

---

---

**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): August 5, 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:

<b>Title of each class</b>	<b>Trading Symbol(s)</b>	<b>Name of exchange on which registered</b>
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. ☐

---

**Item 7.01 Regulation FD Disclosure.**

On August 5, 2020, BioSig Technologies, Inc. (the “Company”) issued a press release, attached hereto as Exhibit 99.1, announcing that Dr. Michael J. Sofia has joined the Scientific Advisory Board (“SAB”) of ViralClear Pharmaceuticals, Inc. (“ViralClear”), the Company’s majority-owned subsidiary. The Company undertakes no obligation to update, supplement or amend the materials attached hereto as Exhibit 99.1.

On August 6, 2020, the Company issued a press release, attached hereto as Exhibit 99.2, announcing that Professor Angus Dalgleish has joined ViralClear’s SAB. The Company undertakes no obligation to update, supplement or amend the materials attached hereto as Exhibit 99.2.

In accordance with General Instruction B.2 of Form 8-K, the information in this Item 7.01 of this Current Report on Form 8-K, including Exhibits 99.1 and 99.2, shall not be deemed “filed” for the purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Exchange Act or the Securities Act of 1933, as amended, except as shall be expressly set forth by reference in such a filing. Furthermore, the furnishing of information under Item 7.01 of this Current Report on Form 8-K is not intended to constitute a determination by the Company that the information contained herein, including the exhibits hereto, is material or that the dissemination of such information is required by Regulation FD.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits

<b>Exhibit Number</b>	<b>Description</b>
99.1	<a href="#"><u>Press Release, dated August 5, 2020</u></a>
99.2	<a href="#"><u>Press Release, dated August 6, 2020</u></a>

---

## **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: August 7, 2020

By: /s/ Kenneth L. Londoner

Name: Kenneth L. Londoner

Title: Executive Chairman



## ViralClear Appoints Dr. Michael Sofia to Its Scientific Advisory Board

Westport, CT, Aug. 05, 2020 (GLOBE NEWSWIRE) --

- **Dr. Sofia is an Inductee in the American Chemical Society Medicinal Chemistry Hall of Fame**
- **ViralClear Pharmaceuticals is Currently in Phase 2 Trials with Merimepodib in the Fight Against COVID-19**

BioSig Technologies, Inc.'s (Nasdaq: BSGM) ("BioSig" or the "Company") subsidiary, ViralClear Pharmaceuticals, Inc. (ViralClear), today announced the addition of Dr. Michael J. Sofia to its Scientific Advisory Board (SAB).

Michael Sofia, Ph.D., has introduced numerous drugs into clinical development for the treatment of infectious diseases and inflammatory diseases. He has authored over 110 publications, 15 book chapters and is an inventor on more than 54 US patents. He is the principal inventor of sofosbuvir, currently marketed as the backbone of hepatitis C curative therapies Sovaldi(R), Harvoni(R), Epclusa(R) and Vosevi (R). Dr. Sofia has received numerous recognitions, including the 2016 Lasker-DeBakey Award in Clinical Medical Research and induction in the American Chemical Society Medicinal Chemistry Hall of Fame. Currently, Dr. Sofia is a Co-founder and Chief Scientific Officer of Arbutus Biopharma, Inc., a company focused on the discovery and development of therapies to cure hepatitis B.

"With Mike's addition to our Scientific Advisory Board, we believe that we have a very well rounded group of experts who can provide ViralClear with guidance on how to best proceed in addressing the current COVID-19 pandemic and other significant viral infections of special interest," stated Jerome B. Zeldis, M.D., Ph.D., the acting Chief Medical Officer and Head of ViralClear Pharmaceuticals, Inc.

Dr. Sofia commented, "I am excited to join the Scientific Advisory Board of ViralClear. The COVID-19 pandemic has called the drug development community to action to find therapies that can combat this disease. ViralClear's merimepodib is a novel agent with a new mechanism of action that has the potential to help address this pandemic now. I am eager to bring my expertise in antiviral drug development to help ViralClear in its efforts to develop merimepodib for the fight against COVID-19."

ViralClear recently announced the formation of its Scientific Advisory Board. The goal of the SAB is to review all aspects of drug discovery and development and advise ViralClear on its mission to control emerging infections and viral diseases of special interest, including COVID-19. Robin Robinson, Ph.D., the former Head of Biomedical Advanced Research and Development Authority (BARDA), and J. Paul Waymack, M.D., ScD, formerly of the Food and Drug Administration, were the first advisors appointed to the SAB.

### 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.

### About Merimepodib and ViralClear

BioSig's Technologies, Inc (Nasdaq: BSGM) subsidiary, ViralClear Pharmaceuticals, Inc. (ViralClear), 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 antiviral 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.

---

## 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  
BioSig Technologies, Inc.  
Vice President, Investor Relations  
54 Wilton Road, 2nd floor  
Westport, CT 06880  
[aballou@biosigtech.com](mailto:aballou@biosigtech.com)  
203-409-5444, x133



## Infectious Diseases Expert Prof. Angus Dalglish Joins ViralClear's Scientific Advisory Board

Westport, CT, Aug. 06, 2020 (GLOBE NEWSWIRE) --

- **Expert in Infectious Diseases and Immune Oncology**
- **ViralClear Pharmaceuticals is Currently in Phase 2 Trials with Merimepodib in the Fight Against COVID-19**

BioSig Technologies, Inc.'s (Nasdaq: BSGM) ("BioSig" or the "Company") subsidiary, ViralClear Pharmaceuticals, Inc. (ViralClear), today announced the addition of Professor Angus Dalglish to its Scientific Advisory Board (SAB).

Angus Dalglish, M.D. is the Foundation Chair of Oncology and Honorary Consultant in Medical Oncology at St. George's University and Hospital in London and Principal of the Institute of Cancer Vaccines and Immunotherapy. He is a fellow of the Royal College of Physicians both in the UK and Australia, the Royal College of Pathologists, and the Academy of Medical Sciences (UK). Prof. Dalglish published seminal papers on human immunodeficiency virus (HIV), its receptor on lymphocytes, and its pathogenesis.

"We are delighted to welcome Gus to our Scientific Advisory Board with his breadth of experience in developing antiviral drugs," stated Jerome B. Zeldis, M.D., Ph.D., the acting Chief Medical Officer and Head of ViralClear Pharmaceuticals, Inc. "Gus' expertise will complement those who are already on the SAB, and we look forward to working with him on this important mission."

Professor Dalglish said, "I am delighted to join the SAB of ViralClear Pharmaceuticals with my long-term collaborator, friend and colleague Jerry Zeldis with whom I worked with for several years successfully developing lenalidomide and pomalidomide, and I look forward to repeating this success against COVID-19."

ViralClear recently announced the formation of its Scientific Advisory Board to review all aspects of drug discovery and development and advise the Company on its mission to control emerging infections and viral diseases of special interest, including COVID-19. Robin Robinson, Ph.D., the former Head of Biomedical Advanced Research and Development Authority (BARDA), J. Paul Waymack, M.D., ScD, formerly of the Food and Drug Administration and a pioneer antiviral drug developer Michael Sofia, Ph.D. were the first advisors appointed to the Board.

### 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.

### About Merimepodib and ViralClear

BioSig's Technologies, Inc (Nasdaq: BSGM) subsidiary, ViralClear Pharmaceuticals, Inc. (ViralClear), 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 antiviral 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.

---

## 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  
BioSig Technologies, Inc.  
Vice President, Investor Relations  
54 Wilton Road, 2nd floor  
Westport, CT 06880  
[aballou@biosigtech.com](mailto:aballou@biosigtech.com)  
203-409-5444, x133