ONCOLYTICS BIOTECH INC
Form 6-K
October 30, 2007

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 October 2007

Commission File Number 000-31062

Oncolytics Biotech Inc.

(Translation of registrant s name into English)

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

(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 o Form 40-F þ

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): o

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): o

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 o No þ

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

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: October 30, 2007

By: /s/ Doug Ball

Doug Ball
Chief Financial Officer

210, 1167 Kensington Crescent
N.W.
Calgary, Alberta
Canada T2N 1X7

FOR IMMEDIATE RELEASE

Oncolytics Biotech Inc. Announces 2007 Third Quarter Results

CALGARY, AB, October 30, 2007 - Oncolytics Biotech Inc. (Oncolytics) (TSX:ONC, NASDAQ:ONCY) today announced its financial results and highlights for the three and nine-month periods ended September 30, 2007.

Third Quarter Highlights

Announced positive interim results from a U.K. Phase Ia/1b combination REOLYSIN® and radiation trial for patients with advanced cancers including partial and remote responses in patients with a variety of advanced cancers;

Commenced patient enrolment in a multi-centre, combination REOLYSIN® and docetaxel (Taxotere®) systemic administration trial in the U.K.;

In October, received approval from the U.K. regulatory authorities to begin a combination REOLYSIN® and cyclophosphamide trial for patients with advanced cancers;

Secured two additional U.S. patents, for a total of more than 150 issued patents worldwide; and,

Presented preclinical work at the National Cancer Research Institute Conference in Birmingham, U.K. demonstrating for the first time how reovirus-infected melanoma cells stimulate dendritic cells to prime the immune system against cancer cells.

With positive results being reported from our clinical trial program in the U.K. and the U.S., seven trials actively enrolling, an additional combination trial approved to begin and an expanding intellectual property portfolio supporting our technology, Oncolytics is looking forward to making substantial progress through the balance of 2007 and 2008, said Dr. Brad Thompson, President and CEO of Oncolytics.

Oncolytics Biotech Inc.
BALANCE SHEETS
(unaudited)

As at,

	September 30, 2007 $	**December 31, 2006** $
ASSETS		
Current		
Cash and cash equivalents	**3,326,374**	3,491,511
Short-term investments	**24,865,090**	24,122,237
Accounts receivable	**36,637**	84,003
Prepaid expenses	**413,811**	638,540
	28,641,912	28,336,291
Property and equipment	**169,226**	149,596
Intellectual property	**5,085,755**	5,079,805
	33,896,893	33,565,692
LIABILITIES AND SHAREHOLDERS EQUITY		
Current		
Accounts payable and accrued liabilities	**2,298,064**	2,616,421
Alberta Heritage Foundation loan		150,000
Shareholders equity		
Share capital		
Authorized: unlimited number of common shares Issued: 41,120,748 (December 31, 2006 36,520,748)	**92,708,665**	83,083,271
Warrants	**6,654,740**	4,216,740
Contributed surplus	**8,672,204**	8,529,326
Deficit	**(76,436,780)**	(65,030,066)
	31,598,829	30,799,271
	33,896,893	33,565,692

Oncolytics Biotech Inc.
STATEMENTS OF LOSS AND COMPREHENSIVE LOSS
(unaudited)

	Nine Month Period Ending September 30, 2007 $	**Nine Month Period Ending September 30, 2006** $	**Three Month Period Ending September 30, 2007** $	**Three Month Period Ending September 30, 2006** $	**Cumulative from inception on April 2, 1998 to September 30, 2007** $
Revenue					
Rights revenue					310,000
					310,000
Expenses					
Research and development	**8,815,255**	6,582,687	**2,890,644**	2,705,746	52,036,449
Operating	**2,798,630**	2,789,647	**880,158**	766,618	19,569,211
Stock-based compensation	**142,878**	293,880	**38,909**	34,671	4,308,527
Foreign exchange loss/(gain)	**2,829**	(2,703)	**18,917**	5,129	651,677
Amortization intellectual property	**713,887**	647,893	**244,299**	220,774	4,750,721
Amortization property and equipment	**30,061**	43,379	**10,197**	12,685	437,744
	12,503,540	10,354,783	**4,083,124**	3,745,623	81,754,329
Loss before the following:	**12,503,540**	10,354,783	**4,083,124**	3,745,623	81,444,329
Interest income	**(946,826)**	(947,364)	**(319,223)**	(320,454)	(5,749,831)
Gain on sale of BCY LifeSciences Inc.					(299,403)
Loss on sale of Transition Therapeutics Inc.					2,156,685
Loss before taxes	**11,556,714**	9,407,419	**3,763,901**	3,425,169	77,551,780
Future income tax recovery					(1,115,000)

Net loss and comprehensive loss for the period	**11,556,714**	9,407,419	**3,763,901**	3,425,169	76,436,780
Basic and diluted loss per share		**0.29**	0.26	**0.09**	0.09
Weighted average number of shares (basic and diluted)		**40,181,777**	36,317,687	**41,120,748**	36,368,270

Oncolytics Biotech Inc.
STATEMENTS OF CASH FLOWS
(unaudited)

	Nine Month Period Ending September 30, 2007 $	**Nine Month Period Ending September 30, 2006** $	**Three Month Period Ending September 30, 2007** $	**Three Month Period Ending September 30, 2006** $	**Cumulative from inception on April 2, 1998 to September 30, 2007** $
OPERATING ACTIVITIES					
Net loss for the period	**(11,556,714)**	(9,407,419)	**(3,763,901)**	(3,425,169)	(76,436,780)
Add/(deduct) non-cash items					
Amortization intellectual property	**713,887**	647,893	**244,299**	220,774	4,750,721
Amortization property and equipment	**30,061**	43,379	**10,197**	12,685	437,744
Stock-based compensation	**142,878**	293,880	**38,909**	34,671	4,308,527
Other non-cash items					1,383,537
Net changes in non-cash working capital	**(179,975)**	(34,485)	**239,121**	261,875	1,724,946
	(10,849,863)	(8,456,752)	**(3,231,375)**	(2,895,164)	(63,831,305)
INVESTING ACTIVITIES					
Intellectual property	**(586,124)**	(552,319)	**(99,066)**	(187,283)	(6,085,404)
Property and equipment	**(49,691)**	(29,342)	**(11,386)**	(8,294)	(673,039)
Purchase of short-term investments	**(742,853)**	(801,358)	**(255,688)**	(261,480)	(48,862,320)
Redemption of short-term investments		10,158,000			23,578,746
Investment in BCY LifeSciences Inc.					464,602
Investment in Transition Therapeutics Inc.					2,532,343
	(1,378,668)	8,774,981	**(366,140)**	(457,057)	(29,045,072)
FINANCING ACTIVITIES					
Proceeds from exercise of warrants and stock options		127,500		85,000	15,208,468

Proceeds from private placements					38,137,385
Proceeds from public offerings	**12,063,394**				42,856,898
	12,063,394	127,500		85,000	96,202,751
Increase (decrease) in cash and cash equivalents during the period	**(165,137)**	445,729	**(3,597,515)**	(3,267,221)	3,326,374
Cash and cash equivalents, beginning of the period	**3,491,511**	3,511,357	**6,923,889**	7,224,307	
Cash and cash equivalents, end of the period	**3,326,374**	3,957,086	**3,326,374**	3,957,086	3,326,374

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. as at and for the three and nine months ended September 30, 2007 and 2006, 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 our annual report for the year ended December 31, 2006. 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 our belief as to the potential of REOLYSIN® as a cancer therapeutic and our expectations as to the success of our research and development and manufacturing programs in 2007 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 our actual results to differ materially from those in the forward-looking statements. Such risks and uncertainties include, among others, the need for and 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, our ability to successfully commercialize REOLYSIN®, uncertainties related to the research, development and manufacturing of pharmaceuticals, uncertainties related to competition, changes in technology, the regulatory process and general changes to the economic environment. Investors should consult our 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. We do not undertake to update these forward-looking statements.

OVERVIEW

Oncolytics Biotech Inc. is a Development Stage Company

Since our inception in April of 1998, Oncolytics Biotech Inc. has been a development stage company and we have focused our research and development efforts on the development of REOLYSIN®, our potential cancer therapeutic. We have not been profitable since our inception and expect to continue to incur substantial losses as we continue research and development efforts. We do not expect to generate significant revenues until, if and when, our cancer product becomes commercially viable.

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 we 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 pharmaceutical product, we rely upon our employees, contractors, consultants and collaborators and other third party relationships, including our 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 Oncolytics. See also *RISK FACTORS AFFECTING FUTURE PERFORMANCE* in our 2006 MD&A.

REOLYSIN® Development Update for the Third Quarter of 2007

We continue to develop our lead product REOLYSIN® as a possible cancer therapy. Our goal each year is to advance REOLYSIN® through the various steps and stages of development required for potential pharmaceutical products. In order to achieve this goal, we actively manage the development of our clinical trial program, our pre-clinical and collaborative programs, our manufacturing process and REOLYSIN® supply, and our intellectual property.

Clinical Trial Program

Our clinical trial program includes eight clinical trials of which seven are being conducted by us and one is being sponsored by the U.S. National Cancer Institute (NCI). In the third quarter of 2007, we announced positive interim results from our U.K. Phase Ia/Ib combination REOLYSIN® and radiation clinical trial. As well, we commenced patient enrollment in our U.K. combination REOLYSIN®/docetaxel clinical trial, increasing our actively enrolling clinical trials to seven.

Clinical Trial Results

In the third quarter of 2007, we announced positive interim results from our U.K. Phase Ia/Ib combination REOLYSIN® and radiation clinical trial for patients with advanced or metastatic cancers. As of September 28, 2007, 22 patients had been treated with 15 having completed the study. Five patients withdrew from the study, and two patients are still on study.

A total of 11 patients in the Ia portion of the trial have received two intratumoural treatments of REOLYSIN® at dosages of $1x10^8$, $1x10^9$, or $1x10^{10}$ $TCID_{50}$ with a constant localized radiation dose of 20 Gy given in five fractions. Of these 11 patients, three patients (oesophageal, squamous skin carcinoma and squamous cell scalp) experienced significant partial responses.

One month following treatment, the oesophageal patient experienced a 28.5% reduction in the target tumour, with stable disease noted in four, non-treated tumours. At two and three months, the target tumour had shrunk 64%, with stable disease continuing in the four non-treated tumours, including a 15% volume reduction in non-treated mediastinal disease that was maintained for more than six months. The squamous skin cancer patient experienced a 50% reduction in the target tumour, as well as stable disease in two, non-treated tumours at one, two and three months post treatment. The squamous cell scalp patient experienced stable disease in the target tumour for two months which then became a partial response at three months. This patient also experienced stable disease in one non-treated tumour measured at three months post-treatment.

Patients in the Ib portion received either two, four or six intratumoural doses of REOLYSIN® at $1x10^{10}$ $TCID_{50}$ with a constant localized radiation dose of 36 Gy given in 12 fractions. Of the six patients who have completed the study to date, three patients (colorectal, melanoma and lung cancer) experienced tumour regression in the target tumour, as well as stable disease in non-treated tumours.

The colorectal patient experienced a partial response with a more than 50% regression in the target tumour as well as stable disease in four, non-treated tumours measured at one month following treatment. A melanoma patient experienced minor regression in the target tumour as well as stable disease in two, non-treated tumours at one and two months following treatment. A lung cancer patient experienced minor regression in the target tumour, as well as stable disease in three, non-treated tumours at two months following treatment.

The treatment has been well tolerated, with mostly Grade 1 or 2 toxicities noted including fatigue, lymphopenia, fever, and neutropenia. Grade 3 toxicities including cellulitis, dysphasia and diarrhoea were related to disease progression and not to the combination treatment. Viral replication was unaffected by cellular irradiation.

The primary objective of the Phase Ia/Ib trial was to determine the maximum tolerated dose (MTD), dose limiting toxicity (DLT), and safety profile of REOLYSIN® when administered intratumourally to patients receiving radiation treatment. A secondary objective is to examine any evidence of anti-tumour activity. Eligible patients include those who have been diagnosed with late stage advanced or metastatic solid tumours that are refractory (have not responded) to standard therapy or for which no curative standard therapy exists.

Clinical Trials Actively Enrolling

At the end of the third quarter of 2007, we were actively enrolling in seven clinical trials. In the third quarter of 2007, we commenced enrollment in the following study:

U.K. Combination REOLYSIN® Docetaxel Clinical Trial

We commenced patient enrolment in our U.K. clinical trial to evaluate the anti-tumour effects of systemic administration of REOLYSIN® in combination with docetaxel (Taxotere®) in patients with advanced cancers including bladder, prostate, lung and upper gastro-intestinal. In preclinical studies, the combination of REOLYSIN® and various taxanes including docetaxel has been shown to be synergistic against a variety of cancer cell lines.

The trial has two components. The first is an open-label, dose-escalating, non-randomized study of REOLYSIN® given intravenously with docetaxel every three weeks. A standard dosage of docetaxel will be delivered with escalating dosages of REOLYSIN® intravenously. A maximum of three cohorts will be enrolled in the REOLYSIN® dose escalation portion. The second component of the trial will immediately follow and will include the enrolment of a further 12 patients at the maximum dosage of REOLYSIN® in combination with a standard dosage of docetaxel. Eligible patients include those who have been diagnosed with advanced or metastatic solid tumours such as bladder, prostate, lung or upper gastro-intestinal cancers that are refractory to standard therapy or for which no curative standard therapy exists. The primary objective of the trial is to determine the MTD, DLT, recommended dose and dosing schedule and safety profile of REOLYSIN® when administered in combination with docetaxel. Secondary objectives include the evaluation of immune response to the drug combination, the body s response to the drug combination compared to chemotherapy alone and any evidence of anti-tumour activity.

Pre-Clinical Trial and Collaborative Program

In the third quarter of 2007, we announced that a poster presentation entitled Reovirus Infection of Human Melanoma Cells Supports Priming of Anti-Tumour Cytotoxic T Cell Immunity was presented by Dr. Robin Prestwich of CR-UK Clinical Centre, Leeds Institute of Molecular Medicine, University of Leeds, U.K at the National Cancer Research Institute Cancer Conference in Birmingham, U.K.

In this study, the investigators infected melanoma cell lines with reovirus. The reovirus-infected cell lines stimulated the maturation of dendritic cells, which in turn educated cancer-killing T cells to attack and kill the melanoma cells.

Manufacturing and Process Development

We continued to have REOLYSIN® manufactured in order to supply our current and future clinical trial program. In the third quarter of 2007, our manufacturing activity was focused on the completion of the vial filling and packaging of the production runs that were completed earlier in 2007. Also in the third quarter of 2007, we continued process development that examined the scale up of our manufacturing process increasing the batch size from our present GMP scale of 20-litres to 40-litres and then to 100-litres.

Intellectual Property

In the third quarter of 2007, two U.S. patents were issued. At the end of the third quarter of 2007, we had been issued over 150 patents including 23 U.S. and six Canadian patents as well as issuances in other jurisdictions. We also have over 180 patent applications filed in the U.S., Canada and other jurisdictions.

Financial Impact

We estimated at the beginning of 2007 that our monthly cash usage would be approximately $1,400,000 for 2007. Our cash usage for the nine months ending September 30, 2007 was $10,849,863 from operating activities and $635,815 for the purchases of intellectual property and capital assets which is in line with our estimate. Our net loss for the nine month period ending September 30, 2007 was $11,556,714.

Cash Resources

We exited the third quarter of 2007 with cash resources totaling $28,191,464 (see *Liquidity and Capital Resources*).

Expected REOLYSIN® Development for the Remainder of 2007

We plan to continue to enroll patients in our seven clinical trials and expect to add an additional clinical co-therapy trial. We believe that the NCI sponsored melanoma clinical trial will receive approval to commence in 2007. We believe we will complete enrollment in our U.K. Phase Ia/Ib clinical trial by the end of 2007 and complete enrollment in our Phase II combination REOLYSIN®/radiation and chemotherapy co-therapy studies in 2008. Also, our process development activity will focus on scale up studies and the examination of a lyophilization process for REOLYSIN®. Based on our expected activity in 2007, we continue to estimate our average monthly cash usage to be $1,400,000 per month (see *Liquidity and Capital Resources*).

Recent 2007 Progress

On October 23, 2007, we announced receipt of a letter of approval to commence our clinical trial using intravenous administration of REOLYSIN® in combination with cyclophosphamide, a chemotherapeutic agent as well as immune modulator, in patients with advanced cancers.

The trial is an open-label, dose-escalating, non-randomized trial of REOLYSIN® given intravenously with escalating doses of cyclophosphamide. A standard dose of REOLYSIN® is administered intravenously over five consecutive days, while an intravenous dose of cyclophosphamide is administered three days before REOLYSIN® treatment and continues through the course of the treatment cycle. The total number of patients studied will depend on the number of dose levels tested, but it is anticipated to be approximately 30 patients.

Eligible patients include those who have been diagnosed with advanced or metastatic solid tumours including pancreatic, lung and ovarian cancers that are refractory to standard therapy or for which no curative standard therapy exists. The primary objectives of the trial include determining the Minimum Effective Immunomodulatory Dose of cyclophosphamide to obtain successful immune modulation. Secondary objectives include the safety profile of the combination and gathering any evidence of anti-tumour activity.

THIRD QUARTER RESULTS OF OPERATIONS

(for the three months ended September 30, 2007 and 2006)

Net loss for the three month period ending September 30, 2007 was $3,763,901 compared to $3,425,169 for the three month period ending September 30, 2006.

Research and Development Expenses (R&D)

	2007 $	**2006 $**
Manufacturing and related process development expenses	**879,937**	1,259,716
Clinical trial expenses	**1,278,175**	688,435
Pre-clinical trial and research collaboration expenses	**293,785**	301,165
Other R&D expenses	**438,747**	456,430
Research and development expenses	**2,890,644**	2,705,746

For the third quarter of 2007, R&D increased to $2,890,644 compared to $2,705,746 for the third quarter of 2006. The increase in R&D was due to the following:

Manufacturing & Related Process Development (M&P)

	2007 $	**2006 $**
Product manufacturing expenses	**610,842**	896,776
Technology transfer expenses		184,761
Process development expenses	**269,095**	178,179

Manufacturing and related process development expenses	**879,937**	1,259,716

During the third quarter of 2007, our M&P expenses decreased to $879,937 compared to $1,259,716 for the third quarter of 2006. In the third quarter of 2007, we continued to fill, test, and package the REOLYSIN® that was produced earlier in the year. During the third quarter of 2006, we commenced a number of cGMP production runs using our improved manufacturing process. The technology transfer of our improved process was successfully completed at the beginning of the third quarter of 2006.

Our process development studies in the third quarter of 2007 focused on increasing the scale of our production runs from batch sizes of 20 litres to 40 and then 100 litres. In the third quarter of 2006, we completed process development studies that were successful in improving virus yields.

Clinical Trial Program

	2007 **$**	**2006** **$**
Direct clinical trial expenses	**1,201,557**	639,719
Other clinical trial expenses	**76,618**	48,716
Clinical trial expenses	**1,278,175**	688,435

During the third quarter of 2007, our direct clinical trial expenses increased to $1,201,557 compared to $639,719 for the third quarter of 2006. In the third quarter of 2007, we incurred direct clinical trial expenses in our seven actively enrolling trials compared to only four enrolling trials in the third quarter of 2006.

Pre-Clinical Trial Expenses and Research Collaborations

	2007 **$**	**2006** **$**
Research collaboration expenses	**293,785**	252,460
Pre-clinical trial expenses		48,705
Pre-clinical trial expenses and research collaborations	**293,785**	301,165

During the third quarter of 2007, our research collaboration expenses were $293,785 compared to $252,460 for the third quarter of 2006. Our research collaboration activity continues to focus on the interaction of the immune system and the reovirus, the use of the reovirus as a co-therapy with existing chemotherapeutics, and to investigate new uses of the reovirus as a therapeutic.

Other Research and Development Expenses

	2007 **$**	**2006** **$**
R&D consulting fees	**38,152**	70,323
R&D salaries and benefits	**342,155**	299,224
Other R&D expenses	**58,440**	86,883
Other research and development expenses	**438,747**	456,430

Our R&D salaries and benefits costs were $342,155 in the third quarter of 2007 compared to $299,224 in the third quarter of 2006. The increase is a result of increases in salary and staff levels along with the addition of our Vice President of Intellectual Property in 2007.

Operating Expenses

	2007 $	2006 $
Public company related expenses	**635,076**	507,828
Office expenses	**245,082**	258,790
Operating expenses	**880,158**	766,618

During the third quarter of 2007, our public company related expenses were $635,076 compared to $507,828 for the third quarter of 2006. In the third quarter of 2007, we increased our professional fees and investor relations activity in the United States and Europe compared to the third quarter of 2006.

Stock Based Compensation

	2007 $	2006 $
Stock based compensation	**38,909**	34,671

Stock based compensation for the third quarter of 2007 was $38,909 compared to $34,671 for the third quarter of 2006. In the third quarters of 2007 and 2006, we incurred stock based compensation associated with the vesting of previously granted options.

YEAR TO DATE RESULTS OF OPERATIONS

(for the nine months ended September 30, 2007 and 2006)

Net loss for the nine month period ending September 30, 2007 was $11,556,714 compared to $9,407,419 for the nine month period ending September 30, 2006.

Research and Development Expenses (R&D)

	2007 $	2006 $
Manufacturing and related process development expenses	**3,546,732**	2,751,207
Clinical trial expenses	**2,983,688**	1,920,467
Pre-clinical trial and research collaboration expenses	**731,445**	691,553
Other R&D expenses	**1,553,390**	1,219,460
Research and development expenses	**8,815,255**	6,582,687

For the nine month period ending September 30, 2007, R&D increased to $8,815,255 compared to $6,582,687 for 2006. The increase in R&D was due to the following:

Manufacturing & Related Process Development (M&P)

	2007 $	2006 $
Product manufacturing expenses	**3,134,143**	1,664,308
Technology transfer expenses		457,975
Process development expenses	**412,589**	628,924

Manufacturing and related process development expenses	**3,546,732**	2,751,207

Our M&P expenses for the nine month period ending September 30, 2007 increased to $3,546,732 compared to $2,751,207 in 2006. For the nine month period ending September 30, 2007, our production and vial filling activity increased compared to 2006. During this period of 2007, we completed production runs that commenced in 2006 and initiated additional production runs to manufacture REOLYSIN® at the beginning of 2007. Also, as a result of the increased viral yields from the process development activity in 2006, we have incurred additional vial filling and packaging costs compared to 2006.

For the nine month period ending September 30, 2006, we completed the production runs that were ongoing at the end of 2005 for our Phase I trials. At the same time, our process development activity helped improve the virus yields from our manufacturing process. These improvements were then transferred to our cGMP manufacturer with additional production runs initiated in the third quarter of 2006.

Our process development expenses for the nine month period ending September 30, 2007 were $412,589 compared to $628,924 for the nine month period ending September 30, 2006. During this period of 2007, our main process development focus has been on the scale up of our production process, which has included scale up studies at 40 and 100 litres. During the nine month period ending September 30, 2006, our process development activity included viral yield and scale up studies along with the validation of our fill process.

We still expect that our overall manufacturing and related process development expenses for 2007 will be in line with 2006. In the fourth quarter of 2007, we plan to initiate a 40-litre technology transfer, complete our 100-litre scale up studies and investigate the lyophilization of REOLYSIN®. We are also examining ways to reduce our economic dependence resulting from having only a single cGMP manufacturer. This might include building up a level of inventory, increasing the scale of each production run, engaging another cGMP manufacturer or manufacturing REOLYSIN® ourselves. Depending on how we mitigate our risk of economic dependence our expectation of our 2007 M&P expenses may change.

Clinical Trial Program

	2007 **$**	**2006** **$**
Direct clinical trial expenses	**2,798,024**	1,783,138
Other clinical trial expenses	**185,664**	137,329
Clinical trial expenses	**2,983,688**	1,920,467

During the nine month period ending September 30, 2007, our direct clinical trial expenses were $2,798,024 compared to $1,783,138 for the nine month period ending September 30, 2006. In this period of 2007, we incurred direct patient costs in our seven ongoing clinical trials. As well, we incurred clinical site start up costs for our three co-therapy trials in the U.K. and our Phase II sarcoma clinical trial in the U.S. During the nine month period ending September 30, 2006, we incurred direct patient costs in four ongoing clinical trials along with clinical site start up costs associated with our U.S. recurrent malignant glioma and U.K. Phase Ia combination REOLYSIN® /radiation therapy trials.

We expect our clinical trial expenses will continue to increase for the remainder of 2007 compared to 2006 as we continue patient enrollment and expand our clinical trial program to include other trial sites and other studies.

Pre-Clinical Trial Expenses and Research Collaborations

	2007 **$**	**2006** **$**
Research collaboration expenses	**694,315**	634,199
Pre-clinical		