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**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
Washington, D.C. 20549**

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**FORM 10-Q**

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(Mark One)

**QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the quarterly period ended **March 31, 2019**

**TRANSITION REPORT UNDER SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the transition period from \_\_\_\_\_ to \_\_\_\_\_

Commission file number: **000-55473**

**BIOSIG TECHNOLOGIES, INC.**

(Exact name of registrant as specified in its charter)

**Delaware**

(State or other jurisdiction of incorporation  
or organization)

**26-433375**

(IRS Employer Identification No.)

**12424 Wilshire Blvd, Suite 745**

**Los Angeles, CA**

(Address of principal executive office)

**90025**

(Zip Code)

**(310) 620-9320**

(Registrant's telephone number, including area code)

Securities registered pursuant to Section 12(g) of the Act: **Common Stock, \$0.001 par value per share**

Indicate by check mark if the registrant is a well-known seasoned issuer, as defined by Rule 405 of the Securities Act. Yes  No

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. Yes  No

Indicate by check mark whether the registrant has submitted electronically every Interactive Data File required to be submitted pursuant to Rule 405 of Regulation S-T (§ 229.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit such files). Yes  No

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, a smaller reporting company, or an emerging growth company. See definitions of "large accelerated filer," "accelerated filer," "smaller reporting company," and "emerging growth company" in Rule 12b-2 of the Exchange Act.

Large accelerated filer	<input type="checkbox"/>	Accelerated filer	<input type="checkbox"/>
Non-accelerated filer	<input checked="" type="checkbox"/>	Smaller reporting company	<input checked="" type="checkbox"/>
Emerging growth company	<input checked="" type="checkbox"/>		

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.

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). Yes  No

As of April 15, 2019, there were 20,211,063 shares of registrant's common stock outstanding.

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## PART I – FINANCIAL INFORMATION

## ITEM 1. FINANCIAL STATEMENTS

**BIOSIG TECHNOLOGIES, INC.**  
**CONDENSED CONSOLIDATED BALANCE SHEETS**

	<b>March 31, 2019</b>	<b>December 31, 2018</b>
	<i>(unaudited)</i>	
<b>ASSETS</b>		
Current assets:		
Cash	\$ 10,933,593	\$ 4,450,160
Prepaid expenses	156,577	178,442
Total current assets	<u>11,090,170</u>	<u>4,628,602</u>
Property and equipment, net	53,559	44,346
Right-to-use assets, net	380,586	-
Other assets:		
Patents, net	324,397	268,796
Trademarks	1,125	850
Deposits	<u>54,238</u>	<u>54,238</u>
Total assets	<u>\$ 11,904,075</u>	<u>\$ 4,996,832</u>
<b>LIABILITIES AND STOCKHOLDERS' EQUITY</b>		
Current liabilities:		
Accounts payable and accrued expenses, including \$14,632 and \$32,366 to related parties as of March 31, 2019 and December 31, 2018, respectively	\$ 597,006	\$ 954,655
Dividends payable	253,449	242,908
Lease liability, short term	<u>158,836</u>	<u>-</u>
Total current liabilities	1,009,291	1,197,563
Lease liability, long term	<u>226,531</u>	<u>-</u>
Total debt	1,235,822	1,197,563
Series C Preferred Stock, 475 shares issued and outstanding; liquidation preference of \$475,000 as of March 31, 2019 and December 31, 2018	<u>475,000</u>	<u>475,000</u>
Stockholders' equity		
Preferred stock, \$0.001 par value, authorized 1,000,000 shares, designated 200 shares of Series A, 600 shares of Series B, 4,200 shares of Series C, 1,400 shares of Series D, 1,000 shares of Series E Preferred Stock. No shares outstanding.	-	-
Common stock, \$0.001 par value, authorized 200,000,000 shares, 20,009,985 and 16,868,783 issued and outstanding as of March 31, 2019 and December 31, 2018, respectively	20,010	16,869
Additional paid in capital	86,465,732	74,039,341
Common stock subscription	309,000	-
Accumulated deficit	<u>(76,601,489)</u>	<u>(70,731,941)</u>
Total stockholders' equity	<u>10,193,253</u>	<u>3,324,269</u>
Total liabilities and stockholders' equity	<u>\$ 11,904,075</u>	<u>\$ 4,996,832</u>

See the accompanying notes to the unaudited condensed consolidated financial statements

**BIOSIG TECHNOLOGIES, INC.**  
**CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS**  
*(unaudited)*

	<b>Three months ended March 31,</b>	
	<b>2019</b>	<b>2018</b>
Operating expenses:		
Research and development	\$ 1,488,839	\$ 862,420
General and administrative	4,378,897	1,771,796
Depreciation and amortization	7,935	2,903
Total operating expenses	<u>5,875,671</u>	<u>2,637,119</u>
Loss from operations	(5,875,671)	(2,637,119)
Other income (expense):		
Interest income	<u>6,123</u>	<u>85</u>
Loss before income taxes	(5,869,548)	(2,637,034)
Income taxes (benefit)	<u>-</u>	<u>-</u>
Net loss	(5,869,548)	(2,637,034)
Preferred stock dividend	<u>(10,541)</u>	<u>(305,046)</u>
<b>NET LOSS AVAILABLE TO COMMON STOCKHOLDERS</b>	<b><u>\$ (5,880,089)</u></b>	<b><u>\$ (2,942,080)</u></b>
Net loss per common share, basic and diluted	<u>\$ (0.33)</u>	<u>\$ (0.25)</u>
Weighted average number of common shares outstanding, basic and diluted	<u>17,848,238</u>	<u>11,977,778</u>

See the accompanying notes to the unaudited condensed consolidated financial statements

**BIOSIG TECHNOLOGIES, INC.**  
**CONDENSED CONSOLIDATED STATEMENT OF STOCKHOLDERS' EQUITY**  
**THREE MONTHS ENDED MARCH 31, 2019**

	Common stock		Additional Paid in Capital	Common stock Subscription	Accumulated Deficit	Total
	Shares	Amount				
Balance, December 31, 2018	16,868,783	\$ 16,869	\$ 74,039,341	\$ -	\$ (70,731,941)	\$ 3,324,269
Common stock issued for services	560,000	560	2,332,640	-	-	2,333,200
Sale of common stock	2,155,127	2,155	8,617,123	-	-	8,619,278
Common stock issued upon exercise of warrants at an average of \$3.86 per share	298,319	298	1,150,482	-	-	1,150,780
Common stock issued upon cashless exercise of warrants	104,424	105	(105)	-	-	-
Warrant subscription exercise received	-	-	-	309,000	-	309,000
Stock based compensation	23,332	23	336,792	-	-	336,815
Preferred stock dividend	-	-	(10,541)	-	-	(10,541)
Net loss	-	-	-	-	(5,869,548)	(5,869,548)
Balance, March 31, 2019 ( <i>unaudited</i> )	<u>20,009,985</u>	<u>\$ 20,010</u>	<u>\$ 86,465,732</u>	<u>\$ 309,000</u>	<u>\$ (76,601,489)</u>	<u>\$ 10,193,253</u>

See the accompanying notes to the unaudited condensed consolidated financial statements

**BIOSIG TECHNOLOGIES, INC.**  
**CONDENSED CONSOLIDATED STATEMENT OF STOCKHOLDERS' DEFICIT**  
**THREE MONTHS ENDED MARCH 31, 2018**

	Series D Preferred stock		Series E Preferred stock		Common stock		Additional	Common	Accumulated	Total
	Shares	Amount	Shares	Amount	Shares	Amount	Paid in Capital	stock Subscription	Deficit	
Balance, December 31, 2017	1,334	\$ 1	-	\$ -	11,728,482	\$ 11,728	\$ 53,233,228	\$ 29,985	\$ (56,524,786)	\$ (3,249,844)
Reclassify fair value of derivative and warrant liabilities to equity upon adoption of ASU 2017-11	-	-	-	-	-	-	-	-	3,044,162	3,044,162
Common stock issued for services	-	-	-	-	40,000	40	141,960	-	-	142,000
Sale of common stock	-	-	-	-	80,000	80	299,905	(29,985)	-	270,000
Common stock issued upon conversion of Series C Preferred Stock at \$1.50 per share	-	-	-	-	5,334	5	19,995	-	-	20,000
Common stock issued settlement of Series C Preferred Stock accrued dividends at \$1.25 per share	-	-	-	-	2,638	3	8,273	-	-	8,276
Common stock issued upon conversion of Series D Preferred Stock at \$1.50 per share	(647)	-	-	-	258,800	259	(259)	-	-	-
Common stock issued settlement of Series D Preferred Stock accrued dividends at \$1.25 per share	-	-	-	-	83,830	84	261,951	-	-	262,035
Sale of Series E Preferred stock	-	-	1,000	1	-	-	1,492,968	-	-	1,492,969
Common stock subscription received	-	-	-	-	-	-	-	115,470	-	115,470
Stock based compensation	-	-	-	-	-	-	246,710	-	-	246,710
Preferred stock dividend	-	-	-	-	-	-	(305,046)	-	-	(305,046)
Net loss	-	-	-	-	-	-	-	-	(2,637,034)	(2,637,034)
Balance, March 31, 2018 <i>(unaudited)</i>	<u>687</u>	<u>\$ 1</u>	<u>1,000</u>	<u>\$ 1</u>	<u>12,199,084</u>	<u>\$ 12,199</u>	<u>\$ 55,399,685</u>	<u>\$ 115,470</u>	<u>\$ (56,117,658)</u>	<u>\$ (590,302)</u>

See the accompanying notes to the unaudited condensed financial statements

**BIOSIG TECHNOLOGIES, INC.**  
**CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS**  
*(unaudited)*

	<b>Three months ended March 31,</b>	
	<b>2019</b>	<b>2018</b>
<b>CASH FLOWS FROM OPERATING ACTIVITIES:</b>		
Net loss	\$ (5,869,548)	\$ (2,637,034)
Adjustments to reconcile net loss to cash used in operating activities:		
Depreciation and amortization	7,935	2,903
Equity based compensation	2,670,015	388,710
Changes in operating assets and liabilities:		
Prepaid expenses	21,865	(2,173)
Security deposit	-	(5,393)
Accounts payable and accrued expenses	(354,272)	(134,714)
Lease liability, net	1,404	-
Deferred rent payable	-	427
Net cash used in operating activities	<u>(3,522,601)</u>	<u>(2,387,274)</u>
<b>CASH FLOWS FROM INVESTING ACTIVITIES:</b>		
Payments of patent costs	(58,327)	-
Payment of trademark costs	(275)	-
Purchase of property and equipment	(14,422)	(3,644)
Net cash used in investing activity	<u>(73,024)</u>	<u>(3,644)</u>
<b>CASH FLOWS FROM FINANCING ACTIVITIES:</b>		
Proceeds from sale of common stock	8,619,278	270,000
Proceeds from sale of Series E preferred stock	-	1,492,969
Proceeds from exercise of warrants	1,459,780	-
Proceeds from common stock subscription	-	115,470
Net cash provided by financing activities	<u>10,079,058</u>	<u>1,878,439</u>
Net increase (decrease) in cash and cash equivalents	6,483,433	(512,479)
Cash and cash equivalents, beginning of the period	4,450,160	1,547,579
Cash and cash equivalents, end of the period	<u>\$ 10,933,593</u>	<u>\$ 1,035,100</u>
<b>Supplemental disclosures of cash flow information:</b>		
Cash paid during the period for interest	<u>\$ -</u>	<u>\$ -</u>
Cash paid during the period for income taxes	<u>\$ -</u>	<u>\$ -</u>
<b>Non cash investing and financing activities:</b>		
Common stock issued upon conversion of Series C Preferred Stock and accrued dividends	<u>\$ -</u>	<u>\$ 28,276</u>
Reclassify initial fair value of derivative and warrant liabilities from equity upon issuance of Series D preferred stock	<u>\$ -</u>	<u>\$ 262,035</u>
Reclassify fair value of derivative and warrant liabilities to equity upon adoption of ASU 2017-11	<u>\$ -</u>	<u>\$ 3,044,162</u>
Dividends payable on preferred stock charged to additional paid in capital	<u>\$ 10,541</u>	<u>\$ 305,046</u>
Right-to-use assets and lease liability recorded upon adoption of ASC 842	<u>\$ 418,838</u>	<u>\$ -</u>

See the accompanying notes to the unaudited condensed consolidated financial statements

**BIOSIG TECHNOLOGIES, INC.**  
**NOTES TO THE CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**  
**MARCH 31, 2019**  
*(unaudited)*

**NOTE 1 – NATURE OF OPERATIONS AND BASIS OF PRESENTATION**

BioSig Technologies Inc. (the “Company”) was initially incorporated on February 24, 2009 under the laws of the State of Nevada and subsequently re-incorporated in the state of Delaware in 2011. The Company is principally devoted to improving the quality of cardiac recordings obtained during EP studies and catheter ablation procedures. The Company has not generated any revenue to date and consequently its operations are subject to all risks inherent in the establishment of a new business enterprise.

On November 7, 2018, the Company formed NeuroClear Technologies, Inc., a Delaware Corporation, for the purpose to pursue additional applications of the PURE EP™ signal processing technology outside of electrophysiology. As of March 31, 2019, there were no significant assets or liabilities in NeuroClear Technologies, Inc, or operations since its formation.

The condensed consolidated financial statements include the accounts of BioSig Technologies, Inc. and its wholly owned subsidiary, NeuroClear Technologies, Inc. to as the “Company” or “BioSig”.

The unaudited condensed interim financial statements of the Company have been prepared in accordance with accounting principles generally accepted in the United States (“GAAP”) for interim financial information and the instructions to Form 10-Q and Rule 8-03 of Regulation S-X. Accordingly, they do not include all of the information and footnotes required by GAAP for complete financial statements. In the opinion of management, all adjustments (consisting of normal recurring accruals) considered necessary for a fair presentation have been included.

The condensed consolidated balance sheet as of December 31, 2018 has been derived from audited financial statements.

Operating results for the three months ended March 31, 2019 are not necessarily indicative of results that may be expected for the year ending December 31, 2019. These condensed consolidated financial statements should be read in conjunction with the audited financial statements for the year ended December 31, 2018 filed with the Company’s Form 10-K with the Securities and Exchange Commission on March 15, 2019.

Effective September 10, 2018, the Company amended its Articles of Incorporation to implement a reverse stock split in the ratio of 1 share for every 2.5 shares of common stock. As a result, 40,333,758 shares of the Company’s common stock were exchanged for 16,133,544 shares of the Company's common stock. These financial statements have been retroactively restated to reflect the reverse stock split. (See Note 8)

**NOTE 2 – GOING CONCERN AND MANAGEMENT’S LIQUIDITY PLANS**

As of March 31, 2019, the Company had cash of \$10,933,593 and working capital of \$10,080,879. The Company raised approximately \$8,600,000 through the sale of common stock and \$1,500,000 from the exercise of previously issued warrants during the three months ended March 31, 2019 and approximately \$417,000 subsequent to March 31, 2019 (See Note 13). During the three months ended March 31, 2019, the Company used net cash in operating activities of \$3,522,601. These conditions raise substantial doubt about the Company’s ability to continue as a going concern. Management believes that the Company has sufficient funds to meet its research and development and other funding requirements for at least the next 10 months.

The Company’s primary source of operating funds since inception has been cash proceeds from private placements of common and preferred stock. The Company has experienced net losses and negative cash flows from operations since inception and expects these conditions to continue for the foreseeable future. The Company will require additional financing to fund future operations. Further, the Company does not have any commercial products available for sale and there is no assurance that the Company will be able to generate cash flow to fund operations. In addition, there can be no assurance that the Company’s research and development will be successfully completed or that any product will be commercially viable.



**BIOSIG TECHNOLOGIES, INC.**  
**NOTES TO THE CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**  
**MARCH 31, 2019**  
*(unaudited)*

Accordingly, the accompanying financial statements have been prepared in conformity with U.S. GAAP, which contemplates continuation of the Company as a going concern and the realization of assets and satisfaction of liabilities in the normal course of business. The carrying amounts of assets and liabilities presented in the financial statements do not necessarily purport to represent realizable or settlement values. The condensed consolidated financial statements do not include any adjustment that might result from the outcome of this uncertainty.

**NOTE 3 – SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES**

*Use of estimates*

The preparation of financial statements in conformity with Generally Accepted Accounting Principles requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities, disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting period. Significant estimates include the recoverability and useful lives of long-lived assets, the fair value of the Company's stock, stock-based compensation and the valuation allowance related to deferred tax assets. Actual results may differ from these estimates.

*Fair Value of Financial Instruments*

Accounting Standards Codification subtopic 825-10, Financial Instruments ("ASC 825-10") requires disclosure of the fair value of certain financial instruments. The carrying value of cash and cash equivalents, accounts payable and accrued liabilities as reflected in the balance sheets, approximate fair value because of the short-term maturity of these instruments. All other significant financial assets, financial liabilities and equity instruments of the Company are either recognized or disclosed in the financial statements together with other information relevant for making a reasonable assessment of future cash flows, interest rate risk and credit risk. Where practicable the fair values of financial assets and financial liabilities have been determined and disclosed; otherwise only available information pertinent to fair value has been disclosed.

The Company follows Accounting Standards Codification subtopic 820-10, Fair Value Measurements and Disclosures ("ASC 820-10") and Accounting Standards Codification subtopic 825-10, Financial Instruments ("ASC 825-10"), which permits entities to choose to measure many financial instruments and certain other items at fair value.

*Derivative Instrument Liability*

The Company accounts for derivative instruments in accordance with ASC 815, which establishes accounting and reporting standards for derivative instruments and hedging activities, including certain derivative instruments embedded in other financial instruments or contracts and requires recognition of all derivatives on the balance sheet at fair value, regardless of hedging relationship designation. Accounting for changes in fair value of the derivative instruments depends on whether the derivatives qualify as hedge relationships and the types of relationships designated are based on the exposures hedged. At March 31, 2019 and December 31, 2018, the Company did not have any derivative instruments that were designated as hedges.

At March 31, 2019 and December 31, 2018, the Company had outstanding preferred stock and warrants that contained embedded derivatives. These embedded derivatives include certain conversion features and reset provisions. On January 1, 2018, the Company adopted ASU 2017-11 and according reclassified the fair value of the reset provisions embedded in previously issued Preferred stock and certain warrants with embedded anti-dilutive provisions from liability to equity.

**BIOSIG TECHNOLOGIES, INC.**  
**NOTES TO THE CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**  
**MARCH 31, 2019**  
*(unaudited)*

*Research and development costs*

The Company accounts for research and development costs in accordance with the Accounting Standards Codification subtopic 730-10, Research and Development (“ASC 730-10”). Under ASC 730-10, all research and development costs must be charged to expense as incurred. Accordingly, internal research and development costs are expensed as incurred. Third-party research and development costs are expensed when the contracted work has been performed or as milestone results have been achieved. Company-sponsored research and development costs related to both present and future products are expensed in the period incurred. The Company incurred research and development expenses of \$1,488,839 and \$862,420 for the three months ended March 31, 2019 and 2018, respectively.

*Concentrations of Credit Risk*

Financial instruments and related items, which potentially subject the Company to concentrations of credit risk, consist primarily of cash and cash equivalents. The Company places its cash and temporary cash investments with credit quality institutions. At times, such amounts may be in excess of the FDIC insurance limit. At March 31, 2019 and December 31, 2018, deposits in excess of FDIC limits were \$10,683,593 and \$4,200,160, respectively.

*Net Income (loss) Per Common Share*

The Company computes earnings (loss) per share under Accounting Standards Codification subtopic 260-10, Earnings Per Share (“ASC 260-10”). Net loss per common share is computed by dividing net loss by the weighted average number of shares of common stock outstanding during the year. Diluted earnings per share, if presented, would include the dilution that would occur upon the exercise or conversion of all potentially dilutive securities into common stock using the “treasury stock” and/or “if converted” methods as applicable.

The computation of basic and diluted loss per share as of March 31, 2019 and 2018 excludes potentially dilutive securities when their inclusion would be anti-dilutive, or if their exercise prices were greater than the average market price of the common stock during the period.

Potentially dilutive securities excluded from the computation of basic and diluted net income (loss) per share are as follows:

	<b>March 31, 2019</b>	<b>March 31, 2018</b>
Series C convertible preferred stock	126,667	257,334
Series D convertible preferred stock	-	274,800
Series E convertible preferred stock	-	400,000
Options to purchase common stock	3,940,828	3,292,128
Warrants to purchase common stock	3,987,088	4,895,051
Totals	<u>8,054,583</u>	<u>9,119,313</u>

*Stock Based Compensation*

The Company measures the cost of services received in exchange for an award of equity instruments based on the fair value of the award. For employees and directors, the fair value of the award is measured on the grant date and for non-employees, the fair value of the award is generally re-measured on vesting dates and interim financial reporting dates until the service period is complete. The fair value amount is then recognized over the period during which services are required to be provided in exchange for the award, usually the vesting period.

As of March 31, 2019, the Company had 3,940,828 options outstanding to purchase shares of common stock, of which 3,042,512 were vested.

As of December 31, 2018, there were outstanding stock options to purchase 3,135,828 shares of common stock, 3,007,946 shares of which were vested.

**BIOSIG TECHNOLOGIES, INC.**  
**NOTES TO THE CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**  
**MARCH 31, 2019**  
*(unaudited)*

*Income Taxes*

The Company follows Accounting Standards Codification subtopic 740-10, Income Taxes (“ASC 740-10”) for recording the provision for income taxes. Deferred tax assets and liabilities are computed based upon the difference between the financial statement and income tax basis of assets and liabilities using the enacted marginal tax rate applicable when the related asset or liability is expected to be realized or settled. Deferred income tax expenses or benefits are based on the changes in the asset or liability during each period. If available evidence suggests that it is more likely than not that some portion or all of the deferred tax assets will not be realized, a valuation allowance is required to reduce the deferred tax assets to the amount that is more likely than not to be realized. Future changes in such valuation allowance are included in the provision for deferred income taxes in the period of change. Deferred income taxes may arise from temporary differences resulting from income and expense items reported for financial accounting and tax purposes in different periods.

*Patents, net*

The Company capitalizes certain initial asset costs in connection with patent applications including registration, documentation and other professional fees associated with the application. Patent costs incurred prior to the Company’s U.S. Food and Drug Administration (“FDA”) 510 (k) application on March 28, 2018 were charged to research and development expense as incurred. Commencing upon first in-man trials on February 18 and 19, 2019, capitalized costs are amortized to expense using the straight-line method over the lesser of the legal patent term or the estimated life of the product of 10 years. During the three months ended March 31, 2019, the Company recorded amortization of \$2,726 to current period operations.

*Registration Rights*

On March 12, 2019, in connection with the Company’s Private Placement of common stock, the Company also agreed on or prior the date that is 45 calendar days after the closing date of the Private Placement, the Company will be required to use commercially reasonable efforts to prepare and file a registration statement on Form S-3 or Form S-1 with the Securities and Exchange Commission (the “SEC”) covering the resale of the common shares. The Company is additionally required to use its commercially reasonable efforts to cause such registration statement to be declared effective by the SEC as soon as practicable thereafter. All expenses related to the filing of such registration statement, including legal fees, will be borne by the Company. The Company has estimated the liability under the registration rights agreement at \$-0- as of March 31, 2019.

*Adoption of Accounting Standards*

In February 2016, the FASB established ASC Topic 842, Leases (Topic 842), by issuing ASU No. 2016-02, which requires lessees to recognize leases on-balance sheet and disclose key information about leasing arrangements. Topic 842 was subsequently amended by ASU No. 2018-01, Land Easement Practical Expedient for Transition to Topic 842; ASU No. 2018-10, Codification Improvements to Topic 842, Leases; and ASU No. 2018-11, Targeted Improvements. The new standard establishes a right-of-use (ROU) model that requires a lessee to recognize a ROU asset and lease liability on the balance sheet. Leases will be classified as finance or operating, with classification affecting the pattern and classification of expense recognition in the statement of operations. The Company adopted the new standard on January 1, 2019.

The new standard provides a number of optional practical expedients in transition. The Company has elected the ‘package of practical expedients’, which permit it not to reassess under the new standard its prior conclusions about lease identification, lease classification and initial direct costs. The Company did not elect the use-of-hindsight or the practical expedient pertaining to land easements; the latter is not applicable to the Company.

The new standard had a material effect on the Company’s financial statements. The most significant effects of adoption relate to (1) the recognition of new ROU assets and lease liabilities on its balance sheet for real estate operating leases; and (2) providing significant new disclosures about its leasing activities.

**BIOSIG TECHNOLOGIES, INC.**  
**NOTES TO THE CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**  
**MARCH 31, 2019**  
*(unaudited)*

Upon adoption, the Company recognized additional operating lease liabilities, net of deferred rent, of approximately \$419,000 based on the present value of the remaining minimum rental payments under current leasing standards for existing operating leases. The Company also recognized corresponding ROU assets of approximately \$419,000.

The new standard also provides practical expedients for an entity's ongoing accounting. The Company elected the short-term lease recognition exemption for all leases that qualify. This means, for those leases that qualify, the Company will not recognize ROU assets or lease liabilities, and this includes not recognizing ROU assets or lease liabilities for existing short-term leases of those assets in transition. Beginning in 2019, the Company changed to its disclosed lease recognition policies and practices, as well as to other related financial statement disclosures due to the adoption of this standard. See Note 5.

*Recent Accounting Pronouncements*

There were various updates recently issued, most of which represented technical corrections to the accounting literature or application to specific industries and are not expected to have a material impact on the Company's financial position, results of operations or cash flows.

*Subsequent Events*

The Company evaluates events that have occurred after the balance sheet date but before the consolidated financial statements are issued. Based upon the evaluation, the Company did not identify any recognized or non-recognized subsequent events that would have required adjustment or disclosure in the consolidated financial statements, except as disclosed.

**NOTE 4 – PROPERTY AND EQUIPMENT**

Property and equipment as of March 31, 2019 and December 31, 2018 is summarized as follows:

	<b>March 31, 2019</b>	<b>December 31, 2018</b>
Computer equipment	\$ 117,469	\$ 105,447
Furniture and fixtures	35,019	32,619
Subtotal	152,488	138,066
Less accumulated depreciation	(98,929)	(93,720)
Property and equipment, net	<u>\$ 53,559</u>	<u>\$ 44,346</u>

Property and equipment are stated at cost and depreciated using the straight-line method over their estimated useful lives of 3 to 5 years. When retired or otherwise disposed, the related carrying value and accumulated depreciation are removed from the respective accounts and the net difference less any amount realized from disposition, is reflected in earnings.

Depreciation expense was \$5,209 and \$2,903 for three months ended March 31, 2019 and 2018, respectively.

**NOTE 5 – RIGHT TO USE ASSETS AND LEASE LIABILITY**

On May 22, 2018, the Company entered into a fifth lease amendment agreement, whereby the Company agreed to extend the lease for the original office space and expand with additional space in Los Angeles, California, commencing June 14, 2018 and expiring on June 30, 2021 at an initial rate of \$14,731 per month with escalating payments. In connection with the lease, the Company is obligated to lease parking spaces at an aggregate approximate cost of \$1,070 per month. In addition, the Company entered into a lease for storage space with the Los Angeles, California building commencing on December 1, 2017 and expiring on August 31, 2019 for approximately \$223 per month. The Company has an option to extend the lease for an additional 3 year (option) term.

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On April 11, 2018, the Company extended a short-term lease agreement whereby the Company leased office space in Austin, Texas commencing on August 1, 2018 and expiring July 31, 2019 for \$979 per month.

On October 1, 2018, the Company entered into a lease agreement whereby the Company leased office space in Norwalk, Connecticut commencing on October 1, 2018 and expiring September 30, 2019 for \$2,000 per month.

In adopting ASC Topic 842, Leases (Topic 842), the Company has elected the 'package of practical expedients', which permit it not to reassess under the new standard its prior conclusions about lease identification, lease classification and initial direct costs. The Company did not elect the use-of-hindsight or the practical expedient pertaining to land easements; the latter is not applicable to the Company. In addition, the Company elected not to apply ASC Topic 842 to arrangements with lease terms of 12 month or less. In determining the length of the lease term to its long term lease, the Company determined not to consider an embedded 3 year option primarily due to i) the renewal rate is at future market rate to be determined and ii) Company does not have significant leasehold improvements that would restrict its ability to consider relocation. At lease commencement date, the Company estimated the lease liability and the right of use assets at present value using the Company's estimated incremental borrowing rate of 8% and determined the initial present value, at inception, of \$496,467. On January 1, 2019, upon adoption of ASC Topic 842, the Company recorded right to use assets of \$418,838, lease liability of \$422,215 and eliminated deferred rent of \$3,377.

Right to use assets is summarized below:

	<b>March 31, 2019</b>
Los Angeles, Suite 740	\$ 218,875
Los Angeles, Suite 745	277,592
Subtotal	496,467
Less accumulated depreciation	(115,881)
Right to use assets, net	\$ 380,586

During the three months ended March 31, 2019, the Company recorded \$46,451 as lease expense to current period operations.

Lease liability is summarized below:

	<b>March 31, 2019</b>
Los Angeles, Suite 740	\$ 169,885
Los Angeles, Suite 745	215,482
Total lease liability	385,367
Less: short term portion	(158,836)
Long term portion	\$ 226,531

Maturity analysis under these lease agreements are as follows:

Nine months ended December 31, 2019	\$ 137,503
Year ended December 31, 2020	189,141
Year ended December 31, 2021	96,196
Total	\$ 422,840
Less: Present value discount	(37,473)
Lease liability	\$ 385,367

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Lease expense for the three months ended March 31, 2019 was comprised of the following:

Operating lease expense	\$	46,451
Short-term lease expense		13,238
Variable lease expense		438
	\$	60,127

**NOTE 6 – ACCOUNTS PAYABLE AND ACCRUED EXPENSES**

Accounts payable and accrued expenses at March 31, 2019 and December 31, 2018 consist of the following:

	<b>March 31,</b>	<b>December 31,</b>
	<b>2019</b>	<b>2018</b>
Accrued accounting and legal	\$ 94,477	\$ 59,439
Accrued reimbursements and travel	26,245	27,853
Accrued consulting	102,967	89,718
Accrued research and development expenses	345,751	351,631
Accrued office and other	3,821	14,304
Accrued payroll and related expenses	10,412	395,000
Deferred rent	-	3,377
Accrued settlement related to arbitration	13,333	13,333
	\$ 597,006	\$ 954,655

**NOTE 7 – SERIES C 9% CONVERTIBLE PREFERRED STOCK**

*Series C 9% Convertible Preferred Stock*

On January 9, 2013, the Board of Directors authorized the issuance of up to 4,200 shares of 9% Series C Convertible Preferred Stock (the “Series C Preferred Stock”).

The Series C Preferred Stock is entitled to preference over holders of junior stock upon liquidation in the amount of \$1,000 plus any accrued and unpaid dividends; entitled to dividends as a preference to holders of junior stock at a rate of 9% per annum of the stated value of \$1,000 per share, payable quarterly beginning on September 30, 2013 and are cumulative. The holders of the Series C Preferred Stock vote together with the holders of our common stock on an as-converted basis, but may not vote the Series C Preferred Stock in excess of the beneficial ownership limitation of the Series C Preferred Stock. The beneficial ownership limitation is 4.99% of our then outstanding shares of common stock following such conversion or exercise, which may be increased to up to 9.99% of our then outstanding shares of common stock following such conversion or exercise upon the request of an individual holder. The beneficial ownership limitation is determined on an individual holder basis, such that the as-converted number of shares of one holder is not included in the shares outstanding when calculating the limitation for a different holder.

In connection with the sale of the Series C preferred stock, the Company issued an aggregate of 532,251 warrants to purchase the Company’s common stock at \$6.53 per share expiring five years from the initial exercise date. The warrants contain full ratchet anti-dilution price protection upon the issuance of equity or equity-linked securities at an effective common stock purchase price of less than \$6.53 per share as well as other customary anti-dilution protection. The warrants are exercisable for cash; or if at any time after six months from the issuance date, there is no effective registration statement registering the resale, or no current prospectus available for the resale, of the shares of common stock underlying the warrants, the warrants may be exercised by means of a “cashless exercise”.

As a result of an amendment to the conversion price of our Series C Preferred Stock, the full-ratchet anti-dilution protection provision of the warrants decreased the exercise price of the warrants from \$6.53 per share to \$3.75 per share and increased the aggregate number of shares issuable under the warrants to 926,121.

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Series C Preferred Stock issued and outstanding totaled 475 as of March 31, 2019 and December 31, 2018, respectively. As of March 31, 2019 and December 31, 2018, the Company has accrued \$253,449 and \$242,908 dividends payable on the Series C Preferred Stock.

**NOTE 8 – STOCKHOLDER EQUITY**

*Preferred stock*

The Company is authorized to issue 1,000,000 shares of \$0.001 par value preferred stock. As of March 31, 2019 and December 31, 2018, the Company has authorized 200 shares of Series A preferred stock, 600 shares of Series B preferred stock, 4,200 shares of Series C Preferred Stock, 1,400 shares of Series D Preferred Stock and 1,000 shares of Series E Preferred Stock. As of March 31, 2019 and December 31, 2018, there were no outstanding shares of Series A, Series B, Series D and Series E preferred stock.

*Common stock*

On September 10, 2018, the Company amended its Articles of Incorporation to implement a reverse stock split in the ratio of 1 share for every 2.5 shares of common stock. No fractional shares were issued from such aggregation of common stock, upon the reverse split; any fractional share was rounded up and converted to the nearest whole share of common stock. As a result, 40,333,758 of the Company's common stock were exchanged for 16,133,544 of the Company's common stock resulting in the transfer of \$24,200 from common stock to additional paid in capital. These consolidated financial statements have been retroactively restated to reflect the reverse stock split.

The Company is authorized to issue 200,000,000 shares of \$0.001 par value common stock. As of March 31, 2019 and December 31, 2018, the Company had 20,009,985 and 16,868,783 shares issued and outstanding, respectively.

During the three months ended March 31, 2019, the Company issued an aggregate of 560,000 shares of its common stock for services totaling \$2,333,200 (\$4.17 per share).

During the three months ended March 31, 2019, the Company issued an aggregate of 23,332 shares of its common stock for vested restricted stock units.

During the three months ended March 31, 2019, the Company entered into securities purchase agreements with investors pursuant to which the Company issued 2,155,127 shares of common stock for aggregate proceeds of \$8,619,278, net of \$1,230 in expenses

During the three months ended March 31, 2019, the Company issued 298,319 shares of common stock in exchange for proceeds of \$1,150,780 from the exercise of warrants.

During the three months ended March 31, 2019, the Company issued 104,424 shares of common stock in exchange for the exercise of 210,144 cashless exercises of warrants.

At March 31, 2019, the Company had received proceeds of \$309,000 for the exercise of warrants.

**NOTE 9 – OPTIONS, RESTRICTED STOCK UNITS AND WARRANTS**

*Options*

On October 19, 2012, the Company's Board of Directors approved the 2012 Equity Incentive Plan ("the "Plan) and terminated the Long-Term Incentive Plan (the "2011 Plan"). The Plan provides for the issuance of options to purchase up to 15,186,123 (as amended) shares of the Company's common stock to officers, directors, employees and consultants of the Company (as amended). Under the terms of the Plan the Company may issue Incentive Stock Options as defined by the Internal Revenue Code to employees of the Company only and nonstatutory options. The Board of Directors of the Company or a committee thereof administers the Plan and determines the exercise price, vesting and expiration period of the grants under the Plan.

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However, the exercise price of an Incentive Stock Option should not be less than 110% of fair value of the common stock at the date of the grant for a 10% or more stockholder and 100% of fair value for a grantee who is not 10% stockholder. The fair value of the common stock is determined based on the quoted market price or in absence of such quoted market price, by the administrator in good faith.

Additionally, the vesting period of the grants under the Plan will be determined by the administrator, in its sole discretion, with an expiration period of not more than ten years. The Company reserved 910,346 shares of its common stock for future issuance under the terms of the Plan.

During the three months ended March 31, 2019, the Company granted an aggregate of 805,000 options to officers, directors and key consultants.

The following table presents information related to stock options at March 31, 2019:

<b>Options Outstanding</b>			<b>Options Exercisable</b>	
<b>Exercise Price</b>	<b>Number of Options</b>	<b>Weighted Average Remaining Life In Years</b>	<b>Exercisable Number of Options</b>	
\$ 2.51-5.00	1,663,218	8.1	1,163,218	
5.01-7.500	2,157,610	4.3	1,759,294	
7.51-10.00	120,000	6.0	120,000	
	<u>3,940,828</u>	6.0	<u>3,042,512</u>	

A summary of the stock option activity and related information for the 2012 Plan for the three months ended March 31, 2019 is as follows:

	<b>Shares</b>	<b>Weighted-Average Exercise Price</b>	<b>Weighted-Average Remaining Contractual Term</b>	<b>Aggregate Intrinsic Value</b>
Outstanding at December 31, 2018	3,135,828	\$ 5.34	5.2	\$ 311,545
Grants	805,000	4.90	10.0	-
Exercised	-			
Canceled	-			
Outstanding at March 31, 2019	<u>3,940,828</u>	\$ 5.25	6.0	\$ 4,162,880
Exercisable at March 31, 2019	3,042,512	\$ 5.36	4.8	\$ 3,020,099

The aggregate intrinsic value in the preceding tables represents the total pretax intrinsic value, based on options with an exercise price less than the Company's stock price of \$6.14 as of March 31, 2019, which would have been received by the option holders had those option holders exercised their options as of that date.

Option valuation models require the input of highly subjective assumptions. The fair value of stock-based payment awards was estimated using the Black-Scholes option model with a volatility figure derived from an index of historical stock prices of comparable entities until sufficient data exists to estimate the volatility using the Company's own historical stock prices. Management determined this assumption to be a more accurate indicator of value. The Company accounts for the expected life of options based on the contractual life of options for non-employees.



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For employees, the Company accounts for the expected life of options in accordance with the “simplified” method, which is used for “plain-vanilla” options, as defined in the accounting standards codification. The risk-free interest rate was determined from the implied yields of U.S. Treasury zero-coupon bonds with a remaining life consistent with the expected term of the options. The fair value of stock-based payment awards during the three months ended March 31, 2019 was estimated using the Black-Scholes pricing model.

On January 22, 2019, the Company granted an aggregate of 460,000 options to purchase the Company stock in connection with the services rendered at the exercise price of \$4.33 per share for a term of ten years with vesting quarterly beginning April 1, 2019 over 3 years

On March 14, 2019, the Company granted an aggregate of 345,000 options to purchase the Company stock in connection with the services rendered at the exercise price of \$5.66 per share for a term of ten years with 150,000 options vesting at anniversary date beginning March 14, 2020 over 3 years, 175,000 options vesting quarterly beginning June 14, 2019 over 3 years and 20,000 options vesting at one year anniversary.

The following assumptions were used in determining the fair value of options during the three months ended March 31, 2019:

Risk-free interest rate	2.53% - 2.74%
Dividend yield	0%
Stock price volatility	90.73% to 91.55%
Expected life	6 – 10 years
Weighted average grant date fair value	\$ 4.244

The fair value of all options vesting during the three months ended March 31, 2019 and 2018 of \$193,234 and \$246,710, respectively, was charged to current period operations. Unrecognized compensation expense of \$3,472,040 and \$173,446 at March 31, 2019 and December 31, 2018, respectively, will be expensed in future periods.

#### *Restricted Stock*

The following table summarizes the restricted stock activity for the three months ended March 31, 2019:

Total restricted shares issued as of December 31, 2018	-
Granted	190,000
Vested and issued	(23,332)
Vested restricted shares as of March 31, 2019	-
Unvested restricted shares as of March 31, 2019	<u>166,668</u>

On February 28, 2019, the Company granted an aggregate of 70,000 restricted stock grants for services with 23,332 vested immediately; 23,334 vesting at one year anniversary and 23,334 vesting at two year anniversary.

On March 20, 2019, the Company granted an aggregate of 120,000 restricted stock grants for services vesting quarterly beginning on April 1, 2019 over one year.

Stock based compensation expense related to restricted stock grants was \$143,581 and \$0 for the three months ended March 31, 2019 and 2018, respectively. As of March 31, 2019, the stock-based compensation relating to restricted stock of \$1,063,717 remains unamortized.

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*Warrants*

The following table summarizes information with respect to outstanding warrants to purchase common stock of the Company at March 31, 2019:

Exercise Price	Number Outstanding	Expiration Date
\$ 0.0025	153,328	January 2020
\$ 3.75	1,977,295	April 2019 to August 2021
\$ 4.375	633,272	May 2021
\$ 4.60	12,294	January 2020
\$ 4.875	352,943	April 2019 to September 2019
\$ 5.05	12,227	January 2020
\$ 6.85	217,958	July 2021 to August 2021
\$ 6.875	91,504	August 2019 to September 2019
\$ 9.375	536,267	April 2019 to March 2020
	<u>3,987,088</u>	

A summary of the warrant activity for the three months ended March 31, 2019 is as follows:

	Shares	Weighted-Average Exercise Price	Weighted-Average Remaining Contractual Term	Aggregate Intrinsic Value
Outstanding at December 31, 2018	4,579,511	\$ 4.73	1.5	\$ 1,924,388
Grants	-			
Exercised	(508,463)	\$ 3.81		
Expired	(83,960)	\$ 6.61	-	-
Outstanding at March 31, 2019	3,987,088	\$ 4.81	1.2	\$ 7,263,244
Vested and expected to vest at March 31, 2019	3,987,088	\$ 4.81	1.2	\$ 7,263,244
Exercisable at March 31, 2019	3,987,088	\$ 4.81	1.2	\$ 7,263,244

The aggregate intrinsic value in the preceding tables represents the total pretax intrinsic value, based on options with an exercise price less than the Company's stock price of \$6.14 of March 31, 2019, which would have been received by the option holders had those option holders exercised their options as of that date.

**NOTE 10 – RELATED PARTY TRANSACTIONS**

The Company's President and shareholders have advanced funds to the Company for working capital purposes since the Company's inception in February 2009. No formal repayment terms or arrangements exist and the Company is not accruing interest on these advances. The net amount of outstanding advances at March 31, 2019 and December 31, 2018 was \$-0-.

At March 31, 2019 and December 31, 2018, the Company had reimbursable travel and other related expenses due related parties of \$14,632 and \$32,366, respectively.

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On November 1, 2017, in connection with Mr. Filler joining the Company's Board of Directors, the Company entered into a Master Services Agreement (the "Agreement") with 3LP Advisors LLC (d/b/a Sherpa Technology Group) ("Sherpa") and an initial statement of work (the "SOW"), pursuant to which Sherpa will develop, execute and expand the Company's intellectual property strategy over the course of the next approximately 18 months by evaluating the business and technology landscape in which the Company operates, and charting and executing a strategy of patent filing and licensing. In connection with the SOW, the Company will pay Sherpa fee of (i) \$200,000 in cash, of which \$25,000 will be paid on January 1, 2018, with the remainder to be paid upon completion of certain objectives, and (ii) a ten-year option to purchase up to 120,000 of the Company's common stock at an exercise of \$3.75 per share of common stock, of which 60,000 options vest immediately and 60,000 options are performance conditioned. Mr. Filler is the general counsel and partner of Sherpa.

During the three months ended March 31, 2019 and 2018, the Company paid \$75,000 and \$175,000 as patent costs, consulting fees and expense reimbursements. As of March 31, 2019 and December 31, 2018, there was an unpaid balance of \$0.

**NOTE 11 – FAIR VALUE MEASUREMENT**

The Company adopted the provisions of Accounting Standards Codification subtopic 825-10, Financial Instruments ("ASC 825-10"). ASC 825-10 defines fair value as the price that would be received from selling an asset or paid to transfer a liability in an orderly transaction between market participants at the measurement date. When determining the fair value measurements for assets and liabilities required or permitted to be recorded at fair value, the Company considers the principal or most advantageous market in which it would transact and considers assumptions that market participants would use when pricing the asset or liability, such as inherent risk, transfer restrictions, and risk of nonperformance. ASC 825-10 establishes a fair value hierarchy that requires an entity to maximize the use of observable inputs and minimize the use of unobservable inputs when measuring fair value. ASC 825-10 establishes three levels of inputs that may be used to measure fair value:

Level 1 – Quoted prices in active markets for identical assets or liabilities.

Level 2 – Observable inputs other than Level 1 prices such as quoted prices for similar assets or liabilities; quoted prices in markets with insufficient volume or infrequent transactions (less active markets); or model-derived valuations in which all significant inputs are observable or can be derived principally from or corroborated by observable market data for substantially the full term of the assets or liabilities.

Level 3 – Unobservable inputs to the valuation methodology that are significant to the measurement of fair value of assets or liabilities.

All items required to be recorded or measured on a recurring basis are based upon level 3 inputs.

To the extent that valuation is based on models or inputs that are less observable or unobservable in the market, the determination of fair value requires more judgment. In certain cases, the inputs used to measure fair value may fall into different levels of the fair value hierarchy. In such cases, for disclosure purposes, the level in the fair value hierarchy within which the fair value measurement is disclosed and is determined based on the lowest level input that is significant to the fair value measurement.

The carrying value of the Company's cash and cash equivalents, accounts payable and other current assets and liabilities approximate fair value because of their short-term maturity.

As of March 31, 2019 and December 31, 2018, the Company did not have any items that would be classified as level 1 or 2 disclosures.

As of March 31, 2019 and December 31, 2018, the Company did not have any derivative instruments that were designated as hedges.

There were no derivative and warrant liability as of March 31, 2019 and December 31, 2018.

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**NOTE 12 – SUBSEQUENT EVENTS**

In April 2019, the Company issued an aggregate of 167,327 shares of its common stock in exchange for warrant subscriptions received of \$309,000 and proceeds of \$416,719 from the exercise of warrants.

On April 2, 2019, the Company issued 244 shares of its common stock in exchange for the cashless exercise of 600 warrants.

In April 2019, the Company issued an aggregate of 30,000 shares of its common stock for vested restricted stock units.

*Series C conversions:*

On April 1, 2019 the Company issued 3,507 shares of its common stock in exchange for 10 shares of Series C preferred stock and accrued dividends.

## ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

*This Management's Discussion and Analysis of Financial Condition and Results of Operations includes a number of forward-looking statements that reflect Management's current views with respect to future events and financial performance. You can identify these statements by forward-looking words such as "may," "will," "expect," "anticipate," "believe," "estimate" and "continue," or similar words. Those statements include statements regarding the intent, belief or current expectations of us and members of our management team as well as the assumptions on which such statements are based. Prospective investors are cautioned that any such forward-looking statements are not guarantees of future performance and involve risk and uncertainties, and that actual results may differ materially from those contemplated by such forward-looking statements.*

*Readers are urged to carefully review and consider the various disclosures made by us in this report and in our other reports filed with the Securities and Exchange Commission. Important factors currently known to Management could cause actual results to differ materially from those in forward-looking statements. We undertake no obligation to update or revise forward-looking statements to reflect changed assumptions, the occurrence of unanticipated events or changes in the future operating results over time. We believe that our assumptions are based upon reasonable data derived from and known about our business and operations. No assurances are made that actual results of operations or the results of our future activities will not differ materially from our assumptions. Factors that could cause differences include, but are not limited to, expected market demand for our products, fluctuations in pricing for materials, and competition.*

### Business Overview

We are a development stage medical device company that is developing a proprietary biomedical signal processing technology platform to extract information from physiologic signals. Our initial emphasis is on providing intracardiac signal information to electrophysiologists during electrophysiology ("EP") studies and cardiac catheter ablation of atrial fibrillation ("AF") and ventricular tachycardia ("VT"). Cardiac catheter ablation is a procedure that involves delivery of energy through the tip of a catheter that scars or destroys heart tissue in order to correct heart rhythm disturbances. Our first product which received FDA 510(k) clearance in August 2018 is the PURE (Precise Uninterrupted Real-time evaluation of Electrograms) EP System.

The PURE EP™ System is a non-invasive computerized system intended for acquiring, digitizing, amplifying, filtering, measuring and calculating, displaying, recording and storing of electrocardiographic and intracardiac signals for patients undergoing EP procedures in an EP laboratory. The system is indicated for use under the supervision of licensed healthcare practitioners who are responsible for interpreting the data. The PURE EP System aims to minimize noise and artifacts and acquire high-fidelity cardiac signals. Improving cardiac signals may potentially increase the diagnostic value of these signals, thereby possibly improving accuracy and efficiency of the EP studies and related procedures. The PURE EP System is intended to be used in addition to existing electrophysiology recorders. We believe that data provided by the PURE EP System will increase the workload ability and enhance the capabilities of the typical electrophysiology laboratory.

Our initial focus is on improving intracardiac signal acquisition and enhance diagnostic information for catheter ablation procedures for the complex arrhythmias, atrial fibrillation, the most common cardiac arrhythmia, and ventricular tachycardia, an arrhythmia evidenced by a fast heart rhythm originating from the lower chambers of the heart, which can be life-threatening. Cardiac catheter ablation is a procedure that corrects conduction of electrical impulses in the heart that cause arrhythmias and is now a preferred treatment for certain arrhythmias. During this procedure, a catheter is usually inserted using a venous access into a specific area of the heart. Cryo or radiofrequency energy is delivered through the catheter to destroy small areas of the heart muscle that cause the abnormal heart rhythm. According to the 2017 HRS/EHRA/ECAS/APHRS/SOLAECE Expert Consensus Statement on Catheter and Surgical Ablation of Atrial Fibrillation, the role of catheter ablation as first-line therapy, prior to a trial of a Class I or III antiarrhythmic agent, is an appropriate indication for catheter ablation of AF in patients with symptomatic paroxysmal or persistent AF.

Catheter ablation for many arrhythmias have high success rates; however, more complex or long-standing examples of the disease (like recurrent AF and VT) often require multiple procedures (each typically lasting from 3-6 hours), evidencing the need for additional research and technology to help diagnose and treat these cases. Consequently, ablating AF and VT is regarded as being more difficult. Therefore, access to these procedures has traditionally been limited to being performed by only the most well-trained electrophysiologists.

Our overall goal is to establish the PURE EP System as a new platform in the EP market. We believe that the PURE EP System and its signal processing tools will possibly improve patient outcomes because we believe that the PURE EP System may have the following advantages over the EP recording systems currently available on the market:

- Precise, uninterrupted, real-time evaluations of electrograms;
- Higher quality cardiac signal acquisition for accurate and more efficient electrophysiology studies and catheter ablation procedures to help reduce costs and length of procedures;
- Reliable display of information to better determine precise ablation targets, strategy and end point of procedures with the objective of reducing the need for multiple procedures;
- A device that can run in parallel with the existing EP lab equipment.

On February 18 and February 19, 2019, we conducted the first clinical cases with our PURE EP™ System which was announced on February 20, 2019. The patient cases were performed by Andrea Natale, M.D., F.A.C.C., F.H.R.S., F.E.S.C., Executive Medical Director, Texas Cardiac Arrhythmia Institute at St. David's Medical Center. Initial results showed improved signal detection and fidelity compared to the data acquired using the existing recording devices in the EP lab. We intend to conduct additional clinical external evaluation at a select number of centers.

We also intend to continue additional research studies of our technology at Mayo Clinic. On November 13, 2018, we announced that we entered into a new advanced research agreement with Mayo Clinic. The program will be run under the leadership of Samuel J. Asirvatham, M.D., Mayo Clinic's Vice-Chair of Innovation and Medical Director, Electrophysiology Laboratory and will consist of a number of two- to three-year projects, which will focus on development of additional advanced features of PURE EP™ System within the field of EP and potential clinical applications of our technology in a new, previously unexplored, field.

To date, we have conducted a total of sixteen pre-clinical studies with the PURE EP™ System prototype, fifteen of which were conducted at Mayo Clinic in Rochester, Minnesota. We also conducted a pre-clinical study at the Mount Sinai Hospital in New York, NY with emphasis on the VT model.

Over the three months ended March 31, 2019, our other significant achievements include:

- On January 21, 2019, Dr. Asirvatham's team at Mayo performed our fourteenth pre-clinical study at Mayo Clinic in Rochester, Minnesota which was part of the new agreement announced November 13, 2018.
- On February 5, 2019, we announced that John Kowalski, former Biosense Webster (a Johnson & Johnson company) Northeast Area Director, joined BioSig as Vice President of Sales.
- On February 12, 2019, we issued a 2019 Shareholder Letter which provided updates on our recent business development and highlighted our plans for future growth.
- On February 26, 2019, we announced expansion of our engineering team to drive R&D and manufacturing efforts.
- On March 6, 2019, we announced the appointment of Barry Keenan Ph.D., MBA, PMP as Vice President of Engineering to head up our advanced product development.

- On April 3, 2019, Dr. Asirvatham's team at Mayo performed our fifteenth pre-clinical study at Mayo Clinic in Rochester, Minnesota.

We continue to work with JK Advisors to scale up operational activities; Quintain Project Solutions LLC as the manufacturing project management leader for the PURE EP System; and with our Regulatory and Compliance Advisor to move the development process forward. These relationships have aided us with our first 510(k) clearance from the U.S. Food and Drug Administration for the PURE EP System in August 2018. Our manufacturing partner, Minnetronix, a medical technology and innovation company, has built initial beta units for our first installations, clinical procedures, and IP proposals.

We are currently working on audit preparation for the International Organization for Standardization ("ISO") and Medical Device Single Audit Program ("MDSAP") to ensure that our product meets specific requirements - audit submission is targeted for Q2 2019. We believe that we will have obtained ISO Certification and product CE Mark by the fourth quarter of 2019 and Japanese approval by the end of 2019.

Because we have not yet entered the commercialization stage, with our initial product beginning clinical trials, we currently do not have any customers. We anticipate that our initial customers will be hospitals and other health care facilities that operate electrophysiology labs.

## Results of Operations

We anticipate that our results of operations will fluctuate for the foreseeable future due to several factors, such as the progress of our commercialization efforts and the timing and outcome of future regulatory submissions. Due to these uncertainties, accurate predictions of future operations are difficult or impossible to make.

### *Three Months Ended March 31, 2019 Compared to Three Months Ended March 31, 2018*

*Revenues and Cost of Goods Sold.* We had no revenues or cost of goods sold during the three months ended March 31, 2019 and 2018.

*Research and Development Expenses.* Research and development expenses for the three months ended March 31, 2019 were \$1,488,839, an increase of \$626,419, or 72.6%, from \$862,420 for the three months ended March 31, 2018. This increase is primarily due to increase in compensation and consulting work with us adding personnel and advisors along with increases in stock based compensation in 2019, as compared to 2018, as we finalize our initial product towards commercialization. Research and development expenses were comprised of the following:

Three months ended:

	March 31, 2019	March 31, 2018
Salaries and equity compensation	\$ 681,632	\$ 287,849
Consulting expenses	230,263	131,601
Research studies and design work	536,196	419,325
Travel, supplies, other	40,748	23,645
Total	<u>\$ 1,488,839</u>	<u>\$ 862,420</u>

Stock based compensation for research and development personnel was \$428,747 and \$98,589 for the three months ended March 31, 2019 and 2018, respectively.

*General and Administrative Expenses.* General and administrative expenses for the three months ended March 31, 2019 were \$4,378,897, an increase of \$2,607,101, or 147.1%, from \$1,771,796 incurred in the three months ended March 31, 2018. This increase is primarily due to an increase in employee performance pay and staff in the current period as compared to the same period in the prior year and additional service provider fees paid.

Payroll related expenses increased to \$2,923,770 in the current period from \$534,627 for the three months ended March 31, 2018, an increase of \$2,389,143. The increase was due to performance pay and added staff in 2019 for commercialization and support personnel. We incurred \$2,241,268 in stock based compensation in connection with the vesting of stock and stock options issued to board members, officers, employees and consultants for the three months ended March 31, 2019 as compared to \$290,121 in stock based compensation for the same period in 2018.

Professional services for the three months ended March 31, 2019 totaled \$157,807, a decrease of \$51,115, or 24.5%, over the \$208,922 recognized for the three months ended March 31, 2018. Of professional services, legal fees totaled \$110,307 for the three months ended March 31, 2019, a decrease of \$51,615, or 31.9%, from \$161,922 incurred for the three months ended March 31, 2018. The primary decrease was due to costs incurred in registration statement and patent filings in 2018 as compared to 2019. Accounting fees incurred in the three months ended March 31, 2019 amounted to \$47,500, an increase of \$500, or 1.1%, from \$47,000 incurred in same period last year.

Consulting, public and investor relations fees for the three months ended March 31, 2019 were \$598,845 as compared to \$532,050 incurred for the three months ended March 31, 2018. The increase in consulting and investor relations fees during the three months ended March 31, 2019 related to our continued efforts to develop our recognition throughout the medical industry in an effective manner.

Travel, meals and entertainment costs for the three months ended March 31, 2019 were \$125,864, an increase of \$34,943, or 38.4%, from \$90,921 incurred in the three months ended March 31, 2018. Travel, meals and entertainment costs include travel related to business development and financing. The increase in 2019 was due to added commercialization and business development efforts as compared to 2018.

Rent for the three months ended March 31, 2019 totaled \$60,127, an increase of \$25,274 or 72.5%, from \$34,853 incurred in three months ended March 31, 2018. The increase in rent for 2019 as compared to 2018 is due primarily expanding our Los Angeles office, adding an administrative center in Austin, Texas and a Norwalk, CT office.

*Depreciation and amortization Expense.* Depreciation and amortization expense for the three months ended March 31, 2019 totaled \$7,935 an increase of \$5,032, or 173.3%, over the expense of \$2,903 incurred in the three months ended March 31, 2018, as a result of the adding additional office computers and other equipment. In addition, we begun amortizing our incurred patent costs during the three months ended March 31, 2019.

*Preferred Stock Dividend.* Preferred stock dividend for the three months ended March 31, 2019 totaled \$10,541, a decrease of \$294,505, or 96.5% from \$305,046 incurred during the three months ended March 31, 2018, 2017. Preferred stock dividends are primarily related to the issuance of our Series C, D and E Preferred Stock from 2013 through 2018. The significant decrease in 2019 as compared to 2018 is the result of conversions in 2018 of the Series D Preferred Stock and the payment, upon conversion, of a required minimum dividend of \$405 per share of Series D Preferred Stock for the first three years of issuance.

*Net Loss available to common shareholders.* As a result of the foregoing, net loss available to common shareholders for the three months ended March 31, 2019 was \$5,880,089 compared to a net loss of \$2,942,080 for the three months ended March 31, 2018.

## **Liquidity and Capital Resources**

### *Three Months Ended March 31, 2019 Compared to three Months Ended March 31, 2018*

As of March 31, 2019, we had a working capital of \$10,080,879, comprised of cash of \$10,933,593 and prepaid expenses of \$156,577, which was offset by \$597,006 of accounts payable and accrued expenses, accrued dividends on preferred stock issuances of \$253,449 and current portion of lease liability of \$158,836. For the three months ended March 31, 2019, we used \$3,522,601 of cash in operating activities and \$73,024 of cash in investing activities.



Cash provided by financing activities totaled \$10,079,058, comprised of proceeds from the sale of our common stock of \$8,619,278 and proceeds from exercise of warrants of \$1,459,780. In the comparable period in 2018, our aggregate cash provided by financing activities totaled \$1,878,439 comprised proceeds from the sale of our common stock and common stock subscriptions of \$385,470 and proceeds from the sale of our Series E preferred stock of \$1,492,969. At March 31, 2019, we had cash of \$10,933,593 compared to \$1,035,100 at March 31, 2018. Our cash is held in bank deposit accounts. At March 31, 2019 and March 31, 2018, we had no convertible debentures outstanding.

Cash used in operations for the three months ended March 31, 2019 and 2018 was \$3,522,601 and \$2,387,274, respectively, which represent cash outlays for research and development and general and administrative expenses in such periods. The increases in cash outlays principally resulted from additional operating costs and general and administrative expenses and a decrease in our operating assets of \$21,865 and decrease our operating liabilities of \$352,868, net of stock based compensation.

We used \$73,024 cash for investing activities for the three months ended March 31, 2019, compared to \$3,644 for the three months ended March 31, 2018. For the current period, we purchased computer and other equipment of \$14,422 and paid \$58,327 and \$275 in patent and trademark costs, respectively, as compared to \$3,644 in 2018 to purchase computer and other equipment.

In their report dated March 15, 2019, our independent registered public accounting firm stated at December 31, 2018, there is substantial doubt about our ability to continue as a going concern. Our ability to continue as a going concern is an issue raised due to our net losses and negative cash flows from operations since inception and our expectation that these conditions will continue for the foreseeable future. In addition, we will require additional financing to fund future operations.

Further, we do not have any commercial products available for sale and have not generated revenues to date, and there is no assurance that we will be able to generate cash flow to fund operations. In addition, there can be no assurance that our research and development will be successfully completed or that any product will be approved or commercially viable. Our ability to continue as a going concern is subject to our ability to obtain necessary funding from outside sources, including obtaining additional funding from the sale of our securities, obtaining loans from various financial institutions or being awarded grants from government agencies, where possible. Our continued net operating losses increase the difficulty in meeting such goals and there can be no assurances that such methods will prove successful.

We expect to incur losses from operations for the near future. We expect to incur increasing research and development expenses, including expenses related to clinical and research trials. We expect that our general and administrative expenses will increase in the future as we expand our business development, add infrastructure and incur additional costs related to being a public company, including incremental audit fees, investor relations programs and increased professional services.

Our future capital requirements will depend on a number of factors, including the progress of our research and development of product candidates, the timing and outcome of regulatory approvals, the costs involved in preparing, filing, prosecuting, maintaining, defending and enforcing patent claims and other intellectual property rights, the status of competitive products, the availability of financing and our success in developing markets for our product candidates. We believe our existing cash will not be sufficient to fund our operating expenses and capital equipment requirements. We anticipate we will need approximately \$4 million in addition to our current cash on hand to fund our operating expenses and capital equipment requirements for the next 12 months.

We will have to raise additional funds to continue our operations and, while we have been successful in doing so in the past, there can be no assurance that we will be able to do so in the future. Our continuation as a going concern is dependent upon our ability to obtain necessary additional funds to continue operations and the attainment of profitable operations.

Future financing may include the issuance of equity or debt securities, obtaining credit facilities, or other financing mechanisms. Even if we are able to raise the funds required, it is possible that we could incur unexpected costs and expenses or experience unexpected cash requirements that would force us to seek alternative financing. Furthermore, if we issue additional equity or debt securities, existing holders of our securities may experience additional dilution or the new equity securities may have rights, preferences or privileges senior to those of existing holders of our securities.

If additional financing is not available or is not available on acceptable terms, we may be required to delay, reduce the scope of or eliminate our research and development programs, reduce our commercialization efforts or obtain funds through arrangements with collaborative partners or others that may require us to relinquish rights to certain product candidates that we might otherwise seek to develop or commercialize independently.

#### **Off-Balance Sheet Arrangements**

We do not have any off-balance sheet arrangements.

#### **Critical Accounting Policies and Estimates**

Our discussion and analysis of our financial condition and results of operations are based on our financial statements, which have been prepared in accordance with accounting principles generally accepted in the United States of America. The preparation of these financial statements requires us to make estimates and judgments that affect the reported amounts of assets, liabilities and expenses. We evaluate our estimates and judgments on an ongoing basis. We base our estimates on historical experience and on assumptions that we believe to be reasonable under the circumstances, the results of which form the basis for making judgments about the carrying values of assets and liabilities that are not readily apparent from other sources. Actual results may differ from these estimates.

We believe the following critical accounting policies affect our more significant judgments and estimates used in the preparation of our financial statements.

##### *Research and Development.*

We account for research and development costs in accordance with the Accounting Standards Codification subtopic 730-10, Research and Development (“ASC 730-10”). Under ASC 730-10, all research and development costs must be charged to expense as incurred. Accordingly, internal research and development costs are expensed as incurred. Third-party research and development costs are expensed when the contracted work has been performed or as milestone results have been achieved. Company-sponsored research and development costs related to both present and future products are expensed in the period incurred.

##### *Stock Based Compensation.*

All stock-based payments to employees and to nonemployee directors for their services as directors consisted of grants of restricted stock and stock options, which are measured at fair value on the grant date and recognized in the statements of operations as compensation expense over the relevant vesting period. Restricted stock payments and stock-based payments to nonemployees are recognized as an expense over the period of performance.

Such payments are measured at fair value at the earlier of the date a performance commitment is reached or the date performance is completed. In addition, for awards that vest immediately and are non-forfeitable, the measurement date is the date the award is issued.

On October 29, 2014, our common stock commenced trading on OTCQB and on September 21, 2018 on the NASDAQ Capital Market under the symbol “BSGM.” Fair value is typically determined by the closing price of our common stock on the date of the award.

*Income Taxes.*

Deferred income tax assets and liabilities are determined based on the estimated future tax effects of net operating loss and credit carryforwards and temporary differences between the tax basis of assets and liabilities and their respective financial reporting amounts measured at the current enacted tax rates. We record an estimated valuation allowance on our deferred income tax assets if it is not more likely than not that these deferred income tax assets will be realized. We recognize a tax benefit from an uncertain tax position only if it is more likely than not that the tax position will be sustained on examination by taxing authorities, based on the technical merits of the position. The tax benefits recognized in the financial statements from such a position are measured based on the largest benefit that has a greater than 50% likelihood of being realized upon ultimate settlement.

**ITEM 3. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK**

Not required under Regulation S-K for “smaller reporting companies.”

**ITEM 4. CONTROLS AND PROCEDURES**

**Evaluation of Disclosure Controls and Procedures**

As required under Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934, as amended (the “Exchange Act”), we carried out an evaluation under the supervision and with the participation of our management, including our Chief Executive Officer and Chief Financial Officer, of the effectiveness of the design and operation of our disclosure controls and procedures as of March 31, 2019. Based upon that evaluation, the Chief Executive Officer and Chief Financial Officer identified material weaknesses in internal control over financial reporting as previously disclosed under Item 9A. “Controls and Procedures” in our Annual Report on Form 10-K for our fiscal year ended December 31, 2018, which have not been remediated, and therefore concluded that our disclosure controls and procedures as of March 31, 2019 were not effective.

**Changes in Internal Controls over Financial Reporting**

There have been no changes in our internal controls over financial reporting (as defined in Rule 13a-15(f) and 15d-15(f) of the Exchange Act) that occurred during the last fiscal quarter to which this report relates that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

**PART II – OTHER INFORMATION**

**ITEM 1. LEGAL PROCEEDINGS**

None.

**ITEM 1A. RISK FACTORS**

Not required under Regulation S-K for “smaller reporting companies.”

**ITEM 2. UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS**

On March 12, 2019, we entered into a Securities Purchase Agreement (the “Purchase Agreement”) with certain accredited investors (the “Investors”), pursuant to which the Company sold to the Investors an aggregate of 2,155,127 shares of the Company’s common stock in exchange for aggregate consideration of \$8,619,278, net of transaction expenses of \$1,230. The securities sold in this offering were not registered under the Securities Act of 1933, as amended, or the securities laws of any state, and were offered and sold in reliance on the exemption from registration under the Securities Act of 1933, as amended, provided by Section 4(2) and Regulation D (Rule 506) under the Securities Act of 1933, as amended.

**ITEM 3. DEFAULTS UPON SENIOR SECURITIES**

None.

**ITEM 4. MINE SAFETY DISCLOSURES**

None.

**ITEM 5. OTHER INFORMATION**

None.

**ITEM 6. EXHIBITS**

10.1	<a href="#">Securities Purchase Agreement dated as of March 12, 2019, by and between BioSig Technologies, Inc. and certain purchasers set forth therein (incorporated by reference to Exhibit 10.1 to the Form 8-K filed on March 14, 2019)</a>
31.01	<a href="#">Certification of Chief Executive Officer pursuant to Exchange Act Rules 13a-14(a) and 15d-14(a), as adopted pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.</a>
31.02	<a href="#">Certification of Chief Financial Officer pursuant to Exchange Act Rules 13a-14(a) and 15d-14(a), as adopted pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.</a>
32.01	<a href="#">Certifications of Chief Executive Officer and Chief Financial Officer pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.</a>
101 INS	XBRL Instance Document
101 SCH	XBRL Taxonomy Extension Schema Document
101 CAL	XBRL Taxonomy Calculation Linkbase Document
101 DEF	XBRL Taxonomy Extension Definition Linkbase Document
101 LAB	XBRL Taxonomy Labels Linkbase Document
101 PRE	XBRL Taxonomy Presentation Linkbase Document

**SIGNATURES**

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned thereunto duly authorized.

**BIOSIG TECHNOLOGIES, INC.**

Date: April 15, 2019

By: /s/ KENNETH L. LONDONER  
Kenneth L. Londoner  
Chairman & Chief Executive Officer (Principal Executive Officer)

Date: April 15, 2019

By: /s/ STEVEN CHAUSSY  
Steven Chaussy  
Chief Financial Officer (Principal Accounting Officer)

CERTIFICATION

I, Kenneth L. Londoner, certify that:

1. I have reviewed this quarterly report on Form 10-Q of BioSig Technologies, Inc.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
  - (a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
  - (b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
  - (c) Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
  - (d) Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
5. The registrant's other certifying officer and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
  - (a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonable likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
  - (b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal controls over financial reporting.

Date: April 15, 2019

/s/ KENNETH L. LONDONER

Kenneth L. Londoner

Chairman & Chief Executive Officer (Principal Executive Officer)

CERTIFICATION

I, Steven Chaussy, certify that:

1. I have reviewed this quarterly report on Form 10-Q of BioSig Technologies, Inc.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
  - (a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
  - (b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
  - (c) Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
  - (d) Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
5. The registrant's other certifying officer and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
  - (a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonable likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
  - (b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal controls over financial reporting.

Date: April 15, 2019

/s/ STEVEN CHAUSSY

Steven Chaussy

Chief Financial Officer (Principal Accounting Officer)

**CERTIFICATIONS OF CHIEF EXECUTIVE OFFICER AND CHIEF FINANCIAL OFFICER  
PURSUANT TO  
18 U.S.C. SECTION 1350,  
AS ADOPTED PURSUANT TO  
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002**

I, Kenneth L. Londoner, certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that the Quarterly Report of BioSig Technologies, Inc. on Form 10-Q for the fiscal quarter ended March 31, 2019 fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934 and that information contained in this Quarterly Report on Form 10-Q fairly presents in all material respects the financial condition and results of operations of BioSig Technologies, Inc.

Date: April 15, 2019

By: /s/ KENNETH L. LONDONER  
Name: Kenneth L. Londoner  
Title: *Chairman & Chief Executive Officer (Principal Executive Officer)*

I, Steven Chaussy, certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that the Quarterly Report of BioSig Technologies, Inc. on Form 10-Q for the fiscal quarter ended March 31, 2019 fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934 and that information contained in this Quarterly Report on Form 10-Q fairly presents in all material respects the financial condition and results of operations of BioSig Technologies, Inc.

Date: April 15, 2019

By: /s/ STEVEN CHAUSSY  
Name: Steven Chaussy  
Title: *Chief Financial Officer (Principal Accounting Officer)*