April 30, 2019



## **BioSig To Participate at Heart Rhythm Society's Scientific Sessions 2019**

Santa Monica, CA, April 30, 2019 (GLOBE NEWSWIRE) -- BioSig Technologies, Inc. (NASDAQ: BSGM), a medical device company developing a proprietary biomedical signal processing platform designed to address unmet needs for the electrophysiology (EP) marketplace, today announced that the Company will be exhibiting at the Heart Rhythm Society's 40<sup>th</sup> Annual Scientific Sessions on May 8-11, 2019 at Moscone Center in San Francisco, CA.

During the event BioSig will be presenting at booth 2056 in the South Hall of Moscone Center.

Previously, BioSig announced that it successfully conducted its first patient cases using PURE EP(tm) System at the <u>Texas Cardiac Arrhythmia Institute</u> in Austin, TX in February 2019 and <u>Greenville Memorial Hospital</u> in Greenville, SC in April 2019. These initial experiences suggested improved cardiac signal detection and fidelity.

The event will convene some of the finest clinicians, scientists, researchers and innovators in the field of cardiac pacing and electrophysiology. More than 800 of the world's most noted experts in cardiac rhythm management are expected to serve as faculty for more than 200 educational sessions covering topic areas such as pacing, defibrillation, clinical arrhythmia management, ablation, pharmacology, genetics, basic science, health policy, and more.

The PURE EP(tm) System is a computerized system intended for acquiring, digitizing, amplifying, filtering, measuring and calculating, displaying, recording and storing of electrocardiographic and intracardiac signals for patients undergoing electrophysiology (EP) procedures in an EP laboratory. The PURE EP(tm) System aims to minimize noise and artifacts and acquire high-fidelity cardiac signals. Improving fidelity of acquired cardiac signals may potentially increase the diagnostic value of these signals, thereby possibly improving accuracy and efficiency of the EP studies and related procedures. The results of pre-clinical studies have been <u>published</u> in a number of journals, including <u>The Journal of Innovations in Cardiac Rhythm Management</u> and the <u>Journal of the American College of Cardiology</u>, a manuscript, which was included in the top 5 most read, discussed and shared articles in 2016.

## About Heart Rhythm Society's Scientific Sessions

The Heart Rhythm Society's Annual Scientific Sessions is a Heart Rhythm Society program. Heart Rhythm Society (HRS) is a 501(c)(3) international nonprofit organization with a mission to improve the care of patients by promoting research, education, and optimal health care policies and standards. Founded in 1979, HRS is a leading resource on cardiac pacing and electrophysiology. This specialty organization represents medical, allied health, and science professionals from more than 70 countries who specialize in cardiac rhythm disorders.

## About BioSig Technologies

BioSig Technologies is a medical technology company developing a proprietary biomedical signal processing platform designed to improve the electrophysiology (EP) marketplace (www.biosig.com). Led by a proven management team and a veteran, independent Board of Directors, Los Angeles-based BioSig Technologies is preparing to commercialize its PURE EP<sup>™</sup> System. The technology has been developed to address an unmet need in a large and growing market.

The Company's first product, PURE EP<sup>™</sup> System, is a novel cardiac signal acquisition and display system which is engineered to assist electrophysiologists in clinical decision-making during procedures to diagnose and treat patients with abnormal heart rates and rhythms. BioSig's main goal is to deliver technology to improve upon catheter ablation treatments for the prevalent and potentially deadly arrhythmias, Atrial Fibrillation and Ventricular Tachycardia. BioSig has partnered with Minnetronix on technology development and received FDA 510(k) clearance for the PURE EP<sup>™</sup> System in August 2018.

## **Forward-looking Statements**

This press release contains "forward-looking statements." Such statements may be preceded by the words "intends," "may," "will," "plans," "expects," "anticipates," "projects," "predicts," "estimates," "aims," "believes," "hopes," "potential" or similar words. Forwardlooking statements are not guarantees of future performance, are based on certain assumptions and are subject to various known and unknown risks and uncertainties, many of which are beyond the Company's control, and cannot be predicted or quantified and consequently, actual results may differ materially from those expressed or implied by such forward-looking statements. Such risks and uncertainties include, without limitation, risks and uncertainties associated with (i) our inability to manufacture our products and product candidates on a commercial scale on our own, or in collaboration with third parties; (ii) difficulties in obtaining financing on commercially reasonable terms; (iii) changes in the size and nature of our competition; (iv) loss of one or more key executives or scientists; and (v) difficulties in securing regulatory approval to market our products and product candidates. More detailed information about the Company and the risk factors that may affect the realization of forward-looking statements is set forth in the Company's filings with the Securities and Exchange Commission (SEC), including the Company's Annual Report on Form 10-K and its Quarterly Reports on Form 10-Q. Investors and security holders are urged to read these documents free of charge on the SEC's website at http://www.sec.gov. The Company assumes no obligation to publicly update or revise its forward-looking statements as a result of new information, future events or otherwise.

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Source: BioSig Technologies, Inc.