

**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): March 23, 2016

BioSig Technologies, Inc.
(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction
of incorporation)

000-55473
(Commission File Number)

26-433375
(IRS Employer
Identification No.)

8441 Wayzata Blvd., Suite 240
Minneapolis, Minnesota
(Address of principal executive offices)

55426
(Zip Code)

Registrant's telephone number, including area code: **(763) 999-7330**

(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:

- 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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Item 8.01 Other Events.

On March 23, 2016, BioSig Technologies, Inc. (the “Company”) issued a press release announcing the establishment of an advanced research program with Dr. Samuel Asirvatham at the Mayo Clinic in Rochester, Minnesota. A copy of the press release is filed as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated by reference herein.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

<u>Exhibit Number</u>	<u>Description</u>
99.1	Press release dated March 23, 2016

SIGNATURES

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.

BIOSIG TECHNOLOGIES, INC.

Date: March 23, 2016

By: /s/ Kenneth Londoner
Name: Kenneth Londoner
Title: Executive Chairman



BioSig Technologies Announces Advanced Research Program with Mayo Clinic

MINNEAPOLIS, MN, March 23, 2016 – BioSigä Technologies, Inc. (OTCQB:BSGM) today announced the establishment of an advanced research program with Dr. Samuel Asirvatham at the Mayo Clinic in Rochester, Minnesota.

This program has been designed to build upon the initial studies conducted at Mayo Clinic in March, June and November 2015 through a tripling of investment to fully characterize and develop novel features discovered during this prior preclinical work. These features have the potential to make significant impact in the treatment of complex arrhythmias.

“We are extremely pleased to continue our relationship with Dr. Asirvatham and his colleagues at the Mayo Clinic.” said Greg Cash, President and CEO of BioSig Technologies. “The previous studies confirmed the potential of PURE EP to improve the clarity of cardiac signals while minimizing electrical noise in the cardiac electrophysiology laboratory.”

About BioSig Technologies

BioSig Technologies is a medical device company that is developing a proprietary technology platform designed to improve the \$3 billion EP marketplace¹ (www.biosigtech.com). Led by a proven management team and a veteran, independent Board of Directors, Minneapolis-based BioSig Technologies is preparing to commercialize its PURE EP System.

The PURE EP System is a surface electrocardiogram and intracardiac multichannel signal acquisition and analysis system designed to assist electrophysiologists in making clinical decisions in real-time by acquiring and displaying high-fidelity cardiac signal recordings and providing clarity of data which may be used to guide the electrophysiologists in identifying ablation targets - areas of tissue to treat that otherwise create a heart rhythm disturbance (arrhythmia).

Analysts forecast the global market for EP devices will grow at a 12.1 percent compound annual growth rate, from \$2.5 billion in 2012 to \$5.5 billion by 2019¹, making it one of the fastest growing medical device segments. Just in the US, the number of Atrial Fibrillation (AF) and Ventricular Tachycardia (VT) arrhythmia ablations is forecast to grow at 10.5 percent from 2012 to 2017².

BioSig intends to seek FDA 510(k) clearance for the PURE EP System. The Company has achieved proof of concept validation through UCLA labs, and has performed pre-clinical studies at the Mayo Clinic in Minnesota. The Company is collaborating with several of the nation's most prestigious cardiac arrhythmia centers including Texas Cardiac Arrhythmia Institute, UCLA Cardiac Arrhythmia Center, and Mayo Clinic.

- (1) Electrophysiology Devices Market - Global Industry Analysis, Size, Share, Growth, Trends and Forecast, 2013 – 2019
- (2) HRI 2013 "Global Opportunities in Medical Devices & Diagnostics" report; triangulation of multiple sources; *AF includes left atrial tachycardia, left WPW, left atrial flutter.

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