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# BioSig to Present at the 25th Annual International AF Symposium

**PURE EP(tm) System to be highlighted in a Spotlight session by Andrea Natale, M.D., Texas Cardiac Arrhythmia Institute**

Westport, CT, Jan. 21, 2020 (GLOBE NEWSWIRE) -- BioSig Technologies, Inc. (NASDAQ: BSGM) ("BioSig" or the "Company"), a medical technology company developing a proprietary biomedical signal processing platform designed to improve signal fidelity and uncover the full range of ECG and intra-cardiac signals, today announced that the Company will be exhibiting at the *25<sup>th</sup> Annual International AF Symposium* on January 23-25, 2020 at Gaylord National Hotel in Washington, DC.

Clinical observations collected with BioSig's PURE EP(tm) System will be presented by Andrea Natale, M.D., Executive Medical Director, Texas Cardiac Arrhythmia Institute at St. David's Medical Center, during *Spotlight Session: Early Stage and Emerging New Technologies and Drugs in Cardiac EP* on January 23, 2020.

BioSig will be represented by its executive, clinical, marketing and commercial teams, and will be performing product demonstrations and hold physician sessions at the Company's booth, as well as feature a new [product video](#) about PURE EP(tm) System.

Previously, BioSig presented initial clinical data at [Heart Rhythm Society's Scientific Sessions 2019](#), [Venice Arrhythmias 2019](#) and [14<sup>th</sup> Annual International Symposium on Ventricular Arrhythmias: Pathophysiology & Therapy 2019](#).

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.

## About 25<sup>th</sup> Annual International AF Symposium

This intensive, highly focused three-day symposium brings together the world's leading medical scientists to share in a highly interactive environment the most recent advances in the field of atrial fibrillation. The primary objective of the meeting is to provide attendees with a thorough and practical course on the current state of the art in the field of atrial fibrillation in a scholarly and collegial atmosphere, as well as an opportunity to network with colleagues and faculty between sessions. More information about the event on [www.afsymposium.com](http://www.afsymposium.com).

## **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](http://www.biosig.com)).

The Company's first product, 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.

## **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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