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): June 29, 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 June 28, 2010, Oramed Pharmaceuticals Inc. issued a press release announcing that its executive management is participating in the 2010 Scientific Sessions of the American Diabetes Association currently being held in Orlando, Florida. Dr. Miriam Kidron, Chief Scientific Officer, is scheduled to present results of two of the company's recently completed clinical trials between 12:00 – 2:00 PM ET on Monday, June 28, 2010 in Poster Hall C at the Orange County Convention Center.

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 June 28, 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: June 29, 2010

By: /s/ Nadav Kidron

Nadav Kidron
President, CEO and Director

Exhibit Index

Exhibit Number	Description
99.1	Press Release dated June 28, 2010.

Oramed Pharmaceuticals to present at the 70th Scientific Sessions of the American Diabetes Association

Results of first exposure of Type I diabetes patients to Oramed's oral insulin and of a first-in-humans study with its oral GLP-1 analog

Jerusalem, IL - June 28, 2010 - Oramed Pharmaceuticals Inc. (OTCBB: ORMP) announced today that its executive management is participating in the 2010 Scientific Sessions of the American Diabetes Association currently being held in Orlando, Florida. Dr. Miriam Kidron, Chief Scientific Officer, is scheduled to present results of two of the company's recently completed clinical trials between 12:00 – 2:00 PM ET on Monday, June 28, 2010 in Poster Hall C at the Orange County Convention Center.

One presentation will provide an overview of the results of a Phase IIa study testing the company's flagship ORMD-0801 oral insulin capsule in Type I diabetes patients when administered just before (10-90 minutes) a standard meal. This first exposure of Oramed's oral insulin established the drug's safety and efficacy in Type I diabetics. ORMD-0801 led to a significant increase in insulin levels in 61% of the treatment sessions, irrespective of the time lapse between ORMD-0801 administration and mealtime. The oral insulin preparation effectively kept glucose levels in check in all sessions. These data suggest both tolerance of Oramed's platform to preprandial dosing regimens and its therapeutic potential in regulating Type I diabetes.

In addition, Dr. Kidron will present a summary of Oramed's first-in-humans clinical trial testing the safety and efficacy of ORMD-0901, an encapsulated oral GLP-1 analog formulation. The study monitored the responses of healthy males to a single dose delivered 60 minutes before a glucose load. ORMD-0901 was well tolerated by all subjects and demonstrated physiological activity, as extrapolated from ensuing subject insulin levels when compared to those observed after treatment with placebo.

The poster entitled "Oral Insulin: Type I Diabetes (T1DM) Patient Response Upon Preprandial Administration" will be featured in a guided tour in the Poster Hall. The tour title is "Innovations in Novel Insulin Preparations" and is scheduled for Monday June 28 at 12:00-1:00 PM.

To view the posters being presented, please visit: www.oramed.com

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 http://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.

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