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BioSig Technologies Begins Trading on Nasdaq

Santa Monica, CA, Sept. 21, 2018 (GLOBE NEWSWIRE) -- BioSig Technologies, Inc. (NASDAQ:BSGM) a medical device company developing a proprietary biomedical signal processing platform designed to address an unmet technology need for the $4.6 billion electrophysiology (EP) marketplace, today announced that shares of its common stock have begun officially trading on the Nasdaq Capital Market, under the ticker “BSGM”.

BioSig’s listing on the Nasdaq is the Company’s most recent operational achievement. In August 2018, the Company received 510(k) clearance from the FDA for their patented advanced signal acquisition and processing technology, PURE EP(tm) System. The non-invasive technology is a computerized system intended to enhance electrocardiographic and intracardiac signals for patients undergoing electrophysiology (EP) procedures in an EP laboratory for the treatments of arrhythmia, or irregular heartbeat. The Company completed 13 pre-clinical studies at Mayo Clinic and Mount Sinai to date and received first units from its contract manufacturer, Minnetronix, in September 2018. BioSig recently made an announcement about their ongoing progress with effective physician engagement and continues to add experienced industry hires to its clinical and commercial teams.

“The Nasdaq stock exchange is home to many of the world’s most innovative medical technology and healthcare companies and is a natural fit for BioSig,” stated Mr. Kenneth Londoner, Founder, Chairman and CEO of BioSig Technologies. “Our listing on the Nasdaq exchange is a milestone and reflects the significant progress we have made in strengthening our corporate governance and expanding our industry footprint.”

Analysts forecast the global market for EP devices will grow at a 10.6 percent compound annual growth rate to more than $8.5 billion by 2024*, making it one of the fastest growing medical device segments. In the United Stated alone, the number of Atrial Fibrillation (AF) and Ventricular Tachycardia (VT) arrhythmia ablations is forecast to grow at 11 percent from 2015 to 2020**.

*EP device growth forecast from *Electrophysiology Market by Devices Analysis*, Market Research Engine, July 2017.
**Cardiac ablations growth forecast from Global Opportunities in *Medical Devices & Diagnostics*, Health Research International, 2016

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.biosigtech.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™ System. The technology has been developed to address an unmet need in a large and growing market.

The Company's first product, PURE EP™ 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 has received FDA 510(k) clearance for the PURE EP™ 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. Forward-looking 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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