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UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
WASHINGTON, D.C. 20549

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**FORM 8-K**

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**CURRENT REPORT**  
**Pursuant to Section 13 or 15(d) of the**  
**Securities Exchange Act of 1934**

Date of Report (Date of earliest event reported): July 10, 2020

**BioSig Technologies, Inc.**

(Exact name of registrant as specified in its charter)

**Delaware**  
(State or other jurisdiction  
of incorporation)

**001-38659**  
(Commission File Number)

**26-4333375**  
(IRS Employer  
Identification No.)

**54 Wilton Road, 2nd Floor**  
**Westport, Connecticut**  
(Address of principal executive offices)

**06880**  
(Zip Code)

**(203) 409-5444**  
(Registrant's telephone number, including area code)

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:

- ☐ 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))

Securities registered pursuant to Section 12(b) of the Act:

<u>Title of each class</u>	<u>Trading Symbol(s)</u>	<u>Name of exchange on which registered</u>
Common Stock, par value \$0.001 per share	BSGM	The NASDAQ Capital Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ☐

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

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**Item 8.01 Other Events**

On July 10, 2020, BioSig Technologies, Inc. (the “Company”) announced that it has submitted its application to Systems for Award Management (SAM). A copy of this press release is filed as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated by reference herein.

On July 13, 2020, the Company and its majority-owned subsidiary, ViralClear Pharmaceuticals, Inc. (“ViralClear”), today announced the enrollment of adult patients for ViralClear’s Phase II trial for merimepodib, a broad- spectrum, orally administered antiviral drug candidate for the treatment of COVID-19, at four key trial sites. A copy of this press release is filed as Exhibit 99.2 to this Current Report on Form 8-K and is incorporated by reference herein.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits

<b>Exhibit Number</b>	<b>Description</b>
99.1	<a href="#">Press Release, dated July 10, 2020</a>
99.2	<a href="#">Press Release, dated July 13, 2020</a>

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## 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.

Date: July 16, 2020

By: /s/ Kenneth L. Londoner

Name: Kenneth L. Londoner

Title: Executive Chairman



## BioSig Submits Application to Systems for Award Management (SAM) with U.S. Government

Registration would provide Company access to business with the federal government, including contract and grant opportunities

Westport, CT, July 10, 2020 (GLOBE NEWSWIRE) -- BioSig Technologies, Inc. (BSGM) ("BioSig" or the "Company"), a medical technology company commercializing 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 has submitted its application to Systems for Award Management (SAM).

Upon completion of its registration with SAM, BioSig, along with its majority owned subsidiary ViralClear, will become eligible to conduct business with the federal government. This includes the ability to apply and be awarded government contracts, assistance and grant programs. Additionally, government agencies will be able to search for BioSig when contracts that fit the Company become available.

BioSig expects to receive its Commercial and Government Entity (CAGE) code, required to begin business operations with the federal government, within the next two weeks.

### About SAM

The System for Award Management (SAM) is a federal government owned and operated free website that consolidates the capabilities in Central Contractor Registration (CCR)/FedReg, Online Representations and Certifications Applications (ORCA) and the Excluded Parties List System (EPLS). Both current and potential government vendors are required to register in SAM in order to be awarded contracts by the government. SAM allows government agencies and contractors to search for companies based on ability, size, location, experience, ownership, and more. SAM allows users to search for firms certified by the SBA under the 8(a) Development and Hubzone Programs. SAM validates the vendor's information and electronically shares the secure and encrypted data with the federal agencies' finance offices to facilitate paperless payments through electronic funds transfer (EFT). Additionally, SAM shares the data with government procurement and electronic business systems.

### About BioSig Technologies

BioSig Technologies is a medical technology company commercializing a proprietary biomedical signal processing platform designed to improve signal fidelity and uncover the full range of ECG and intra-cardiac signals ([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) the geographic, social and economic impact of COVID-19 on our ability to conduct our business and raise capital in the future when needed, (ii) our inability to manufacture our products and product candidates on a commercial scale on our own, or in collaboration with third parties; (iii) difficulties in obtaining financing on commercially reasonable terms; (iv) changes in the size and nature of our competition; (v) loss of one or more key executives or scientists; and (vi) 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.

Andrew Ballou  
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## Phase II Human Trial of ViralClear's Anti-Viral for SARS-CoV-2 Coronavirus Underway at 4 Key Trial Sites

Adult patients hospitalized with COVID-19 are enrolled in Austin, TX, Rochester, MN, Jacksonville, FL, and Scottsdale, AZ

Westport, CT, July 13, 2020 (GLOBE NEWSWIRE) -- BioSig Technologies, Inc. (Nasdaq: BSGM) ("BioSig" or the "Company") and its subsidiary, ViralClear Pharmaceuticals, Inc., today announced the enrollment of adult patients for its Phase II trial for merimepodib, a broad- spectrum, orally administered antiviral drug candidate for the treatment of COVID-19, at four key trial sites. The trial sites are located in Austin, TX, Rochester, MN, Jacksonville, FL, and Scottsdale, AZ.

Merimepodib is intended to be orally administered and has demonstrated broad-spectrum in-vitro antiviral activity, including strong activity against SARS-CoV-2 in cell cultures.

"Three of our four trial sites where our partners are enrolling patients are in the recognized hot spots of Texas, Arizona, and Florida. The patients in this trial are hospitalized and requiring oxygen; therefore, providing effective treatment is of the utmost importance to all," commented Kenneth L. Londoner, Chairman and CEO of BioSig Technologies, Inc., the majority shareholder of ViralClear Pharmaceuticals, Inc. "Arizona, Florida, and Texas are unfortunately seeing spikes in COVID-19 cases resulting in hospitals reaching near capacity. We would like to thank our clinical partners for the steady progress with the trial and expect to report the trial data later this summer".

The Phase II randomized, double-blind, placebo-controlled study is designed to enroll adult patients with advanced Coronavirus Disease 2019 (COVID-19). A description of this clinical trial can be accessed via [www.clinicaltrials.gov](http://www.clinicaltrials.gov).

Preclinical in vitro laboratory studies performed by the Galveston National Laboratory at The University of Texas Medical Branch demonstrated that merimepodib, provided in combination with remdesivir, showed a reduction in SARS-CoV-2 replication to undetectable levels. Peer-reviewed publication of these findings can be found at F1000 Research: <https://f1000research.com/articles/9-361>

### About BioSig Technologies

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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.

### About ViralClear Pharmaceuticals and Merimepodib (MMPD)

BioSig's subsidiary, ViralClear Pharmaceuticals, Inc., is seeking to develop a novel pharmaceutical called merimepodib to treat patients with COVID-19. Merimepodib is intended to be orally administered and has demonstrated broad-spectrum in vitro anti-viral activity, including strong activity against SARS-CoV-2 in cell cultures. Merimepodib was previously in development as a treatment for chronic hepatitis C and psoriasis by Vertex Pharmaceuticals Incorporated (Vertex), with 12 clinical trials (7 in phase 1 and 5 in phase 2) with over 400 subjects and patients and an extensive preclinical safety package was completed. A manuscript titled, "The IMPDH inhibitor merimepodib provided in combination with the adenosine analogue remdesivir reduces SARS-CoV-2 replication to undetectable levels in vitro", was submitted to an online peer-reviewed life sciences journal. This manuscript is authored by Natalya Bukreyeva, Rachel A. Sattler, Emily K. Mantlo, John T. Manning, Cheng Huang and Slobodan Paessler of the UTMB Galveston National Laboratory and Dr. Jerome Zeldis of ViralClear Pharmaceuticals, Inc. ("ViralClear") as a corresponding author. This article highlights pre-clinical data generated under contract with Galveston National Laboratory at The University of Texas Medical Branch.

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## 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) the geographic, social and economic impact of COVID-19 on our ability to conduct our business and raise capital in the future when needed, (ii) our inability to manufacture our products and product candidates on a commercial scale on our own, or in collaboration with third parties; (iii) difficulties in obtaining financing on commercially reasonable terms; (iv) changes in the size and nature of our competition; (v) loss of one or more key executives or scientists; and (vi) 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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