FORM 6-K

Current report of foreign issuer pursuant to Rules 13a-16 and 15d-16 Amendments

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ONCOLYTICS BIOTECH INC

CIK:**1129928**| IRS No.: **000000000** | Fiscal Year End: **1231**
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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 January 2013

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 ☑ 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 - _____

EXHIBIT NUMBER	DESCRIPTION
99.1	News Release Dated January 28, 2013: Oncolytics Biotech® Inc. Announces Positive REOLSYIN® Clinical Trial Data Presented at ASCO Gastrointestinal Cancers Symposium

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: January 28, 2013

By: /s/ Brad Thompson
Brad Thompson
President and Chief Executive Officer

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

FOR IMMEDIATE RELEASE

Oncolytics Biotech® Inc. Announces Positive REOLSYIN® Clinical Trial Data Presented at ASCO Gastrointestinal Cancers Symposium

CALGARY, AB, --- January 28, 2013 - Oncolytics Biotech Inc. ("Oncolytics") (TSX:ONC, NASDAQ:ONCY) today announced a poster presentation covering positive preliminary results from a Phase I study examining the intravenous administration of REOLYSIN in combination with FOLFIRI in patients with metastatic colorectal cancer (REO 022). The results were presented at the ASCO Gastrointestinal Cancers Symposium in San Francisco, CA, which took place from January 24-26, 2013.

The poster presentation, titled: "A Multicenter Phase I Study of Intravenous Administration of REOLYSIN in combination with Irinotecan/Fluorouracil/Leucovorin (FOLFIRI) in Patients (pts) with Oxaliplatin-Refractory/Intolerant KRAS-Mutant Metastatic Colorectal Cancer (mCRC)," was authored by Ocean et al. Twenty-one patients were enrolled in the study, including nine that were FOLFIRI-naïve. Of the 18 patients evaluable for response there was one partial response and nine had stable disease. The combined overall progression free survival (PFS) of FOLFIRI-naïve and FOLFIRI-failed patients was 7.4 months. The authors concluded that the combination of REOLYSIN and FOLFIRI was safe and well tolerated and resulted in disease control in the majority of evaluable patients, including patients that had previously progressed on Irinotecan.

"These results are intriguing, particularly with respect to patients who have failed prior FOLFIRI treatment," said Dr. Brad Thompson, President and CEO of Oncolytics. "This study, in conjunction with an earlier study examining REOLYSIN monotherapy in metastatic colorectal patients (REO 013), formed the basis for our decision to proceed into a randomized study in colorectal cancer (IND 211), which is now enrolling."

The trial was a 21-patient, single arm dose escalation study designed to determine a maximum tolerated dose and dose-limiting toxicities for the combination of REOLYSIN and FOLFIRI. Eligible patients included those with histologically confirmed cancer of the colon or rectum with Kras mutation and measurable disease. They must have progressed on or within 190 after the last dose of an oxaliplatin regimen in the metastatic setting, or be intolerant to oxaliplatin.

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 human trials including a Phase III trial in head and neck cancers using REOLYSIN®, its proprietary formulation of the human reovirus. For further information about Oncolytics, please visit: www.oncolyticsbiotech.com.

This press release contains forward-looking statements within the meaning of the U.S. Securities Act of 1933, as amended, and U.S. Securities Exchange Act of 1934, as amended, and forward-looking information within the meaning of Canadian securities laws. Statements, other than statements of historical facts, included in this press release that address activities, events or developments that Oncolytics expects or anticipates will or may occur in the future, including such things as, the Company's expectations related to the Phase I colorectal cancer trial of REOLYSIN in combination with FOLFIRI, and the Company's belief as to the potential of REOLYSIN as a cancer therapeutic, and other such matters are forward-looking statements and forward-looking information and 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 and forward-looking information. Such risks and uncertainties include, among others, risks related to the statistical sufficiency of patient enrollment numbers in separate patient groups, 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 statement and forward-looking information. Investors are cautioned against placing undue reliance on forward-looking statements and forward-looking information. The Company does not undertake to update these forward-looking statements and forward-looking information, except as required by applicable laws.

FOR FURTHER INFORMATION PLEASE CONTACT:

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