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

June 14, 2011
Date of Report (Date of Earliest Event Reported)

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

Delaware	**000-31187**	**87-0638336**
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

6425 Abrams, Ville St- Laurent, Quebec, Canada	**H4S 1X9**
(Address of principal executive offices)	(Zip Code)

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

Check the appropriate box below if the Form 8K fining 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))

Item 7.01 Regulation FD Disclosure - News Release

IntelGenx Corp. ("IntelGenx", or “the Company”), announced today that the U.S. Food and Drug Administration (“FDA”) has accepted the Company's resubmission of its antidepressant CPI-300 New Drug Application 505(b)(2) in response to the February 2010 Complete Response Letter (“CRL”) as a complete, Class 2 response. In addition, the FDA has established November 13, 2011 as its target action date under the Prescription Drug User Fee Act (“PDUFA”). CPI-300 is a novel, high strength of Bupropion Hydrochloride (HCl), the active ingredient in Wellbutrin XL®

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: June 14,, 2011

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

Exhibit 99.1

IntelGenx Announces New PDUFA Date for its Single Dose 450 mg Bupropion Hydrochloride Tablet

SAINT LAURENT, QUEBEC, June 14, 2011 – IntelGenx Corp. (TSX VENTURE: IGX) (OTCBB: IGXT) ("IntelGenx", or "the Company"), announced that the U.S. Food and Drug Administration ("FDA") has accepted the Company's resubmission of its antidepressant CPI-300 New Drug Application 505(b)(2) in response to the February 2010 Complete Response Letter ("CRL") as a complete, Class 2 response. In addition, the FDA has established November 13, 2011 as its target action date under the Prescription Drug User Fee Act ("PDUFA"). CPI-300 is a novel, high strength of Bupropion Hydrochloride (HCl), the active ingredient in Wellbutrin XL®

"We are very pleased to note that FDA considers our response to be complete," said Dr. Horst G. Zerbe, President and Chief Executive Officer of IntelGenx. "As expected, FDA considers our resubmission a class 2 response to the CRL and, as such, has issued a PDUFA target date of 6 months from the date of our resubmission. We look forward to working with the FDA during the review process and, following approval, making this proprietary product available to patients who suffer from major depressive disorder."

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' research and development pipeline includes products for the treatment of pain, hypertension, erectile dysfunction, sleep disorders, allergies and depressive disorders. 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, 2010, 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 OTC Bulletin Board has neither approved nor disapproved the contents of this press release.

CONTACT:

Dr. Horst G. Zerbe,
President and CEO
IntelGenx Corp.

T: +1 514-331-7440 (ext. 201)
F: +1 514-331-0436

horst@intelgenx.com
www.intelgenx.com