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): April 12, 2021

BioSig Technologies, Inc.

(Exact name of registrant as specified in its charter)

<u>Delaware</u> (State or other jurisdiction of incorporation) <u>001-38659</u> (Commission File Number)

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

26-433375 (IRS Employer Identification No.)

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

financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. \Box

<u>06880</u> (Zip Code)

(ICE	sistiant's telephone number, including area code	"
(Former	N/A name or former address, if changed since last re	eport)
Check the appropriate box below if the Form 8-K filing is intend	ded to simultaneously satisfy the filing obligation	n of the registrant under any of the following provisions:
\square Written communications pursuant to Rule 425 under the Secu	rrities Act (17 CFR 230.425)	
☐ Soliciting material pursuant to Rule 14a-12 under the Exchange	ge 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))
$\hfill \square$ Pre-commencement communications pursuant to Rule 13e-4 ((c) under the Exchange Act (17 CFR 240.13e-4((c))
Securi	ities registered pursuant to Section 12(b) of the A	Act:
Title of each class	Trading Symbol(s)	Name of exchange on which registered
Common Stock, par value \$0.001 per share	BSGM	The NASDAQ Capital Market
Indicate by check mark whether the registrant is an emerging groof the Securities Exchange Act of 1934 (§240.12b-2 of this chap	1 7	curities Act of 1933 (§230.405 of this chapter) or Rule 12b-2
Emerging growth company □		
If an emerging growth company, indicate by check mark if the re-	egistrant has elected not to use the extended tran	nsition period for complying with any new or revised

Item 7.01 Regulation FD Disclosure.

On April 12, 2021, Kenneth L. Londoner, Chairman and Chief Executive Officer of BioSig Technologies, Inc. (the "Company"), is expected to make a presentation regarding the Company at the Annual Needham Virtual Healthcare Conference at 3:00 PM ET. The slide presentation is attached hereto as Exhibit 99.1 and incorporated herein by reference. Additionally, the Company may, from time to time, to present and/or distribute to the investment community and utilize the slide presentation at various industry and other conferences. The Company undertakes no obligation to update, supplement or amend the materials attached hereto as Exhibit 99.1.

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 Exhibit 99.1, 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

Exhibit Number	Description
99.1	Slide Presentation of BioSig Technologies, Inc., April 2021
104	Cover Page Interactive Data File (formatted as Inline XBRL)

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: April 12, 2021

BIOSIG TECHNOLOGIES, INC.

By:

/s/ Kenneth L. Londoner Name: Kenneth L. Londoner Title: Executive Chairman



Disclaimer

This presentation contains forward-looking statements including statements that address activities, events or developments that BioSig expects, believes or anticipates will or may occur in the future, such as predictions of financial performance, approvals and launches by BioSig of new products, market acceptance of BioSig's products, market and procedure projections, financing plans, and related documents. Forward-looking statements are based on BioSig's experience and perception of current conditions, trends, expected future developments and other factors it believes are appropriate under the circumstances and are subject to numerous risks and uncertainties, many of which are beyond BioSig's control.

These risks and uncertainties include the timing of approvals for BioSig products, rate and degree of market acceptance of products, BioSig's ability to develop and market new and enhanced products, the timing of and ability to obtain and maintain regulatory clearances and approvals for its products and the impact of failure to obtain such clearances and approvals on its ability to promote its products and train doctors and operators in the use of its products, the timing of and ability to obtain reimbursement if required of procedures utilizing BioSig's products and the potential impact of current healthcare reform initiatives thereon, competition from existing and new products and procedures or BioSig's ability to effectively react to other risks and uncertainties described from time to time in BioSig's SEC filings, such as fluctuation of financial results, reliance on third party manufacturers and suppliers, litigation or other proceedings, government regulation, negative publicity, current worldwide economic conditions and share price volatility.

BioSig does not guarantee any forward-looking statements, and actual results may differ materially from those projected. Unless required by law, BioSig undertakes no obligation to update any forward-looking statement, whether as a result of new information, future events or otherwise.



Who We Are - NASDAQ: BSGM



- EP Tech Market estimated 13.5% CAGR
- Global EP Market Expected to be \$12.2 Billion by 2026
- 34 Patents Issued or Allowed
- First customer #1 center by volume in the U.S.
- Second customer voted #1 hospital in the world
- Razor & Blade Business Model
- FDA Cleared
- · Unblinding of clinical data in 2021 expected



Data source: MarketWatchGlobal Electrophysiology Device Market Could Exceed \$12.2 Billion By 2026. Definitive Healthcare, U.S. News & World Report and Newsweek.

Our Solution



More information, better decisions™

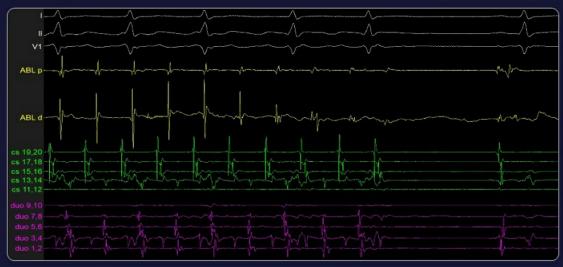
- Signal processing platform
- Intracardiac signals
- Customizable interface
- Artificial intelligence



"Secret Sauce"

Optimize signals for mapping, signal interpretation and during therapy delivery

• Right Atrial Typical Flutter signals during ablation with optimized settings showing no RF noise and detailed signal attenuation





Source: BioSig Spotlight Cases

Market Opportunity – PURE EP™

Global EP Procedures Growing Rapidly

Cardiac Ablation Procedures Are Key Hospital Revenue Drivers

Complex Cardiac Ablation:

13.5% Growth Rate

(Estimated)

U.S.

OUS:

3,915 EP labs

Total Global Addressable Market

\$2.0 Billion



BioSig* Published by Cardiovascular Rounds, Guggenhem Analyst - Chris Pasquale technologies Data source: 2018 MD&D report Advisory Board and BioSig Technologies, Inc. e:

Expanded Product Adoption, Experience, and a Growing Database

# PURE EP Procedures by Account				
Texas Cardiac Arrythmia Institute - Austin	293			
Mayo Clinic	120			
UNIV of Penn	99			
Mass General	13			
Deborah Heart & Lung	61			
Overland Park - Kansas	62			
Memorial – South Bend, IN	28			
Houston Methodist	31			
First in-human - TX, IN, SC	20			
TOTAL	727			

# Physician Users			
TCAI	10		
Mayo	3		
Penn	13		
MGH	1		
Deborah	2		
Overland Park	4		
Memorial	2		
Houston Methodist	2		
First in-human - TX, IN, SC	4		
TOTAL	41		

Diagnosis	# of Cases
Paroxysmal AF	222
Persistent AF	183
Ventricular Tachycardias and PVCs	132
SVTs	68

*Data as of 04/02/21





PURE EP™ System – Customer Installations



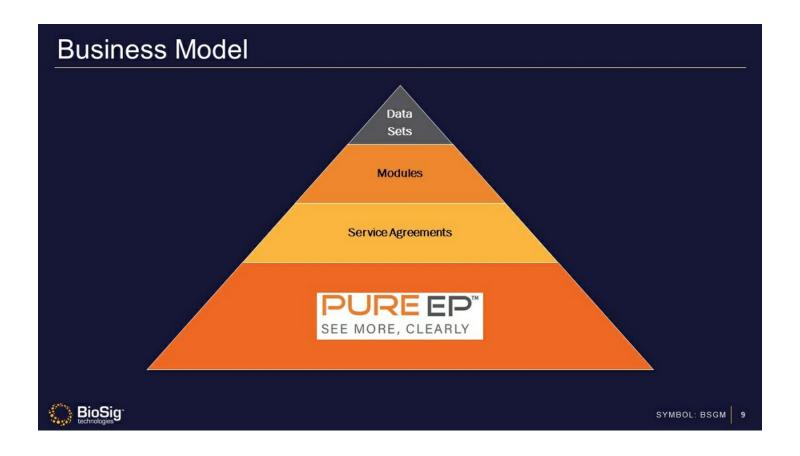
Hospital of the University of Pennsylvania

BioSig*

PURE EP™



Texas Cardiac Arrhythmia Institute



Commercialization – Regions and Centers of Excellence



Florida

120 EP Programs 240 Labs

Texas

220 EP Programs 440 Labs

Northeast

175 EP Programs 400 Labs

Revenue Potential \$286 Million



BioSig*

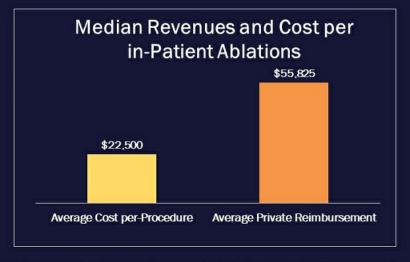
Highly Profitable Elective Procedure

EP procedures are clinically urgent

 Delaying procedure increases stroke risk and worsens outcomes

EP is revenue generating

 CV surgery and invasive cardiology have the highest net annual revenue compared to all other service lines





*Data Source: Advisory Board, Published by Cardiovascular Rounds, Cardiovascular rounds for DRG 273, UCLA Health

Clinical Data Catalysts

Further establish the safety and effectiveness of the PURE EP™ System

Multi-center: Mayo Clinic, Mass General, TCAI

Enrollment Completion April 13, 2021*

Abstract Acceptance Q2, 2021*

Unblinding data to the public July 15, 2021*

Heart Rhythm Society Presentation July 27, 2021*



*These are expected dates. ClinicalTrials.gov Identifier. NCT04112433

SYMBOL: BSGM

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Physician Testimonies



"The PURE EP System does provide innovative design with greater resolution and greater bandwidth, which should allow for safer and more efficacious ablations in the future."

- K. Venkatachalam, M.D., Cardiac Electrophysiologist, Mayo Clinic



"The quality of the recording that we can get in the lab can make a big difference [during the procedure]. . . The PURE EP System clearly gave us better quality of intracardiac recording [compared to conventional systems.]"

- Andrea Natale, M.D., Texas Cardiac Arrhythmia Institute







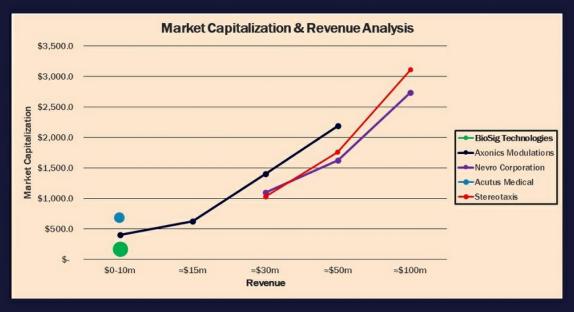
Market Value Inflection

Company	Market Capitalization (in millions USD)				
BioSig Technologies Formed 2009/IPO 2014	\$140				
Axonics Modulation Formed 2012/IPO 2018	\$405	\$620	х	\$2,200	
Acutus Medical Formed 2011/IPO 2020	х	\$670			
Nevro Corp Formed 2006/IPO 2014	х	х	\$1,090	\$1,623	\$2,740
Stereotaxis Formed 1990/IPO 2004	Х	Х	\$1,028	\$1,756	\$3,105
Revenues (millions USD)	≈\$1m	≈\$15m	≈\$30m	≈\$50m	≈\$100m



Source: FactSet Systems and BioSig Technologies' estimates. Market cap and revenues in millions

Revenue Increase = Market Capitalization Increase





Source: FactSet Systems and BioSig Technologies' estimates. Market cap and revenues in millions

SYMBOL: BSGM

1:

Management Team



Kenneth L. Londoner, MBA
Founder, Chairman, CEO, Director
NeuroClear Technologies; Endicott Management Partners;



Steve Chaussy, CPA

NeuroClearTechnologies; Liberski Inc; Anna & Co; Penske Automotive; Ford Hogg and Cobbe



Brenda Castrodad

Director of Human Resources TissueTech, Inc.; HeartWare, Inc., Schering-Plough Corp



Andy Ballou

Vice President, Investor Relations
Janney Montgomery Scott; RBC Capital Markets



Julie Stephenson, BSN, MBA

Vice President, Clinical Affairs Medtronic; Boston Scientific; Guidant Corporation



Manasi Patwardhan

Director of Strategic Planning Verily Life Sciences, Boston Scientific - Neuromodulation; Medtronic























Ewald Riechert

Natasha Drapeau

Todd Wiltshire

John Kowalski

Vice President, Sales

Olivier Chaudoir

Biosense Webster; DePuy Synthes

Biosense Webster (Johnson & Johnson)

Executive Vice President
NeuroClear Technologies; Alliance for Advancing Bioelectronic
Medicine; Institute of Directors, UK; IG Group Plc, UK

Senior Vice President, Corporate Development Fidelity Investments; Credit Agricole; UBS; Morgan Stanley

Director of Regulatory Affairs / Quality Assurance TUV Rheinland, Siemens, Acclarent, Inc. (J&J Company)

Medtronic; Nexeon MedSystems; Alfred Mann Institute for Biomedical Engineering; Alfred Mann Foundation for Scientific Research

Barry Keenan, Ph.D, MBA, PMP





Value Drivers

- \$28 million* in cash on the balance sheet and no debt
- Expect to triple the customer base in 2021
- Targeting sequential revenue growth in 2021
- Expect CE mark approval / EU entry 2022
- Launched Al program Mayo partnership
- Expect presentation of data Q3 2021





