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): December 1, 2010

ORAMED PHARMACEUTICALS INC.

(Exact name of registrant as specified in its charter)

Nevada	**000-50298**	**98-0376008**
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

Hi-Tech Park 2/5 Givat Ram
PO Box 39098
Jerusalem, Israel 91390
(Address of principal executive offices and zip code)

Registrant's telephone number, including area code: 972-2-566-0001

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

o Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

o Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

o Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

o Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

ITEM 7.01 REGULATION FD DISCLOSURE

On December 1, 2010, Oramed Pharmaceuticals Inc. issued a press release announcing a publication in the Journal of Diabetes Science and Technology's November issue describing evaluation of the company's glucagon-like peptide- (GLP-1) analog formulation (ORMD-0901) in regulating glucose excursions in animal models.

A copy of the press release is attached to this Current Report on Form 8-K as Exhibit 99.1 and is incorporated by reference herein.

ITEM 9.01 FINANCIAL STATEMENTS AND EXHIBITS.

(d) Exhibits

Exhibit Number	Description
99.1	Press Release dated December 1, 2010.

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.

ORAMED PHARMACEUTICALS INC.

Dated: December 1, 2010

By: /s/ Nadav Kidron
Nadav Kidron
President, CEO and Director

Exhibit Index

Exhibit Number	Description
99.1	Press Release dated December 1, 2010.

Oramed Pharmaceuticals Announces Publication in the Journal of Diabetes Science and Technology

Results describe preclinical trials demonstrating retained GLP-1 analog activity when delivered with Oramed's oral drug delivery formulations

JERUSALEM, Israel – December 1, 2010 – Oramed Pharmaceuticals Inc. (OTCBB: ORMP.OB; www.oramed.com), a developer of oral drug delivery solutions, announces a publication in the Journal of Diabetes Science and Technology's November issue describing evaluation of the company's glucagon-like peptide- (GLP-1) analog formulation (ORMD-0901) in regulating glucose excursions in animal models.

GLP-1 and its analogs harbor significant therapeutic potential for management of Type 2 diabetes mellitus through their broad physiological impact on glucose regulation-related mechanisms. However, the naturally secreted GLP-1 is degraded within minutes and in cases of impaired release or activity, is typically substituted by exogenously supplied, long-lived analogues. To date, these drugs are only available in injectable forms, leading to systemic effects upon administration as well as unsatisfactory patient compliance and adherence.

With the objective of offering a safer, more practical and physiologically relevant alternative, Oramed has applied its oral drug delivery platform to the GLP-1 analog family. High sugar-content meals were delivered to animals in the presence or absence of ORMD-0901, followed by close monitoring of blood glucose levels. Specific formulations were found to effectively blunt expected glucose surges in both ORMD-0901-treated porcines and canines. The two animal models used for testing the performance of ORMD-0901, indicated retained drug activity when delivered using Oramed's oral drug delivery platform.

Oramed has already embarked on clinical testing of its ORMD-0901 product to assess the formulation's safety and efficacy in healthy individuals. Further testing and development are expected to significantly contribute to current diabetes control options and success rates.

In addition to an oral GLP1 analog capsule, the Company’s flagship product is an oral insulin capsule. The Company is currently working towards submission of an IND to the FDA for a Phase 2 clinical trial.

To view the article abstract please click on the link below: http://www.journalofdst.org/November2010/PDF/Abstract/VOL-4-6-ORG20-ELDOR-ABSTRACT.pdf

About Oramed Pharmaceuticals

Oramed Pharmaceuticals is a technology pioneer in the field of oral delivery solutions for drugs and vaccines presently delivered via injection. Oramed is seeking to revolutionize the treatment of diabetes through its patented flagship product, an orally ingestible insulin capsule currently in Phase 2 clinical trials. Established in 2006, Oramed's technology is based on over 25 years of research performed by top research scientists at Jerusalem's Hadassah Medical Center. The Company's corporate and R&D headquarters are based in Jerusalem.
For more information, please visit www.oramed.com

Safe Harbor Statement

Some of the statements contained in this press release are forward-looking statements which involve known and unknown risks, uncertainties and other factors which may cause the actual results, performance or achievements of the company, or industry results, to be materially different from any future results, performance or achievements expressed or implied by such forward looking statements, including the risks and uncertainties related to the progress, timing, cost, and results of clinical trials and product development programs; difficulties or delays in obtaining regulatory approval for our product candidates; competition from other pharmaceutical or biotechnology companies; and the company's ability to obtain additional funding required to conduct its research, development and commercialization activities. Please refer to the company's filings with the Securities and Exchange Commission for a comprehensive list of risk factors that could cause actual results, performance or achievements of the company to differ materially from those expressed or implied in such forward looking statements. The company undertakes no obligation to update or revise any forward-looking statements.

Company and Investor Relation Contacts:
Oramed Pharmaceuticals
Tara Horn
Cell: + 972-54-334-318
Office: + 972-2-566-0001
Email: tara@oramed.com