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

Date of Report (Date of earliest event reported): November 26, 2007

Idera Pharmaceuticals, Inc.

(Exact name of Registrant as Specified in Charter)

Delaware	001-31918	04-30/2298
(State or Other Jurisdiction of Incorporation)	(Commission File Number)	(IRS Employer Identification No.)
167 Sidney Street, Cambridge, Massac	nusetts	02139
(Address of Principal Executive Offi	ces)	(Zip Code)
Registrat	at's telephone number, including area code: (617) 6	79-5500
(Forme	r Name or Former Address, if Changed Since Last R	Leport)
Check the appropriate box below if the Form 8-K filing provisions:	g is intended to simultaneously satisfy the filing o	bligation of the registrant under any of the following
☐ Written communications pursuant to Rule 425 un	der the Securities Act (17 CFR 230.425)	
☐ Soliciting material pursuant to Rule 14a-12 under	the Exchange Act (17 CFR 240.14a-12)	
☐ Pre-commencement communications pursuant to l	Rule 14d-2(b) under the Exchange Act (17 CFR 240	0.14d-2(b))
☐ Pre-commencement communications pursuant to	Rule 13e-4(c) under the Exchange Act (17 CFR 240	1.13e-4(c))

Item 3.01(d). Transfer of Listing.

On November 26, 2007, Idera Pharmaceuticals, Inc. (the "Company") issued a press release announcing that it has received approval for the listing of its common stock on the NASDAQ Global Market ("NASDAQ") and that it anticipates that its common stock will commence trading on NASDAQ on December 10, 2007. The Company further anticipates that its common stock will cease trading on the American Stock Exchange ("AMEX") prior to the open of business on December 10, 2007. The Company has notified AMEX of its intention to voluntarily withdraw its common stock from listing on AMEX. The Company's transfer of listing of its common stock to NASDAQ from AMEX was authorized by the Company's Board of Directors. A copy of the press release is attached as Exhibit 99.1 and is incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits

(d) Exhibit(s)

99.1 Press Release, dated November 26, 2007

SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, as amended, the Registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Date: November 26, 2007

IDERA PHARMACEUTICALS, INC.

By: /s/ Sudhir Agrawal
Sudhir Agrawal

Chief Executive Officer



FOR IMMEDIATE RELEASE

Contacts:

Idera Pharmaceuticals, Inc. Kelly Luethje 617-679-5519 E-mail: kluethje@iderapharma.com MacDougall Biomedical Communications Chris Erdman 508-647-0209 E-mail: cerdman@macbiocom.com

Idera Pharmaceuticals Common Stock Approved for Listing on NASDAQ Global Market

Cambridge, MA, November 26, 2007 — Idera Pharmaceuticals, Inc. (AMEX: IDP) today announced that it has received notification from The NASDAQ Stock Market® that its common stock has been approved for listing on NASDAQ Global Market. The Company is currently going through all the required regulatory processes associated with the transition and expects its common stock to begin trading on NASDAQ under its new symbol IDRA on Monday, December 10, 2007. The Company's common stock will continue to trade on the American Stock Exchange under its current symbol IDP until such time.

"We believe the move to NASDAQ's electronic market will provide our stock with improved visibility and liquidity and give us more exposure to institutional investors," said Sudhir Agrawal, D. Phil., Chief Executive Officer and Chief Scientific Officer. "We thank the American Stock Exchange and our specialists, Cohen Specialists, for their support over the last few years."

About Idera Pharmaceuticals, Inc.

Idera Pharmaceuticals is a drug discovery and development company that is developing drug candidates to treat cancer and infectious, respiratory, and autoimmune diseases, and for use as vaccine adjuvants. Idera's proprietary drug candidates are designed to modulate specific TLRs, which are a family of immune system receptors. Idera's pioneering DNA chemistry expertise enables it to identify drug candidates for internal development and creates opportunities for multiple collaborative alliances. Idera's most advanced clinical candidate, IMO-2055, is an agonist of TLR9 and is currently in a Phase 2 trial in oncology and in a Phase 1/2 chemotherapy combination trial in oncology. Idera's second TLR9 agonist, IMO-2125, is currently in a Phase 1 trial for the treatment of hepatitis C virus infection. Idera is collaborating with Novartis International Pharmaceutical, Ltd. for the discovery, development, and commercialization of TLR9 agonists for the treatment of asthma and allergy indications. Idera is also collaborating with Merck & Co., Inc. for the use of Idera's TLR7, 8 and 9 agonists in combination

with Merck's therapeutic and prophylactic vaccines in the areas of oncology, infectious diseases, and Alzheimer's disease. For more information, visit www.iderapharma.com.

Forward Looking Statements

This press release contains forward-looking statements concerning Idera Pharmaceuticals, Inc. that involve a number of risks and uncertainties. For this purpose, any statements contained herein that are not statements of historical fact may be deemed to be forward-looking statements. Without limiting the foregoing, the words "believes," "anticipates," "plans," "expects," "estimates," "intends," "should," "could," "will," "may," and similar expressions are intended to identify forward-looking statements. There are a number of important factors that could cause Idera's actual results to differ materially from those indicated by such forward-looking statements, including whether products based on Idera's technology will advance into or through the clinical trial process on a timely basis or at all and receive approval from the United States Food and Drug Administration or equivalent foreign regulatory agencies; whether, if the Company's products receive approval, they will be successfully distributed and marketed; whether the Company's collaborations with Novartis and Merck will be successful; whether Idera's cash resources will be sufficient to fund product development and clinical trials; and such other important factors as are set forth under the caption "Risk Factors" in Idera's Quarterly Report on Form 10-Q filed on November 13, 2007, which important factors are incorporated herein by reference. Idera disclaims any intention or obligation to update any forward-looking statements.

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