
**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): May 11, 2016

GALECTIN THERAPEUTICS INC.

(Exact name of registrant as specified in its charter)

Nevada
(State or Other Jurisdiction
of Incorporation)

001-31791
(Commission
File Number)

04-3562325
(IRS Employer
Identification No.)

**4960 PEACHTREE INDUSTRIAL BOULEVARD, Ste 240
NORCROSS, GA 30071**
(Address of principal executive office) (zip code)

Registrant's telephone number, including area code: (678) 620-3186

N/A
(Former name or former address, if changed since last report)

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):

- Written communications pursuant to Rule 425 under 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 Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
 - Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))
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SECTION 7 – REGULATION FD

Item 7.01 Regulation FD Disclosure.

On May 11, 2016, Galectin Therapeutics Inc. (the “Company”) issued the press release attached hereto as Exhibit 99.1.

The information in this report is being furnished pursuant to this Item 7.01 and shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934 (the “Exchange Act”) or otherwise subject to the liabilities of that section, and it shall not be deemed incorporated by reference in any filing under the Securities Act of 1933 or under the Exchange Act, whether made before or after the date hereof, except as expressly set forth by specific reference in such filing to this report.

SECTION 9 – FINANCIAL STATEMENTS AND EXHIBITS

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

<u>Exhibit Number</u>	<u>Description</u>
99.1	Press release

SIGNATURES

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

Galectin Therapeutics Inc.

Date: May 11, 2016

By: /s/ Jack W. Callicutt
Jack W. Callicutt
Chief Financial Officer



**Galectin Therapeutics Announces Complete Enrollment in its Phase 2 Trial
with GR-MD-02 in NASH Patients with Advanced Fibrosis**

Top line results of the NASH-FX trial to be reported in September 2016

NORCROSS, GA. (May 11, 2016) – **Galectin Therapeutics Inc. (NASDAQ: GALT)**, the leading developer of therapeutics that target galectin proteins to treat fibrosis and cancer, announces the completion of enrollment in its Phase 2 clinical trial with GR-MD-02 in patients with non-alcoholic steatohepatitis (NASH) with advanced fibrosis (stage 3) (the NASH-FX trial). The Company expects to report the top line results of this trial by the end of September 2016, as previously planned.

“We are pleased that enrollment in our NASH-FX trial was completed right on schedule,” said Peter G. Traber, M.D., Galectin’s president, chief executive officer and chief medical officer. “This is one of two Phase 2 trials we are conducting in subjects with NASH, and is designed to assess the efficacy of our lead compound GR-MD-02 in patients with NASH with advanced fibrosis (stage 3). Since the diagnosis of NASH and the monitoring of the disease are hampered by the need for liver biopsy, in addition to providing proof-of-concept on the efficacy of GR-MD-02, this study explores the utility of three non-invasive measures of fibrosis which may be useful in the design and execution of later stage clinical studies and perhaps have relevance in commercial clinical settings.”

The NASH-FX trial has enrolled a total of 30 liver biopsy-confirmed NASH patients with advanced fibrosis (stage 3) -15 patients to receive 8 mg/kg of GR-MD-02 and 15 patients to receive placebo every other week for 16 weeks, for a total of nine doses. The effect of GR-MD-02 on liver fibrosis will be assessed by three independent non-invasive tests following the treatment period. The primary endpoint will be an assessment of fibrosis using multi-parametric magnetic resonance imaging (LiverMultiScan®), which is a validated and proprietary MRI protocol developed by Perspectum Diagnostics. Secondary endpoints will evaluate liver stiffness, which correlates to the degree of liver fibrosis, as assessed by magnetic resonance-elastography and by FibroScan®. More information on the NASH-FX trial may be found in a post on Dr. Traber’s blog, [CEO Perspectives](#) and at www.clinicaltrials.gov.

This single-site study is being conducted by Stephen A. Harrison, M.D., FACP, FAASLD, Colonel, Medical Corps U.S.A., Director, Medical Education, Associate Dean, San Antonio Uniformed Services Health Education Consortium, Professor of Medicine, Uniformed Services University of the Health Sciences and Consultant to The Army Surgeon General for Gastroenterological Diseases, San Antonio Military Medical Center. Dr. Harrison is also the co-lead investigator for Galectin’s Phase 2 NASH-CX trial, which is studying two different doses of GR-MD-02 against placebo in 156 patients with NASH with cirrhosis (the NASH-CX trial).

About GR-MD-02

GR-MD-02 is a complex carbohydrate drug that targets galectin-3, a critical protein in the pathogenesis of fatty liver disease and fibrosis. Galectin-3 plays a major role in diseases that involve scarring of organs including fibrotic disorders of the liver, lung, kidney, heart and

vascular system. The drug binds to galectin proteins and disrupts their function. Preclinical data in animals have shown that GR-MD-02 has robust treatment effects in reversing liver fibrosis and cirrhosis.

About Fatty Liver Disease with Advanced Fibrosis and Cirrhosis

Non-alcoholic steatohepatitis (NASH), also known as fatty liver disease, has become a common disease of the liver with the rise in obesity rates. NASH is estimated to affect up to 28 million people in the U.S. Fatty liver disease is characterized by the presence of fat in the liver along with inflammation and damage in people who consume little or no alcohol. Over time, patients with fatty liver disease can develop fibrosis, or scarring of the liver, and it is estimated that as many as 1-2 million individuals in the U.S. have cirrhosis, a severe liver disease for which liver transplant is the only treatment available. Approximately 6,300 liver transplants are performed annually in the U.S. There are no drug therapies approved for the treatment of NASH or liver fibrosis. A recent analyst estimate indicated that by 2025 the worldwide market for NASH treatments could approach \$35 billion.

About Galectin Therapeutics

Galectin Therapeutics is developing promising carbohydrate-based therapies for the treatment of fibrotic liver disease and cancer based on the Company's unique understanding of galectin proteins, which are key mediators of biologic function. Galectin seeks to leverage extensive scientific and development expertise as well as established relationships with external sources to achieve cost-effective and efficient development. The Company is pursuing a development pathway to clinical enhancement and commercialization for its lead compounds in liver fibrosis and cancer. Additional information is available at www.galectintherapeutics.com.

Forward Looking Statements

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. These statements relate to future events or future financial performance, and use words such as "may," "estimate," "could," "expect" and others. They are based on management's current expectations and are subject to factors and uncertainties that could cause actual results to differ materially from those described in the statements. These statements include those regarding the hope that Galectin's development program for GR-MD-02 will lead to the first therapy for the treatment of fatty liver disease with advanced fibrosis and/or cirrhosis and/or moderate to severe psoriasis. Factors that could cause actual performance to differ materially from those discussed in the forward-looking statements include, among others, that Galectin may not be successful in developing effective treatments and/or obtaining the requisite approvals for the use of GR-MD-02 or any of its other drugs in development. The Company's current clinical trial and any future clinical studies may not produce positive results in a timely fashion, if at all, and could prove time consuming and costly. Plans regarding development, approval and marketing of any of Galectin's drugs are subject to change at any time based on the changing needs of the Company as determined by management and regulatory agencies. Regardless of the results of any of its development programs, Galectin may be unsuccessful in developing partnerships with other companies or raising additional capital that would allow it to further develop and/or fund any studies or trials. Galectin has incurred operating losses since inception, and its ability to successfully develop and market drugs may be impacted by its ability to manage costs and finance continuing operations. For a discussion of additional factors impacting Galectin's business, see the Company's Annual Report on Form 10-K for the year ended December 31, 2015, and subsequent filings with the SEC. You should not place undue reliance on forward-looking statements. Although subsequent events may cause its views to change, management disclaims any obligation to update forward-looking statements.

Contacts:
Jack Callicutt, Chief Financial Officer
(678) 620-3186
ir@galectintherapeutics.com.

LHA
Kim Golodetz
(212) 838-3777
kgolodetz@lhai.com

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