
**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION**
WASHINGTON, D.C. 20549

FORM 8-K

Current Report
Pursuant to Section 13 or 15(d) of the
Securities Exchange Act of 1934

November 8, 2012
Date of Report (Date of Earliest Event Reported)

IntelGenx Technologies Corp.
(Exact Name of Registrant as Specified in its Charter)

Delaware
*(State or other jurisdiction of
incorporation)*

000-31187
*(Commission File
Number)*

870299034
*(IRS Employer Identification
No.)*

6425 Abrams, Ville St- Laurent, Quebec, Canada
(Address of principal executive offices)

H4S 1X9
(Zip Code)

Registrant's telephone number, including area code: (514) 331-7440

Check the appropriate box below if the Form 8K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17CFR230.425)
 - Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17CFR 240.14a -12)
 - Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d -2(b))
 - Pre-commencement communication pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e -4(c))
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Item 7.01 Regulation FD Disclosure - News Release

IntelGenx Announces Successful Pre-NDA meeting for Anti-Migraine Film

Exhibit	Description
99.1	Press Release

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 hereunto duly authorized.

INTELGENX TECHNOLOGIES CORP.

Dated: November 8, 2012

By: /s/ *Horst Zerbe*
Horst G. Zerbe
President and Chief
Executive Officer

IntelGenx Announces Successful Pre-NDA meeting for Anti-Migraine Film

SAINT LAURENT, QUEBEC , November 8, 2012 - IntelGenx Corp. (TSX-V: IGX) (OTCQX: IGXT) ("IntelGenx", "the Company") today announced that the Company concluded a pre-New Drug Application ("NDA") meeting with the U.S. Food and Drug Administration ("FDA") related to its novel oral thin-film formulation of Rizatriptan, the active drug in Maxalt-MLT® orally disintegrating tablets.

The purpose of the meeting was to confirm the adequacy of the clinical, non-clinical and CMC data for the Company's proposed 505(b)(2) NDA submission, which the Company intends to file in the first quarter of 2013, as previously announced.

Maxalt-MLT® is a leading branded anti-migraine product manufactured by Merck & Co. According to Merck's most recent annual report, sales of Maxalt® grew 16% to \$639 million in 2011. The thin-film formulation of Rizatriptan has been developed in accordance with the co-development and commercialisation agreement with RedHill Biopharma Ltd. (TASE:RDHL) using IntelGenx' proprietary immediate release "VersaFilm" drug delivery technology.

About IntelGenx:

IntelGenx is a drug delivery company focused on the development of oral controlled-release products as well as novel rapidly disintegrating delivery systems. IntelGenx uses its unique multiple layer delivery system to provide zero-order release of active drugs in the gastrointestinal tract. IntelGenx has also developed novel delivery technologies for the rapid delivery of pharmaceutically active substances in the oral cavity based on its experience with rapidly disintegrating films. IntelGenx' development pipeline includes products for the treatment of severe depression, hypertension, erectile dysfunction, benign prostatic hyperplasia, migraine, insomnia, idiopathic pulmonary fibrosis, allergies and pain management. More information is available about the company at www.intelgenx.com.

Forward Looking Statements:

This document may contain forward-looking information about IntelGenx' operating results and business prospects that involve substantial risks and uncertainties. Statements that are not purely historical are forward-looking statements within the meaning of Section 21E of the Securities Exchange Act of 1934, as amended, and Section 27A of the Securities Act of 1933, as amended. These statements include, but are not limited to, statements about IntelGenx' plans, objectives, expectations, strategies, intentions or other characterizations of future events or circumstances and are generally identified by the words "may," "expects," "anticipates," "intends," "plans," "believes," "seeks," "estimates," "could," "would," and similar expressions. All forward looking statements are expressly qualified in their entirety by this cautionary statement. Because these forward-looking statements are subject to a number of risks and uncertainties, IntelGenx' actual results could differ materially from those expressed or implied by these forward looking statements. Factors that could cause or contribute to such differences include, but are not limited to, those discussed under the heading "Risk Factors" in IntelGenx' annual report on Form 10-K for the fiscal year ended December 31, 2011, filed with the United States Securities and Exchange Commission and available at www.sec.gov, and also filed with Canadian securities regulatory authorities and www.sedar.com. IntelGenx assumes no obligation to update any such forward-looking statements.

Each of the TSX Venture Exchange and OTCQX has neither approved nor disapproved the contents of this press release.

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