

PALADIN LABS INC.
CONSOLIDATED FINANCIAL STATEMENTS

March 31, 2010



Management Discussion and Analysis:

All numbers are in thousands of Canadian dollars except for share and per share amounts

This management's discussion and analysis provides our overview of the Company's operations, performance and financial condition for the quarter ended March 31, 2010 and compares these unaudited quarterly results to those of the quarter ended March 31, 2009. It is intended to complement and supplement financial information included in the interim and annual consolidated financial statements, related notes, other financial information found elsewhere in our annual report and in our annual information form or other documents filed on SEDAR at www.sedar.com. As a result, it should be read in conjunction with such financial information. This management's discussion and analysis is current as at May 7, 2010 and as at this date 18,647,453 shares and 1,443,117 options were issued and outstanding. Reference to "Paladin" or the "Company" includes Paladin Labs Inc. and all its subsidiaries including the acquisition of Isotechnika Inc. ("Isotechnika") as of June 18, 2009, the effective date of the acquisition, further described in note 4 to these unaudited interim consolidated financial statements.

Forward-Looking Statements

This document contains forward-looking statements for the Company and its subsidiaries. These forward looking statements, by their nature, necessarily involve risks and uncertainties that could cause actual results to differ materially from those contemplated by the forward-looking statements. The Company considers the assumptions on which these forward-looking statements are based to be reasonable at the time they were prepared, but cautions the reader that these assumptions regarding future events, many of which are beyond the control of the Company and its subsidiaries, may ultimately prove to be incorrect. Factors and risks, which could cause actual results to differ materially from current expectations, are discussed in the Company's Annual Report as well as in the Company's Annual Information Form for the year ended December 31, 2009. The Company disclaims any intention or obligation to update or revise any forward-looking statements whether as a result of new information, future events, except as required by law. For additional information on risks and uncertainties relating to these forward-looking statements, investors should consult the Company's ongoing quarterly filings, Annual Report and Annual Information Form and other filings found on SEDAR at www.sedar.com.

Overview

Paladin is a specialty pharmaceutical company focused on developing, acquiring, in-licensing, marketing, and distributing innovative pharmaceutical products.

First quarter highlights:

- Revenues reached \$30,837, an increase of 19% over the same period last year
- Net income was \$3,045, remaining steady over the same period last year
- Cash flows from operations reached \$10,365, a 25% increase over the same period last year
- EBITDA¹ was \$11,546, an increase of 5% over the same period last year
- Filed a new drug submission for Abstral[®] sublingual fentanyl, which has been accepted for priority review by Health Canada
- Amended its existing agreements with IsoPharma (TSX: ISA) giving Paladin full share of future net profits of the Isodiagnostika line of diagnostic products
- Entered into a distribution agreement with Protherics Inc., a wholly-owned subsidiary of BTG plc (LSE: BGC), for the exclusive Canadian rights to market and sell DigiFab[™] (Digoxin Immune Fab (Ovine))
- Received approval from the Toronto Stock Exchange ("TSX") on March 1, 2010 to carry out a normal course issuer bid to purchase up to 1,102,000 common shares
- Completed a strategic investment in SpePharm Holding B.V., a rapidly-growing, specialty pharmaceutical company focused on acquiring, registering and marketing high medical value specialty medicines throughout Europe
- Completed a strategic investment to acquire an initial 34.99% ownership interest in PharmaPlan (Pty) Ltd., a leading privately owned specialty pharmaceutical company, based in Johannesburg, South Africa, dedicated to the execution of a search-acquire-commercialize business model in the South African and broader sub-Saharan African region

- Subsequent to the quarter, obtained approval from Health Canada for Seasonique[®], a next generation extended-cycle oral contraceptive for the prevention of pregnancy

Paladin's revenues are principally derived from sales of pharmaceutical products to large pharmaceutical wholesalers and large chain pharmacies. The Company's expenses have been comprised primarily of cost of goods sold (including royalty payments to those companies from whom Paladin licenses its products), selling, marketing and administrative expenses and research and development expenses. In addition, a substantial portion of the Company's expenses are related to amortization of the pharmaceutical product licenses and rights the Company acquires.

Paladin's annual and quarterly operating results are primarily affected by the level of acceptance of Paladin's products by physicians and their patients, and the timing and number of product launches. The level of patient and physician acceptance of Paladin's products, the acceptance of provincial government reimbursement on such products, market access, as well as the availability of similar therapies, impact Paladin's revenues by driving the level and timing of prescriptions for its products. Each new product launch requires significant promotional investment during the first three to five years from launch.

¹ EBITDA – Non-GAAP financial measures

The term EBITDA (earnings before interest, taxes, depreciation and amortization) does not have any standardized meaning under Canadian GAAP and therefore may not be comparable to similar measures presented by other companies. The Company defines EBITDA as earnings before interest expense, taxes, amortization, foreign exchange gains (losses) and unusual items; such as write-downs and gains (losses) on intellectual property and investments. EBITDA is calculated and presented consistently from period to period and agrees, on a consolidated basis, with the amount disclosed as "*Earnings before under-noted items*" on the consolidated statements of income. The Company believes EBITDA to be an important measurement that allows it to assess the operating performance of its ongoing business on a consistent basis without the impact of amortization expenses. The Company excludes amortization expenses because their level depends substantially on non-operating factors such as the historical cost of intangible and capital assets. The Company's method for calculating EBITDA may differ from that used by other issuers and, accordingly, this measure may not be comparable to EBITDA used by other issuers.

Critical Accounting Estimates

Paladin's consolidated financial statements are prepared in accordance with Canadian GAAP, applied on a consistent basis. Paladin's critical accounting estimates include revenue recognition, inventory valuation, the recording of research and development expenses and related tax credits, the useful lives and fair value of pharmaceutical product licenses and rights, stock-based compensation expense, income taxes and the determination of fair value of financial instruments. For a more detailed discussion of the Company's critical accounting estimates, please refer to the management's discussion & analysis included in the Company's 2009 Annual Report. There have been no material changes to accounting estimates since December 31, 2009.

New Accounting Standards and Disclosure Changes

Effective January 1, 2010, the Company adopted the following recently introduced Canadian Institute of Chartered Accountants ("CICA") Handbook Sections:

Section 1582, *Business Combinations*. This section establishes the standards for the accounting of business combinations, and states that all assets and liabilities of an acquired business will be recorded at fair value. Obligations for contingent considerations and contingencies will also be recorded at fair value at the acquisition date. The standard also states that acquisition-related costs will be expensed as incurred and that restructuring charges will be expensed in the periods after the acquisition date. The initial application of this standard did not have a significant impact on the Company's consolidated financial statements.

Section 1601, *Consolidated Financial Statements*. This section establishes the standards for preparing consolidated financial statements. The initial application of this standard did not have a significant impact on the Company's consolidated financial statements.

Section 1602, *Non-controlling Interests*, which establishes standards for the accounting of non-controlling interests of a subsidiary in the preparation of consolidated financial statements subsequent to a business combination. These changes did not have any impact on the Company's consolidated financial statements as the Company owns 100% of its current subsidiaries.

IFRS Changeover Plan

In February 2008 the Canadian Accounting Standards Board ("AcSB") confirmed that the use of International Financial Reporting Standards ("IFRS") would be required for Canadian publicly accountable enterprises for interim and annual financial statements effective for fiscal years beginning on or after January 1, 2011. The Company will implement these standards on January 1, 2011. The AcSB also stated that, during the transition period, companies will be required to provide comparative figures in accordance with IFRS. For the year ended December 31, 2010, the Company will be providing readers a reconciliation of current CICA GAAP to IFRS accounting standards. It is important to note that under IFRS there is significantly more disclosure required. Further, while IFRS uses a conceptual framework similar to Canadian GAAP, there are significant differences in accounting policy that must be addressed.

The Company's IFRS changeover plan is progressing according to expectations. The Company continues to monitor standards to be issued by the IASB but it is difficult to predict the IFRS that will be in effect at the end of the Company's first IFRS reporting period as the IASB work plan anticipates the completion of several projects in calendar years 2010 and 2011. The Company will provide updates as further progress is achieved and conclusions are reached.

IFRS 1 "First time adoption of international financial reporting standards" requires that first-time adopters select accounting policies that comply with each IFRS effective at the end of its first IFRS reporting period, December 31, 2011 for the Company, and apply those policies to all periods presented in its first IFRS financial statements. IFRS 1 also provides certain optional exemptions to the full retrospective application. The following are the Company's conclusions with respect to the key IFRS 1 optional exemptions:

- Business combinations – the Company will apply the new standard only to business combinations that have occurred after the transition date, without restatement of prior business combinations.
- Property, plant and equipment and pharmaceutical product licenses and rights – the Company will use the historical cost method for capital assets and pharmaceutical product licenses and rights.
- Share-based payments - the Company will not restate any of its existing stock options which have been fully vested before January 1, 2010.

Based on the Company's assessment of current IFRS, the Company has not identified any significant balance sheet items or any significant differences in the recording of the Company's revenues and expenses that would be materially impacted by the application of IFRS standards. Continued progress is necessary before we can increase the specificity of the disclosure of the impacts of IFRS.

Pursuant to the Canadian Securities Administrators Staff Notice 52-320, Disclosure of Expected Changes in Accounting Policies Relating to Changeover to IFRS, the Company is presenting below the progress towards the completion to our changeover plan.

	KEY ACTIVITIES	MILESTONES / DEADLINES	PROGRESS TO DATE
Accounting policies and financial statement preparation	Identify relevant differences between IFRS and the Company's accounting policies and practices and design and implement solutions; Evaluate and select one-time and ongoing accounting policy alternatives; Benchmark findings with peer companies; Prepare financial statements and related note disclosures to comply with IFRS; Quantify the effects of changeover to IFRS;	Assessment and quantification of the significant effects of the changeover to be completed by approximately the second quarter of 2010; Final selection of accounting policy alternatives by the changeover date;	Preliminary identification of IFRS differences completed by management and reviewed by third-party experts under way; Evaluation and selection of accounting policy alternatives is ongoing; Third-party experts are assisting in the transition; Analysis, subject to the impact of the proposed IFRS standards, is underway; Impact quantification is underway;
Information technology and data systems	Identify and address IFRS differences that require changes to financial systems; Evaluate and select methods to address need for dual record-keeping during 2010 (i.e. IFRS and Canadian GAAP) for comparatives, budget and planning purposes in 2011;	Changes to significant systems and dual record-keeping process completed in time for the first quarter of 2010;	No IFRS differences with significant system impacts have been identified to date Dual record-keeping solution design is underway;
Internal control over financial reporting (ICFR)	Revise existing internal control processes and procedures to address significant changes to existing accounting policies and practices, including the need for dual record-keeping in 2010; Design and implement internal controls with respect to one-time changeover adjustments and related communications;	Changes completed by the first quarter of 2010. Conduct management evaluation of new or revised controls throughout 2010; Update the Chief Executive Officer/Chief Financial Officer certification process by fourth quarter of 2010;	Monitoring design of solutions to address IFRS differences to permit concurrent design or revision and implementation of necessary internal controls;
Disclosure controls and procedures (DC&P)	For changes to accounting policies and practices identified, assess the DC&P design and effective implementation;	See ICFR deadlines above;	MD&A disclosures have begun;
Training and communication	Provide training to employees and management; Communicate progress of changeover plan to internal and external stakeholders;	Timely training provided to align with work under changeover – to be completed during 2010; Communicate effects of changeover for 2011 financial planning process, by the third quarter of 2010;	Selected training for employees directly engaged in the changeover and general awareness to broader group of finance employees; Periodic internal and external communications about our progress are ongoing;
Business activities	Identify impact of changeover on contractual arrangements, including customer and supplier agreements and employee compensation plans; Make any required changes to arrangements and plans;	Changes to be completed by third quarter of 2010;	No potential impact identified to date;

Results of Operations

Three-month period ended March 31, 2010 compared to three-month period ended March 31, 2009.

Revenues

Revenues increased \$5,022 or 19% to \$30,837 for the three-month period ended March 31, 2010 from \$25,815 for the three-month period ended March 31, 2009. Revenues from products acquired after the comparative quarter, March 31, 2009, contributed \$1,559 to the quarter ended March 31, 2010. Furthermore, during the quarter, an amount of \$1,668 (2009 - \$1,693) previously recorded in deferred revenue was recognized into revenue in accordance with the Company's revenue recognition policy.

The increase in revenues for the quarter ended March 31, 2010 is also attributable to the sales growth of certain significant promoted products, including Tridural[®], Trelstar[®], Testim[®], Metadol[®] and Plan B[®], which combined increased by 26% compared to the quarter ended March 31, 2009.

Product revenues highlights for the Company's most significant promoted products using IMS Canada data² for the quarter ended March 31, 2010 compared to the quarter ended March 31, 2009 are as follows:

Promoted Products	Three-month period ended March 31, 2010	
	Sales per IMS Canada	% change vs. 2009
	\$	
Tridural [®]	2,414	20%
Trelstar [®]	1,041	106%
Testim [®]	712	53%
Metadol [®]	2,217	17%
Plan B [®]	2,252	15%
Total	8,636	26%

Gross Profit

Total gross profit increased \$2,612 or 13% to \$22,057 for the three-month period ended March 31, 2010 from \$19,445 for the three-month period ended March 31, 2009. Gross profit, as a percentage of revenues, decreased 3% to 72% for the quarter ended March 31, 2010 from 75% from the same period last year. The decrease in gross profit as a percentage of revenues for the quarter ended March 31, 2010 is mainly the result of the effect of reduced margins on the Company's product mix as a result of growth in certain promoted products.

Selling and Marketing Expense

Selling and marketing expense increased \$276 or 5% to \$5,788 for the quarter ended March 31, 2010 from \$5,512 for the same comparative period last year. Selling and marketing expense, as percentage of revenues, decreased to 19% for the quarter ended March 31, 2010 compared to 21% for the same period last year. The decrease in the current quarter's selling and marketing expenses as a percentage of revenues is the result of certain streamlining efforts and the Company's growth in non-promoted product revenue during the first quarter of 2010, shifting selling and marketing expense as a percentage of revenues downward for the quarter. The promotional activities driving selling and marketing costs primarily relate to Paladin's continued promotional activities for Tridural[®], Plan B[®], Metadol[®], Twinject[®], Trelstar[®], and Testim[®].

General and Administrative Expense

General and administrative expense increased \$372 or 19% to \$2,287 for the three-month period ended March 31, 2010 from \$1,915 for the three-month period ended March 31, 2009. General and administrative expense, as percentage of revenues, remained steady at 7% for the quarters ended March 31, 2010 and 2009, respectively. The increase in the current quarter's general and administrative

² The Company has chosen not to disclose detailed product by product revenue information for competitive reasons, however, does include detailed IMS Canada sales data, essentially end-user pharmacy purchase volume data, to allow the reader to better understand revenue changes from period to period on certain significant products. It is important that readers of this sales data note that IMS Canada sales data may not necessarily correspond to the Company's recording of revenue in accordance with GAAP.

expenses amount is primarily the result of the Pharmaplan (Pty) Ltd. acquisition related costs expensed in accordance with recently adopted accounting standards.

Research and Development Expense

Research and development expense increased \$1,475 or 117% to \$2,737 for the three-month period ended March 31, 2010 from \$1,262 for the three-month period ended March 31, 2009. Research and development expense, as percentage of revenues, increased to 9% for the quarter ended March 31, 2010 compared to 5% for the quarter ended March 31, 2009. The increase in the current quarter's research and development expenses primarily relates to the recently initiated and on-going research and development efforts at IsoPharma, as further described in note 4 to the quarterly unaudited consolidated financial statements and incremental research and development expenses related to Paladin Biosciences advancements. In addition to the above, the research activities driving research and development expense include managing development projects with licensors and preparing new drug submissions to strengthen the Company's pipeline as well as to search and explore potential product opportunities for internal development.

Interest Income

Interest income increased \$102 or 51% to \$301 for the three-month period ended March 31, 2010 from \$199 for the three-month period ended March 31, 2009. This increase is primarily the result of the additional interest earned on the Company's investment in a portfolio company in the form of a convertible debenture acquired during the quarter ended March 31, 2010. Furthermore, the Company held higher average daily cash and marketable securities balances over the quarter ended March 31, 2009, mainly as a result of the Company's share offering, partially offset by a lower effective rate of return during the quarter ended March 31, 2010, compared to the same comparative period last year.

Amortization of Pharmaceutical Product Licenses, Rights

Amortization expense increased \$174 or 3% to \$6,270 for the three-month period ended March 31, 2010 from \$6,096 for the three-month period ended March 31, 2009. The increase in amortization expense is the result of the amortization related to the Company's recently acquired pharmaceutical product licenses and rights partially offset by a reduction in amortization expense for certain pharmaceutical product licenses and rights having reached full amortization during the quarter.

Unrealized Gain on Derivative Financial Instruments

In accordance with Section 3856, the Company used the Black-Scholes option pricing model to re-measure the fair value of a conversion option on a secured convertible term note investment in a portfolio company recognizing an unrealized gain on the conversion option in the amount of \$nil for the quarter ended March 31, 2010 [2009 - \$344] and an unrealized gain on warrants in a portfolio company in the amount of \$nil [2009 - \$15].

Net (Gain) Loss on Investments

During the three-month period ended March 31, 2010, the Company disposed of certain shares held in a portfolio company for proceeds of \$27, representing a gain of \$7.

During the three month period ended March 31, 2009, the Company disposed of certain shares held in a portfolio company for proceeds of \$568, representing a gain of \$62. In addition, during the quarter ended March 31, 2009, Endo Pharmaceuticals Inc. ("Endo") acquired Indevus Pharmaceuticals Inc. ("Indevus") for US\$4.50 per Indevus share in cash and up to an additional US\$3.00 per share in cash upon achievement of certain regulatory and sales milestones. The Company received proceeds in the amount of \$2,167 (US\$1,720) for the investment it held in 382,253 common shares of Indevus, resulting in a realized loss on disposal in the amount of \$414. Furthermore, the Company recorded \$416 representing the estimated fair value of a contingent right to cash receivable in relation to the achievement of certain regulatory and sales milestones, determined to represent the fair value upon receipt and measured based upon the average incremental Indevus common stock trading price over the US\$4.50 cash payment received over a 31 day trading period on NASDAQ.

Foreign Exchange Loss

During the three-month period ended March 31, 2010, the Company recorded a foreign exchange loss of \$320, compared to a loss of \$22 for the same period last year, mainly as a result of the strengthening of the Canadian dollar relative to the US dollar and Euro and its impact on Company's net monetary position in these currencies during the quarter.

Share of Net Income in Companies Subject to Significant Influence

On March 16, 2010, the Company entered into a strategic investment to acquire an initial 34.99% ownership interest in Pharmaplan (Pty) Ltd. ("Pharmaplan"), a privately-owned specialty pharmaceutical company based in Johannesburg, South Africa. The equity interest acquired in Pharmaplan represents an investment subject to significant influence which is accounted for using the equity method from the date of the transaction, March 16, 2010. The investment was initially recorded at cost and adjustments are made to include the Company's share of Pharmaplan's net income. The Company's share of net income is adjusted to reflect the amortization of the fair value adjustments related to the Company's acquired share of Pharmaplan's net identifiable assets. Paladin's share of Pharmaplan's net income for the 15 days from the acquisition date, March 16, 2010 and March 31, 2010, amounted to \$131 (net of \$70 amortization of fair value adjustments).

Income Tax Expense

Income tax expense increased \$257 or 14% to \$2,049 for the three-month period ended March 31, 2010 from \$1,792 for the three-month period ended March 31, 2009. For the quarter ended March 31, 2010, the effective tax rate was 40% compared to 37%, for the quarter ended March 31, 2009. The increase in effective tax rates in the current quarter is principally due to increases in permanent differences in comparison to the previous quarter, mainly capital losses not deductible for tax purposes. The Company has the following tax pools detailed below which may be applied against taxable income:

	Available \$	Recognized \$	Expires in
Non-capital tax losses			
Federal	65,255	30,166	2013-2028
Provincial	65,255	30,166	2013-2028
Scientific Research and Experimental Development expenditures			
Federal	83,455	63,524	N/A
Provincial	80,008	63,524	N/A
Investment tax credits			
Federal	23,208	15,699	2017-2030

The amount of the tax benefit claimed in the current and prior years, is subject to audit by the taxation authorities and could be reduced by a material amount in the future.

During the quarter ended March 31, 2010, in connection with the Company's previously disclosed tax contingency, the Company received notices of re-assessment from the Canada Revenue Agency ("CRA") and the Ontario Minister of Finance ("OMF") reversing their original position on the use of certain non-capital losses acquired as part of the Dimethaid Health Care Ltd. (subsequently renamed Squire Pharmaceuticals Inc. "Squire") acquisition from Nuvo Research Inc. ("Nuvo").

As previously disclosed, on various dates during fiscal 2008 and 2009 the Company had received notices of re-assessment from the CRA relating to the taxation years ending August 16, 2005, July 31, 2006, July 31, 2007, and December 31, 2008 and from the OMF for the taxation year ended August 16, 2005, containing adjustments relating to the use of certain non-capital losses. The notices of assessment and re-assessment, if they had stood as a result of the CRA's position, amounted to a total tax liability exposure to the federal and relevant provincial governments of approximately \$11,625 including interest and penalties. The Company filed a Notice of Objection through the CRA appeals process on October 23, 2008. Furthermore, the Company, under the terms of the Share Purchase Agreement ("SPA") for Squire with Nuvo holds indemnities with respect to the status of the Squire tax accounts and certain tax asset values the Company as well as all costs relating to reassessment including advisory fees, interest and penalties, as

applicable. In addition, Nuvo had issued additional security over the indemnity obligations by entitling the Company to the benefit of security over certain assets and product revenue streams of Nuvo and certain of its subsidiaries.

In connection with the appeals process, during the years ended December 31, 2009 and 2008, the Company had posted a deposit of \$3,752 to the CRA and \$500 to the OMF, representing up to one half of the tax and interest assessed. In addition, during the year ended December 31, 2009, the Company issued from its revolving unsecured credit facility, a bank guarantee in the amount of \$720 to the OMF. As a result of the Company's success in the appeal process, an amount of \$3,936 was received from the CRA on January 20, 2010 and an amount of \$504 was received from OMF on April 6, 2010, representing a refund for the full amount of the deposits above, along with accrued interest in the amount of \$188. In addition, the bank guarantee previously issued to the OMF expired February 1, 2010 without being drawn-down by the OMF.

Net Income

Due to the factors set forth above, net income essentially remained steady decreasing by \$7 or nil% to \$3,045 for the three-month period ended March 31, 2010 compared to net income of \$3,052 for the same period ended last year.

Liquidity and Capital Resources

The Company's Investment Policy regulates the investment activities relating to cash resources. An Investment Committee composed of representatives from management and the Board of Directors monitors compliance with said policy. The Company invests strictly in liquid, high-grade investment securities with varying terms to maturity, selected with regard to the expected timing of investments and expenditures for continuing operations and prevailing interest rates. As at March 31, 2010, there were no restrictions on the flow of these funds nor have any of these funds been committed in any way, except as set out in Note 7 to the unaudited interim consolidated financial statements.

The Company has a one-year \$5,000 revolving unsecured credit facility in place with one of the Company's bankers maturing on July 29, 2010. The credit facility may be used for general corporate purposes including financing acquisitions and has been drawn down in the form of security in the amount of \$2,000 mainly to enter into foreign currency derivative transactions in order to hedge certain foreign exchange positions.

The Company believes that its existing cash, cash equivalents and short-term marketable securities, as well as cash generated from operations, are sufficient to finance its current operations, working capital needs and future product acquisitions. At present, the Company is actively pursuing product acquisitions that may require the use of substantial capital resources. There are no present agreements or commitments with respect to any such acquisitions.

	Three-month period ended March 31	
	2010	2009
	\$	\$
Cash flows from operating activities	10,365	8,266
Cash flows from (used in) investing activities	4,359	(661)
Cash flows from financing activities	771	188
Foreign exchange rate change on cash and cash equivalents	(18)	(57)
Increase in cash position	15,495	7,793
Cash and cash equivalents, beginning of period	31,227	4,646
Cash and cash equivalents, end of period	46,722	12,439
Short and long-term marketable securities, end of period	43,519	20,052
Cash, cash equivalents and marketable securities, end of period	90,241	32,491

Paladin's cash, cash equivalents and marketable securities decreased \$15,128 to \$90,241 at March 31, 2010 from \$105,369 at December 31, 2009. This decrease is primarily a result of the Company's acquisition of investments in the amount of \$24,612 and by a partial repayment of the balance of sale

payable in the amount of \$1,650 partially offset by cash flows generated from operating activities in the amount of \$10,365 and common shares issued for cash in the amount of \$771. Working capital (current assets less current liabilities) decreased \$12,613 to \$97,843 at March 31, 2010 from \$110,456 at December 31, 2009 primarily due to the decrease in the cash, cash equivalents and marketable securities explained above.

Cash flows from operating activities increased 25% or \$2,099 to \$10,365 for the three-month period ended March 31, 2010 from \$8,266 for the three-month period ended March 31, 2009. Cash flows from operating activities represent the cash flows from net earnings, excluding revenues and expenses not affecting cash, principally amortization, future income taxes, stock based compensation expense, foreign exchange (gains) losses, gains (losses) on investments and derivative instruments and accreted interest.

Cash flows from investing activities were \$4,359 compared to cash flows used in investing activities of \$661 for the three-month period ended March 31, 2010 and 2009, respectively. During the three-month period ended March 31, 2010, the Company generated proceeds from maturing marketable securities net of the acquisition of short and long-term marketable securities in the amount of \$30,616 and proceeds from disposal of investments in the amount of \$27, partially offset by \$24,612 used for the acquisition of investments, \$1,650 towards a partial repayment of the balance of sale payable and \$22 for the acquisition of capital assets. During the three-month period ended March 31, 2009, the Company invested \$3,367 towards the acquisition of short-term and long-term marketable securities net of cash flows generated by maturing marketable securities, \$29 for the acquisition of capital assets, partially offset by proceeds from disposal of investments in the amount of \$2,735.

Cash flows from financing activities were \$771 compared to \$188 for the three-month periods ended March 31, 2010 and 2009, respectively, and represent proceeds from common stock options exercised and the issuance of common shares under the stock purchase plan for cash.

Acquisition of Isotechnika Inc.

On June 18, 2009, the Company acquired all the issued and outstanding shares of Isotechnika (TSX:ISA) in accordance with a court supervised Plan of Arrangement. As part of the transaction the Company paid \$7,594 in cash and Isotechnika entered into a collaborative research and development agreement with IsoPharma in exchange for supporting research and development services for the commercialization of voclosporin, Isotechnika's next-generation calcineurin inhibitor, in Canada, Mexico, Central & South America, Israel and South Africa ("Paladin-acquired territories"). The research and development services extend for a period of seven years and included an amount of \$4,350 payable by the Company to IsoPharma over 12 months following the acquisition and expensed as incurred over this same 12 month period. Furthermore, the Research and Development Agreement in conjunction with the Company's Licence Agreement for voclosporin in the Paladin acquired territories, contains certain other voclosporin research, development and commercialization payment arrangements including possible licensing and royalty revenue payments over the remaining period. During the quarter ended March 31, 2010, the Company has expensed \$988 with respect to these research and development services.

As part of the acquisition, the Company also received the international rights to a portfolio of products under development and a commercialized diagnostic product portfolio. The Company had also assumed an obligation to pay out certain future contractually pre-defined amounts relating to the commercialization of the diagnostic product portfolio over a period of seven years, estimated to amount to approximately \$5,950 at the time of acquisition (the "contingent balance of sale payable"). This estimated amount was determined based upon the historical sales of the products, current expenditure levels and market conditions and was payable over a seven-year period pursuant to its Research and Development Agreement and represented management's best estimate of this liability at the time of acquisition. It was considered reasonably possible that changes in future conditions could require a material change in the recognized amount. On December 31, 2009, the Company entered into an Amended Agreement with IsoPharma to settle the contingent balance of sale payable for a total amount of \$1,991, of which \$1,650 was paid on February 26, 2010 and the remainder in the amount of \$341, representing the discounted present value, is payable on January 31, 2011 if certain conditions are met. This settlement resulted in an adjustment to the extraordinary gain recorded upon acquisition in the amount of \$3,458 which has been recorded in accordance with related party accounting standards (see note 6 to the the interim unaudited consolidated financial statements for additional information). The Company had originally recorded an extraordinary

gain in the amount of \$25,959 upon the acquisition of Isotechnika which in conjunction with the settlement above results in an aggregate extraordinary gain of \$29,417 for the year ended December 31, 2009.

As at March 31, 2010, the Company is carrying a balance of sale payable to Isotechnika in the amount of \$343 and is committed to research and development services expected to be paid up to June 30, 2010 in the amount of \$988.

Isotechnika is an international biopharmaceutical company dedicated to the discovery, development and commercialization of novel immunosuppressive therapeutics for the treatment of autoimmune diseases and for use in the prevention of organ rejection in transplantation. In addition to the Company's drug pipeline, Isotechnika also has commercialized diagnostic products. Isotechnika was subsequently wound up into the Company on October 1, 2009. Furthermore, as part of the purchase price, the Company received 24,921,312 common shares, representing a 19 percent interest in IsoPharma as at the date of acquisition, with an approximate value of \$4,348 using the weighted average trading price of the common shares on the TSX for the 20 trading days pre and post acquisition (\$9,221 as at March 31, 2010).

The results of Isotechnika's operations have been included in the Company's results since June 18, 2009, the effective date of acquisition. The Company accounts for its investment in IsoPharma using the equity method of accounting. As per applicable accounting standards, the Company eliminated the value assigned to its investment in common shares of IsoPharma against the excess ("negative goodwill") of the amounts assigned to assets acquired and undiscounted liabilities assumed over the cost of the total purchase price. As a result the Company's investment in IsoPharma has a carrying value of \$nil effective June 18, 2009. Since the Company's acquisition, IsoPharma has incurred net losses from operations. As the Company is not committed to make further capital contributions to IsoPharma, the Company has not recorded its share of IsoPharma's net loss since acquisition as per applicable accounting standards. The Company refers the reader to note 4 of the interim unaudited consolidated financial statements, for further details regarding the acquisition.

Equity investment in Pharmaplan

On March 16, 2010, the Company entered into a strategic investment to acquire an initial 34.99% ownership interest in Pharmaplan (Pty) Ltd. ("Pharmaplan"), a privately-owned specialty pharmaceutical company based in Johannesburg, South Africa. The Company paid \$18,861 including a non-interest bearing loan in the amount of \$2,879. In addition, the Company has committed to additional future consideration by increasing its ownership position by 5% per year over the next 3 years to 49.99%, with such additional consideration based upon Pharmaplan's future financial results.

The equity interest acquired in Pharmaplan represents an investment subject to significant influence which is accounted for using the equity method from the date of the acquisition, March 16, 2010. The investment was initially recorded at cost and adjustments are made to include the Company's share of Pharmaplan's net income. The Company's share of net income is adjusted to reflect the amortization of the fair value adjustments related to the Company's share of net identifiable assets of Pharmaplan acquired.

The total cost was allocated to the Company's share of net identifiable assets acquired on the basis of their fair values using the purchase method of accounting. The purchase price allocations are preliminary and are subject to changes once the final valuation of the net identifiable assets acquired has been made. The preliminary allocation of the cost of the investment in Pharmaplan over the underlying net book value of assets acquired amounted to \$13,496 as at March 16, 2010, and represents intangible assets (consisting mainly of exclusive distribution licenses), future income tax liabilities and incremental goodwill.

The Company is in the process of finalizing the purchase price allocation which will be completed during 2010.

	\$
Investment in Pharmaplan, March 16, 2010	15,982
Share of net income for the period	201
Amortization of fair value adjustments	(70)
Investment in Pharmaplan, March 31, 2010	16,113

The Company is presenting selected financial information derived from Pharmaplan's unaudited financial statements using South African GAAP translated into Canadian GAAP for information purposes.

Pharmaplan's statement of income data (unaudited)	15 days ended March 31, 2010	
	\$	
Revenues	2,509	
Cost of sales	1,146	
Gross profit	1,363	
Operating expenses	562	
Earnings before under-noted items	801	
Interest, depreciation and income taxes	226	
Net income	575	

Pharmaplan's balance sheet data (unaudited)	March 31, 2010	March 16, 2010
	\$	\$
Total assets	15,299	16,117
Total liabilities	7,648	9,009

Related Party Transactions

Joddes Limited ["Joddes"], a private Canadian corporation, is a significant shareholder holding approximately 37% of the outstanding shares of the Company, and one director of the Company, the Company's President and CEO, is related to Joddes.

The Company engages a wholly-owned subsidiary of Joddes to provide logistics services including: customer service, warehousing, shipping, invoicing, collection services and certain manufacturing and selling services on behalf of the Company. The Company also engages this affiliate to perform certain research and development and selling services on a contractual pay-for-use basis. The Company also leases its office facilities from another wholly owned subsidiary of Joddes. This lease is for a period of 10 years, ending in 2013 and includes minimum annual payments for a total remaining committed amount of \$663 as at March 31, 2010 and is included in the purchase and service based commitments amount in Contractual Obligations and Commitments below.

The Company has also entered into contractual royalty agreements with a wholly-owned subsidiary of Joddes for certain legacy and over-the-counter products. The terms of these arrangements vary whereby the Company may earn a royalty fee based on certain established terms relating to the net sales of the respective products such as through a percentage of net sales, certain guaranteed minimum annual payments, or as a percentage of a defined product contribution.

Effective November 1, 2006, the Company acquired the Canadian distribution rights to Metadol[®] from a wholly-owned subsidiary of Joddes for cash consideration of \$15,000. Under the terms of the agreement, the Company can purchase the Canadian license for Metadol[®] on the fourth anniversary of the agreement for \$1 and can receive a reimbursement of up to \$3,750 subject to certain acquisition related conditions. As at March 31, 2010, the Company has not received or earned any reimbursement. The acquisition of the Canadian distribution rights to Metadol[®] was not in the normal course of operations and was recorded at an agreed upon exchange amount in accordance with the requirements of accounting standard CICA 3840.

The Company owns a 19 percent interest in the common shares of IsoPharma and considers this investment a related party. The Company, in conjunction with the business combination discussed in note 4 to the interim unaudited consolidated financial statements, settled with IsoPharma on December 31, 2009 the balance of sale contingently payable over a seven-year period for an amount of \$1,991, of which \$1,650 was paid on February 26, 2010 and \$341, representing the discounted present value, is payable on January 31, 2011 if certain conditions are met. This transaction was not in the normal course of operations and was recorded at an agreed upon exchange amount in accordance with the requirements of accounting standard CICA 3840. Furthermore, the Company is committed to research and development services

payments and a milestone payment with IsoPharma, as further discussed in note 4, in the amount of \$1,388 as at March 31, 2010 that are included in the contractual obligations and commitments paragraph below. All transactions with related parties, except for the Metadol[®] and IsoPharma transactions described above, are carried out in the normal course of operations, and are recorded at an agreed upon exchange amount. The accounts payable to related parties are on normal commercial terms and conditions and are non-interest bearing.

The table below reflects all transactions and services with related parties carried in the normal course of operations, which include those referred to in the agreements described above, as well as revenues from a wholly-owned subsidiary of Joddes:

	Three-month period ended	
	March 31	
	2010	2009
	\$	\$
Revenues	1,064	889
Purchases	4,162	1,570
Sales and marketing	1,539	1,239
Research and development	1,131	133
General and administrative	162	121

Quarterly Information (unaudited)

(In thousands of Canadian dollars except per share information)

	Q1	Q4	Q3	Q2	Q1	Q4	Q3	Q2
	F2010	F2009	F2009	F2009	F2009	F2008	F2008	F2008
Revenues	30,837	29,279	28,374	26,255	25,815	23,051	22,191	20,668
EBITDA ¹	11,546	9,661	10,161	8,410	10,955	7,415	9,189	6,997
Earnings before income taxes	5,094	2,501	4,212	3,051	4,844	3,512	5,975	4,067
Net Income before extraordinary gain	3,045	934	2,564	1,771	3,052	2,044	3,617	2,588
Net Income	3,045	4,392	2,564	27,730	3,052	6,116	3,617	2,588
Earnings per share before extraordinary gain	\$0.16	\$0.05	\$0.14	\$0.11	\$0.20	\$0.14	\$0.24	\$0.17
Earnings per share	\$0.16	\$0.24	\$0.13	\$1.77	\$0.20	\$0.41	\$0.24	\$0.17
Diluted earnings per share before extraordinary gain	\$0.16	\$0.05	\$0.14	\$0.11	\$0.20	\$0.14	\$0.24	\$0.17
Diluted earnings per share	\$0.16	\$0.23	\$0.13	\$1.71	\$0.20	\$0.41	\$0.24	\$0.17

Paladin's annual and quarterly operating results are primarily affected by the level of acceptance of Paladin's products by physicians and their patients, and the timing and number of product launches. The level of patient and physician acceptance of Paladin's products, the acceptance of provincial government reimbursement on such products, market access, as well as the availability of similar therapies, impact Paladin's revenues by driving the level and timing of prescriptions for its products. Each new product launch requires significant promotional investment during the first three to five years from launch.

Off-Balance Sheet Arrangements

The Company's off balance sheet arrangements consist of contractual obligations and agreements for development, sales, marketing and distribution rights to innovative drug products for the Canadian market. The effect of terminating these arrangements under normal operating circumstances consists of an effective transition of the remaining responsibilities and obligations to the licensor under agreed upon time frames and conditions. Please refer to this section below or note 7 of the Company's quarterly unaudited consolidated financial statements for additional details. Other than these contractual obligations and commitments, the Company does not have any off-balance sheet arrangements that have, or are reasonably likely to have, a current or future effect on the Company's financial condition, changes in revenues or

expenses, results of operations, liquidity, capital expenditures, or capital resources that are material to investors.

The Company does not issue guarantees contemplated by the applicable CICA Guidelines.

Concentration of Credit Risk and Major Customers

The Company's cash and cash equivalents, short-term and long-term investments are held through various institutions. Marketable securities are mainly investments in liquid, high-grade investment securities. They are subject to minimal risk of changes in value and generally have an original maturity from three months to twenty-four months from the date of purchase. Marketable securities are substantially all invested with large Canadian financial institutions, with a minor proportion of securities invested in one large U.S. financial institution.

The Company is exposed to credit risk from its customers and continually monitors its customers' credit. It establishes the provision for doubtful accounts based upon the credit risk applicable to each customer. For the three-month period ended March 31, 2010, two customers, a major wholesale distributor and a major retail chain, represented 29% and 16% of revenues, respectively [2009 – 28% and 16%]. As at March 31, 2010, two customers, a major wholesale distributor and a major retail chain, represented 14% and 18% of trade accounts receivable, respectively [2009 – 32% and 13%]. For a more detailed analysis and disclosure of credit risk please refer to note 8 to the quarterly unaudited consolidated financial statements.

Liquidity risk

The Company generates sufficient cash from operating activities to fund its operations and fulfill its obligations as they become due. The Company has sufficient funds available through its cash and marketable securities, should its cash requirements exceed cash generated from operations to cover all financial liability obligations. As at March 31, 2010, there were no restrictions on the flow of these funds nor have any of these funds been committed in any way, except as set out in the "Contractual Obligations and Commitments" section below. All financial liabilities are short term in nature except for the long-term portion of the balance of sale payable, which is payable to the extent of future product sales.

Foreign exchange risk

The Company principally operates within Canada, however, a portion of the Company's revenues, expenses, and current assets and liabilities, are denominated in United States dollars ("USD"), EURO and South African Rand ("ZAR"). This results in financial risk due to fluctuations in the value of the USD, EURO and ZAR relative to the Canadian dollar ("CAD"). The Company has significant monetary assets denominated in EURO and ZAR that are required to be revalued in CAD at each period end. As of March 31, 2010, the Company has entered into a forward foreign exchange contract to cover the foreign exchange exposure in a certain investment denominated in EURO. With the exception of the forward contract described above, the Company does not currently use derivative financial instruments to reduce its foreign exchange exposure and often relies on natural hedges to mitigate foreign currency risk. Based on the net exposure described in note 8 to the unaudited interim consolidated financial statements, and assuming that all other variables remain constant, a ten-point increase or decrease in the CAD/USD, CAD/EURO and CAD/ZAR exchange rates would have an effect of \$291 on net earnings.

Interest rate risk

The Company is subject to interest rate risk on its cash and marketable securities. The Company does not believe that the results of operations or cash flows would be materially affected to any significant degree by a sudden change in market interest rates relative to interest rates on the investments, owing to the relative short-term nature of the marketable securities.

Risk Factors

For a more detailed discussion of the risk factors that could materially affect the results of operations and the financial condition of the Company, please refer to the Company's Annual Information Form.

Internal Control Over Financial Reporting

No changes were made in our internal control over financial reporting during the quarter ended March 31, 2010 that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

Contractual Obligations and Commitments

In the normal course of business, Paladin secures development, sales, marketing and distribution rights to innovative drug products and has entered into various agreements, which include contractual obligations extending beyond the current year. The Company is committed to making minimum purchases of inventory, and minimum expenditures for regulatory, selling and marketing services in the amount of \$28,709, including €4,062, to retain exclusive distribution agreements for certain products. The Company, as further discussed in note 5 to the unaudited interim consolidated financial statements, is also committed to purchase an additional 15% of Pharmaplan's common shares evenly over 3 years, currently estimated to amount to ZAR62,816 and subject to change based upon Pharmaplan's future operating results. These commitments end in 2015 and annual commitments are as follows:

Contractual Obligations	Total	Less than 1 year	1-3 years	4-5 years	After 5 years
Purchase and service based commitments	28,709	10,618	16,159	1,933	—

In addition, under certain agreements, Paladin may have to pay additional consideration should the Company achieve certain sales volumes or if certain milestones are met, such as regulatory approval in Canada. The Company may have to pay up to \$16,241 including US\$11,561, €975 and GBP£500 over a maximum period of 15 years if it achieves certain product, regulatory or sales milestones on specific products in the future. The Company has the following commitments related to product license, trademark and distribution agreements:

Commitments	Total	Less than 1 year	1-3 years	4-5 years	After 5 years
Milestone based commitments	9,892	2,317	3,717	508	3,350
Revenues based commitments	6,349	508	1,524	508	3,809

**NOTICE TO READER OF THE INTERIM
CONSOLIDATED FINANCIAL STATEMENTS**

The consolidated financial statements of Paladin Labs Inc. (the “**Company**”) and the accompanying interim consolidated balance sheet as at March 31, 2010 and the interim consolidated statements of income, cash flows, comprehensive income, accumulated other comprehensive income (loss) and retained earnings for the three-month period then ended are the responsibility of the Company’s management. These consolidated financial statements have not been audited or reviewed on behalf of the shareholders by the independent external auditors, Ernst & Young LLP.

The interim consolidated financial statements have been prepared by management and include the selection of appropriate accounting principles, judgments and estimates necessary to prepare these financial statements in accordance with Canadian generally accepted accounting principles. Management has determined such amounts on a reasonable basis in order to ensure that the consolidated financial statements are presented fairly in all material respects. The Company’s accounting procedures and related systems of internal controls are designed to provide a reasonable assurance that its assets are safeguarded and its financial records are reliable. Readers are cautioned that these interim consolidated statements may not be appropriate for their purposes.

(signed) Jonathan Ross Goodman

Jonathan Ross Goodman, B.A., LL.B, M.B.A.
President and Chief Executive Officer

Montreal, Canada
May 7, 2010

(signed) Samira Sakhia

Samira Sakhia C.A., M.B.A.
Chief Financial Officer

Montreal, Canada
May 7, 2010

CONSOLIDATED BALANCE SHEETS

[In thousands of Canadian dollars]

	March 31 2010 \$ (unaudited)	December 31 2009 \$ (audited ³)
ASSETS		
Current		
Cash and cash equivalents	46,722	31,227
Marketable securities	39,185	73,274
Accounts receivable <i>[note 8]</i>	12,075	14,167
Inventories	13,664	12,361
Other current assets	5,378	2,668
Income taxes receivable <i>[note 10]</i>	690	4,630
Investment tax credits recoverable	776	776
Future income tax assets	6,751	6,196
Total current assets	125,241	145,299
Long-term marketable securities	4,334	868
Property, plant and equipment	493	691
Pharmaceutical product licences and rights	36,277	42,543
Investments <i>[note 5]</i>	21,701	62
Investments tax credits recoverable	14,923	14,903
Future income tax assets	28,826	31,029
Total assets	231,795	235,395
LIABILITIES AND SHAREHOLDERS' EQUITY		
Current		
Accounts payable and accrued liabilities	18,238	22,934
Accounts payable to related parties	962	1,122
Deferred revenue	93	1,776
Income taxes payable	7,512	7,109
Balance of sale payable	343	1,650
Future income tax liabilities	250	252
Total current liabilities	27,398	34,843
Long-term		
Balance of sale payable	1,402	1,743
Future income tax liabilities	4,013	4,007
Total liabilities	32,813	40,593
Shareholders' equity		
Capital stock <i>[note 3]</i>	120,828	119,652
Other paid-in capital	4,364	4,408
Accumulated other comprehensive income	101	98
Retained earnings	73,689	70,644
Total shareholders' equity	198,982	194,802
Total liabilities and shareholders' equity	231,795	235,395

See accompanying notes

³ Derived from the audited annual financial statements filed on SEDAR at www.sedar.com

CONSOLIDATED STATEMENTS OF INCOME

[In thousands of Canadian dollars except for share and per share amounts]
[unaudited]

	Three-month period ended March 31	
	2010	2009
	\$	\$
Revenues	30,837	25,815
Cost of sales	8,780	6,370
Gross profit	22,057	19,445
Expenses (income)		
Selling and marketing	5,788	5,512
General and administrative	2,287	1,915
Research and development	2,737	1,262
Interest income	(301)	(199)
Earnings before under-noted items	11,546	10,955
Amortization of pharmaceutical product licenses and rights	6,270	6,096
Unrealized gain on derivative financial instruments	—	(359)
Net (gain) loss on of investments	(7)	352
Foreign exchange loss	320	22
Share of net income in companies subject to significant influence <i>[note 5]</i>	(131)	—
Income before income taxes	5,094	4,844
Provision for income taxes		
Current	422	194
Future	1,627	1,598
	2,049	1,792
Net income for the period	3,045	3,052
Earnings per share		
Basic	0.16	0.20
Diluted	0.16	0.20
Weighted average number of shares outstanding		
Basic	18,595,616	14,929,672
Diluted	19,160,825	15,212,910

See accompanying notes

CONSOLIDATED STATEMENTS OF CASH FLOWS

[In thousands of Canadian dollars]

[unaudited]

	Three-month period ended	
	2010	March 31 2009
	\$	\$
Operating activities		
Net income	3,045	3,052
Add items not affecting cash		
Amortization	6,485	6,148
Future income taxes	1,625	1,598
Stock based compensation expense <i>[note 3]</i>	353	402
Unrealized foreign exchange loss	338	57
Unrealized gain on derivative financial instruments	—	(359)
Net accreted interest income	(18)	(25)
Net (gain) loss on investments	(7)	352
Share of net income in companies subject to significant influence <i>[note 5]</i>	(131)	—
	11,690	11,225
Net change in non-cash balances relating to operations	(1,307)	(2,902)
Cash flows from operating activities	10,383	8,323
Investing activities		
Acquisition of investments	(24,612)	—
Repayment of balance of sale payable	(1,650)	—
Acquisition of property, plant and equipment	(22)	(29)
Purchases of short-term marketable securities	(39,415)	(10,362)
Maturities of marketable securities	74,404	8,964
Purchases of long-term marketable securities	(4,373)	(1,969)
Proceeds from disposal of investments	27	2,735
Cash flows from (used in) investing activities	4,359	(661)
Financing activities		
Common shares issued for cash	771	188
Cash flows from financing activities	771	188
Foreign exchange rate change on cash and cash equivalents	(18)	(57)
Net change in cash and cash equivalents during the period	15,495	7,793
Cash and cash equivalents, beginning of period	31,227	4,646
Cash and cash equivalents, end of period	46,722	12,439
Cash and cash equivalents	46,722	12,439
Short-term marketable securities	39,185	18,093
Long-term marketable securities	4,334	1,959
	90,241	32,439

See accompanying notes

CONSOLIDATED STATEMENTS OF COMPREHENSIVE INCOME, ACCUMULATED OTHER COMPREHENSIVE INCOME (LOSS), AND RETAINED EARNINGS

[In thousands of Canadian dollars except for share and per share amounts]
[unaudited]

	Three-month period ended March 31	
	2010	2009
	\$	\$
Net income for the period	3,045	3,052
Other comprehensive income (loss):		
Change in fair value of available-for-sale financial instruments [net of (\$2) taxes [2009 – \$29]]	13	185
Reclassification adjustment for losses (gains) on available-for-sale financial instruments included in net income in the current period [net of \$1 taxes [2009 – (\$219)]]	(10)	1,399
Other comprehensive income for the period	3	1,584
Comprehensive income for the period	3,048	4,636
Accumulated other comprehensive income (loss), beginning of period	98	(1,420)
Other comprehensive income for the period	3	1,584
Accumulated other comprehensive income, end of period	101	164
Retained earnings, beginning of period	70,644	32,949
Net income for the period	3,045	3,052
Retained earnings, end of period	73,689	36,001
Retained earnings and accumulated other comprehensive income, end of period	73,790	36,165

See accompanying notes

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

[In thousands of Canadian dollars except for share and per share amounts]

1. Governing Statute and Nature of Operations

Paladin Labs Inc. is a specialty pharmaceutical public company continued under the Canada Business Corporations Act, focusing on developing, acquiring, marketing and distributing innovative pharmaceutical products. Paladin Labs Inc., together with its subsidiaries, is hereinafter referred to as the “Company”.

2. Basis of Presentation and Accounting policies

The unaudited interim consolidated financial statements have been prepared by the Company in accordance with Canadian generally accepted accounting principles [“GAAP”] applicable to interim financial statements and include the accounts of all its subsidiaries. The Company’s 35% ownership in Pharmaplan Ltd. [“Pharmaplan”] is accounted for using the equity method as the Company exercises significant influence. These financial statements do not include all the information and disclosures required according to GAAP for annual financial statements and should be read in conjunction with the Company’s audited consolidated financial statements and notes thereto in the Company’s Annual Report for the year ended December 31, 2009.

Information with respect to the December 31, 2009 balance sheet is derived from the Company’s complete audited consolidated financial statements. The accounting policies underlying these unaudited interim financial statements are those set forth in note 2 of the audited consolidated financial statements for the year ended December 31, 2009.

Changes in accounting policies

Effective January 1, 2010, the Company adopted the following recently introduced Canadian Institute of Chartered Accountants [“CICA”] Handbook Sections:

Section 1582, Business Combinations. This section establishes the standards for the accounting of business combinations, and states that all assets and liabilities of an acquired business will be recorded at fair value. Obligations for contingent considerations and contingencies will also be recorded at fair value at the acquisition date. The standard also states that acquisition-related costs will be expensed as incurred and that restructuring charges will be expensed in the periods after the acquisition date. The initial application of this standard did not have a significant impact on the Company’s consolidated financial statements.

Section 1601, Consolidated Financial Statements. This section establishes the standards for preparing consolidated financial statements. The initial application of this standard did not have a significant impact on the Company’s consolidated financial statements.

Section 1602, Non-controlling Interests, which establishes standards for the accounting of non-controlling interests of a subsidiary in the preparation of consolidated financial statements subsequent to a business combination. These changes did not have any impact on the Company’s consolidated financial statements as the Company owns 100% of its current subsidiaries.

3. Capital Stock

Authorized

100,000,000 common shares without nominal or par value.

Issued and outstanding

	Number of shares	Amount
Balance at beginning of year	18,563,250	\$119,652
Issued upon exercise of stock options	72,411	1,107
Issued under employee share purchase plan	3,430	69
Balance at March 31, 2010	18,639,091	\$120,828

Stock option plan

The changes to the number of stock options granted by the Company and their weighted average exercise price are as follows:

	2010		2009	
	#	Weighted average exercise price \$	#	Weighted average exercise price \$
Balance at beginning of year	1,246,518	10.65	1,067,948	9.03
Granted	330,201	19.67	342,750	12.19
Exercised	(72,411)	9.80	(15,453)	9.12
Expired or forfeited	(50,329)	13.72	(1,925)	10.58
Balance at March 31	1,453,979	12.63	1,393,320	9.81
Options exercisable at March 31	541,272	8.95	438,661	6.86

The Company recorded stock option compensation expense with a corresponding credit to other paid-in-capital and determined the fair value of stock options under the Black-Scholes option-pricing model using the following assumptions:

	Three-month period ended March 31	
	2010	2009
Stock-based compensation expense	\$353	\$402
Weighted average fair value of options	\$6.21	\$5.74
Weighted average risk-free interest rate	2.29%	2.44%
Dividend yield	Nil	Nil
Weighted average volatility factor	34%	42%
Weighted average expected life	4 years	7 years

4. Business acquisition

On June 18, 2009, the Company acquired all the issued and outstanding shares of Isotechnika Inc. ["Isotechnika"] [TSX:ISA] in accordance with a court supervised Plan of Arrangement. As part of the transaction the Company paid \$7,594 in cash and Isotechnika entered into a collaborative research and development agreement with Isotechnika Pharma Inc. ["IsoPharma"] in exchange for supporting research and development services for the commercialization of voclosporin, Isotechnika's next-generation calcineurin inhibitor, in Canada, Mexico, Central & South America, Israel and South Africa ["Paladin-acquired territories"]. The research and development services extend for a period of seven years and included an amount of \$4,350 payable by the Company to IsoPharma over the 12 months following the acquisition and expensed as incurred over this same 12 month period. Furthermore, the Research and Development Agreement in conjunction with the Company's Licence Agreement for voclosporin in the Paladin acquired territories, contains certain other voclosporin research, development and commercialization payment arrangements including possible licensing and royalty revenue payments over the remaining period. During the quarter ended March 31, 2010, the Company has expensed \$988 with respect to these research and development services.

As part of the acquisition, the Company also received the international rights to a portfolio of products under development and a commercialized diagnostic product portfolio. The Company had also assumed an obligation to pay out certain future contractually pre-defined amounts relating to the commercialization of the diagnostic product portfolio over a period of seven years, estimated to amount to approximately \$5,950 at the time of acquisition [the "contingent balance of sale payable"]. This estimated amount was determined based upon the historical sales of the products, current expenditure levels and market conditions and was payable over a seven-year period pursuant to its Research and Development Agreement and represented management's best estimate of this liability at the time of acquisition. It was considered reasonably possible that changes in future conditions could require a material change in the recognized amount. On December 31, 2009, the Company entered into an Amended Agreement with IsoPharma to settle the contingent balance of sale payable for a total amount of \$1,991, of which \$1,650 was paid on February 26, 2010 and the remainder in the amount of \$341, representing the discounted present value, is payable on January 31, 2011 if certain conditions are met. This settlement resulted in an adjustment to the extraordinary gain recorded upon acquisition in the amount of \$3,458 which has been recorded in accordance with related party accounting standards [see note 6 for additional information]. The Company had originally recorded an extraordinary gain in the amount of \$25,959 upon the acquisition of Isotechnika which in conjunction with the settlement above results in an aggregate extraordinary gain of \$29,417 for the year ended December 31, 2009.

As at March 31, 2010, the Company is carrying a balance of sale payable to Isotechnika in the amount of \$343 and is committed to research and development services expected to be paid up to June 30, 2010 in the amount of \$988.

Isotechnika is an international biopharmaceutical company dedicated to the discovery, development and commercialization of novel immunosuppressive therapeutics for the treatment of autoimmune diseases and for use in the prevention of organ rejection in transplantation. In addition to the Company's drug pipeline, Isotechnika also has commercialized diagnostic products. Isotechnika was subsequently wound up into the Company on October 1, 2009.

4. Business acquisition (cont'd)

Furthermore, as part of the purchase price, the Company received 24,921,312 common shares, representing a 19 percent interest in IsoPharma as at the date of acquisition, with an approximate value of \$4,348 using the weighted average trading price of the common shares on the TSX for the 20 trading days pre and post acquisition. In connection with the acquisition, the Company had incurred transaction costs in the amount of \$530 included in the purchase price below.

The acquisition was accounted for using the purchase method. The results of Isotechnika's operations have been included in the Company's results since June 18, 2009, the effective date of acquisition. The total purchase price of \$10,689 was allocated to the fair value of the net assets acquired in the amount of \$47,204, representing negative goodwill in the amount of the excess of \$36,515. The Company, as per applicable accounting standards, eliminated the value previously assigned to certain prescribed assets in the amount of \$7,098 against the excess of the amounts assigned to assets acquired and undiscounted liabilities assumed over the cost of the purchase above. The remaining excess is presented as an extraordinary gain in the amount of \$29,417. The purchase price was preliminarily allocated as follows:

Purchase price allocation	\$
Cash	1,565
Current assets	627
Future income tax assets	39,440
Current liabilities	(1,526)
	<u>40,106</u>
 Consideration represented by:	
Cash paid	7,594
Balance of sale payable	2,565
Acquisition costs	530
	<u>10,689</u>
 Extraordinary gain [net of \$nil taxes]	 <u>29,417</u>

The Company is in the process of finalizing the purchase price allocation which will be completed during 2010.

5. Investments

	March 31, 2010	December 31, 2009
	\$	\$
Companies subject to significant influence	16,113	—
Portfolio investments	5,588	62
Total investments	<u>21,701</u>	<u>62</u>

5. Investments (cont'd)

Investment in Pharmaplan

On March 16, 2010, the Company entered into a strategic investment to acquire an initial 34.99% ownership interest in Pharmaplan (Pty) Ltd. ["Pharmaplan"], a privately-owned specialty pharmaceutical company based in Johannesburg, South Africa. The Company paid \$18,861 including a non-interest bearing loan in the amount of \$2,879. In addition, the Company has committed to additional future consideration by increasing its ownership position by 5% per year over the next 3 years to 49.99%, with such additional consideration based upon Pharmaplan's future financial results.

The equity interest acquired in Pharmaplan represents an investment subject to significant influence which is accounted for using the equity method from the date of the acquisition, March 16, 2010. The investment was initially recorded at cost and adjustments are made to include the Company's share of Pharmaplan's net income. The Company's share of net income is adjusted to reflect the amortization of the fair value adjustments related to the Company's share of net identifiable assets of Pharmaplan acquired.

The total cost was allocated to the Company's share of net identifiable assets acquired on the basis of their fair values using the purchase method of accounting. The purchase price allocations are preliminary and are subject to changes once the final valuation of the net identifiable assets acquired has been made. The preliminary allocation of the cost of the investment in Pharmaplan over the underlying net book value of assets acquired amounted to \$13,496 as at March 16, 2010, and represents intangible assets [consisting mainly of exclusive distribution licenses], future income tax liabilities and incremental goodwill.

The Company is in the process of finalizing the purchase price allocation which will be completed during 2010.

	\$
Investment in Pharmaplan, March 16, 2010	15,982
Share of net income for the period	201
Amortization of fair value adjustments	(70)
Investment in Pharmaplan, March 31, 2010	16,113

The Company is presenting selected financial information derived from Pharmaplan's unaudited financial statements using South African GAAP translated into Canadian GAAP for information purposes.

Pharmaplan's statement of income data (unaudited)	15 days ended March 31, 2010
	\$
Revenues	2,509
Cost of sales	1,146
Gross profit	1,363
Operating expenses	562
Earnings before under-noted items	801
Interest, depreciation and income taxes	226
Net income for the period	575

5. Investments (cont'd)

Pharmaplan's balance sheet data (unaudited)	March 31, 2010	March 16, 2010
	\$	\$
Total assets	15,299	16,117
Total liabilities	7,648	9,009

Investment in Isotechnika

On June 18, 2009, as part of the Isotechnika acquisition, the Company received 24,921,312 common shares, representing a 19 percent interest in IsoPharma, with an approximate value of \$4,348 using the weighted average trading price of the common shares on the TSX for the 20 trading days pre and post acquisition [\$9,221 as at March 31, 2010]. The Company accounts for its investment in IsoPharma using the equity method of accounting. As per applicable accounting standards, the Company eliminated the value assigned to its investment in common shares of IsoPharma against the excess ["negative goodwill"] of the amounts assigned to assets acquired and undiscounted liabilities assumed over the cost of the total purchase price. As a result the Company's investment in IsoPharma has a carrying value of \$nil effective June 18, 2009. Since the Company's acquisition, IsoPharma has incurred net losses from operations. As the Company is not committed to make further capital contributions to IsoPharma, the Company has not recorded its share of IsoPharma's net loss since acquisition as per applicable accounting standards. The Company refers the reader to note 4 for further details regarding the acquisition.

6. Related party transactions

Joddes Limited ["Joddes"], a private Canadian corporation, together with its affiliates own in aggregate approximately 37% of the outstanding shares of the Company, and one director of the Company, the Company's President and CEO, is related to this group.

The Company engages a wholly-owned subsidiary of Joddes to provide logistics services including: customer service, warehousing, shipping, invoicing, collection services and certain manufacturing and selling services on behalf of the Company. The Company also engages this affiliate to perform certain research and development and selling services on a contractual pay-for-use basis. In addition, the Company leases its office facilities from another wholly-owned subsidiary of Joddes. This lease is for a period of 10 years, ending in 2013 and includes minimum annual payments for a total remaining committed amount of \$663 as at March 31, 2010 and is included in the purchase and service based commitments in Note 7.

The Company has also entered into contractual royalty agreements with a wholly-owned subsidiary of Joddes for certain legacy and over-the-counter products. The terms of these arrangements vary whereby the Company may earn a royalty fee based on certain established terms relating to the net sales of the respective products such as through a percentage of net sales, certain guaranteed minimum annual payments, or as a percentage of a defined product contribution.

Effective November 1, 2006, the Company acquired the Canadian distribution rights to Metadol[®] from a wholly-owned subsidiary of Joddes for cash consideration of \$15,000. Under the terms of the agreement, the Company can purchase the Canadian license for Metadol[®] on the fourth anniversary of the agreement for \$1 and can receive a reimbursement of up to \$3,750 subject to certain acquisition related conditions. As at March 31, 2010, the Company has not received or earned any reimbursement. The acquisition of the Canadian distribution rights to Metadol[®] was

6. Related party transactions (cont'd)

not in the normal course of operations and was recorded at an agreed upon exchange amount in accordance with the requirements of accounting standard CICA 3840.

The Company owns a 19 percent interest in the common shares of IsoPharma and considers this investment a related party. The Company, in conjunction with the business combination discussed in note 4, settled with IsoPharma on December 31, 2009 the balance of sale contingently payable over a seven-year period for an amount of \$1,991, of which \$1,650 was paid on February 26, 2010 and \$341, representing the discounted present value, is payable on January 31, 2011 if certain conditions are met. This transaction was not in the normal course of operations and was recorded at an agreed upon exchange amount in accordance with the requirements of accounting standard CICA 3840. Furthermore, the Company is committed to research and development service payments and a milestone payment with IsoPharma, as further discussed in note 4, in the amount of \$1,388 as at March 31, 2010 that are included in the purchase and service based commitments in note 7.

All transactions with related parties, except for the Metadol[®] and IsoPharma transactions described above, are carried out in the normal course of operations, and are recorded at an agreed upon exchange amount. The accounts payable to related parties are on normal commercial terms and conditions and are non-interest bearing.

The table below reflects all transactions and services with related parties carried in the normal course of operations, which include those referred to in the agreements described above, as well as revenues from a wholly-owned subsidiary of Joddes:

	Three-month period ended March 31	
	2010	2009
	\$	\$
Revenues	1,064	889
Purchases	4,162	1,570
Selling and marketing	1,539	1,239
Research and development	1,131	133
General and administrative	162	121

7. Commitments

In the normal course of business, the Company secures development, sales, marketing and distribution rights to innovative drug products and has entered into various agreements which include contractual obligations extending beyond the current year. These obligations are classified into three major categories: revenue based, milestone based, and purchase and services based commitments.

Revenue based commitments

Most pharmaceutical product license agreements require that the Company make royalty payments ranging from 10% to 20% of sales, or generally require payments for products at rates ranging from 20% to 40% of the net selling price and in certain cases require revenue sharing at various rates over and above a pre-established net sales threshold.

7. Commitments (cont'd)

In addition, the Company may have to pay up to \$6,349 [US\$6,250] if it achieves specific sales volumes on specific products in the future, over a maximum of ten years.

Milestone based commitments

The Company has also committed to fund certain research and development expenditures of third parties upon milestone attainment in the amount of \$2,826, including €975, over the next five years. In addition, certain additional payments may be required under these agreements if milestones are met, such as regulatory approval in Canada. Based on the outcome of these milestones, the Company may have to pay up to \$7,066, including US\$5,311 and GBP£500, over a maximum period of 15 years.

Purchase and service based commitments

The Company is committed to making minimum purchases of inventory, and minimum expenditures for regulatory, selling and marketing services in the amount of \$28,709, including €4,062, to retain exclusive distribution agreements for certain products. The Company, as further discussed in note 5, is also committed to purchase an additional 15% of Pharmaplan's common shares evenly over 3 years, currently estimated to amount to ZAR62,816 and subject to change based upon Pharmaplan's future operating results. These commitments end in 2015 and annual commitments are as follows:

	\$
2010	10,618
2011	7,976
2012	4,109
2013	4,073
2014	1,003
2015	930

8. Financial Instruments

The Company's Investment Policy regulates the investment activities relating to cash resources. An Investment Committee composed of representatives from management and the Board of Directors monitors compliance with said policy. The Company invests in strategic investments in the form of equity or strictly in liquid, high-grade investment securities with varying terms to maturity, selected with regard to the expected timing of investments and expenditures for continuing operations and prevailing interest rates.

8. Financial Instruments (cont'd)

March 31, 2010	Available- for-sale	Loans and receivables	Other financial liabilities	Derivatives	Carrying value	Fair value
	\$	\$	\$	\$	\$	\$
Financial assets						
Cash and cash equivalents	46,722				46,722	46,722
Marketable securities	39,185				39,185	39,185
Accounts receivable		12,075			12,075	12,075
Other current assets		4,346			4,346	4,346
Long term marketable securities	4,334				4,334	4,334
Investments ⁴	52	4,960		576	5,588	5,588
Total financial assets	90,293	21,381	—	576	112,250	112,250
Financial liabilities						
Accounts payable and accrued liabilities			18,238		18,238	18,238
Accounts payable to related parties			962		962	962
Balance of sale payable			343		343	343
Long-term balance of sale payable			1,402		1,402	1,402
Total financial liabilities	—	—	20,945	—	20,945	20,945
December 31, 2009						
	Available- for-sale	Loans and receivables	Other financial liabilities	Derivatives	Carrying value	Fair value
	\$	\$	\$	\$	\$	\$
Financial assets						
Cash and cash equivalents	31,227				31,227	31,227
Marketable securities	73,274				73,274	73,274
Accounts receivable		14,167			14,167	14,167
Other current assets		901			901	901
Long-term marketable securities	868				868	868
Investments ⁴	62				62	4,299
Total financial assets	105,431	15,068	—	—	120,499	124,736
Financial liabilities						
Accounts payable and accrued liabilities			22,934		22,934	22,934
Accounts payable to related parties			1,122		1,122	1,122
Balance of sale payable			1,650		1,650	1,650
Long-term balance of sale payable			1,743		1,743	1,743
Total financial liabilities	—	—	27,449	—	27,449	27,449

⁴ In accordance with Section 3856, certain investments in private companies included in the balance above are carried at cost as there are no quoted market prices in an active market for such equity instruments. Fair value has not been disclosed because fair value cannot be measured reliably.

8. Financial Instruments (cont'd)

Fair Value

Fair value is the amount of consideration that would be agreed upon in an arm's length transaction between knowledgeable, willing parties who are under no compulsion to act. Fair values for marketable securities and investments classified "Available for sale" are obtained using quoted active market prices as such securities. In accordance with CICA Section 3856, investments in private companies are carried at cost unless evidence of an other than temporary impairment exists in which case they are written down to their recoverable amount.

The carrying amounts of cash and cash equivalents, accounts receivable, other current assets, accounts payable and accrued liabilities, accounts payable to related parties and the short-term portion of the balance of sale payable are a reasonable estimate of their fair values because of the short maturity of these instruments.

In accordance with Section 3856, certain investments in private companies included in the balance above are carried at cost as there are no quoted market prices in an active market for such equity instruments. Fair value has not been disclosed because fair value cannot be measured reliably.

The carrying values of all financial instruments approximate their fair values, except for the fair values for the bifurcated conversion option of a secured convertible term note in a portfolio company classified as a derivative and certain warrants in a portfolio company that were obtained using the residual amount method. The allocated loan portion described above classified as "Loans and receivables" and the long-term portion of the balance of sale payable classified as "Other financial liabilities" are being discounted using a 20% and 2.5% discount rates, respectively, such approximating market value.

These estimates are affected by assumptions the Company makes about the amount and timing of estimated future cash flows and discount rates, all of which reflect varying degrees of risk. Income taxes and other expenses that would be incurred on disposition of these financial instruments are not reflected in the fair values. As a result, the fair values are not necessarily the net amounts that would be realized if these instruments were settled.

Cash, cash equivalents, marketable securities and investments were measured using level 1 in the fair value hierarchy. The balance of sale payable and a secured convertible term note were measured using level 2 in the fair value hierarchy.

Risk arising from financial instruments

The Company does not use derivative financial instruments for speculative or trading purposes. Since the Company does not trade actively in derivative instruments it is not exposed to any significant liquidity risks relating to them.

Concentration of credit risk and major customers

The Company's cash and cash equivalents, short-term and long-term investments are held through various institutions. Marketable securities are mainly investments in liquid, high-grade investment securities. They are subject to minimal risk of changes in value and generally have an original maturity from three months to twenty-four months from the date of purchase. Marketable securities are substantially all invested with large Canadian financial institutions, with a minor proportion of securities invested in one large U.S. financial institution.

8. Financial Instruments (cont'd)

The Company is exposed to credit risk from our customers and continually monitors its customers' credit. It establishes the provision for doubtful accounts based upon the credit risk applicable to each customer. For the three-month period ended March 31, 2010, two customers, a major wholesale distributor and a major retail chain, represented 29% and 16% of revenues, respectively [2009 – 28% and 16%]. As at March 31, 2010, two customers, a major wholesale distributor and a major retail chain, represented 14% and 18% of trade accounts receivable, respectively [2009 – 32% and 13%].

The following table provides further details on the Company's accounts receivable balances:

	March 31, 2010	December 31, 2009
	\$	\$
Accounts receivable	16,929	19,188
Allowance for product returns	(4,751)	(4,918)
Allowance for doubtful accounts	(103)	(103)
Total accounts receivable	12,075	14,167

The following table provides the change in the allowance for doubtful accounts and product returns for trade accounts receivable:

	2010	2009
	\$	\$
Balance, beginning of the year	5,021	3,980
Change in provision for doubtful accounts	—	(27)
Change in provision for product returns	(167)	37
Balance at March 31	4,854	3,990

The following table provides further details on trade accounts receivable past due but not provisioned:

	March 31, 2010	December 31, 2009
	\$	\$
Trade accounts receivable not passed due	12,641	13,212
Trade accounts receivable passed due and not provisioned		
Under 30 days	3,719	4,402
31 to 60 days	466	1,269
Over 60 days	—	202
Allowance for product returns	(4,751)	(4,918)
Total accounts receivable, net of allowance for doubtful accounts and product returns	12,075	14,167

Liquidity risk

The Company generates sufficient cash from operating activities to fund its operations and fulfill its obligations as they become due. The Company has sufficient funds available through its cash, cash equivalents and marketable securities to cover all financial liability obligations should its cash requirements exceed cash generated from operations. As at March 31, 2010, there were no restrictions on the flow of these funds nor have any of these funds been committed in any way, except as set out in note 7.

8. Financial Instruments (cont'd)

All financial liabilities are short term in nature except for the long-term portion of the balance of sale payable, which is payable to the extent of future product sales.

Foreign exchange risk

The Company principally operates within Canada, however, a portion of the Company's revenues, expenses, and current assets and liabilities, are denominated in United States dollars ["USD"], EURO and South African Rand ["ZAR"]. This results in financial risk due to fluctuations in the value of the USD, EURO and ZAR relative to the Canadian dollar ["CAD"]. The Company has significant monetary assets denominated in EURO and ZAR that are required to be revalued in CAD at each period end. As of March 31, 2010, the Company has entered into a forward foreign exchange contract to cover the foreign exchange exposure in a certain investment denominated in EURO. With the exception of the forward contract described above, the Company does not currently use derivative financial instruments to reduce its foreign exchange exposure and often relies on natural hedges to mitigate foreign currency risk. Fluctuations in foreign exchange rates could cause unanticipated fluctuations in the Company's operating results, financial position or cash flows. The significant balances in foreign currencies as at March 31, 2010 are as follows:

	U.S. dollars	EURO	ZAR
	\$	€	
Cash, cash equivalents and deposits	1,089	412	21,000
Accounts receivable	316	319	—
Accounts payable and accrued liabilities	(1,286)	(142)	—
Net Exposure	119	589	21,000

Based on the aforementioned net exposure as at March 31, 2010, and assuming that all other variables remain constant, a ten-point increase or decrease in the CAD/USD, CAD/EURO and CAD/ZAR exchange rates would have an effect of \$291 on net earnings.

Interest rate risk

The Company is subject to interest rate risk on its cash and marketable securities. The Company does not believe that the results of operations or cash flows would be materially affected to any significant degree by a sudden change in market interest rates relative to interest rates on the investments, owing to the relative short-term nature of the marketable securities.

9. Comparative figures

Certain of the comparative figures have been reclassified to conform to the presentation adopted in the current year.

10. Contingencies

During the quarter ended March 31, 2010, in connection with the Company's previously disclosed tax contingency, the Company received notices of re-assessment from the Canada Revenue Agency ["CRA"] and the Ontario Minister of Finance ["OMF"] reversing their original position on the use of certain non-capital losses acquired as part of the Dimethaid Health Care Ltd. [subsequently renamed Squire Pharmaceuticals Inc. "Squire"] acquisition from Nuvo Research Inc. ["Nuvo"].

10. Contingencies (cont'd)

As previously disclosed, on various dates during fiscal 2008 and 2009 the Company had received notices of re-assessment from the CRA relating to the taxation years ending August 16, 2005, July 31, 2006, July 31, 2007, and December 31, 2008 and from the OMF for the taxation year ended August 16, 2005, containing adjustments relating to the use of certain non-capital losses. The notices of assessment and re-assessment, if they had stood as a result of the CRA's position, amounted to a total tax liability exposure to the federal and relevant provincial governments of approximately \$11,625 including interest and penalties. The Company filed a Notice of Objection through the CRA appeals process on October 23, 2008. Furthermore, the Company, under the terms of the Share Purchase Agreement ["SPA"] for Squire with Nuvo holds indemnities with respect to the status of the Squire tax accounts and certain tax asset values the Company as well as all costs relating to reassessment including advisory fees, interest and penalties, as applicable. In addition, Nuvo had issued additional security over the indemnity obligations by entitling the Company to the benefit of security over certain assets and product revenue streams of Nuvo and certain of its subsidiaries.

In connection with the appeals process, during the years ended December 31, 2009 and 2008, the Company had posted a deposit of \$3,752 to the CRA and \$500 to the OMF, representing up to one half of the tax and interest assessed. In addition, during the year ended December 31, 2009, the Company issued from its revolving unsecured credit facility, a bank guarantee in the amount of \$720 to the OMF. As a result of the Company's success in the appeal process, an amount of \$3,936 was received from the CRA on January 20, 2010 and an amount of \$504 was received from OMF on April 6, 2010, representing a refund for the full amount of the deposits above, along with accrued interest in the amount of \$188. In addition, the bank guarantee previously issued to the OMF expired February 1, 2010 without being drawn-down by the OMF.

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