expressed or implied by those forward-looking statements, such as significant changes in market conditions, the inability of the Company to close sales and the inability of the Corporation to attract sufficient financing and including the risk factors summarized below under the heading "Risk Factors". New risk factors may arise from time to time and it is not possible for management of the Company to predict all of those risk factors or the extent to which any factor or combination of factors may cause actual results, performance or achievements of the Corporation to be materially different from those expressed or implied in such forward-looking statements. Given these risks and uncertainties, investors should not place undue reliance on forward-looking statements as a prediction of actual results. Although the forward-looking statements contained in this MD&A are based upon what management believes to be reasonable assumptions, the Company cannot assure investors that actual results will be consistent with these forward-looking statements. The forward-looking statements contained in this MD&A speak only as of the date hereof. The Corporation does not undertake or assume any obligation to release publicly any revisions to these forward-looking statements to reflect events or circumstances after the date hereof or to reflect the occurrence of unanticipated events, except as required by securities legislation.

Description of Operations and Outlook

Hamilton Thorne Ltd. is a Ontario corporation currently trading on the TSX Venture Exchange, as a Tier 2 Corporation, under the stock symbol "HTL". The Company was formed via a reverse takeover of Calotto Capital Inc. on October 28, 2009 ("RTO") and commenced trading on November 5, 2009. The Company's principal business is the development, manufacture and sale of advanced laser systems and instruments for living cell applications in the regenerative medical research and fertility markets.

The Company sells its products both direct and through distributors to pharmaceutical companies, biotechnology companies, fertility clinics, university research centers and other commercial and academic research establishments worldwide.

Hamilton Thorne's ZILOS-tk and XYClone laser systems attach to standard inverted microscopes and operate as robotic micro-surgeons, reducing time and increasing efficiency in key stem cell, embryo and other living cell procedures. Currently the Company has three laser products that have been introduced, one that is pending introduction and three more that are in development. Each member of the laser family is built on the same architecture but serves different market applications. The miniature size, high performance and reasonable price permit laboratories to mount three or four of the Company's lasers onto one microscope and thus have different applications at the researcher's fingertips.

Hamilton Thorne's CASA (Computer Assisted Sperm Analysis) systems are designed to bring quality, efficiency and reliability to studies of reproductive cells in the animal, human infertility and reproductive toxicology fields. These systems assist researchers and clinicians in analyzing sperm motility and other characteristics in human fertility, toxicology and animal applications. These "legacy" products generate meaningful cash flow with relatively little investment in a stable market and are sold through the same distribution channels as the laser family.

The Company is relying for its growth primarily on the growth of the market for instruments in the regenerative medicine (stem cell) field. While this field is rapidly growing, it is also continually evolving and demand for the Company's products has periodically been impacted as certain procedures and research paths increase and decline in laboratory usage. In addition, as the field evolves, the Company's spending priorities may change or be accelerated to address new opportunities in the market.

During the first quarter Hamilton Thorne generated sales increases from its existing products, supported by continued strong demand in Asia and the initial installation of its new Hawk-i product in the US. Given the current instability and currency fluctuations in Europe, the Company is re-focusing its major sales efforts in Asia, particularly in China where the economy continues to be strong, and in North America, where demand is growing, in part as a result of the release of additional NIH funding for stem cell research.

Key Financial Data and Comparative Figures

Income Statements	Quarters Ended March 31							
	2010	2009						
Sales	\$ 1,180,687	\$ 1,115,554						
Gross profit	\$ 710,201	\$ 696,693						
Operating expenses	\$ 1,189,377	\$ 1,001,773						
(Loss) from continuing operations	\$ (479,176)	\$ (389,476)						
(Loss) per share, basic and diluted	\$ (0.02)	\$ (0.02)						
Net income (loss)	\$ (549,125)	\$ (389,476)						
Basic and diluted (loss) per share	\$ (0.02)	\$ (0.02)						
Balance Sheets as at:	31-Mar-10	31-Dec-09						
Cash	\$ 696,343	\$ 1,356,371						
Working capital (deficiency)	\$ 576,718	\$ 1,168,815						
Total assets	\$ 2,098,252	\$ 2,627,983						
Non-current liabilities	\$ 5,054,251	\$ 5,057,904						
Shareholders' (deficiency)	\$ (4,235,529)	\$ (3,726,154)						
Quarterly Data	Mar. 31 10	Dec. 31 09	Sep. 30 09	Jun. 30 09	Mar. 31 09	Dec. 31 08	Sep. 30 08	Jun. 30 08
Sales	\$ 1,180,687	\$ 1,197,160	\$ 1,463,109	\$ 1,018,870	\$ 1,115,554	\$ 1,655,813	\$ 1,426,310	\$ 1,438,163
Gross profit	\$ 710,201	\$ 743,713	\$ 969,971	\$ 614,658	\$ 696,693	\$ 1,111,146	\$ 942,922	\$ 918,286
Operating expenses	\$ 1,189,377	\$ 1,241,584	\$ 1,042,708	\$ 1,047,581	\$ 1,001,773	\$ 1,077,676	\$ 1,150,120	\$ 1,203,086
(Loss) from continuing operations	\$ (479,176)	\$ (577,877)	\$ (185,998)	\$ (521,584)	\$ (389,476)	\$ (73,305)	\$ (319,138)	\$ (400,538)
(Loss) per share, basic and diluted	\$ (0.02)	\$ (0.03)	\$ (0.01)	\$ (0.03)	\$ (0.02)	\$ -	\$ (0.02)	\$ (0.02)
Net income (loss)	\$ (549,125)	\$ (577,877)	\$ (185,998)	\$ (521,584)	\$ (389,476)	\$ (73,324)	\$ (319,138)	\$ (218,398)
Basic and diluted (loss) per share	\$ (0.02)	\$ (0.03)	\$ (0.01)	\$ (0.03)	\$ (0.02)	\$ -	\$ (0.02)	\$ (0.01)

The above financial information has been prepared in accordance with Canadian generally accepted accounting principles and is stated in US dollars.

Results of Operations for quarter ended March 31, 2010

The Company had sales of \$1,180,687 during the quarter ended March 31, 2010 which was up from \$1,115,554 during the previous year. The sales increase was attributable primarily to the general improvement in worldwide economic conditions. The Company's customers in the regenerative medicine field primarily use the Company's products in research applications. During 2009, the Company's sales, particularly in North America and Europe, were adversely impacted by budget freezes and extended purchasing cycles for research products.

Cost of sales for the quarter ended March 31, 2010 totaled \$470,486 compared to \$418,861 for the same period in 2009. The gross profit as a percent of sales declined slightly from 62.5% for 2009 to 60.2% for 2010, primarily due to certain lower margin accessories sold in connection with its initial Hawk-I installation, offset by the fact that manufacturing costs are largely fixed and did not increase proportionately with the increase in sales.

Operating expenses increased to \$1,189,377 for the quarter ended March 31, 2010 as compared with \$1,001,773 for the prior year quarter. Research and development expenses increased to \$250,307 from \$176,773 due primarily to a general increase in third party development contracts and patent costs. General and administrative expenses increased from \$382,465 to \$453,383 due primarily to the increase in stock compensation costs and public company expenses. Sales and marketing expenses increased from \$442,535 to \$485,687, primarily due to additional compensation expense.

Interest expense decreased from \$84,605 to \$70,159. The decrease was due to the elimination of interest accreted on the redeemable preferred stock offset by the higher average interest rate on the bank line of credit during 2010.

The net loss for the quarter ended March 31, 2010 was \$549,125, an increase from the net loss of \$389,476 for the same period of the previous year. The increase was due to the increase in operating expenses, partially offset by increased gross profit from the increased sales.

Liquidity

The Company's cash balance at March 31, 2010 was \$696,343 as compared with \$1,356,371 at December 31, 2009. Working capital decreased from \$1,168,815 at December 31, 2009 to \$576,718 at March 31, 2010. The declines in cash and working capital are as a result of the net losses incurred by the Company during the quarter ended March 31, 2010. Cash used by operations was \$617,291 for the quarter ended March 31, 2010 compared to \$5,656 provided in the prior year due primarily to high accounts receivable collections in the 2009 quarter. The Company's continued strong gross margins provide funds to help cover expanding operating expenses, provided sales continue to grow and the planned new products are brought online as scheduled.

Although the Company currently has positive working capital, it should be noted that the Company has incurred recurring losses over its history and has been able to meet its obligations through equity and debt financings. The Company's bank line of credit matures on October 1, 2011 and the Company is now examining options to meet that obligation, including reviewing the possibilities of refinancing/extending the note, raising additional equity and other financing alternatives.

The Company is expected to continue to use cash in funding its operations during 2010. The Company believes that its current cash position, reduced by cash used in operations, will be sufficient to support operations through all of 2010.

Capital Resources

As the Company is primarily an assembler, as opposed to a "manufacturer from scratch", it does not have significant capital expenditures as part of its financial plans and outlook. It is expected that its capital expenditures in the coming year will be limited to new computers, software and fixtures and will total approximately \$120,000. Most of these expenditures will be purchased with current funds and some may be financed through leasing arrangements.

The purpose of the RTO and private placement in 2009 was to provide financing to undertake the continued development of existing laser products, expansion of the laser product family and to get these products to market in 2010 and ensuing years. It is expected that current funds will be used to pay for these expenses.

Share Capital

As of March 31, 2010, there were 24,415,157 common shares issued and outstanding.

As of March 31, 2010, there were 5,500,005 warrants outstanding to purchase common shares at a price of Cdn \$0.60. The warrants all expire in April 2011.

Stock options issued to employees and directors outstanding at March 31, 2010 totaled 3,532,756 at prices ranging from Cdn \$0.2176 to Cdn \$0.7712255. Options for 1,518,695 shares are exercisable as of March 31, 2010. Options expire at varying times from November 2010 through March 2020.

Agent compensation options outstanding at March 31, 2010 totaled 440,001 options to purchase units at a price of Cdn \$0.40. Each unit consists of one common share and a warrant to purchase a common share at a price of Cdn \$0.60. The options and warrants will expire in April 2011.

A subordinated convertible note for \$50,000 was outstanding at March 31, 2010. The note was issued in September 2009 to a then director. The note was convertible, at the option of the holder, in the next round of equity financing that raised a minimum of \$1.5 million (exclusive of the noteholders participation in the equity financing) at the same terms and conditions as other investors participating. The noteholder chose not to convert this note in the financing. Had the noteholder participated in the financing, the noteholder would have received 132,500 common shares. It is not possible to determine the number of shares issuable as of March 31, 2010 to this now shareholder.

Related Parties

At March 31, 2010, the Company was indebted to certain officers under various unsecured notes payable bearing interest at 7%. In December 2008 an officer of the Company lent \$50,000 on a promissory note payable, subordinated to the bank line of credit, with interest at the prime rate plus 1%. Total indebtedness to officers at March 31, 2010 amounted to \$56,049.

In January 2007, the Company lent \$20,000 to an officer on an unsecured promissory note payable at maturity on December 31, 2009, bearing interest at the prime lending rate plus 1%. At March 31, 2010, total indebtedness to the Company totaled \$24,023.

As of March 31, 2010 an officer owed the Company approximately \$24,000 for advances and an advance against future pay.

In April 2010 the Board of Directors, with the permission of the bank providing the Company's line of credit, allowed the offset of the above \$20,000 note to an officer, along with accrued interest of \$4,023, and \$24,042 of advances to the same officer against the \$50,000 note due to that officer and the accrued interest due on the note.

New Accounting Pronouncements

Business Combinations

Section 1582, Business Combinations, replaces Section 1581, Business Combinations. The section establishes standards for the accounting for a business combination. It provides the Canadian equivalent to the IFRS standard, IFRS 3 (Revised), Business Combinations. The section applies prospectively to business combinations for which the acquisition date is on or after the beginning of the first annual reporting period beginning on or after January 1, 2011. The new standard is not expected to have a material effect on the Company's consolidated financial statements.

Consolidated Financial Statements

Section 1601, Consolidated Financial Statements and Section 1602, Non-Controlling Interests, together replace Section 1600, Consolidated Financial Statements. Section 1601 establishes standards for the preparation of consolidated financial statements. Section 1602 establishes standards for accounting for a non-controlling interest in a subsidiary in consolidated financial statements subsequent to a business combination. It is equivalent to the corresponding provisions of IFRS standard, IAS 27 (Revised), Consolidated and Separate Financial Statements. The sections apply to interim and annual consolidated financial statements relating to fiscal years beginning on or after January 1, 2011. Earlier adoption is permitted as of the beginning of a fiscal year. The Company is currently evaluating the impact of the adoption of these new sections on the consolidated financial statements.

International Financial Reporting Standards

In February 2008, Canada's Accounting Standards Board (AcSB) confirmed that Canadian GAAP, as used by publicly accountable enterprises, will be fully converged into IFRS, as issued by the International Accounting Standards Board (IASB). The changeover date is for interim and annual financial statements relating to fiscal years beginning on or after January 1, 2011. Therefore, the Company will be required to report under International Financial Reporting Standards ("IFRS") no later than in the first quarter of 2011, with restatement of comparative information. The conversion to IFRS will impact the Company's accounting policies, systems, disclosures and controls. The Company has identified the major differences between its current accounting policies and those required or expected to apply in preparing IFRS financial statements and has begun the process of developing a conversion plan. During the second quarter of 2010, the Company expects to quantify the differences between GAAP and IFRS reporting and determine their materiality. Updates regarding the progress of the conversion plan will be provided to the Company's Audit Committee on a quarterly basis.

EIC-175

In December 2009, the CICA issued EIC 175, Multiple Deliverable Revenue Arrangements, replacing EIC 142, Revenue Arrangements with Multiple Deliverables. The accounting changes summarized in EIC-175 are effective for fiscal years beginning on or after January 1, 2011, with early adoption permitted. Adoption may either be on a prospective basis or by retrospective application. If the Abstract is adopted early, in a reporting period that is not the first reporting period in the entity's fiscal year, it must be applied retroactively from the beginning of the Company's fiscal period of adoption. The Company is currently assessing the future impact of these amendments on its consolidated financial statements and has not yet determined the timing and method of adoption.

Risk Factors

An investment in the Company must be considered highly speculative due to the relatively early stage of the development of its current operations. There are trends and factors that may be beyond the Company's control which affect its operations and business. Such trends and factors include adverse changes in the conditions in the specific markets for the Company's products and services, the conditions in the broader market of laboratory instruments, consumables and accessories and conditions in the domestic or global economy generally. It is not possible for management to predict economic fluctuations and the impact of such fluctuations on its performance.

1. General Economic Conditions – The demand for capital asset purchases declines in the face of difficult economic conditions such as those experienced in the United States and much of the rest of the world during 2008 and 2009. The Company's customers include pharmaceutical and chemical companies, laboratories, universities, IVF labs, government agencies and public and private research institutions. Many factors, including public policy spending priorities, available resources and product and economic cycles, have a significant effect on the capital spending policies of these entities. These policies in turn can have a significant effect on the demand for its products.

2. History of Losses – The Company has a history of losses and cannot predict the extent of future losses. The Company may not achieve profitability in the foreseeable future, if at all. Its ability to generate profits in the future will depend on a number of factors, including: (i) its ability to grow sales based on the continued market demand for its existing products and expected demand for additional products; (ii) costs relating to the commercialization, sale and marketing of its products; (iii) general and administrative costs relating to its operations; (iv) research and development costs; and (v) charges related to purchases of technology or other assets.

3. Limited Operating History in Certain Markets - The Company will be a small company focused on commercializing, marketing and selling products in the in vitro fertilization, transgenic and regenerative medical research markets. Its operating history in some of these markets is extremely limited. Its laser products for the regenerative medical market are in the early stages of commercialization. Other products are only in the early stages of development. An investor should evaluate the likelihood of financial and operational success in light of the uncertainties and complexities present in an early-stage company, many of which are beyond the Company's control, including: (i) the Company's potential inability to distribute, sell and market its products; and (ii) the significant investment to achieve its commercialization, marketing and sales objectives. The regenerative medicine market is relatively new and its long-term growth prospects are uncertain. Should the regenerative medicine market fail to expand, it could have a materially adverse effect on the Company's business and financial condition.

4. Product Development - The development of additional products is subject to the risks of failure inherent in the development of new, state of the art products, laboratory devices and products based on new technologies. These risks include: (i) delays in product development or manufacturing; (ii) unplanned expenditures for product development or manufacturing; (iii) failure of new products to have the desired effect or an acceptable accuracy profile; (iv) emergence of superior or equivalent products; (v) failure by any potential collaborative partners to successfully develop products; and (vi) the dependence on third parties for the manufacture, development and sale of the Reporting Issuer's products. Because of these risks, the Company's research and development efforts or those of potential collaborative partners may not result in any commercially viable products. If a significant portion of these development efforts is not successfully completed, or any products are not commercially successful, the Company is less likely to business, financial condition and results of its operations.

5. Technological Advancement - The areas in which the Company is commercializing, distributing, and/or selling products involve rapidly developing technology. There can be no assurance that the Company will be able to establish itself in such fields, or, if established, that it will be able to maintain its position. There can be no assurance that the development by others of new or improved products will not make the Company's present and future products, if any, superfluous or obsolete.

6. Intellectual Property Rights - Although three patents have now been issued in the United States, all applications are still pending in all overseas jurisdictions as well as continuations in part and new applications still pending in the United States. The Company's success and ability to compete are substantially dependent on these patents. Although management of Hamilton Thorne believes that the patents and associated trademarks and licenses are valid, there can be no assurance that they will not be challenged and subsequently invalidated and/or canceled. The invalidation or cancellation of any one or all of the patents or trademarks would significantly damage the Company's commercial prospects. Further, the Company may find it necessary to legally challenge parties infringing its patents or trademarks or licensed trademarks to enforce its rights thereto. There can be no assurance that any of the patents would ultimately be held valid or that efforts to defend any of the patents, trade secrets, know-how or other intellectual property rights would be successful.

The Company's future success will depend, in part, on its ability to obtain patents for newly developed products, maintain trade secrets protection, and operate without infringing on the proprietary rights of third parties or having third parties circumvent its rights. The patent position of regenerative medicine firms is uncertain and involves complex legal and financial questions for which, in some cases, certain important legal principles remain unresolved. There can be no assurance that the patent applications made in respect of the owned products will result in the issuance of patents, that the term of a patent will be extendable after it expires in due course, that any patent issued to the Company will provide it with any competitive advantages, that the patents of others will not impede the Company's ability to do business or that third parties will not be able to circumvent or successfully challenge the patents obtained in respect of the products. The cost of obtaining and maintaining patents is high. Furthermore, there can be no assurance that others will not independently develop similar products which duplicate any of the products, or, if patents are issued, design around the patent for the product. There can be no assurance that the Company's processes or products do not or will not infringe upon the patents of third parties, or that the scope of the Company's patents will successfully prevent third parties from developing similar and competitive products.

Much of the Company's know-how and technology may not be patentable, though they may constitute trade secrets. There can be no assurance, however, that the Company will be able to meaningfully protect its trade secrets. To help protect its intellectual property rights and proprietary technology, the Company requires employees, consultants, advisors and collaborators to enter into confidentiality agreements. There can be no assurance that these agreements will provide meaningful protection for the Company's trade secrets, know-how or other proprietary information in the event of any unauthorized use or disclosure.

The Company's commercial success will also depend, in part, on not infringing on the patents or proprietary rights of others. There can be no assurance that the technologies and products used or developed by the Company will not infringe such rights. If such infringement occurs and the Company is not able to obtain a license from the relevant third party, it will not be able to continue the development, manufacture, use, or sale of any such infringing technology or product. There can be no assurance that necessary licenses to third-party technology will be available at all or on commercially reasonable terms. In some cases, litigation or other proceedings may be necessary to defend against or assert claims of infringement or to determine the scope and validity of the proprietary rights of third parties. Any potential litigation could result in substantial costs to, and diversion of, its resources and could have a material and adverse impact on the Company. An adverse outcome in any such litigation or proceeding could subject the Company to significant liabilities, require it to cease using the subject technology or require it to license the subject technology from the third party, all of which could have a material adverse effect on the Company's business.

The Company's future success and competitive position depends in part upon its ability to maintain its intellectual property portfolio. There can be no assurance that any patents will be issued on any existing or future patent applications. Even if such patents are issued, there can be no assurance that any patents issued or licensed to the Company will not be challenged. The Company's ability to establish and maintain a competitive position may be achieved in part by prosecuting claims against others who it believes to be infringing its rights. In addition, enforcement of the Company's patents in foreign jurisdictions will depend on the legal procedures in those jurisdictions. Even if such claims are found to be invalid, the Company's involvement in intellectual property litigation could have a material adverse effect on its ability to distribute any products that are the subject of such litigation. In addition, the Company's involvement in intellectual property litigation could result in significant expense, which could materially adversely affect the use or distribution of related intellectual property and divert the efforts of the Company's technical and management personnel from their principal responsibilities, whether or not such litigation is resolved in the Company's favor.

7. Competition - The Company is engaged in a rapidly evolving field. Competition for its laser products from numerous companies is expected to increase. Competition from other unknown entities and competition from research and academic institutions is also expected to increase. The market for solutions to the many regenerative medical research problems is growing rapidly and is likely to attract new entrants. Numerous biotechnology companies have focused on developing new media or devices and most, if not all, of these companies have greater financial and other resources and development capabilities than the Company. The Company's future success depends in part on its ability to maintain a competitive position, including its ability to further progress and develop its products for sale and commercialization. Other companies may succeed in commercializing products earlier than the Company or they may succeed in developing products that are more effective than the Company's products. While the Company will seek to expand its technological capabilities in order to remain competitive, there can be no assurance that developments by others will not render its products non-competitive or that the Company will be able to keep pace with technological developments. The success of the Company's competitors and their products relative to the Company's products could have a material adverse effect on the future operations of the Company.

In addition to competing with universities and other research institutions in the development of products, technologies and processes, the Company may compete with other companies in acquiring rights to products or technologies from universities. There can be no assurance that the Company's products, existing or to be developed, will be more effective or achieve greater market acceptance than competitive products, or that its competitors will not succeed in developing products and technologies that are more effective than those being developed by or that would render its products and technologies less competitive or obsolete.

8. Product Liability - The sale of the Company's products may expose it to potential liability resulting from the sale and use of such products. Liability might result from claims made directly by consumers or by laboratory companies. Hamilton Thorne currently maintains US\$3 million of product liability insurance. There can be no assurance that the Company will be able to renew its current insurance, renew it at a rate comparable to what it now pays, or that the coverage will be adequate to protect it against liability. If it were held liable for a claim or claims exceeding the limits of its current or future insurance coverage, or if coverage was discontinued for any reason, it could have a materially adverse effect on the Company's business and financial condition.

9. Government Regulation - The growth of the regenerative medicine market in the United States and certain other countries has been hampered by government regulations regarding the use of embryonic stem cells. The current political administration in the United States is more supportive of stem cell research than the prior administration, however, there is no assurance that this support will continue and the support could conceivably be reversed if a new administration is elected in 2012 or thereafter. Should these regulations be readopted, stiffened, or extended to other jurisdictions, the market for the Company's products could be adversely affected which could have a materially adverse effect on the Company's business and financial condition.

The Company's business may also be affected in varying degrees by changes to government regulation of intellectual property or export controls. Such changes are beyond the control of the Company and the effect of any such changes cannot be predicted.

The Company conducts its business internationally and is subject to laws and regulations of several countries which may affect its ability to access regulatory agencies and may affect the enforceability and value of its intellectual property rights. There can be no assurance that any sovereign government, including Canada's or the United States', will not establish laws or regulations that will be deleterious to the Company's interests. There is no assurance that the Company, as a Canadian corporation, will continue to have access to the regulatory agencies in any jurisdiction where it might want to obtain final regulatory approval, and there can be no assurance that the Company will be able to enforce its intellectual property rights in foreign jurisdictions. Governments have, from time to time, established foreign exchange controls which could have a material adverse effect on the Company's business and financial condition, since such controls may limit its ability to flow funds or products into a particular country to meet its obligations under distribution agreements and to flow funds which the Company is entitled to, in the form of sales proceeds, out of a particular country.

10. Dependence upon Management - The Company is substantially dependent upon the services of a few key personnel. The loss of the services of any of these personnel could have a material adverse effect on the business of the Company. The Company maintains key man insurance on certain management personnel. In order to pursue its marketing and commercialization plans, the Company will need to hire additional personnel with experience in marketing and finance. The Company may not be able to attract and retain personnel on acceptable terms given the intense competition for such personnel among high technology enterprises, including biotechnology, and healthcare companies, universities and non-profit research institutions. If it loses any of these persons, or is unable to attract and retain qualified personnel, its business, financial condition and results of operations may be materially and adversely affected.

11. Financing - The Company currently maintains a \$5,000,000 secured line of credit with a US bank that currently matures October 1, 2011. This line of credit is secured by two letters of credit, issued by shareholders, which expire on November 1, 2011. If the Company is unable to extend such letters of credit or refinance or pay off the debt to the Bank, or if there were otherwise an event of default, the Company's business, financial condition and results of operations may be materially and adversely affected.

The Company may need to issue common shares to raise additional capital and/or as full or partial consideration in connection with future acquisitions. To the extent that it does so, existing shareholders will be diluted and the trading price of its shares may also decrease.

12. Dividends - The Company intends to retain any future earnings to finance the growth and development of its business and does not plan to pay cash dividends in the foreseeable future, if ever.

13. Principal Stockholder Influence - The Company's principal stockholders own or control approximately 60% of the shares outstanding and therefore have significant ability to control the outcome of stockholder votes, including votes concerning the election of directors, the adoption or amendment to provisions in the Company's articles or by-laws, the approval of mergers and/or acquisitions, decisions affecting capital structure and other significant