

Management Discussion and Analysis of Financial Condition and Results of Operations (As of June 28, 2007)

This MD&A contains projections and other forward-looking statements regarding future events. Such statements are predictions, which may involve known and unknown risks, uncertainties and other factors, which could cause the actual events or results and company plans and objectives to differ materially from those expressed. For information concerning factors affecting the Company's business, the reader is referred to the documents that the Company files from time to time with applicable Canadian securities and regulatory authorities.

This discussion and analysis of the results of operations of Quest PharmaTech Inc. (“Quest” or the “Company”) should be read in conjunction with the unaudited consolidated financial statements and accompanying notes for the three months ended April 30, 2007 and the audited consolidated financial statements for the years ended January 31, 2007 and January 31, 2006. This discussion and analysis provides an update to the discussion and analysis prepared for the year ended January 31, 2007. The unaudited consolidated financial statements have been prepared in accordance with generally accepted accounting principles in Canada (Canadian GAAP) and have not been reviewed by the Company’s auditors. This discussion and analysis provides information on the operations of Quest on a consolidated basis. All amounts are expressed in Canadian dollars unless otherwise noted and references to the term “year” refer to the fiscal year ended January 31st. Additional information related to the Company is on SEDAR at www.sedar.com.

Overview

Quest is committed to the development and commercialization of new pharmaceutical products. It is developing a series of products for the treatment of cancer and dermatological conditions based on its unique photodynamic and sonodynamic therapy platform.

A summary of the Company’s projects follows.

HYPOCRELLIN-BASED TECHNOLOGIES

This technology platform is based on a unique, non-toxic family of photosensitizing and sonosensitizing compounds. The active ingredient is a derivative of Hypocrellin B (“HB”), a small molecular compound isolated from parasitic fungi on bamboo. Quest has formulated HB derivatives into a topical gel (SL017), and an injectable solution (SL052). The drug is delivered to the target tissue and then the target tissue is illuminated. This activates the photosensitizer which has the potential to destroy the diseased tissue, predominately through the action of singlet oxygen.

HB Topical - SL017

This gel penetrates skin and can potentially be used to treat various skin conditions such as acne, actinic keratosis, psoriasis, etc. HB gels target a large patient population and will face comparatively less stringent regulatory requirements than injectible HB compounds. The Company has reported positive results from a phase I clinical trial for acne and hair removal, and has initiated a 50 patient clinical trial to determine the appropriate light dose to be used with SL017 for cosmetic hair removal applications. Additionally, the Company recently completed the enrolment of 12 patients in a Phase I clinical study for PDT treatment of actinic keratosis.

HB Injectable – SL052

Quest has developed SL052, which may have utility in the photodynamic therapy of prostate cancer. During 2004, the Company entered into an agreement with Dr. Ronald B. Moore, the Alberta Cancer Board and the Cross Cancer Institute to conduct preclinical studies on SL052. Over the next year, the Company will continue to focus on the development of HB Injectable for prostate cancer and will incur additional costs associated with accumulating preclinical data in order to advance this technology into clinical trials.

Quest is also developing SL052 (SDT) for peritoneal carcinomatosis (cancer of the abdominal cavity and linings) to demonstrate the proof-of-principle for a sonodynamic therapy approach for the treatment of cancer. The Company intends to seek a partner to undertake development of an ultrasound transducer suitable for abdominal and thoracic activation of the sonosensitizer, SL052, previously administered to the ascites fluid or pleural effusion of patients with advanced carcinomatosis.

OTHER DRUG DEVELOPMENT PROGRAMS

2127 (formerly CDK Immunomodulator)

2127 is a novel immunomodulator which inhibits cycline-dependent kinases (“CDKs”) and prevents the growth of cancer cells. CDKs are the proteins that control the growth cycle of cancer cells. Initial studies demonstrated unique advantages of 2127 over other CDK inhibitors, and the Company has filed patent applications to protect the intellectual property generated to date. Further development of 2127 is on hold pending receipt of additional funding.

Results of Operations

Net consolidated loss for the three months ended April 30, 2007 was \$137,373 or \$0.00 per share on a fully diluted basis as compared to a consolidated loss of \$472,863 or \$0.01 per share for the three months ended April 30, 2006. After adjusting for non-cash items, cash flows provided by (used in) continuing operations for the three months ended April 30, 2007 were \$83,568 as compared to (\$71,482) for the three months ended April 30, 2006.

Expenses

The following table identifies the changes in General and Administrative expense for the three month period ended April 30, 2007 compared to the three month period ended April 30, 2006.

General and administrative expense	2007	2006	Increase (decrease)
	\$	\$	\$
Salaries, wages and benefits	47,341	52,076	(4,735)
Other support costs	14,522	7,740	6,782
Consulting	81,000	44,500	36,500
Legal fees	4,633	2,879	1,754
Public company related costs	17,181	9,465	7,716
Rent	1,797	2,762	(965)
Travel	747	5,346	(4,599)
Insurance	4,610	14,747	(10,137)
Total general and administrative expense	171,831	139,515	32,316

Salaries, wages and benefits, rent and travel expenses have decreased due to the Company's continued cost containment efforts. Consulting costs have increased due to an increase in consulting and business development activities in Q1, fiscal 2008. Public Company related costs have increased as a result of the Company's activity related to the issuance of private placement units during the period. Insurance costs have decreased due to the allocation of a portion of insurance expense to R&D activities in Q1, fiscal 2008.

The following table identifies the changes in research and development (R&D) expense for the three month period ended April 30, 2007 compared to the three month period ended April 30, 2006.

Research and development expense	2007	2006	Increase (decrease)
	\$	\$	\$
Salaries, wages and benefits	83,551	105,571	(22,020)
Sub-contract and consulting	27,132	24,792	2,340
Rent	2,695	17,773	(15,078)
Legal (patent prosecution)	8,819	44,172	(35,353)
Supplies	4,756	25,414	(20,658)
Other R&D costs	10,417	22,896	(12,479)
Gross research and development expense	137,370	240,618	(103,248)
Less			
NRC-IRAP funding	(13,340)	-	(13,340)
Alberta Ingenuity funding	(12,000)	(12,000)	-
Research and development expense (net)	112,030	228,618	(116,588)

R&D expenses have decreased due to the Company's continuing efforts to cut costs where possible in order to conserve resources.

Discontinued Operations

As part of the Company's decision in February, 2005 to divest itself of its non-core assets, the Company has presented the activities related to non-core assets as discontinued operations.

The following table identifies the activity in connection with the Company's discontinued operations for the three month period ended April 30, 2007 compared to the three month period ended April 30, 2006.

Discontinued operations	2007	2006	Increase (decrease)
	\$	\$	\$
Revenue	9,262	25,032	(20,032)
Direct costs	-	9,354	(9,354)
Gross Margin	9,262	15,678	(10,678)
General and administrative expenses	-	31,229	(35,491)
Amortization expense	-	32,266	(32,266)
Interest expense	-	12,924	(12,924)
Interest income	-	(17)	17
Income / (loss) from discontinued operations	9,262	(60,724)	(69,896)

During the three month period ended April 30, 2007, the Company received incidental revenue of \$5,000 in connection with ACP-HIP and also recognized \$4,262 as a recovery of a bad debt in connection with Accu-MAB.

Revenues

For the three month period ended April 30, 2007, the Company recognized revenue related to licensing fees and marketing distribution rights. The following table identifies the changes in revenue for the three month period ended April 30, 2007 compared to the three month period ended April 30, 2006.

Revenue	2007	2006	Increase (decrease)
	\$	\$	\$
Continuing operations			
Licensing fees and market distribution rights	168,005	-	168,005
Total revenue from continuing operations	168,005	-	168,005
Discontinued operations			
Accu-MAB TM	4,262	23,020	(23,020)
Pharmaceutical pellet core	-	2,012	(2,012)
ACP-HIP	5,000	-	5,000
Total revenue from discontinued operations	9,262	25,032	(20,032)
Total revenue	177,267	25,032	147,973

Stock-Based Compensation Expense

During the three month period ended April 30, 2007, the Company granted a total of nil (for the three month period ended April 30, 2006 – 150,000) stock options, as per the Company's Stock Option Plan, including nil (2006 – nil) to employees and nil (2006 – 150,000) to non-employees. The fair value of these options, nil (2006 - \$4,500), was recognized as an expense and credited to contributed surplus for the three month period ended April 30, 2007 and 2006 respectively.

Liquidity and Capital Resources

As noted in Risks and Uncertainties below, the Company's ability to continue as a going concern is uncertain and is dependent upon its ability to raise additional capital to successfully complete its research and development programs, commercialize its technologies, conduct clinical trials and receive regulatory approval for its products.

At April 30, 2007, cash and cash equivalents was \$202,589 as compared to \$123,022 at January 31, 2007.

During the quarter, the Company completed an equity offering which provided gross proceeds of \$960,000. Also during the quarter, the Company recognized \$25,340 in R&D grant funding from the Alberta Ingenuity Fund and NRC-IRAP.

Funds used in operating activities was \$693,432 for the three month period ended April 30, 2007 compared to \$221,482 for the three month period ended April 30, 2006. The large increase was primarily due to a pay-down of accounts payable and accrued liabilities.

Subsequent to quarter end, the Company completed two additional equity offerings which provided gross proceeds of \$300,000 and \$750,000, respectively. The Company has used a portion of these offering proceeds to make a \$200,000 principal payment against the \$1,000,000 convertible debenture and a \$50,000 principal payment against the \$230,000 demand notes.

The Company continues to implement a disciplined approach to containing costs and is focusing on programs aimed at achieving near-term goals.

Quest's funding needs will vary as its drug development products move into and through clinical trials.

Based on current operating budgets and assuming the ongoing divestiture of non-core assets, management believes that the capital resources of the Company should be sufficient to fund operations to the fourth quarter of fiscal 2008.

The Company will seek additional capital through the sale of the remaining non-core assets, further equity financings, licensing arrangements involving its core technologies, strategic partnerships and/or financings from directors.

Demand Notes and Related Party Transactions

In June, 2007, the Company made a principal payment of \$50,000 towards the demand note held by a company controlled by Dr. Donald Rix, a Director of the Company.

Under the Company's March, 2007 private placement, \$165,000 was raised from Officers and Directors of the Company.

Outstanding Share Data

As at June 28, 2007, there were 60,962,580 common shares issued and outstanding, 3,273,000 stock options and 4,800,000 share purchase warrants.

Disclosure Controls and Procedures

The management of Quest is responsible for establishing and maintaining disclosure controls and procedures for the Company and has designed such disclosure controls and procedures, or caused them to be designed under Quest management's supervision, to provide reasonable assurance that material information relating to the Company, including its consolidated subsidiaries, is made known to Quest management by others within those entities particularly during the period covered by this MD&A.

Quest management has evaluated the effectiveness of the Company's disclosure controls and procedures for the period covered by this MD&A and based on that evaluation, management has concluded that the disclosure controls and procedures are effective.

Internal Control Over Financial Reporting

The Company's management is responsible for establishing and maintaining adequate internal control over financial reporting. Management has considered the effectiveness of design of the Company's internal controls and procedures over financial reporting and has noted several weaknesses in internal controls over financial reporting such as a lack of segregation of duties because of limited staff members.

Management intends to initiate steps to remedy the noted shortcomings over the next fiscal year by carrying out a management assessment of the weaknesses with a view to improving areas where weaknesses exist and implementing procedures aimed at minimizing the risk of material error in its financial reporting.

Risks and Uncertainties

Going concern uncertainty

The Company's financial statements have been prepared on a going concern basis which presumes the realization of assets and discharge of liabilities in the normal course of business for the foreseeable future. The Company has experienced significant operating losses and cash outflows from operations since its inception and it is expected to continue to experience negative cash flows from operations in the coming fiscal year. The Company had a working capital deficiency of \$1,043,614 and a shareholders' deficiency of \$1,069,587 as at April 30, 2007. The Company's ability to continue as a going concern is uncertain and is dependent upon its ability to raise additional capital to successfully complete its research and development programs, commercialize its technologies and conduct clinical trials and receive regulatory approvals for its products. It is not possible at this time to predict the outcome of these matters.

Quest's proprietary technologies are in various stages of development and some technologies have not received regulatory approval to begin clinical trials. It will be necessary for the Company to produce sufficient preclinical data in order to receive regulatory approval to begin clinical trials. There is no assurance that regulatory approval will be received to begin clinical trials. For the proprietary technologies that have received regulatory approval to begin clinical trials, future success will depend upon the ability of the Company to move the products through clinical trials, the effect and safety of these products, the timing and cost to receive regulatory and marketing approvals and the filing and maintenance of patent claims.

Quest's proprietary technologies have exposure to risks associated with commercialization. Even after product approval is obtained, there is no assurance that the Company will have a sufficient market for its products or the working capital required for commercialization.

The Company maintains clinical trial liability and product liability insurance; however, it is possible that this coverage may not provide full protection against all risks.

The Company may be exposed to risks associated with malfunctioning equipment, catastrophic events and other events within and outside of the Company's control. The Company maintains insurance believed to be adequate to cover any eventuality, but there is no guarantee that coverage will be sufficient for all purposes.

To a large degree, the Company's success is dependant upon attracting and retaining key management and scientific personnel to further the Company's drug development programs. There is a risk that required personnel may not be available to the Company when needed and, as a result, this may have a negative impact on the Company.

Quest must continue to raise additional capital through the exercise of stock options and warrants, issuing new share capital through equity financing, licensing arrangements and/or strategic partnerships. The Company's ability to raise additional capital will depend upon the progress of moving its drug development products into and through clinical trials and the strength of the equity markets, which are uncertain. There can be no assurance that additional capital will be available.