

ONCOLYTICS BIOTECH INC

Form 6-K

August 11, 2004

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

**FORM 6-K**

**Report of Foreign Private Issuer**

**Pursuant to Rule 13a-16 or 15d-16  
of the Securities Exchange Act of 1934**

For the month of August 2004

Commission File Number 000-31062

**Oncolytics Biotech Inc.**

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*(Translation of registrant's name into English)*

**Suite 210, 1167 Kensington Crescent NW  
Calgary, Alberta, Canada T2N 1X7**

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*(Address of principal executive offices)*

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F.

Form 20-F

Form 40-F

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):

**Note:** Regulation S-T Rule 101(b)(1) only permits the submission in paper of a Form 6-K if submitted solely to provide an attached annual report to security holders.

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):

**Note:** Regulation S-T Rule 101(b)(7) only permits the submission in paper of a Form 6-K if submitted to furnish a report or other document that the registrant foreign private issuer must furnish and make public under the laws of the jurisdiction in which the registrant is incorporated, domiciled or legally organized (the registrant's home country), or under the rules of the home country exchange on which the registrant's securities are traded, as long as the report or other document is not a press release, is not required to be and has not been distributed to the registrant's security holders, and, if discussing a material event, has already been the subject of a Form 6-K submission or other Commission filing on EDGAR.

Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes

No

If Yes is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b):  
82-\_\_\_\_\_

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Signatures

Second Quarter Report

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**Signatures**

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

Oncolytics Biotech Inc.  
(Registrant)

Date August 11, 2004

By: /s/ Douglas A. Ball  
Douglas A. Ball  
Chief Financial Officer

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Second Quarter Report

June 30, 2004

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Oncolytics Biotech Inc.  
TSX: ONC  
NASDAQ: ONCY

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**SECOND QUARTER REPORT**

*For the quarter ended June 30, 2004*

**Letter to Shareholders**

In the second quarter of 2004, Oncolytics announced the commencement of enrollment in its first systemic administration clinical trial of REOLYSIN®; completed a private placement financing that raised gross proceeds of \$6.73 million; and strengthened the Board of Directors with the addition of Mr. J. Mark Lievonen.

**Clinical Studies**

In May, Oncolytics started patient treatment in its UK Phase I clinical trial investigating the systemic delivery of REOLYSIN® as a treatment for up to 40 patients with advanced or metastatic solid tumours. This clinical trial is an open-label, dose escalation Phase I study in which REOLYSIN® is administered intravenously to patients diagnosed with advanced or metastatic solid tumours that have not responded to conventional therapy, or for which no curative standard therapy exists. This trial is another important step in the development of REOLYSIN®.

**Board of Directors**

The Company also strengthened its board of directors through the addition of Mr. J. Mark Lievonen, President of Aventis Pasteur Limited. Mr. Lievonen is responsible for Aventis Pasteur's operations in Canada.

**Financing**

In April, the Company announced the successful completion of a \$6.73 million private placement with a European institutional investor. In addition, during the second quarter, the Company received proceeds from the exercise of warrants and options of \$4.0 million. The Company now believes it has sufficient reserves to fund its present plans for research and development and operational activities into 2007.

Thank you for your support.

Brad Thompson, PhD  
Chairman, President and  
CEO August 5, 2004

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**August 5, 2004**

**MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL  
CONDITION AND RESULTS OF OPERATIONS**

This discussion and analysis should be read in conjunction with the unaudited financial statements of Oncolytics Biotech Inc. (Oncolytics or the Company) as at and for the three and six months ended June 30, 2004 and 2003, and should also be read in conjunction with the audited financial statements and Management's Discussion and Analysis of Financial Condition and Results of Operations (MD&A) contained in Oncolytics' annual report for the year ended December 31, 2003. The financial statements have been prepared in accordance with Canadian generally accepted accounting principles (GAAP).

**FORWARD-LOOKING STATEMENTS**

*The following discussion contains forward-looking statements, within the meaning of Section 21E of the Securities Exchange Act of 1934, as amended. Forward-looking statements, including the Company's belief as to the potential of REOLYSIN® as a cancer therapeutic, the Company's expectation regarding the adequacy of its existing capital resources, and the Company's expectations as to the success of its research and development programs in 2004 and beyond, future financial position, business strategy and plans for future operations, and statements that are not historical facts, involve known and unknown risks and uncertainties, which could cause the Company's actual results to differ materially from those in the forward-looking statements. Such risks and uncertainties include, among others, the availability of funds and resources to pursue research and development projects, the efficacy of REOLYSIN® as a cancer treatment, the success and timely completion of clinical studies and trials, the Company's ability to successfully commercialize REOLYSIN®, uncertainties related to the research and development of pharmaceuticals, uncertainties related to competition, changes in technology, the regulatory process and general changes to the economic environment. Investors should consult the Company's quarterly and annual filings with the Canadian and U.S. securities commissions for additional information on risks and uncertainties relating to the forward-looking statements. Forward-looking statements are based on assumptions, projections, estimates and expectations of management at the time such forward looking statements are made, and such assumptions, projections, estimates and/or expectations could change or prove to be incorrect or inaccurate. Investors are cautioned against placing undue reliance on forward-looking statements. The Company does not undertake to update these forward-looking statements.*

**OVERVIEW**

**Oncolytics Biotech Inc. is a Development Stage Company**

Since its inception in April of 1998, Oncolytics Biotech Inc. (the Company) has been a development stage company and has focused its research and development efforts on the development of REOLYSIN®, its potential cancer therapeutic. The Company has not been profitable since its inception and expects to continue to incur substantial losses from its research and development. The Company does not expect to generate significant revenues until, if and when, its cancer product becomes commercially viable.



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**General Risk Factors**

Prospects for biotechnology companies in the research and development stage should generally be regarded as speculative. It is not possible to predict, based upon studies in animals, or early studies in humans, whether a new therapeutic will ultimately prove to be safe and effective in humans, or whether necessary and sufficient data can be developed through the clinical trial process to support a successful product application and approval.

If a product is approved for sale, product manufacturing at a commercial scale and significant sales to end users at a commercially reasonable price may not be successful. There can be no assurance that the Company will generate adequate funds to continue development, or will ever achieve significant revenues or profitable operations. Many factors (e.g. competition, patent protection, appropriate regulatory approvals) can influence the revenue and product profitability potential.

In developing a product for approval, the Company will rely upon its employees, contractors, consultants and collaborators and other third party relationships, including the ability to obtain appropriate product liability insurance. There can be no assurance that these reliances and relationships will continue as required.

In addition to developmental and operational considerations, market prices for securities of biotechnology companies generally are volatile, and may or may not move in a manner consistent with the progress being made by the Company.

**Highlights**

During the second quarter of 2004, the Company's net loss was \$3,191,888 compared to \$3,911,473 for the second quarter of 2003. The decrease in the Company's net loss is mainly due to the loss on sale of investment in Transition Therapeutics Inc. ( TTH ) of \$2,156,685 that occurred in 2003. This decrease was offset by an increase in expenses for the three month period ending June 30, 2004 associated with the Company's operations. Specifically, manufacturing and related process development expenses increased in the second quarter of 2004 compared to 2003 as the Company has increased its production of REOLYSIN® in order to supply its clinical trial program. As well, the Company's systemic (intravenous) delivery clinical trial in the United Kingdom ( U.K. ) commenced patient enrolment in the second quarter of 2004 increasing clinical trial costs in the second quarter of 2004 compared to 2003. Also, the Company's pre-clinical trial expenses increased in support of future clinical trial applications that include other jurisdictions and methods of application. Finally, the Company recorded a non-cash expense for stock based compensation related to stock options granted in the second quarter of 2004.

The Company's cash balance continued to improve through the closing of a private placement and the exercise of warrants and options. During the six months ended June 30, 2004, the Company received additional cash proceeds from financing activities of \$10,264,784. The Company exited the second quarter of 2004 with cash and short-term investments of \$25,522,728 compared to \$20,752,735 as at December 31, 2003.

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**Table of Contents****SECOND QUARTER RESULTS OF OPERATIONS***(for the three months ended June 30, 2004 and 2003)*

Net loss for the three month period ended June 30, 2004 was \$3,191,888 compared to \$3,911,473 for 2003. The decrease in the Company's net loss in the second quarter of 2004 was mainly due to the loss from sale of investment in TTH of \$2,156,685 that occurred in 2003 and increases during the second quarter of 2004 in the Company's operating activities as follows:

**Research and Development Expenses ( R&D )**

	<b>2004</b>	<b>2003</b>
	<b>\$</b>	<b>\$</b>
Manufacturing and related process development expenses	<b>810,748</b>	422,626
Clinical trial expenses	<b>167,051</b>	78,145
Pre-clinical trial expenses	<b>334,603</b>	80,141
Other R&D expenses	<b>183,532</b>	267,809
	<hr/>	<hr/>
Research and development expenses	<b>1,495,934</b>	848,721
	<hr/>	<hr/>

For the second quarter of 2004, R&D increased to \$1,495,934 compared to \$848,721 for the second quarter of 2003. The increase in R&D was due to the following:

***Manufacturing & Related Process Development***

During the second quarter of 2004, the Company continued to focus on the production of REOLYSIN® in order to supply its R&D activity. As well, additional production costs were incurred relating to the technology transfer and set up costs associated with the Company's second manufacturer.

***Clinical Trial Programs***

The Company's clinical trial expenses increased to \$167,051 in the second quarter of 2004 compared to \$78,145 for the second quarter of 2003. The increase in the second quarter of 2004 relates to the Company commencing patient enrolment in and supporting its systemic (intravenous) delivery clinical trial in the United Kingdom.

***Pre-Clinical Trial Expenses***

During the second quarter of 2004, the Company incurred pre-clinical trial costs associated with toxicology and equivalency studies being performed in support of future clinical trial applications. These types of studies were limited in the second quarter of 2003.

**Operating Expenses**

<b>2004</b>	<b>2003</b>
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	<u>\$</u>	<u>\$</u>
Salary, insurance and other office expenses	<b>344,624</b>	290,616
Public company and other operating expenses	<b>615,702</b>	424,143
	<b>960,326</b>	714,759

For the second quarter of 2004, the Company's operating expenses increased to \$960,326 compared to \$714,759 for the second quarter of 2003. Public company and other operating costs increased in the second quarter of 2004 compared to 2003 reflecting the increased costs associated with the preparation of the Company's annual filings, annual general meeting and shareholder mail outs plus additional expenses incurred in investor relations and business development.

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**Table of Contents****Stock Based Compensation**

	<b>2004</b>	<b>2003</b>
	<b>\$</b>	<b>\$</b>
	<hr/>	<hr/>
Stock based compensation	<b>734,670</b>	68,318

During the second quarter of 2004, the Company recorded stock based compensation of \$734,670 associated with the granting of stock options to its employees, directors, and certain consultants. Stock based compensation recorded in 2003 related to previously granted options that vested in the second quarter of 2003 and options granted to consultants.

**YEAR TO DATE RESULTS OF OPERATIONS**  
*(for the six months ended June 30, 2004 and 2003)*

Net loss for the six month period ended June 30, 2004 was \$5,868,124 compared to \$5,025,787 for 2003. The increase in the Company's net loss was due to the following:

**Research and Development Expenses ( R&D )**

	<b>2004</b>	<b>2003</b>
	<b>\$</b>	<b>\$</b>
	<hr/>	<hr/>
Manufacturing and related process development expenses	<b>2,187,178</b>	488,735
Clinical trial expenses	<b>292,696</b>	88,787
Pre-clinical trial expenses	<b>513,562</b>	137,966
Other R&D expenses	<b>456,895</b>	612,868
	<hr/>	<hr/>
Research and development expenses	<b>3,450,331</b>	1,328,356
	<hr/>	<hr/>

For the six month period ending June 30, 2004, R&D increased to \$3,450,331 compared to \$1,328,356 for 2003. The increase in R&D was due to the following:

***Manufacturing & Related Process Development***

In 2004 the Company continued to focus on the production of REOLYSIN® in order to supply its existing and planned R&D activity. As well, the Company took steps to mitigate the risk of economic dependence as a result of having only one supplier of REOLYSIN®. Consequently, almost 75% of the Company's manufacturing and related process development expenses incurred in 2004 related to the production of REOLYSIN® compared to only 8% in 2003. The Company's manufacturing expenses in 2004 also include technology transfer and set up costs associated with the addition of a second supplier.

The remaining manufacturing and related process development costs incurred in 2004 and almost all of these costs

incurred in 2003 relate to process development. During the first six months of 2003 the Company was completing the development of its manufacturing process and also commenced the development of its viral and cell banks.

Consequently, 85% of the Company's manufacturing and related process development expenses incurred in 2003 related to these activities compared to only 19% in 2004.

For the remainder of 2004, the Company expects that it will continue to produce REOLYSIN® and that a majority of these costs will relate directly to manufacturing. As well, future manufacturing costs may be impacted by the need to supply the clinical trials to be run in accordance with the agreement between the U.S. National Cancer Institute ( NCI ) and the Company.

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**Table of Contents*****Clinical Trial Programs***

The Company's clinical trial expenses increased to \$292,696 for the six month period ending June 30, 2004 compared to \$88,787 in 2003. The increase in clinical trial expenses relates mainly to the costs associated with the Company's systemic (intravenous) delivery clinical trial in the United Kingdom. The Company also continues to incur expenses related to the Canadian malignant glioma clinical trial.

For the remainder of 2004, the Company expects that clinical trial expenses will continue to be incurred as enrolment continues in the systemic (intravenous) delivery clinical trial. As well, the Company expects that its clinical trial costs may increase as it continues to try and expand its clinical trial program into other jurisdictions.

***Pre-Clinical Trial Expenses***

The Company's pre-clinical trial expenses increased to \$513,562 for the six month period ending June 30, 2004 compared to \$137,966 in 2003. Pre-clinical costs include toxicology studies and are incurred by the Company in support of expanding its clinical trial program into other jurisdictions and other applications.

**Operating Expenses**

	<b>2004</b>	<b>2003</b>
	<b>\$</b>	<b>\$</b>
Salary, insurance and other office expenses	<b>723,040</b>	505,893
Public company and other operating expenses	<b>938,460</b>	730,420
	<b>1,661,500</b>	1,236,313

For the six month period ending June 30, 2004, the Company's operating expenses increased to \$1,661,500 compared to \$1,236,313 for the six month period ending June 30, 2003. Salary, insurance and other office expenses increased to \$723,040 for the six month period ending June 30, 2004 from \$505,893 in 2003 due to the increase in staff levels and insurance premiums that commenced in the second quarter of 2003. Public company and other operating costs increased to \$938,460 for the six month period ending June 30, 2004 from \$730,420 in the first six months of 2003 reflecting the increased costs associated with the preparation of the Company's annual filings, annual general meeting and shareholder mail outs plus additional expenses incurred in investor relations and business development.

**Sale of Investments**

	<b>2004</b>	<b>2003</b>
	<b>\$</b>	<b>\$</b>
Gain on sale of investment in BCY LifeSciences Inc. ( BCY )	<b>47,002</b>	
Loss on sale of investment in TTH		2,156,685

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For the six month period ending June 30, 2004 the Company sold 697,945 common shares of BCY for net cash proceeds of \$133,609. This resulted in an accounting gain of \$47,002. As at June 30, 2004, the Company owned 200,000 common share of BCY with an estimated market value of \$20,000. These remaining shares are held in escrow and will be released over the next two years.

For the six month period ending June 30, 2003, the Company sold its investment in TTH for net cash proceeds of \$2,552,695 resulting in a recorded loss of \$2,156,685.

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**Commitments**

As at June 30, 2004, the Company has committed to payments totaling \$766,500 for activities primarily related to product manufacturing and continued toxicology related work. The Company anticipates that these committed payments will occur in 2004. All of these committed payments are considered to be part of the Company's normal course of business.

**LIQUIDITY AND CAPITAL RESOURCES**

**Liquidity**

As at June 30, 2004, the Company had cash of \$25,522,728 (including cash, cash equivalents and short-term investments) and a working capital position of \$25,234,266 compared to \$20,752,735 and \$20,088,868 respectively as at December 31, 2003. During the second quarter of 2004, the Company continued to improve its cash position through a private placement of 1,077,100 units at an average price of \$6.25 per unit. Net cash proceeds after issue costs were \$6,223,763 and each unit was comprised of 1,077,100 common shares and 538,550 common share purchase warrants. Each whole common share purchase warrant entitles the holder to acquire one common share of the capital of the Company upon payment of \$7.75 per share until October 7, 2005. The Company also received cash proceeds from the exercise of warrants from previously closed financings of \$3,300,038 and from the exercise of stock options of \$740,983. Consequently, for the six month period ending June 30, 2004, the Company has received a net amount of \$10,264,784. This increase in the Company's cash position has been offset by cash outflows from operating activities of \$5,194,579 and purchases of intellectual property and other assets of \$433,821.

The Company desires to maintain adequate cash and short-term investment reserves to support its planned activities which include its clinical trial program, production manufacturing, and its intellectual property expansion and protection as well as administrative activities. The Company believes that its existing capital resources are adequate to fund its current plans for research and development activities into 2007 without presuming the further exercise of outstanding warrants and options. In the event that the Company chooses to seek additional capital, the Company will look to fund additional capital requirements through the issue of additional equity as well as potential partnering or licensing opportunities. The Company recognizes the challenges and uncertainty inherent in the capital markets and the potential difficulties it might face in today's environment. Market prices for securities in biotechnology companies are volatile and the ability to raise funds will be dependent on a number of factors, including the progress of R&D, availability of clinical trial information, and general market conditions.

**Capital Expenditures**

During the six month period ending June 30, 2004, the Company spent \$425,928 on intellectual property compared to \$595,147 in 2003. The difference relates to variances in filing fees on existing patent applications.

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**Table of Contents****SUMMARY OF QUARTERLY RESULTS**

The following unaudited quarterly information is presented in thousands of dollars except for per share amounts:

	2004		2003				2002	
	June <sup>(2)</sup>	March <sup>(2)</sup>	Dec. <sup>(2)</sup>	Sept.	June <sup>(2)</sup>	March	Dec.	Sept.
<b>Revenue<sup>(1)</sup></b>	<b>183</b>	<b>117</b>	<b>127</b>	<b>102</b>	<b>41</b>	<b>43</b>	<b>44</b>	<b>53</b>
<b>Net loss<sup>(3)</sup></b>	<b>3,192</b>	<b>2,676</b>	<b>1,696</b>	<b>1,823</b>	<b>3,911</b>	<b>1,114</b>	<b>1,542</b>	<b>1,990</b>
<b>Loss per common share<sup>(3)</sup></b>	<b>\$ 0.11</b>	<b>\$ 0.10</b>	<b>\$ 0.06</b>	<b>\$ 0.07</b>	<b>\$ 0.17</b>	<b>\$ 0.05</b>	<b>\$ 0.07</b>	<b>\$ 0.09</b>
<b>Total assets<sup>(4), (6)</sup></b>	<b>31,221</b>	<b>25,435</b>	<b>26,051</b>	<b>21,532</b>	<b>18,815</b>	<b>16,702</b>	<b>17,968</b>	<b>17,331</b>
<b>Total cash<sup>(5), (6)</sup></b>	<b>25,522</b>	<b>20,298</b>	<b>20,753</b>	<b>15,843</b>	<b>13,486</b>	<b>6,887</b>	<b>8,319</b>	<b>7,746</b>
<b>Total long-term debt<sup>(7)</sup></b>	<b>150</b>	<b>150</b>	<b>150</b>	<b>150</b>	<b>150</b>	<b>150</b>	<b>150</b>	<b>150</b>
<b>Cash dividends declared<sup>(8)</sup></b>	<b>Nil</b>	<b>Nil</b>	<b>Nil</b>	<b>Nil</b>	<b>Nil</b>	<b>Nil</b>	<b>Nil</b>	<b>Nil</b>

- (1) Revenue is comprised of interest income.
- (2) Included in net loss and net loss per share in March 2004 and December 2003 is a gain on sale of investment of \$47,648 and \$264,453 respectively and in June 2004 and 2003 is a loss from sale of investments of \$646 and \$2,156,685 respectively.
- (3) Included in net loss and net loss per share for 2002 is a future income tax recovery of \$647,618 (2004 and 2003 nil).
- (4) Subsequent to the acquisition of the Company by SYNSORB in April 1999, the Company applied push down accounting. See note 2 to the audited financial statements for 2003.
- (5) Included in total cash are cash, cash equivalents and short-term investments.
- (6) The Company issued 2,163,709 common shares for cash proceeds of \$10,264,784 in 2004 (2003 5,062,978 common shares for \$16,004,981 and 2002 1,040,000 common shares for \$1,803,877).
- (7) The long-term debt recorded in 2004, 2003 and 2002 represents repayable loans from the Alberta Heritage Foundation.
- (8) The Company has not declared or paid any dividends since incorporation.

**OTHER MD&A REQUIREMENTS**

The Company has 29,452,618 common shares outstanding at July 31, 2004. If all of the Company's warrants and options were exercised the Company would have 35,234,077 common shares outstanding.

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Financial Statements

**Oncolytics Biotech Inc.**

June 30, 2004

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**Table of Contents****Oncolytics Biotech Inc.****BALANCE SHEETS**

As at,

	<b>June 30, 2004 \$ (unaudited)</b>	<b>December 31, 2003 \$ (audited)*</b>
<b>ASSETS</b>		
<b>Current</b>		
Cash and cash equivalents	<b>3,058,642</b>	2,641,127
Short-term investments	<b>22,464,086</b>	18,111,608
Accounts receivable	<b>63,618</b>	64,224
Prepaid expenses	<b>566,331</b>	156,837
	<b>26,152,677</b>	20,973,796
<b>Capital assets</b>	<b>5,043,344</b>	4,965,379
<b>Investments [note 2]</b>	<b>24,818</b>	111,425
	<b>31,220,839</b>	26,050,600
<b>LIABILITIES AND SHAREHOLDERS EQUITY</b>		
<b>Current</b>		
Accounts payable and accrued liabilities	<b>918,411</b>	884,928
<b>Alberta Heritage Foundation loan</b>	<b>150,000</b>	150,000
<b>Shareholders equity</b>		
Share capital [note 3]		
Authorized: unlimited		
Issued: 29,371,971 (2003 27,208,262)	<b>54,295,023</b>	44,712,589
Warrants [note 3]	<b>2,280,600</b>	1,598,250
Contributed surplus	<b>4,439,521</b>	3,699,425
Deficit	<b>(30,862,716)</b>	(24,994,592)
	<b>30,152,428</b>	25,015,672

**31,220,839**

**26,050,600**

*See accompanying notes*

\* Derived from the December 31, 2003 audited financial statements

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## Oncolytics Biotech Inc.

## STATEMENTS OF LOSS AND DEFICIT

	Six Month Period Ending June 30, 2004 \$ (unaudited)	Six Month Period Ending June 30, 2003 \$ (unaudited)	Three Month Period Ending June 30, 2004 \$ (unaudited)	Three Month Period Ending June 30, 2003 \$ (unaudited)	Cumulative from inception on April 2, 1998 to June 30, 2004 \$ (unaudited)
<b>Revenue</b>					
Rights revenue					310,000
Interest income	<b>300,815</b>	84,526	<b>183,459</b>	41,356	2,386,798
	<b>300,815</b>	84,526	<b>183,459</b>	41,356	2,696,798
<b>Expenses</b>					
Research and development	<b>3,450,331</b>	1,328,356	<b>1,495,934</b>	848,721	19,868,861
Operating	<b>1,661,500</b>	1,236,313	<b>960,326</b>	714,759	8,865,527
Stock based compensation	<b>740,096</b>	68,789	<b>734,670</b>	68,318	1,769,521
Amortization	<b>363,856</b>	318,940	<b>184,833</b>	163,716	2,273,946
	<b>6,215,783</b>	2,952,398	<b>3,375,763</b>	1,795,514	32,777,855
<b>Loss before the following:</b>	<b>5,914,968</b>	2,867,872	<b>3,192,304</b>	1,754,158	30,081,057
<b>(Gain) loss on sale of BCY LifeSciences Inc. [note 2]</b>	<b>(47,002)</b>		<b>646</b>		(311,455)
<b>Loss on sale of Transition Therapeutics Inc.</b>		2,156,685		2,156,685	2,156,685
<b>Loss before taxes</b>	<b>5,867,966</b>	5,024,557	<b>3,192,950</b>	3,910,843	31,926,287
<b>Capital tax (recovery)</b>	<b>158</b>	1,230	<b>(1,062)</b>	630	51,429
<b>Future income tax recovery</b>					(1,115,000)
<b>Net loss for the period</b>	<b>5,868,124</b>	5,025,787	<b>3,191,888</b>	3,911,473	30,862,716
<b>Deficit, beginning of period</b>	<b>24,994,592</b>	16,450,561	<b>27,670,828</b>	17,564,875	
<b>Deficit, end of period</b>	<b>30,862,716</b>	21,476,348	<b>30,862,716</b>	21,476,348	30,862,716

	<u>          </u>	<u>          </u>	<u>          </u>	<u>          </u>	<u>          </u>
<b>Basic and diluted loss per share</b>	<b>0.21</b>	0.22	<b>0.11</b>	0.17	
	<u>          </u>	<u>          </u>	<u>          </u>	<u>          </u>	
<b>Weighted average number of shares</b>	<b>28,100,033</b>	22,396,218	<b>28,944,326</b>	22,569,011	
	<u>          </u>	<u>          </u>	<u>          </u>	<u>          </u>	

*See accompanying notes*



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## Oncolytics Biotech Inc.

## STATEMENTS OF CASH FLOWS

	Six Month Period Ending June 30, 2004 \$ (unaudited)	Six Month Period Ending June 30, 2003 \$ (unaudited)	Three Month Period Ending June 30, 2004 \$ (unaudited)	Three Month Period Ending June 30, 2003 \$ (unaudited)	Cumulative from inception on April 2, 1998 to June 30, 2004 \$ (unaudited)
<b>OPERATING ACTIVITIES</b>					
Net loss for the period	(5,868,124)	(5,025,787)	(3,191,888)	(3,911,473)	(30,862,716)
Deduct non-cash items					
Amortization	363,856	318,940	184,833	163,716	2,273,946
Non-cash compensation	740,096	68,789	734,670	68,318	1,769,521
(Gain) loss on sale of BCY LifeSciences Inc.	(47,002)		646		(311,455)
Loss on sale of Transition Therapeutics Inc.		2,156,685		2,156,685	2,156,685
Future income tax recovery					(1,115,000)
Net changes in non-cash working capital	(383,405)	(669,452)	(1,523,988)	(412,316)	195,795
	<u>(5,194,579)</u>	<u>(3,150,825)</u>	<u>(3,795,727)</u>	<u>(1,935,070)</u>	<u>(25,893,224)</u>
<b>INVESTING ACTIVITIES</b>					
Intellectual property	(425,928)	(595,147)	(295,388)	(135,487)	(3,090,754)
Other capital assets	(7,893)	(41,431)	(6,295)	(40,809)	(518,865)
Purchase of short-term investments	(6,352,478)		(6,107,212)		(24,464,086)
Redemption of short-term investments	2,000,000		1,000,000		2,000,000
Investment in BCY LifeSciences Inc.	133,609		1,959		456,637
Investment in Transition Therapeutics Inc.		2,552,695		2,552,695	2,532,343
	<u>(4,652,690)</u>	<u>1,916,117</u>	<u>(5,406,936)</u>	<u>2,376,399</u>	<u>(23,084,725)</u>
<b>FINANCING ACTIVITIES</b>					
Alberta Heritage Foundation loan					150,000

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Proceeds from exercise of warrants and stock options	<b>4,041,021</b>	339,975	<b>3,096,276</b>	339,975	7,502,006
Proceeds from private placements	<b>6,223,763</b>	6,061,585	<b>6,223,763</b>	5,817,414	22,741,983
Proceeds from public offerings					21,642,602
	<b>10,264,784</b>	6,401,560	<b>9,320,039</b>	6,157,389	52,036,591
<b>Increase in cash and cash equivalents during the period</b>	<b>417,515</b>	5,166,852	<b>117,376</b>	6,598,718	3,058,642
<b>Cash and cash equivalents, beginning of the period</b>	<b>2,641,127</b>	8,319,244	<b>2,941,266</b>	6,887,378	
<b>Cash and cash equivalents, end of the period</b>	<b>3,058,642</b>	13,486,096	<b>3,058,642</b>	13,486,096	3,058,642

*See accompanying notes*



**Table of Contents****Oncolytics Biotech Inc.****NOTES TO FINANCIAL STATEMENTS**June 30, 2004 and 2003 (*unaudited*)**1. ACCOUNTING POLICIES**

These unaudited interim financial statements do not include all of the disclosures included in the Company's annual financial statements. Accordingly, these unaudited interim financial statements should be read in conjunction with the Company's most recent annual financial statements. The information for the year ended December 31, 2003 has been derived from the Company's audited financial statements for the year then ended.

The accounting policies used in the preparation of these unaudited interim financial statements conform with those used in the Company's most recent annual financial statements.

**2. INVESTMENTS**

During the six month period ending June 30, 2004, the Company sold 697,945 of its BCY shares for net cash proceeds of \$133,609 recording a gain on sale of investment of \$47,002. As at June 30, 2004, the Company's remaining ownership in BCY was 200,000 common shares with a book value of \$24,818 and an estimated market value of \$20,000 based on the trading price at June 30, 2004.

**3. SHARE CAPITAL****Authorized:**

Unlimited number of common shares

**Issued:**

	Shares		Warrants	
	Number	Amount \$	Number	Amount \$
Balance, December 31, 2002	22,145,284	30,191,572	550,000	114,286
Issued for cash pursuant to February 10, 2003 private placement	140,000	265,540	77,000	16,000
Issued for cash pursuant to June 19, 2003 private placement	2,120,000	5,912,113	1,272,000	543,287
Issued for cash pursuant to August 21, 2003 private placement	1,363,900	3,801,778	813,533	349,176
Issued for cash pursuant to October 14, 2003 public offering	1,200,000	5,528,972	720,000	617,428
Exercise of options	64,700	149,615		
Exercise of warrants	174,378	593,194	(174,378)	(41,927)
Share issue costs		(1,730,195)		
Balance, December 31, 2003	27,208,262	44,712,589	3,258,155	1,598,250
Issued for cash pursuant to April 7, 2004 private placement (i)	1,077,100	5,924,050	646,260	1,028,631
Exercise of warrants	890,359	3,646,319	(890,359)	(346,281)
Exercise of options	196,250	740,983		

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Share issue costs		(728,918)		
	<u>                    </u>	<u>                    </u>	<u>                    </u>	<u>                    </u>
Balance, June 30, 2004	<u>29,371,971</u>	<u>54,295,023</u>	<u>3,014,056</u>	<u>2,280,600</u>

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- (i) Pursuant to a private placement, the Company sold 1,077,100 units at an average price of \$6.25 per unit for gross cash proceeds of \$6,731,875. The units were comprised of 1,077,100 common shares and 538,550 common share purchase warrants and have ascribed values of \$5.50 and \$1.50 respectively. Each common share purchase warrant entitles the holder to acquire one common share in the capital of the Company upon payment of \$7.75 per share until October 7, 2005. Share issue costs related to the private placement were \$728,918. In addition, the Company issued 107,710 common share purchase warrants to its advisor entitling the holder to acquire one common share of the capital of the Company upon payment of \$7.00 per share until October 7, 2005. The ascribed value of these additional warrants was \$220,806 (\$2.05 per additional warrant) and has been included in the share issue costs above. The ascribed values of the warrants were based on the Black Scholes Option Pricing Model.

The following table summarizes the Company's outstanding warrants as at June 30, 2004:

<b>Exercise Price</b>	<b>Outstanding, December 31, 2003</b>	<b>Granted During the Period</b>	<b>Exercised During the Period</b>	<b>Outstanding, End of Period</b>	<b>Weighted Average Remaining Contractual Life (years)</b>
\$3.00	480,755		463,255	17,500	0.11
\$4.00	2,057,400		313,060	1,744,340	0.54
\$5.00	120,000		43,794	76,206	0.79
\$6.25	600,000		70,250	529,750	0.79
\$7.00		107,710		107,710	1.25
\$7.75		538,550		538,550	1.25
	<b>3,258,155</b>	<b>646,260</b>	<b>890,359</b>	<b>3,014,056</b>	<b>0.74</b>

**Stock Option Plan**

The Company has issued stock options to acquire common stock through its stock option plan of which the following are outstanding at:

<b>June 30, 2004</b>		<b>December 31, 2003</b>	
<b>Stock</b>	<b>Weighted Average Share</b>	<b>Stock</b>	<b>Weighted Average Share</b>

	<b>Options</b>	<b>Price \$</b>	<b>Options</b>	<b>Price \$</b>
Outstanding at beginning of period	<b>2,800,800</b>	<b>3.81</b>	2,653,500	4.40
Granted during period	<b>243,500</b>	<b>8.12</b>	599,000	3.71
Cancelled during period			(387,000)	7.97
Exercised during period	<b>(196,250)</b>	<b>3.78</b>	(64,700)	2.31
	<b>2,848,050</b>	<b>4.18</b>	2,800,800	3.81
Outstanding at end of period				
Options exercisable at end of period	<b>2,768,633</b>	<b>4.21</b>	2,720,383	3.87

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June 30, 2004 and 2003 (*unaudited*)

As the Company is following the fair value based method of accounting for stock option awards, compensation expense related to options granted to employees and consultants was \$717,276 and \$22,820, respectively for the six month period ending June 30, 2004 (June 30, 2003 \$5,271 and \$63,518, respectively) with an offsetting credit to contributed surplus.

**4. COMPARATIVE FIGURES**

Certain comparative figures have been reclassified to conform to the current period's presentation.

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**Shareholder Information**

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**Officers**

**Brad Thompson, PhD**

Chairman, President and CEO

**Doug Ball, CA**

Chief Financial Officer

**George Gill, MD**

Senior Vice President, Clinical and Regulatory Affairs

**Matt Coffey, PhD**

Vice President, Product Development

**Directors**

**Brad Thompson, PhD**

Chairman, President and CEO of Oncolytics Biotech Inc.

**Doug Ball, CA**

CFO, Oncolytics Biotech Inc.

**William. A. Cochrane, OC, MD**

Biotech Consultant

**Jim Dinning**

Executive Vice President, TransAlta Corporation

**J. Mark Lievonen**

President, Aventis Pasteur Limited

**Tony Noujaim, PhD**

President and CEO of Virexx Research Inc.

**Bob Schultz, FCA**

Chairman of Rockwater Capital Corporation

**Fred Stewart, QC**

President of Fred Stewart and Associates Inc.