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D> For the Three-Month Period Ended March 31, **2014** 2013 **Revenues** Royalties**$ 97** $ 80 License and other revenue **125** 77 **Total Revenues 222** 157 **Expenses** Research and development expense **188** 167 Selling, general and administrative expense **460** 456 Depreciation of tangible assets **7** 10 Amortization of intangible assets **9** 10 **Total Costs and Expenses 664** 643 **Net Loss (442)** (486)**Other Comprehensive Loss** Foreign currency translation adjustment **(231)** (36)**Comprehensive Loss$ (673)**$ (522) **Basic and Diluted Weighted Average Number of Shares Outstanding 62,064,139** 50,236,255 Basic and Diluted Loss Per Common Share (note 9)**$ (0.01)**$ (0.01)

See accompanying notes

IntelGenx Technologies Corp.

Consolidated Statement of Cash Flows
(Expressed in thousands of U.S. Dollars ($000 s) Except Share and Per Share Data)
(Unaudited)

	For the Three-Month Period Ended March 31,	
	2014	2013
Funds Provided (Used) -		
Operating Activities		
Net loss	**$ (442)**	$ (486)
Amortization and depreciation	**16**	20
Stock-based compensation	**32**	18
	(394)	(448)
Changes in assets and liabilities:		
Accounts receivable	**118**	1,112
Prepaid expenses	**41**	13
Investment tax credits receivable	**(17)**	(30)
Accounts payable and accrued liabilities	**(309)**	(560)
Deferred revenue	**(27)**	(77)
Net change in assets and liabilities	**(194)**	458
Net cash (used) / provided by operating activities	**(588)**	10
Financing Activities		
Proceeds from exercise of warrants	**1,064**	195
Net cash provided by financing activities	**1,064**	195
Investing Activities		
Additions to property and equipment	**(105)**	(69)
Net cash used in investing activities	**(105)**	(69)
Increase in Cash and Cash Equivalents	**371**	136
Effect of Foreign Exchange on Cash and Cash Equivalents	**(210)**	(27)
Cash and Cash Equivalents		
Beginning of Period	**5,005**	2,059
End of Period	**$ 5,166**	$ 2,168

See accompanying notes

IntelGenx Technologies Corp.

Notes to Consolidated Interim Financial Statements
March 31, 2014
(Expressed in U.S. Funds)
(Unaudited)

1. Basis of Presentation

The accompanying unaudited consolidated financial statements have been prepared in accordance with generally accepted accounting principles for interim financial information and with the instructions to Form 10-Q and Article 10 of Regulation S-X. Accordingly, they do not include all of the information and footnotes required by generally accepted accounting principles for complete consolidated financial statements. In the opinion of management, all adjustments considered necessary for a fair presentation have been included. All such adjustments are of a normal and recurring nature.

These financial statements should be read in conjunction with the audited consolidated financial statements at December 31, 2013. Operating results for the three months ended March 31, 2014 are not necessarily indicative of the results that may be expected for the year ending December 31, 2014. The Company prepares its financial statements in accordance with accounting principles generally accepted in the United States (U.S. GAAP). This basis of accounting involves the application of accrual accounting and consequently, revenues and gains are recognized when earned, and expenses and losses are recognized when incurred.

The consolidated financial statements include the accounts of the Company and its subsidiary companies. On consolidation, all inter-entity transactions and balances have been eliminated.

The financial statements are expressed in U.S. funds.

Management has performed an evaluation of the Company s activities through the date and time these financial statements were issued and concluded that there are no additional significant events requiring recognition or disclosure.

2. Adoption of New Accounting Standards

The FASB issued Update No. 2013-04, Liabilities (Topic 405) Obligations Resulting from Joint and Several Liability Arrangements for Which the Total Amount of the Obligation Is Fixed at the Reporting Date . The amendments in this Update provide guidance for the recognition, measurement, and disclosure of obligations resulting from joint and several liability arrangements for which the total amount of the obligation within the scope of this Update is fixed at the reporting date, except for obligations addressed within existing guidance in U.S. GAAP. The guidance requires an entity to measure those obligations as the sum of the amount the reporting entity agreed to pay on the basis of its arrangement among its co-obligors and any additional amount the reporting entity expects to pay on behalf of its co-obligors. The guidance in this Update also requires an entity to disclose the nature and amount of the obligation as well as other information about those obligations. For public entities, the amendments in this ASU were applicable for fiscal years, and interim periods within those years, beginning after December 15, 2013. The adoption of this Statement did not have a material effect on the Company s financial position or results of operations.

IntelGenx Technologies Corp.

Notes to Consolidated Interim Financial Statements
March 31, 2014
(Expressed in U.S. Funds)
(Unaudited)

2. Adoption of New Accounting Standards (Cont d)

The FASB issued Update No. 2013-05, Foreign Currency Matters (Topic 830) Parent s Accounting for the Cumulative Translation Adjustment upon Derecognition of Certain Subsidiaries or Groups of Assets within a Foreign Entity or of an Investment in a Foreign Entity . The amendments in this Update resolve the diversity in practice about whether Subtopic 810-10, Consolidation Overall, or Subtopic 830-30, Foreign Currency Matters Translation of Financial Statements, applies to the release of the cumulative translation adjustment into net income when a parent either sells a part or all of its investment in a foreign entity or no longer holds a controlling financial interest in a subsidiary or group of assets that is a nonprofit activity or a business (other than a sale of in substance real estate or conveyance of oil and gas mineral rights) within a foreign entity. In addition, the amendments in this Update resolve the diversity in practice for the treatment of business combinations achieved in stages (sometimes also referred to as step acquisitions) involving a foreign entity. For public entities, the amendments in this ASU were effective prospectively for fiscal years, and interim reporting periods within those years, beginning after December 15, 2013. The adoption of this Statement did not have a material effect on the Company s financial position or results of operations.

The FASB issued Update No. 2013-07, Presentation of Financial Statements Liquidation Basis of Accounting . The objective of this Update is to clarify when an entity should apply the liquidation basis of accounting and to provide principles for the measurement of assets and liabilities under the liquidation basis of accounting, as well as any required disclosures. These amendments were effective for entities that determine liquidation is imminent during annual reporting periods beginning after December 15, 2013, and interim reporting periods therein. Entitles should apply the requirements prospectively from the day that liquidation becomes imminent. The adoption of this Statement did not have a material effect on the Company s financial position or results of operations.

The FASB issued Update No. 2013-11, Income Taxes (Topic 740) Presentation of an Unrecognized Tax Benefit When a Net Operating Loss Carryforward, a Similar Tax Loss, or a Tax Credit Carryforward Exists . The amendments in this ASU provide guidance on the financial statement presentation of an unrecognized tax benefit when a net operating loss carryforward, similar tax loss, or tax credit carryforward exists. The amendments were effective for fiscal years, and interim periods within those years, beginning after December 15, 2013 and should be applied prospectively to all unrecognized tax benefits that exist at the effective date. The adoption of this Statement did not have a material effect on the Company s financial position or results of operations.

3. Significant Accounting Policies

Recently Issued Accounting Pronouncements

None of the new pronouncements issued by the FASB but not yet effective as of March 31, 2014 are applicable to the Company.

IntelGenx Technologies Corp.

Notes to Consolidated Interim Financial Statements
March 31, 2014
(Expressed in U.S. Funds)
(Unaudited)

4. Intangible Assets

As of March 31, 2014 NDA acquisition costs of $70 thousand (December 31, 2013 - $79 thousand) were recorded as intangible assets on the Company s balance sheet and represent the net book value of the final progress payment related to the acquisition of 100% ownership of Forfivo XL®. The asset is being amortized over its estimated useful life of 39 months. The Company commenced amortization upon commercial launch of the product in October 2012.

5. Deferred License Revenue

Deferred license revenue represents upfront payments received for the granting of licenses to the Company s patents, intellectual property, and proprietary technology, for commercialization. Deferred license revenue is recognized in income over the period where sales of the licensed products will occur.

Upon entering into the licensing agreement with Edgemont Pharmaceuticals the Company received an upfront fee of $1 million, which the Company recognized as deferred license revenue. The deferred license revenue is being amortized in income over a period of 39 months, which is the minimum period where sales of Forfivo XL® are expected to be exclusive.

In January, 2014 IntelGenx entered into a development and commercialization agreement with Par Pharmaceutical, Inc. for two products. The Company received $100 thousand upon execution of the agreement, of which $50 thousand has been recognized as deferred revenue until certain development milestones have been achieved.

As a result of this policy, the Company has a deferred revenue balance of $589 thousand at March 31, 2014 (December 31, 2013 - $616 thousand) that has not been recognized as revenue.

6. Capital Stock

	March 31, 2014	**December 31, 2013**
Authorized -		
100,000,000 common shares of $0.00001 par value		
20,000,000 preferred shares of $0.00001 par value		
Issued -		
62,650,655 (December 31, 2013 - 60,984,267) common shares	**$ 627**	$ 610

IntelGenx Technologies Corp.

Notes to Consolidated Interim Financial Statements
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7. Additional Paid-In Capital

Stock options

No stock options were exercised during the three month period ended March 31, 2014. During the three month period ended March 31, 2013 a total of 50,000 stock options were exercised for 50,000 common shares having a par value of $0 thousand in aggregate, for cash consideration of $23 thousand, resulting in an increase in additional paid-in capital of $23 thousand.

Compensation expenses for stock-based compensation of $32 thousand and $18 thousand were recorded during the three-month periods ended March 31, 2014 and 2013 respectively. The entire amount expensed in 2014 relates to stock options granted to employees and directors. Of the amount expensed in 2013, $13 thousand relates to stock options granted to employees and directors and $5 thousand relates to options granted to independent third party consultants. As at March 31, 2014 the Company has $194 thousand (2013 - $50 thousand) of unrecognized stock-based compensation.

Warrants

During the three month period ended March 31, 2014 a total of 1,666,388 (2013 - 362,500) warrants were exercised for 1,666,388 (2013 - 362,500) common shares having a par value of $0 thousand in aggregate, for cash consideration of $1,064 thousand (2013 - $172 thousand), resulting in an increase in additional paid-in capital of $1,064 thousand (2013 - $171 thousand).

8. Related Party Transactions

Included in management salaries are $15 thousand (2013 - $3 thousand) for options granted to the Chief Executive Officer and $11 thousand (2013 - $2 thousand) for options granted to the Chief Financial Officer under the 2006 Stock Option Plan and $3 thousand (2013 - $4 thousand) for options granted to non-employee directors. In addition, included in management salaries during the first three months of 2013 are $1 thousand for options granted to a director, who is also an employee of the Company.

Also included in management salaries are director fees of $22 thousand (2013 - $22 thousand) for attendance to board meetings and audit committee meetings. In addition, during the first three months of 2013 the Company paid $54 thousand in fees to a director under a management consultancy agreement.

The above related party transactions have been measured at the exchange amount which is the amount of the consideration established and agreed to by the related parties.

IntelGenx Technologies Corp.

Notes to Consolidated Interim Financial Statements
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(Expressed in U.S. Funds)
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9. Basic and Diluted Loss Per Common Share

Basic and diluted loss per common share is calculated based on the weighted average number of shares outstanding during the period. The warrants, share-based compensation and convertible notes have been excluded from the calculation of diluted loss per share since they are anti-dilutive.

10. Subsequent Events

Subsequent to the end of the quarter 565,000 warrants were exercised for 565,000 common shares having a par value of $0 thousand for cash consideration of approximately $370 thousand, resulting in an increase in additional paid-in capital of approximately $370 thousand.

Item 2: MANAGEMENT S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

Introduction to management s discussion and analysis

The purpose of this section, Management s Discussion and Analysis of Financial Condition and Results of Operations, is to provide a narrative explanation of the financial statements that enables investors to better understand the business of the Company, to enhance the Company s overall financial disclosures, to provide the context within which the Company s financial information may be analyzed, and to provide information about the quality of, and potential variability of, the Company s financial condition, results of operations and cash flows. Unless otherwise indicated, all financial and statistical information included herein relates to continuing operations of the Company. Unless otherwise indicated or the context otherwise requires, the words, IntelGenx, Company , we , us , and our refer to IntelG Technologies Corp. and its subsidiaries, including IntelGenx Corp. This information should be read in conjunction with the accompanying unaudited Consolidated Financial Statements and Notes thereto.

Company background

We are a drug delivery company established in 2003 and headquartered in Montreal, Quebec, Canada. Our focus is on the development of novel oral immediate-release and controlled-release products for the pharmaceutical market. Our business strategy is to develop pharmaceutical products based on our proprietary drug delivery technologies and, once the viability of a product has been demonstrated, to license the commercial rights to partners in the pharmaceutical industry. In certain cases, we rely upon partners in the pharmaceutical industry to fund development of the licensed products, complete the regulatory approval process with the U.S. Food and Drug Administration (FDA) or other regulatory agencies relating to the licensed products, and assume responsibility for marketing and distributing such products.

In addition, we may choose to pursue the development of certain products until the project reaches the marketing and distribution stage. We will assess the potential for successful development of a product and associated costs, and then determine at which stage it is most prudent to seek a partner, balancing such costs against the potential for additional returns earned by partnering later in the development process.

We have also undertaken a strategy under which we will work with pharmaceutical companies in order to develop new dosage forms for pharmaceutical products for which patent protection is nearing expiration. Under §(505)(b)(2) of the Food, Drug, and Cosmetics Act, the FDA may grant market exclusivity for a term of up to three years of exclusivity following approval of a listed drug that contains previously approved active ingredients but is approved in a new dosage, dosage form, route of administration or combination, or for a new use, the approval of which was required to be supported by new clinical trials, other than bioavailability studies, conducted by or for the sponsor.

We are currently continuing to develop the existing products in our pipeline and may also perform research and development on other potential products as opportunities arise.

We currently purchase and/or lease, on an as-needed basis, the equipment necessary for performing research and development activities related to our products.

We plan to hire new personnel, primarily in the areas of research and development, manufacturing, and administration on an as-needed basis as we enter into partnership agreements, establish our pilot plant VersaFilm manufacturing capability, and increase our research and development activities.

Key developments

Par Pharmaceutical, Inc.

On January 13, 2014 we announced the execution of a second development and commercialization agreement with Par Pharmaceutical, Inc. ("Par") for two new products utilizing our proprietary oral drug delivery platforms.

Under the terms of the agreement, Par has obtained certain exclusive rights to market and sell our products in the USA. In exchange we will receive upfront and milestone payments, together with a share of the profits upon commercialization. In accordance with confidentiality clauses contained in the agreement, the specifics of the products and financial terms remain confidential.

Anti-migraine VersaFilm product

On February 4, 2014 we, together with our co-development partner RedHill Biopharma Ltd. ("RedHill"), announced receipt of a Complete Response Letter ("CRL") from the U.S. Food and Drug Administration ("FDA") regarding the New Drug Application ("NDA") for our VersaFilm product for the treatment of acute migraines. The anti-migraine VersaFilm product is a proprietary oral thin film formulation of rizatriptan benzoate, a 5-HT1 receptor agonist and the active drug in Merck & Co.'s Maxalt®.

A CRL is issued by the FDA's Center for Drug Evaluation and Research to inform companies that certain questions and deficiencies remain that preclude the approval of the application in its present form. The questions raised by the FDA in the CRL regarding the NDA for the anti-migraine VersaFilm product primarily relate to Chemistry, Manufacturing and Controls ("CMC") and to the packaging and labeling of the product. No questions or deficiencies were raised relating to the product's safety and the FDA's CRL does not require additional clinical studies.

On March 3, 2014 we, together with RedHill, announced the submission of a response to the FDA s CRL and, subsequent to the end of the quarter, on April 24, 2014 IntelGenx and RedHill (the Companies), reported that the FDA had acknowledged receipt of our response and has requested additional CMC data, which the Companies believe they can supply within several weeks based on available information.

The Companies further reported that a supplier of raw material for the anti-migraine VersaFilm product is currently holding compliance discussions with the FDA, which are independent of RedHill and IntelGenx and are not specific to our anti-migraine VersaFilm product. The Companies are diligently working on a variety of options to ensure continued supply of the raw material regardless of the result of these compliance discussions.

The Companies believe that FDA approval of the anti-migraine VersaFilm product NDA is subject to the satisfactory resolution of the remaining CMC questions, as well as securing a compliant source of the raw material. Therefore, IntelGenx and RedHill will continue to work with the FDA in order to submit all the data requested, and will provide an update as and when applicable.

Subsequent to the end of the quarter, on April 28, 2014 the Companies announced the commencement of a comparative bioavailability clinical study comparing the anti-migraine VersaFilm product to the European reference drug. The study is intended to support the planned submission of a European Marketing Authorization Application ("MAA") and follows a positive scientific advice meeting with the German Federal Institute for Drugs and Medical Devices ("BfArM") announced by RedHill in November 2013. This single-dose, crossover, comparative bioavailability study includes 26 healthy volunteers and is intended to evaluate and compare the relative bioavailability and to assess the bioequivalence of the anti-migraine VersaFilm product and the reference drug, Maxalt® lingua, marketed in Germany by MSD SHARP & DOHME GMBH.

Results of the bioavailability study are anticipated by June 2014. Subject to the results of the study and to the required regulatory process, and in light of the data from prior successful studies conducted with the anti-migraine VersaFilm product, the Companies plan to submit a European MAA in the third quarter of 2014, with Germany as the reference member state, under the European Mutual Recognition Procedure ("MRP").

Erectile Dysfunction VersaFilm product

On February 24, 2014 we announced the completion of a pilot biostudy with our proprietary VersaFilm tadalafil product for erectile dysfunction that indicated bioequivalence with the leading brand reference listed drug (RLD) tadalafil product.

This was a randomized, two-period, two-way crossover study in healthy male subjects. The study was designed to determine whether VersaFilm tadalafil was bioequivalent as measured by industry standard pharmacokinetic measures of peak plasma concentration (Cmax) and area under the curve (AUC). The study results demonstrated that VersaFilm tadalafil was within an acceptable range of bioequivalency with the RLD on both of these measures.

Government Funding for CNS VersaFilm product

Subsequent to the end of the quarter, on April 30, 2014 we announced financial support from the National Research Council of Canada Industrial Research Assistance Program (NRC-IRAP). In addition to advisory services and technological expertise, the funding provided by NRC-IRAP will support further development of a product for the treatment of central nervous system (CNS) diseases and disorders. The product will be based upon our proprietary, oral thin film, VersaFilm , technology.

In order to maintain our competitive advantage, no specific details related to this project are being disclosed at this time.

U.S. patent allowances

On February 26, 2014 we announced receipt of a Notice of Allowance ("NOA") from the United States Patent and Trademark Office ("USPTO") for U.S. Patent Application Serial No. 11/647,033 entitled "Multilayer tablet" which covers the technology used in our hypertension product currently under development. A second NOA has been received for U.S. Patent Application Serial No. 11/782,838 entitled "Controlled-release pharmaceutical tablets" which is related to the drug delivery technology used in Forfivo XL®, our first FDA-approved product currently commercialized in the U.S. These two NOA's conclude the examination of each U.S. patent application and will result in the issuance of two U.S. patents after administrative processes are completed.

Subsequent to the end of the quarter, on April 16, 2014 we announced receipt of a further NOA from the USPTO for U.S. Patent Application Serial No. 12/836,810 entitled "Oral mucoadhesive dosage form" which covers IntelGenx' proprietary AdVersa mucoadhesive drug delivery technology. This NOA concludes the examination of the U.S. patent application and will result in the issuance of a U.S. patent after the administrative process is completed.

Currency rate fluctuations

Our operating currency is Canadian dollars, while its reporting currency is U.S. dollars. Accordingly, our results of operations and balance sheet position have been affected by currency rate fluctuations. The following management discussion and analysis takes this into consideration whenever material.

Results of operations for the three month period ended March 31, 2014 compared with the three month period ended March 31, 2013.

In U.S.$ thousands	2014	2013	Increase/ (Decrease)	Percentage Increase/ (Decrease)
Revenue	$ 222	$ 157	$ 65	41%
Research and Development Expenses	188	167	21	13%
Selling, General and Administrative Expenses	460	456	4	1%
Depreciation of tangible assets	7	10	(3)	(30%)
Amortization of intangible assets	9	10	(1)	(10%)
Net Loss	**(442)**	**(486)**	**(44)**	**(9%)**

Revenue

Total revenue in the first three months of 2014 increased to $222 thousand from $157 thousand in the same period of 2013.

Of the total revenue recorded during the first quarter of 2014, $173 thousand (2013: $157 thousand) relates to Forfivo XL®, our first FDA approved product, which was launched in October 2012 under a licensing partnership with Edgemont Pharmaceuticals LLP (Edgemont). Upon entering into the licensing agreement, Edgemont paid us an upfront fee of $1 million, which we recognized as deferred license revenue. The deferred license revenue is amortized in income over the period where sales of Forfivo XL® are expected to be exclusive. As a result of this policy, we recognized $77 thousand in income during the first quarter of 2014 (2013: $77 thousand). In addition, we recognized approximately $96 thousand of royalty income earned from the sale of Forfivo XL® in the first quarter of 2014. This compares with approximately $80 thousand in the same period of 2013, the majority of which related to initial supplies for the launch of the product in Q4, 2012. Pursuant to the contractual terms, royalty income relates to sales of Forfivo XL® recorded by Edgemont during the quarter preceding receipt of the royalty income by us. Forfivo XL® is indicated for the treatment of Major Depressive Disorder (MDD) and is the only extended-release bupropion HCl product to provide a once-daily, 450mg dose in a single tablet.

The level of sales achieved for Forfivo XL® to date has been considerably lower than anticipated, resulting in a proportionately lower level of royalty income. Management continues to work diligently with the commercialization partner to explore options to accelerate sales growth of Forfivo XL®, which have grown by an average of approximately 97% per quarter for the last three quarters ending December 31, 2013.

Revenue for the three months ended March 31, 2014 also includes a $50 thousand milestone payment in respect of one of the two new projects being developed in accordance with our new development and commercialization agreement with Par, utilizing our proprietary oral drug delivery platforms. As previously stated, in accordance with confidentiality clauses contained in the agreement, the specifics of the product and financial terms remain confidential.

Research and development (R&D) expenses

R&D expenses, net of R&D investment tax credits, totaled $188 thousand in the three months ended March 31, 2014, representing an increase of $21 thousand, or 13%, to the amount of $167 thousand expensed in the same period of last year.

The increase in R&D expenses primarily relates to the costs of a pilot clinical study for our VersaFilm product for erectile dysfunction that indicated bioequivalence with the leading brand reference listed drug tadalafil product, partly offset by a reduction in R&D staff salaries, as described below.

Included within R&D expenses for the first three months of 2014 are R&D Salaries of $115 thousand, of which approximately $2 thousand represents non-cash compensation. This compares to R&D salaries of $153 thousand in the first three months of 2013, of which approximately $3 thousand represented non-cash compensation. The reduction in R&D salaries is primarily attributable to the retirement, effective December 31, 2013, of Dr. Horst Zerbe, our founder, and former President and CEO. 50% of Dr. Zerbe s expenses were previously apportioned to the R&D department. Dr. Zerbe remains available to us to consult on R&D activities.

In the three months ended March 31, 2014 we recorded estimated Research and Development Tax Credits and refunds of $27 thousand, compared with $35 thousand that was recorded in the same period of the previous year.

Selling, general and administrative (SG&A) expenses

SG&A expenses increased by $4 thousand, to $460 thousand, in the first three months of 2014 compared with $456 thousand in the first three months of 2013.

Included in SG&A expenses are approximately $27 thousand (2013: $5 thousand) in non-cash compensation from options granted to management employees in 2012 and 2013, $3 thousand (2013: $4 thousand) in non-cash compensation from options granted to non-employee directors in 2013, and $Nil (2013: $6 thousand) in non-cash compensation from options granted to consultants in 2012.

Depreciation of tangible assets

In the three months ended March 31, 2014 we recorded an expense of $7 thousand for the depreciation of tangible assets, compared with an expense of $10 thousand for the same period of the previous year.

Amortization of intangible assets

The amortization of intangible assets expense for the first three months of 2014 totaled $9 thousand, compared with $10 thousand in the same period of last year. The expense relates to the amortization of NDA acquisition costs in respect of the final progress payment to acquire 100% ownership of Forfivo XL®. Commercialization of Forfivo XL® in October 2012 triggered amortization of the asset over its estimated useful life of 39 months.

Share-based compensation expense, warrants and stock based payments

Share-based compensation expense, warrants and share-based payments totaled $32 thousand for the three months ended March 31, 2014, compared with $18 thousand for the three months ended March 31, 2013.

We expensed approximately $29 thousand in the first three months of 2014 for options granted to our employees in 2011 and 2012 under the 2006 Stock Option Plan, and approximately $3 thousand for options granted to non-employee directors in 2013, compared with $8 thousand and $4 respectively that was expensed in the same period of the previous year.

We also expensed $6 thousand in the first three months of 2013 for options granted to consultants.

There remains approximately $194 thousand in stock based compensation to be expensed in fiscal 2014 and 2015, all of which relates to the issuance of options to our employees and directors during 2012 and 2013. We anticipate the issuance of additional options and warrants in the future, which will continue to result in stock-based compensation

expense.

Key items from the balance sheet.

In U.S.$ thousands	March 31, 2014	December 31, 2013	Increase/ (Decrease)	Percentage Increase/ (Decrease)
Current Assets	$ 5,569	$ 5,550	$ 19	0%
Leasehold improvements and Equipment	664	588	76	13%
Intangible Assets	70	79	(9)	(11%)
Current Liabilities	642	901	(259)	(29%)
Deferred License Revenue	231	308	(77)	(25%)
Capital Stock	1	1	0	0%
Additional Paid-in-Capital	22,030	20,934	1,094	5%

Current assets

Current assets totaled $5,569 thousand at March 31, 2014 compared with $5,550 thousand at December 31, 2013. The increase of $19 thousand is attributable to an increase in cash and cash equivalents of approximately $161 thousand and an increase in investment tax credits receivable of approximately $17 thousand, partly offset by a decrease in accounts receivable of approximately $118 thousand and a decrease in prepaid expenses of approximately $41 thousand.

Cash and cash equivalents

Cash and cash equivalents totaled $5,166 thousand as at March 31, 2014 representing an increase of $161 thousand compared with the balance of $5,005 thousand as at December 31, 2013. The increase in cash on hand relates to net cash provided by financing activities of $1,064 thousand, partly offset with net cash used by operating activities of $588 thousand, net cash used in investing activities of $105 thousand and an unrealized foreign exchange loss of $210 thousand.

The cash provided by financing activities derives from the exercise of 1,666,388 warrants that were exercised for 1,666,388 common shares for cash consideration of $1,064 thousand.

Accounts receivable

Accounts receivable totaled $26 thousand as at March 31, 2014 representing a decrease of $118 thousand compared with the balance of $144 thousand as at December 31, 2013. The decreased balance relates to the payment of client invoices in Q1, 2014 that were issued and outstanding at December 31, 2013.

Prepaid expenses

As of March 31, 2014 prepaid expenses totaled $92 thousand compared with $133 thousand as of December 31, 2013. The decrease in prepaid expenses relates to a deposit paid in December 2013 for a biostudy undertaken in the first quarter of 2014, and a deposit paid in 2013 for R&D machinery to be supplied and installed in 2014.

Investment tax credits receivable

R&D investment tax credits receivable totaled approximately $285 thousand as at March 31, 2014 compared with $268 thousand as at December 31, 2013. The increase relates to the accrual recorded for the first quarter of 2014, partly offset by the unrealized foreign exchange loss of converting our operating currency of CAD$ into our reporting currency of US$.

Leasehold improvements and equipment

As at March 31, 2014, the net book value of leasehold improvements and equipment amounted to $664 thousand, compared to $588 thousand at December 31, 2013. In the three months ended March 31, 2014 additions to assets totaled $105 thousand and comprised $49 thousand for pilot plant manufacturing equipment for our VersaFilm products, $38 thousand for R&D equipment, $16 thousand for leasehold improvements and $2 thousand for computer equipment. In the three months ended March 31, 2014 we recorded depreciation on leasehold improvements and equipment of $7 thousand and incurred an unrealized foreign exchange loss of $22 thousand.

Intangible assets

As at March 31, 2014 NDA acquisition costs of $70 thousand (December 31, 2013 - $79 thousand) were recorded as intangible assets on our balance sheet and are related to the acquisition of 100% ownership of Forfivo XL®. The asset is being amortized over its expected useful life of 39 months and amortization commenced upon commercial launch of Forfivo XL® in the fourth quarter of 2012.

Current liabilities

Current liabilities totaled $642 thousand as at March 31, 2014 (December 31, 2013 - $901 thousand) and consisted of accounts payable and accrued liabilities of $284 thousand (December 31, 2013 - $593 thousand) and the current portion of deferred license revenue of $358 thousand (December 31, 2013 - $308 thousand).

Included in the accounts payable and accrued liabilities balance of $284 thousand as at March 31, 2014 is approximately $22 thousand relating to research and development activities, approximately $69 thousand relating to professional fees, and approximately $144 thousand relates to accrued payroll liabilities. This compares with approximately $100 thousand relating to research and development activities, approximately $180 thousand relating to professional fees, of which approximately $87 thousand relates to the public offering completed in December, 2013, and approximately $301 thousand relates to accrued payroll liabilities, that was included in the accounts payable and accrued liabilities balance as at December 31, 2013.

Deferred license revenue

Pursuant to the execution of a licensing agreement for Forfivo XL®, we received an upfront fee from Edgemont Pharmaceuticals in the first quarter of 2012, which we recognized as deferred license revenue. The deferred license revenue is amortized in income over the period where sales of Forfivo XL® are expected to be exclusive. As a result of this policy, we have a deferred revenue balance related to this upfront fee of $539 thousand at March 31, 2014 (December 31, 2013: $616 thousand) that has not been recognized as revenue, with $231 thousand recognized as the non-current portion and $308 thousand recognized in current assets as the current portion, versus $308 thousand and $308 thousand respectively as at December 31, 2013.

In January, 2014 IntelGenx entered into a development and commercialization agreement with Par Pharmaceutical, Inc. for two products. The Company received $100 thousand upon execution of the agreement, of which $50 thousand has been recognized in current liabilities as deferred revenue until certain development milestones that are expected to be achieved in 2014 have been realised.

Shareholders equity

As at March 31, 2014 we had accumulated a deficit of $16,544 thousand compared with an accumulated deficit of $16,102 thousand as at December 31, 2013. Total assets amounted to $6,303 thousand and shareholders equity totaled $5,431 thousand as at March 31, 2014, compared with total assets and shareholders equity of $6,217 thousand and $5,008 thousand respectively, as at December 31, 2013.

Capital stock

As at March 31, 2014 capital stock amounted to $627 compared to $610 at December 31, 2013. The increase reflects the issuance of 1,666,388 shares related to the exercise of warrants, with all shares issued at par value of $0.00001. Capital stock is disclosed at its par value with the excess of proceeds shown in Additional paid-in-capital.

Additional paid-in-capital

Additional paid-in capital totaled $22,030 thousand at March 31, 2014, compared with $20,934 thousand at December 31, 2013. Additional paid-in capital increased by $1,064 thousand for warrants exercised during the first quarter, and by $32 thousand for stock based compensation attributable to the amortization of stock options granted to employees and directors.

Key items from the statement of cash flows

In U.S.$ thousands	March 31, 2014	March 31, 2013	Increase/ (Decrease)	Percentage Increase/ (Decrease)
Operating Activities	$ (588)	$ 10	$ (598)	(5,980%)
Financing Activities	1,064	$ 195	$ 869	446%
Investing Activities	(105)	(69)	36	52%
Cash and cash equivalents - end of period	5,166	2,168	2,998	138%

Statement of cash flows

Net cash used by operating activities was $588 thousand in the three months ended March 31, 2014, compared with net cash generated of $10 thousand for the three months ended March 31, 2013. In the first quarter of 2014, net cash used by operating activities consisted of an operating loss of $394 thousand (2013: $448 thousand) net of non-cash related expenses of approximately $48 thousand (2013: $38 thousand), and a decrease in non-cash operating elements of working capital of $194 thousand, compared with an increase in non-cash operating elements of working capital of $458 thousand in the same period of the previous year.

Operating activities will continue to consume our available funds until we are able to generate increased revenues.

The net cash provided by financing activities was $1,064 thousand in the first three months of 2014, compared with $195 thousand provided in the same period of the previous year. The net cash provided in the first quarter of 2014 resulted from the exercise of 1,666,388 warrants, whereas the cash provided in the first quarter of 2013 resulted from the exercise of 362,500 warrants and 50,000 options.

Net cash used in investing activities amounted to $105 thousand in the three months ended March 31, 2014 compared with $69 thousand in the three months ended March 31, 2013. Included within the use of funds in the first quarter of 2014 is an investment of approximately $49 thousand (2013: $68 thousand) for pilot plant manufacturing equipment for our VersaFilm products, $38 thousand (2013: $Nil) for R&D equipment, $16 thousand (2013: $Nil) for leasehold improvements and $2 thousand (2013: $1 thousand) for computer equipment.

The balance of cash and cash equivalents as at March 31, 2014 amounted to $5,166 thousand, compared with $2,168 thousand at March 31, 2013.

Off-balance sheet arrangements

We have no off-balance sheet arrangements.

Item 3. Controls and Procedures.

As of the end of the period covered by this report, we carried out an evaluation, under the supervision and with the participation of management, including our chief executive officer and principal financial officer, of the effectiveness of the design and operation of our disclosure controls and procedures as defined in Rules 13a-15(e) and 15d-15(e) of the Securities Exchange Act of 1934. Based upon that evaluation, our chief executive officer and principal financial officer concluded that our disclosure controls and procedures are effective to cause the material information required to be disclosed by us in the reports that we file or submit under the Exchange Act to be recorded, processed, summarized and reported within the time periods specified in the SEC's rules and forms. There have been no significant changes in our internal controls or in other factors which could significantly affect internal controls subsequent to the date we carried out our evaluation.

PART II

Item 1. Legal Proceedings

This Item is not applicable

Item 2. Unregistered Sales of Equity Securities and Use of Proceeds

This Item is not applicable.

Item 3. Defaults Upon Senior Securities

This Item is not applicable.

Item 4. (Reserved)

Item 5. Other Information

This Item is not applicable.

Item 6. Exhibits

Exhibit 31.1
Certification of C.E.O. Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.

Exhibit 31.2
Certification of Principal Accounting Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.

Exhibit 32.1
Certification of C.E.O. pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.

Exhibit 32.2
Certification of Principal Accounting Officer pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.

SIGNATURES

In accordance with the requirements of the Securities Exchange Act of 1934, the Registrant caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

INTELGENX TECHNOLOGIES CORPORATION

Date: May 13, 2014

By: /s/ *Rajiv Khosla*
Rajiv Khosla
President, C.E.O. and
Director

Date: May 13, 2014

By: /s/ *Paul Simmons*
Paul A. Simmons
Principal Accounting Officer

Exhibit 31.1

CERTIFICATION PURSUANT TO
SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Rajiv Khosla, Chief Executive Officer of IntelGenx Technologies Corp. (the "registrant"), certify that:

1. I have reviewed this quarterly report on Form 10-Q of IntelGenx Technologies Corp.;

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

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

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

a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

c) Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and

d) Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

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

a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and

b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: May 13, 2014

/s/ *Rajiv Khosla*
Rajiv Khosla
Chief Executive Officer

Exhibit 31.2

CERTIFICATION PURSUANT TO
SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Paul A. Simmons, Principal Accounting Officer of IntelGenx Technologies Corp. (the "registrant"), certify that:

1. I have reviewed this quarterly report on Form 10-Q of IntelGenx Technologies Corp. ;

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

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

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

a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

c) Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and

d) Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

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

a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and

b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: May 13, 2014

/s/ *Paul A. Simmons*
Paul A. Simmons
Principal Accounting Officer

Exhibit 32.1

CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350,
AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

In connection with the Quarterly Report of IntelGenx Technologies Corporation (the "Company") on Form 10-Q for the period ending March 31, 2014, as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I, Rajiv Khosla, Chief Executive Officer of the Company, certify, pursuant to 18 U.S.C. Sec. 1350, as adopted pursuant to Sec. 906 of the Sarbanes-Oxley Act of 2002, that, to the best of my knowledge and belief:

(1) The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and

(2) The information contained in the Report fairly presents, in all material respects, the financial condition and result of operations of the Company.

/s/ Rajiv Khosla
Rajiv Khosla
Chief Executive Officer
May 13, 2014

A signed original of this written statement required by Section 906, or other document authenticating, acknowledging, or otherwise adopting the signature that appears in typed form within the electronic version of this written statement has been provided to the Company and will be retained by the Company and furnished to the Securities and Exchange Commission or its staff upon request. The foregoing certifications are accompanying the Company's Form 10-Q solely pursuant to section 906 of the Sarbanes-Oxley Act of 2002 (subsections (a) and (b) of section 1350, chapter 63 of title 18, United States Code) and is not being filed as part of the Form 10-Q or as a separate disclosure document.

Exhibit 32.2

CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350,
AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

In connection with the Quarterly Report of IntelGenx Technologies Corporation(the "Company") on Form 10-Q for the period ending March 31, 2014, as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I, Paul A. Simmons, Principal Accounting Officer of the Company, certify, pursuant to 18 U.S.C. Sec. 1350, as adopted pursuant to Sec. 906 of the Sarbanes-Oxley Act of 2002, that, to the best of my knowledge and belief:

(1) The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and

(2) The information contained in the Report fairly presents, in all material respects, the financial condition and result of operations of the Company.

/s/ *Paul A. Simmons*
Paul A. Simmons
Principal Accounting Officer
May 13, 2014

A signed original of this written statement required by Section 906, or other document authenticating, acknowledging, or otherwise adopting the signature that appears in typed form within the electronic version of this written statement has been provided to the Company and will be retained by the Company and furnished to the Securities and Exchange Commission or its staff upon request. The foregoing certifications are accompanying the Company's Form 10-Q solely pursuant to section 906 of the Sarbanes-Oxley Act of 2002 (subsections (a) and (b) of section 1350, chapter 63 of title 18, United States Code) and is not being filed as part of the Form 10-Q or as a separate disclosure document.