# 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 December 2008

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 b

Form 40-F o

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.

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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	0	No þ
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**

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: December 18, 2008 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. Completes Patient Enrolment in Two U.K. REOLYSIN® Combination Therapy Clinical Trials

Company to Host Conference Call to Update Clinical Program

**CALGARY, AB,** December 18, 2008 - Oncolytics Biotech Inc. (Oncolytics) (TSX:ONC, NASDAQ:ONCY) announced today that it has completed patient enrolment in two U.K. clinical trials using REOLYSIN® in combination with chemotherapy or radiation.

The combination REOLYSIN® and docetaxel trial, (REO 010) was designed to evaluate the anti-tumour effects of systemic administration of REOLYSIN® in combination with docetaxel (Taxotere®) in patients with advanced cancers. The principal investigator is Professor Hardev Pandha of the Royal Surrey County Hospital, U.K. The combination REOLYSIN® and radiation trial (REO 008) was designed to evaluate the anti-tumour effects of direct injection of REOLYSIN® in combination with low-dose radiation in patients with advanced cancers. The Principal Investigator for the trial is Dr. Kevin Harrington of the Targeted Therapy Laboratory, Cancer Research UK Centre for Cell and Molecular Biology at The Institute of Cancer Research and Honorary Consultant in Clinical Oncology at The Royal Marsden NHS Foundation Trust.

### **Conference Call Details**

Dr. Brad Thompson, President and CEO of Oncolytics, will host a conference call and webcast on **Friday**, **December 19, 2008 at 8:00 a.m. MT (10:00 a.m. ET)** to update investors on the current and planned clinical trial program for REOLYSIN®.

To access the conference call by telephone, dial 1-416-644-3415 or 1-800-732-9303. A live audio webcast will also be available at the following link: <a href="http://www.newswire.ca/en/webcast/viewEvent.cgi?eventID=2510860">http://www.newswire.ca/en/webcast/viewEvent.cgi?eventID=2510860</a> or through the Company s website at <a href="http://www.oncolyticsbiotech.com">www.oncolyticsbiotech.com</a>. Please connect at least 15 minutes prior to the webcast to ensure adequate time for any software download that may be needed. A replay of the webcast will be available at <a href="http://www.oncolyticsbiotech.com">www.oncolyticsbiotech.com</a> and will also be available by telephone through December 26, 2008. To access the telephone replay, dial 1-416-640-1917 or 1-877-289-8525 and enter reservation number 21292813#.

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### **About Oncolytics Biotech Inc.**

Oncolytics is a Calgary-based biotechnology company focused on the development of oncolytic viruses as potential cancer therapeutics. Oncolytics clinical program includes a variety of Phase I/II and Phase II human trials using REOLYSIN®, its proprietary formulation of the human reovirus, alone and in combination with radiation or chemotherapy. For further information about Oncolytics, please visit www.oncolyticsbiotech.com. This press release 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 expectations related to the completed and ongoing clinical trials with REOLYSIN®, and the Company s belief as to the potential of REOLYSIN® as a cancer therapeutic, 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 tolerability of REOLYSIN® outside a controlled test, 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 and uncertainties related to the regulatory process. 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. Investors are cautioned against placing undue reliance on forward-looking statements. The Company does not undertake to update these forward-looking statements, except as required by applicable laws.

## FOR FURTHER INFORMATION PLEASE CONTACT:

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