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

FORM 10-Q

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

For the quarterly period ended June 30, 2015

OR

[ ] TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

For the transition period to

Commission File No. 000-55364

HELIUS MEDICAL TECHNOLOGIES, INC.
(Exact name of Registrant as specified in its charter)

Wyoming	**36-4787690**
(State or other jurisdiction of incorporation or organization)	(I.R.S. Employer Identification Number)

Suite 400, 41 University Drive
Newton, Pennsylvania, 18940
(Address of principal executive office) (Zip Code)

(215) 809-2018
(Registrant's telephone number, including area code)

N/A
(Former name, former address and former fiscal year, if changed since last report)

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 [X]

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Web site, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit and post such files).
Yes [X] No [ ]

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

Large accelerated filer [ ]	Accelerated filer [ ]
Non-accelerated filer [X] (Do not check if a smaller reporting company)	Smaller reporting company [ ]

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

Indicate the number of shares outstanding of each of the issuer's classes of common stock, as of the latest practicable date.

Class	**Outstanding at August 14, 2015**
Class A Common Stock	64,969,775

HELIUS MEDICAL TECHNOLOGIES, INC.

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PART I — FINANCIAL INFORMATION

Item 1. Consolidated Financial Statements

HELIUS MEDICAL TECHNOLOGIES, INC.
INTERIM CONDENSED CONSOLIDATED FINANCIAL STATEMENTS
June 30, 2015
(Unaudited)
(Expressed in United States Dollars)

Helius Medical Technologies, Inc.
Interim Condensed Consolidated Balance Sheets
June 30, 2015 and March 31, 2015
(Unaudited)
(Expressed in United States Dollars)

	June 30, 2015 $	March 31, 2015 $
ASSETS		
Current assets		
Cash and cash equivalents	971,863	418,893
Short-term investment	-	378,000
Receivables	24,231	8,833
Prepaid expenses	434,245	410,621
Total current assets	1,430,339	1,216,347
TOTAL ASSETS	1,430,339	1,216,347
LIABILITIES		
Current liabilities		
Accounts payable and accrued liabilities	1,583,499	1,197,804
Total current liabilities	1,583,499	1,197,804
Derivative liability (Notes 2 and 5)	1,846,083	1,581,444
TOTAL LIABILITIES	3,429,582	2,779,248
CAPITAL DEFICIT		
Common stock (Unlimited Class A common shares authorized); (64,303,924 shares outstanding at June 30, 2015 and 63,104,788 shares outstanding at March 31, 2015) (Note 4)	18,792,322	16,358,093
Additional paid-in capital	1,713,992	1,490,790
Shares to be issued	-	39,545
Accumulated other comprehensive income	(931,014)	(971,640)
Accumulated deficit	(21,574,543)	(18,479,689)
TOTALCAPITAL DEFICIT	(1,999,243)	(1,562,901)
TOTAL LIABILITIES & CAPITAL DEFICIT	1,430,339	1,216,347

"Philippe Deschamps " Director *"Savio Chiu "* Director

(The accompanying notes are an integral part of these financial statements.)

Helius Medical Technologies Inc.
Interim Condensed Consolidated Statements of Comprehensive Loss
for the three months ended June 30, 2015 and 2014
(Unaudited)
(Expressed in United States Dollars)

	June 30, 2015 $	June 30, 2014 $
Operating Expenses		
Advertising, marketing & investor relations	308,213	46,226
Audit & accounting	80,462	10,645
Consulting fees (Note 7)	142,822	200,849
Insurance	29,539	7,486
Legal fees	293,736	219,934
Meals & travel	95,867	29,441
Office & general	26,039	21,424
Professional fees	-	13,125
Research & development	1,386,676	653,259
Transfer agent & regulatory	32,579	15,260
Wages and salaries	439,065	170,981
Loss from operations	(2,834,998)	(1,388,630)
Other items		
Interest expense	(53)	(176,488)
Interest income	-	2,810
Change in fair value of derivative liability (Note 2)	(178,378)	-
Foreign exchange loss	(81,425)	(6,446)
	(259,856)	(180,124)
Net loss for the period	(3,094,854)	(1,568,754)
Other comprehensive income		
Translation adjustments	40,626	106,059
Comprehensive loss for the period	(3,054,228)	(1,462,695)
Basic and diluted loss per common stock	(0.05)	(0.04)
Weighted average number of common stock outstanding – basic & diluted	63,722,378	38,812,706

(The accompanying notes are an integral part of these financial statements.)

Helius Medical Technologies Inc.
Interim Condensed Consolidated Statements of Capital Deficit
for the three months ended June 30, 2015
(Unaudited)
(Expressed in United States Dollars)

	Common Stock	Amount $	Additional Paid-In Capital $	Shares to be Issued $	Accumulated Deficit $	Accumulated other comprehensive income (loss) $	Capital (Deficit) $
Balance – March 31, 2014	**32,070,052**	**8,510,000**	**807,157**	-	**(9,585,134)**	-	**(267,977)**
Stock-based compensation on 2,300,000 options granted			50,303	-	-	-	50,303
Shares issued to consultant for option exercise (Note 6)	2,300,000	717	-	-	-	-	717
Shares issued to consultant for option exercise (Note 6)	930,031	290	-	-	-	-	290
Fair value of options allocated to share capital on exercise of options	-	857,460	(857,460)	-	-	-	-
Recapitalization of Helius Medical Technologies, Inc. (Note 3)	10,000,000	-	162,890	-	-	-	162,890
Issuance of common stock for private placement (Note 5)	15,240,000	6,437,041	578,961	-	-	-	7,016,002
Share issuance cost (Note 5)	-	(447,515)	67,709	-	-	-	(379,806)
Beneficial conversion feature (Note 4)	-	-	176,488	-	-	-	176,488
Stock-based compensation on 2,570,000 options granted (Note 6)	-	-	283,962	-	-	-	283,962
Conversion of debenture (Note 4)	2,564,705	1,000,100	-	-	-	-	1,000,100
Stock-based compensation on 100,000 options granted (Note 6)	-	-	74,190	-	-	-	74,190
Stock-based compensation on 100,000 options granted (Note 6)	-	-	43,229	-	-	-	43,229
Stock-based compensation on 400,000 options granted (Note 6)	-	-	135,564	-	-	-	135,564
Stock-based compensation on 100,000 options granted (Note 6)	-	-	41,987	-	-	-	41,987
Fair value of vested non-employee options reallocated to derivative liability	-	-	(74,190)	-	-	-	(74,190)
Private placement proceeds	-	-	-	39,545	-	-	39,545
Net loss for the year	-	-	-	-	(8,894,555)	-	(8,894,555)
Translation adjustments	-	-	-	-	-	(971,640)	(971,640)
Balance – March 31, 2015	**63,104,788**	**16,358,093**	**1,490,790**	**39,545**	**(18,479,689)**	**(971,640)**	**(1,562,901)**
Exercise of finder's warrants	14,400	11,926	-	-	-	-	11,926
Issuance of common stock for private placement	849,273	1,825,937	-	-	-	-	1,825,937
Issuance of common stock for private placement	335,463	721,243	-	(39,545)	-	-	681,698
Share issuance cost	-	(124,877)	-	-	-	-	(124,877)
Stock-based compensation on 2,570,000 options granted	-	-	173,899	-	-	-	173,899
Stock-based compensation on 400,000 options granted	-	-	135,564	-	-	-	135,564
Fair value of non-employee vested options reallocated to derivative liability	-	-	(86,261)	-	-	-	(86,261)
Net loss for the period	-	-	-	-	(3,094,854)	-	(3,094,854)
Translation adjustments	-	-	-	-	-	40,626	40,626
Balance – June 30, 2015	**64,303,924**	**18,792,322**	**1,713,992**	-	**(21,574,543)**	**(931,014)**	**(1,999,243)**

(The accompanying notes are an integral part of these financial statements.)

Helius Medical Technologies, Inc.
Interim Condensed Consolidated Statements of Cash Flows
for the three months ended June 30, 2015 and 2014
(Unaudited)
(Expressed in United States Dollars)

	June 30, 2015 $	June 30, 2014 $
Cash flows from operating activities		
Net loss for the period	(3,094,854)	(1,568,754)
Items not involving cash:		
Change in fair value of derivative liability	178,378	-
Stock-based compensation	309,463	338,291
Accretion	-	176,488
Changes in non-cash working capital items:		
Receivables	(15,264)	-
Prepaid expenses	(22,205)	(108,895)
Accounts payable and accrued liabilities	382,844	622,002
Foreign exchange re-measurement	115,638	-
Net cash used in operating activities	**(2,146,000)**	**(540,868)**
Cash flows from investing activities		
Short-term investment	378,000	-
Cash acquired on recapitalization	-	23,904
Proceeds from bridge loan	-	150,000
Net cash provided by investing activities	**378,000**	**173,904**
Cash flows from financing activities		
Proceeds from the issuance of shares	2,519,561	6,607,402
Share issuance costs	(124,877)	-
Proceeds from the issuance of convertible debt	-	633,195
Net cash provided by financing activities	**2,394,684**	**7,240,597**
Effect of foreign exchange rate changes on cash	(73,714)	140,658
Net change in cash	552,970	7,014,291
Cash, beginning of the period	418,893	15,968
Cash, end of the period	**971,863**	**7,030,259**
Supplemental cash flow information		
Interest paid in cash	-	-
Income taxes paid in cash	-	-

(The accompanying notes are an integral part of these financial statements.)

1. BASIS OF PRESENTATION

The accompanying unaudited condensed financial statements of Helius Medical Technologies Inc. (the "Company") have been prepared in accordance with U.S. generally accepted accounting principles ("GAAP") for interim financial information and in accordance with the instructions to Form 10-Q and Regulation S-X. Accordingly, they should be read in conjunction with the audited consolidated financial statements and notes thereto for the fiscal year ended March 31, 2015 included in the Annual Report on Form 10-K filed with the Securities and Exchange Commission ("SEC") on June 29, 2015. The unaudited condensed consolidated interim financial statements contain all normal recurring accruals and adjustments that, in the opinion of management, are necessary to present fairly the consolidated financial position of the Company at June 30, 2015, and the consolidated results of operations and cash flows for the three months ended June 30, 2015. All intercompany accounts and transactions have been eliminated. It should be understood that accounting measures at interim dates inherently involve greater reliance on estimates than at year end. The results of operations for the three ended June 30, 2015 are not necessarily indicative of the results to be expected for the full year or any future interim periods.

2. SIGNIFICANT ACCOUNTING POLICIES

Basis of Presentation and Liquidity

The Company has incurred a net loss of $3,094,854 for the three months ended June 30, 2015 (June 30, 2014 - $1,568,754) and, as of June 30, 2015, the Company has an accumulated deficit of $21,574,543 (March 31, 2015 - $18,479,689). Until the Company generates a level of revenue to support its cost structure, the Company expects to continue to incur substantial operating losses and net cash outflows. While the Company had cash and cash equivalents of $971,863 as of June 30, 2015 (March 31, 2015 - $418,893), management does not believe these resources will be sufficient to meet the Company's operating and capital needs for the ensuing fiscal year.

The Company intends to fund ongoing activities by utilizing current cash and cash equivalents and by raising additional capital through equity or debt financings. There can be no assurance that the Company will be successful in raising additional capital or that such capital, if available, will be on terms that are acceptable to the Company. If the Company is unable to raise sufficient additional capital, the Company may be compelled to reduce the scope of its operations and planned capital expenditure or sell certain assets, including intellectual property assets. This material uncertainty gives rise to substantial doubt about the Company's ability to continue as a going concern.

Fair Value of Financial Assets and Liabilities

The Company's financial instruments consist primarily of cash and cash equivalents and accounts payable and accrued liabilities. The book values of these instruments approximate their fair values due to the immediate or short-term nature of those instruments.

ASC 820 establishes a fair value hierarchy based on the level of independent, objective evidence surrounding the inputs used to measure fair value. A financial instrument's categorization within the fair value hierarchy is based upon the lowest level of input that is significant to the fair value measurement. ASC 820 prioritizes the inputs into three levels that may be used to measure fair value;

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

Level 2 – Inputs other than quoted prices included within Level 1 that are either directly or indirectly observable; and

Level 3 – Unobservable inputs that are supported by little or no market activity, therefore requiring an entity to develop its own assumptions about the assumptions that market participants would use in pricing.

Cash and cash equivalents and short-term investment are measured using Level 1 inputs.

(Expressed in United States Dollars)

The Company had certain Level 3 liabilities required to be recorded at fair value on a recurring basis in accordance with US GAAP as at June 30, 2015. As at June 30, 2015, the Company's Level 3 liabilities consisted of the grant of share Purchase options to non-employees. The resulting Level 3 liabilities have no active market and are required to be measured at their fair value each reporting period based on information that is unobservable.

A summary of the Company's Level 3 liabilities for the periods ended June 30, 2015 and March 31, 2015 are as follows:

	Three months ended June 30, 2015 $	Year ended March 31, 2015 $
Non-employee options (Note 5(a))		
Beginning fair value	1,581,444	-
Issuance	-	767,879
Reallocation of vested non-employee options	86,261	74,190
Change in fair value	178,378	739,375
Ending fair value of Level 3 liability	1,846,083	1,581,444

Certain assets and liabilities are measured at fair value on a nonrecurring basis; that is, the instruments are not measured at fair value on an ongoing basis but are subject to fair value adjustments only in certain circumstances (for example, when there is evidence of impairment). There were no assets or liabilities measured at fair value on a nonrecurring basis during the periods ended June 30, 2015 and 2014.

Recent Accounting Pronouncements

In April 2015, the FASB issued Accounting Standards Update No. 2015-03 Interest – Imputation of Interest (Subtopic 835-30). This guidance is to simplify the presentation of debt issuance costs by recognizing debt issuance costs in the balance sheet as a direct deduction from that debt liability consistent with the presentation of a debt discount. The amendments in this Update are effective for financial statements issued for fiscal years beginning after December 15, 2015, and interim periods within those fiscal years. The Company is currently evaluating the impact of adoption of this new accounting pronouncement on its financial statements.

3. CONVERTIBLE DEBENTURE

On February 19, 2014, the Company entered into a securities purchase agreement where the Company agreed to sell and issue a note with annual simple interest at 8% (the "Debenture"). A total of $1,000,100 in principal had been received.

On June 13, 2014, the Debenture matured on the closing of the Company's qualified financing. Upon completion of the qualified financing, the Debenture automatically converted into equity securities of the Company at a price per share equal to 85% of the price per share of the qualified financing. The conversion option of the Debenture was accounted for as a contingent beneficial conversion feature valued at $176,488 which was recorded as interest expense in the Statement of Comprehensive Loss on settlement of the contingency.

4. COMMON STOCK

Authorized:

Unlimited Class A common shares without par value.

Each Class A common share is entitled to have the right to vote at any shareholder meeting on the basis of one vote per share. Each Class A share held entitles the holder to receive dividends as declared by the directors. In the event of the liquidation, dissolution or winding-up of the Company other distribution of assets of the Company among its shareholders for the purposes of winding-up its affairs or upon a reduction of capital the holders of the Class A common shares shall, share equally, share for share, in the remaining assets and property of the Company.

Class B common shares and Class A preferred shares were deleted from the list of classes of shares the Company is authorized to issue by way of amendment to the Company's articles effective June 12, 2014.

The Company is subject to a stockholders agreement, which places certain restrictions on the Company's stock and its stockholders. These restrictions include approvals prior to sale or transfer of stock, a right of first refusal to purchase stock held by the Company and a secondary right of refusal to stockholders, right of co-sale whereby certain stockholders may be enabled to participate in a sale of other stockholders to obtain the same price, term and conditions on a pro-rata basis, rights of first offer of new security issuances to current stockholders on a pro-rata basis and certain other restrictions.

On April 30, 2015 the Company closed a non-brokered private placement (the "First Financing") raising gross proceeds of $1,825,937 by the issuance of 849,273 units (each a "First Financing Unit") at a price of $2.15 per First Financing Unit. Each First Financing Unit consists of one (1) common share and one half of one (1/2) common share purchase warrant (each a "First Financing Warrant"). Each whole First Financing Warrant entitles the holder thereof to purchase one additional common share of the Company at a price of $3.00 per share for a period of thirty-six (36) months from the closing date of the Financing. The Company paid a cash finder's fee of $84,074 in connection with this First Financing, as well as 27,396 finder's warrants (the "First Financing Finder's Warrants"). Each First Financing Finder's Warrant entitles the holder thereof to purchase one additional common share of the Company at a price of $3.00 per share for a period of thirty-six (36) months from the closing date of the First Financing.

On June 26, 2015 the Company closed a non-brokered private placement (the "Second Financing") raising gross proceeds of $721,243 by the issuance of 335,463 units (each a "Second Financing Unit") at a price of $2.15 per Second Financing Unit. Each Second Financing Unit consists of one (1) common share and one half of one (1/2) common share purchase warrant (each a "Second Financing Warrant"). Each whole Second Financing Warrant entitles the holder thereof to purchase one additional common share of the Company at a price of $3.00 per share for a period of thirty-six (36) months from the closing date of the Second Financing. The Company paid a cash finder's fee of $40,803 in connection with this Second Financing, as well as 18,978 finder's warrants (the "Second Financing Finder's Warrants"). Each Second Financing Finder's Warrant entitles the holder thereof to purchase one additional common share of the Company at a price of $2.15 per share for a period of sixty (60) months from the closing date of the Second Financing.

(Expressed in United States Dollars)

5. SHARE BASED PAYMENTS

(a) Stock options

The Company has a stock option plan whereby the Company is authorized to grant up to 12,108,016 options. Vesting and the term of an option is determined at the discretion of the Board of Directors of the Company.

The continuity of stock options for the periods as at June 30 and March 31, 2015, are as follows:

	Number	Weighted Average Exercise Price (CAD)	Aggregate Intrinsic Value (CAD)
Balance outstanding at March 31, 2015	4,920,000	$ 1.14	$ 10,120,000
Balance outstanding at June 30, 2015	4,920,000	$ 1.14	$ 6,676,000
Balance exercisable at June 30, 2015	3,319,587	$ 1.16	$ 4,435,212

The options outstanding and exercisable at June 30, 2015 are as follows:

Number of options	Expiry date	Options outstanding remaining contractual life (years)	Exercise Price (CAD)	Grant date fair value (CAD)	Number of options exercisable
3,520,000	June 18, 2019	3.97	$ 0.60	$ 0.23	2,346,669
250,000	June 20, 2019	3.98	$ 0.60	$ 0.23	156,250
100,000	July 14, 2017	2.04	$ 2.52	$ 1.06	100,000
450,000	December 8, 2019	4.44	$ 2.92	$ 1.65	450,000
100,000	December 8, 2019	4.44	$ 2.92	$ 1.49	33,334
400,000	December 8, 2019	4.44	$ 2.96	$ 1.56	200,000
100,000	March 16, 2020	4.72	$ 3.20	$ 1.61	33,334
4,920,000					3,319,587

The weighted average grant date fair value of stock options vested during the period ended June 30, 2015 of CAD$0.54 was estimated using the Black-Scholes option pricing model with the following weighted average assumptions: stock price – CAD$1.06; exercise price – CAD$1.14; expected risk-free interest rate – 1.08%; expected life – 4.1 years; expected volatility – 67.85% and expected dividend rate – 0%. The Company has adopted the simplified method prescribed by the SEC in SAB Topic 14 in respect of estimating the expected term of its stock options as its limited share purchase option history does not provide a reasonable basis to estimate the expected terms. Expected volatility was determined by reference to the average volatility rates of other companies in the same industry due to the Company's limited trading history.

(Expressed in United States Dollars)

Non-Employee Stock Options

In accordance with the guidance of ASC 815-40-15, stock options awarded to non-employees that are performing services for Neuro are required to be accounted for as derivative liabilities once the services have been performed and the options have vested because they are considered not to be indexed to the Company's stock due to their exercise price being denominated in a currency other than Neuro's functional currency. Stock options awarded to non-employees that are not vested are accounted for as equity awards until the terms associated with their vesting requirements have been met.

The non-employee stock options are accounted for at their respective fair values and are summarized as follows for the three months ended June 30, 2015 and year ended March 31, 2015:

	Three months ended June 30, 2015 $	Year ended March 31, 2015 $
Fair value of non-employee options, beginning of the period	1,581,444	-
Issuance	-	767,879
Reallocation of vested non-employee options	86,261	74,190
Change in fair value of non-employee stock options during the period	178,378	739,375
Fair value of non-employee options, end of the period	1,846,083	1,581,444

The non-employee options are required to be re-valued with the change in fair value of the liability recorded as a gain or loss on the change of fair value of derivative liability and included in other items in the Company's Consolidated Statements of Loss at the end of each reporting period. The fair value of the options will continue to be classified as a liability until such time as they are exercised, expire or there is an amendment to the respective agreements that renders these financial instruments to be no longer classified as a liability.

Share-based payments are classified in the Company's Statement of Loss as follows for the period ended June 30, 2015 and 2014:

	June 30, 2015 $	June 30, 2014 $
Consulting fees	85,330	193,698
Research and development	-	50,303
Wages and salaries	224,133	94,290
	309,463	338,291

(Expressed in United States Dollars)

The Company used the Black Scholes option pricing model to estimate the fair value of the options as the fair value of the services provided could not be reliably calculated. The following assumptions were used:

		March 31, 2015	March 31, 2014
Stock Price	CAD$	0.50 – 3.18	0.27
Exercise Price	CAD$	0.60 – 3.20	0.0003
Risk-free interest rate (%)		0.53 – 1.42	1.20 – 1.65
Dividend yield (%)		0.00	0.00
Expected volatility (%)		67.85	97.73 – 116.82
Expected option life (years)		3.00 – 5.00	4.33 – 5.00

The Black Scholes option pricing model was developed for use in estimating the fair value of share options that have no vesting provisions and are fully transferable. Also, option-pricing models require the use of estimates and assumptions including the expected volatility. The Company uses expected volatility rates which are based upon the average volatility rates of other companies in the same industry, due to the Company's limited history. The Company based the current stock price on the value per shares issued to date. Changes in the underlying assumptions can materially affect the fair value estimates.

(b) **Share Purchase Warrants**

The Company closed its First Financing at $2.15 per First Financing Unit of 849,273 First Financing Units raising $1,825,937 on April 30, 2015. Each First Financing Unit consists of one common stock of the Company and one half of a First Financing Warrant of the Company where one full First Financing Warrant is exercisable for 3 years at $3.00 into one common share. The Company also issued 27,396 First Financing Finder's Warrants. Each First Financing Finder's Warrant entitles the holder thereof to purchase one additional common share of the Company at a price of $3.00 per share for a period of thirty-six (36) months from the closing date of the First Financing.

The Company closed its Second Financing at $2.15 per Second Financing Unit of 335,463 Second Financing Units raising $721,243 on June 26, 2015. Each Second Financing Units consists of one common stock of the Company and one half of a Second Financing Warrant of the Company where one full Second Financing Warrant is exercisable for 3 years at $3.00 into one common share. The Company also issued 18,978 Second Financing Finder's Warrants. Each Second Financing Finder's Warrant entitles the holder thereof to purchase one additional common share of the Company at a price of $2.15 per share for a period of sixty (60) months from the closing date of the Second Financing.

The continuity of warrants for the three months ended June 30, 2015 is as follows:

	Number of warrants		Weighted Average Exercise Price	
	CAD	US	CAD $	US $
Balance, March 31, 2015	8,444,400	-	$ 1.00	-
Granted	-	638,741	$ -	2.97
Exercised	(14,400)	-	$ 1.00	-
Balance, June 30, 2015	8,430,000	638,741	$ 1.00	2.97

The warrants outstanding and exercisable at June 30, 2015 are as follows:

Number of warrants outstanding	Exercise Price	Expiry Date
8,430,000	CAD $1.00	May 30, 2016
452,032	US $ 3.00	April 30, 2018
167,731	US $ 3.00	June 26, 2018
18,978	US $ 2.15	June 26, 2020

6. COMMITMENTS AND CONTINGENCIES

(a) The Company entered into a license agreement with ANR for an exclusive right on ANR's patent pending technology, claims and knowhow. In addition to the issuance of 16,035,026 shares, the Company agreed to pay a 4% royalty on net revenue on the sales of devices covered by the patent-pending technology and services related to the therapy or use of devices covered by the patent-pending technology.

(b) On March 7, 2014, the Company entered into a commercial development-to-supply program with Ximedica where Ximedica will design, develop and produce PoNS product solution suitable for clinical trial and commercial sale. Under the program, the Company is responsible for ensuring the device is in compliance with relevant laws and regulations. The agreed budget for phase 1B of development is $499,000; phase 2 is $1,065,000; Phase 3 and 4 is $1,389,000 and 2nd software development cycle is $586,000, of which $3,918,018 was expensed as research and development since inception to June 30, 2015. The estimated duration of the project is 10 months. Invoices are to be issued monthly for work in progress. The Company can cancel the project at anytime with a written notice at least 30 days prior to the intended date of cancellation. As of June 30, 2015, the Company recorded a prepaid expense of $300,000 to Ximedica which will be applied at the end of the project. During the period ended June 30, 2015, the Company incurred charges of $818,020 (June 30, 2014 - $584,446) pursuant to this agreement.

(c) On January 27, 2015 we received a demand letter containing allegations that we had entered into a consulting arrangement with the complainants and breached certain of its terms, and used certain intellectual property in the form of business and marketing plans allegedly prepared by the complainants, and seeking damages. On May 7, 2015, Mr. Rainer Maas and Dr. Jochen Scheld filed a complaint in the U.S. District Court for the Eastern District of Pennsylvania seeking monetary damages. At this point management is unable to determine the outcome of this matter.

7. RELATED PARTY TRANSACTIONS

For the period ended June 30, 2015, the Company was a party to the following related party transactions:

During the period ended June 30, 2015, the Company paid $40,160 (June 30, 2014 - $1,000) in consulting fees to directors of the Company.

During the period ended June 30, 2015, the Company paid $30,491 (June 30, 2014 - $nil) to a company acting as the Company's corporate advisor and Chief Financial Officer.

8. SUBSEQUENT EVENTS

On July 17, 2015 the Company closed a non-brokered private placement (the "Third Financing") raising gross proceeds of $270,375 by the issuance of 125,756 units (each a "Third Financing Unit") at a price of $2.15 per Third Financing Unit. Each Third Financing Unit consists of one (1) common share and one half of one (1/2) common share purchase warrant (each a "Third Financing Warrant"). Each whole Third Financing Warrant entitles the holder thereof to purchase one additional common share of the Company at a price of $3.00 per share for a period of thirty-six (36) months from the closing date of the Third Financing. The Company paid a cash finder's fee of $16,223 in connection with this Third Financing, as well as 7,545 finder's warrants (the "Third Financing Finder's Warrants"). Each Third Financing Finder's Warrant entitles the holder thereof to purchase one additional common share of the Company at a price of $2.15 per share for a period of sixty (60) months from the closing date of the Third Financing.

Item 2. Management's Discussion And Analysis Of Financial Condition And Results Of Operations

In this quarterly report on Form 10-Q, unless otherwise specified, references to “we”, “us” or “our” mean Helius Medical Technologies, Inc. and its wholly-owned subsidiaries, NeuroHabilitation Corporation, or NHC, and Helius Medical Technologies (Canada), Inc., unless the context otherwise requires. All financial information is stated in U.S. dollars unless otherwise specified. Our financial statements are prepared in accordance with accounting principles generally accepted in the United States, or U.S. GAAP.

FORWARD-LOOKING STATEMENTS

This quarterly report on Form 10-Q contains forward-looking statements that involve risks and uncertainties, including statements regarding our market, strategy, competition, capital needs, business plans and expectations. Such forward-looking statements involve risks and uncertainties regarding the success of our business plan, availability of funds, government regulations, operating costs, our ability to achieve significant revenues and other factors. Forward-looking statements are made, without limitation, in relation to operating plans, availability of funds and operating costs. Any statements contained herein that are not statements of historical facts may be deemed to be forward-looking statements. In some cases, you can identify forward-looking statements by terminology such as “may”, “will”, “should”, “expect”, “plan”, “intend”, “anticipate”, “believe”, “estimate”, “predict”, “potential” or “continue”, the negative of such terms or other comparable terminology. Actual events or results may differ materially. In evaluating these statements, you should consider various factors, including the risks outlined in this annual report. These factors may cause our actual results to differ materially from any forward-looking statements. While these forward-looking statements, and any assumptions upon which they are based, are made in good faith, based on information available to us as of the date hereof, and reflect our current judgment regarding our business plans, our actual results will almost always vary, sometimes materially, from any estimates, predictions, projections, assumptions or other future performance suggested herein. We do not intend to update any of the forward-looking statements to conform these statements to actual results, except as required by applicable law, including the securities laws of the United States.

INDUSTRY AND MARKET DATA

Within this quarterly report on Form 10-Q, we reference information, statistics and estimates regarding the medical devices and healthcare industries. We have obtained this information from various independent third-party sources, including independent industry publications, reports by market research firms and other independent sources. This information involves a number of assumptions and limitations, and we have not independently verified the accuracy or completeness of this information. Some data and other information are also based on the good faith estimates of management, which are derived from our review of internal surveys and independent sources. We believe that these external sources and estimates are reliable but have not independently verified them. The industries in which we operate are subject to a high degree of uncertainty and risk due to a variety of factors, including those described in “Item 1A. Risk Factors” in our annual report on Form 10-K for the fiscal year ended March 31, 2015. These and other factors could cause results to differ materially from those expressed in these publications and reports.

Overview

We are a medical technology company focused on neurological wellness. We seek to develop, license or acquire unique and non-invasive platform technologies that amplify the brain’s ability to heal itself.

Our mission is to develop, license and acquire non-invasive treatments designed to help patients affected by neurological symptoms caused by disease or trauma. Applying the principles of neuroplasticity, our patented PoNS™ device induces Cranial Nerve Non Invasive Neuromodulation that utilizes the brain’s innate ability to achieve neuroplastic change to aid persons with neurological, cognitive, sensory, and motor disorders when combined with the rehabilitation process.

The following discussion and analysis of our results of operations, financial condition and plan of operations should be read in conjunction with (i) our audited financial statements for the year ended March 31, 2015 and (ii) the audited financial statements of NHC for the year ended March 31, 2014 and for the period from January 22, 2013 (inception) to March 31, 2013. The discussion below contains forward-looking statements that involve risks, uncertainties and assumptions. Our actual results may differ materially from those anticipated in these forward-looking statements as a result of many factors, including, but not limited to, those set forth under “Item 1. Business – Business Uncertainties and Going Concern Risk” and elsewhere in our annual report on Form 10-K for the fiscal year ended March 31, 2015.

Results of Operations

Three Months Ended June 30, 2015 Compared to the Three Months Ended June 30, 2014

Revenues

During the three months ended June, 2015 and 2014, we did not generate any revenues.

Operating Expenses

Operating expenses incurred during the three months ended June 30, 2015 were $2,834,998 as compared to $1,388,630 during the three months ended June 30, 2014. Significant changes and expenditures are outlined as follows:

- Advertising, marketing, and IR expenses were $308,213 for the three months ended June 30, 2015, and $46,226 for the three months ended June 30, 2014. The increase relates to advertising and promotion expenses and investor relation consulting fees. We have engaged both investor relations and public relations professionals in Canada and the US to help develop corporate material as well as arranging and participating in conferences and road shows to increase the public's awareness of our activities and the PoNS™ device.

- Audit and accounting fees were $80,462 for the three months ended June 30, 2015, and $10,645 for the three months ended June 30, 2014. The increase of $69,817 was mainly due to the requirement to review and audit the Company’s financial statements since it became a reporting issuer.

- Consulting fees were $142,822 for the three months ended June 30, 2015 and $200,849 for the three months ended June 30, 2014. The increase of $58,027 was mainly due to the expense during the three months ended June 30, 2014, associated with the granting of options to consultants for providing services in design and manufacturing and our strategic growth plan.

- Insurance expenses were $29,539 for the three months ended June 30, 2015, and $7,486 for the three months ended June 30, 2014. The increase of $22,053 was mainly due to the need for general liability, directors’ and officers’, and product insurance as the Company continues its research and development plans.

- Legal fees were $293,736 for the three months ended June 30, 2015 as compared to $219,934 for the three months ended June 30, 2014. The increase of $73,802 was primarily composed of fees incurred for general corporate matters. In addition, our legal activity to ensure current and quality regulatory filings has increased significantly since becoming a public company in Canada. Furthermore, the engagement of various specialized legal counsels for the development of our intellectual properties and the commercialization of the PoNS™ device is carried out to secure our intellectual property, including the issuance of two patents.

- Meals and travel expenses were $95,867 for the three months ended June 30, 2015 as compared to $29,441 for the three months ended June 30, 2014. The increase of $66,426 was primarily due to travel to and from various investor and medical conferences as well as required travel for personnel to coordinate the clinical trials.

- Office expenses were $26,039 for the three months ended June 30, 2015 as compared to $21,424 for the three months ended June 30, 2014. The increase of $4,615 was mainly due to general and administrative expenses but also includes computer and internet expenses, telephone expenses, and rent expenses. These expenses increased significantly as we ramped up our operations.

- Professional fees were $nil for the three months ended June 30, 2015 as compared to $13,125 for the three months ended June 30, 2014. Corporate communications and industry research fees incurred in the three months ended June 30, 2014, were not required again.

- Research and development expenses were $1,386,676 for the three months ended June 30, 2015 as compared to $653,259 for the three months ended June 30, 2014. The increase was primarily due to the continuous efforts on research and development activities of the PoNS™ device, especially activities relating to preparation of clinical trials which mostly includes our commercial development-to-supply program with Ximedica, LLC (“Ximedica”), a contract manufacturer, and the NeuroFeedback's 12- month pilot clinical trial.

- Transfer agent and regulatory fees were $32,579 for the three months ended June 30, 2015, as compared to $15,260 for the three months ended June 30, 2014. The increase of $17,319 stems from the Company’s requirement as a public company to retain a transfer agent, as well as the associated filing fees.

- Wages and salaries expenses were $439,065 for the three months ended June 30, 2015 as compared to $170,981 for the three months ended June 30, 2014. The increase of $268,084 was mainly due to the expense during the three months ended June 30, 2015, associated with the granting of options to employees.

Non-Operating Items

We recorded a loss of $259,856 in respect of non-operating items during the three months ended June 30, 2015 as compared to a loss of $180,124 for the three months ended June 30, 2014. Significant changes are outlined as follows:

- Interest income for the three months ended June 30, 2015 was $nil as compared to $2,810 for the three months ended June 30, 2014. The decrease stems from the cashing and closing of a number of interest- bearing short-term investment accounts with our banking institutions.

- Accreted interest expenses were $nil for the three months ended June 30, 2015, and $176,488 for the three months ended June 30, 2014. The decrease relates to the convertible debenture which was converted and settled in fiscal 2015. As such, no amounts were charge during fiscal 2016.

- Change in fair value of derivative liability for the three months ended June 30, 2015 was $(178,378) as compared to $nil for the three months ended June 30, 2014. The change in fair value of derivative liability is based on the change of the remaining term of our options granted to non-employees providing services for NHC and the change in our stock price. The derivative liabilities do not represent cash liabilities.

- Foreign exchange losses for the three months ended June 30, 2015 were $81,425 as compared to losses of $6,446 for the three months ended June 30, 2014. The losses stem from our Canadian dollar holdings as well as translating the balance of the Canadian dollar intercompany to the reporting currency.

Net Loss

The net loss was $3,094,854for the three months ended June 30, 2015 and $1,568,754 for the three months ended June 30, 2014. The increase in net loss of $1,526,100 resulted primarily from an increase in most operating expenses, especially advertising, marketing and IR, consulting fees, insurance expenses, legal fees, research and development, and wages and salaries.

Liquidity and Capital Resources

Our financial statements have been prepared assuming that we will continue as a going concern and, accordingly, does not include adjustments relating to the recoverability and realization of assets and classification of liabilities that might be necessary should we be unable to continue in operation.

The following table sets out our cash and working capital as of June 30, 2015 and March 31, 2015:

	June 30, 2015	March 31, 2015
Cash and cash equivalents	$ 971,863	$ 418,893
Working capital (deficit)	$ (153,160)	$ 18,543

As of June 30, 2015, our current assets were $1,430,339 (March 31, 2015 - $1,216,347), which increased mostly due to the closing of multiple private placements during the three months ended June 30, 2015. Current liabilities were $1,583,499 (March 31, 2015 - $1,197,804), which increased due to an increase in our operations since the closing of a private placement and our acquisition of NHC. Working capital was $(153,160) (March 31, 2015 – $18,543). Our current assets as of June 30, 2015 consisted of cash and cash equivalents of $971,863 (March 31, 2015 - $418,893), which increased mostly due to the closing of multiple private placements, short-term investment of $nil (March 31, 2015 - $378,000), which decreased as a result of cashing and closing certain term deposits with our banking institution, receivables of $24,231 (March 31, 2015 - $8,833), which increased due to the larger amount of refundable Canadian commodity tax based on the Company’s increase in Canadian operations, and prepaid expenses of $434,245 (March 31, 2015 - $410,621), which mostly include a prepayment to Ximedica and insurance expenses. Our current liabilities as of June 30, 2015 consisted of accounts payable and accrued liabilities of $1,583,499 (March 31, 2015 - $1,197,804), which increased due to our increased operations.

As a result of our increased activity, the accumulated deficit increased from $18,479,689 as at March 31, 2015 to $21,574,543 as at June 30, 2015.

We currently have limited working capital and liquid assets. Our cash and cash equivalents as of June 30, 2015 were $971,863. To date we have not generated any revenue from the sales of products or services. There are a number of conditions that we must satisfy before we will be able to generate revenue, including but not limited to successful completion of the design of the PoNS™ device, FDA clearance of the PoNS™ device for treating balance disorder associated with mild to moderate TBI and MS, manufacturing of a commercially-viable version of the PoNS™ device and demonstration of effectiveness sufficient to generate commercial orders by customers for our product. While we are currently seeking additional funding, we do not currently have sufficient resources to accomplish any of these conditions necessary for us to generate revenue. We will therefore require substantial additional funds in order to continue to conduct the research and development and regulatory clearance and approval activities necessary to bring our product to market, to establish effective marketing and sales capabilities and to develop other product candidates.

We will have to continue to rely on equity and debt financing. There can be no assurance that financing, whether debt or equity, will always be available to us in the amount required at any particular time or for any particular period or, if available, that it can be obtained on terms satisfactory to us. Without additional financing, we do not believe our resources will be sufficient to meet our operating and capital needs through the third quarter of 2015.

Statement of Cash Flows

Quarter ended June 30, 2015 compared to quarter ended June 30, 2014

During the quarter ended June 30, 2015, our net cash increased by $552,970 (June 30, 2014 - $7,014,291), which included net cash used in operating activities of $2,146,000 (June 30, 2014 - $540,868) stemming from our increase in operations, net cash provided by investing activities of $378,000 (June 30, 2014 - $173,904) stemming from the closing of a number of short-term investment accounts and net cash provided by financing activities of $2,434,229 (June 30, 2014 - $7,240,597) stemming mainly from the closing of multiple private placements.

Cash Used in Operating Activities

Operating activities in the three months ended June 30, 2015 used cash of $2,146,000 (June 30, 2014 - $540,868). This was made up of a net loss of $3,094,854 (June 30, 2014 - $1,568,754) less adjustments for non-cash items such as accretion of beneficial conversion feature of $nil (June 30, 2014 – $176,488), change in fair value of derivative liability of $178,378 (June 30, 2014 – $nil), stock based compensation of $309,463 (June 30, 2014 - $338,291), receivables of ($15,264) (June 30, 2014 - $nil), accounts payable of $382,844 (June 30, 2014 – $622,002), prepaid expenses of ($22,205) (June 30, 2014 – $108,895) and foreign exchange on re-measurement of $115,638 (June 30, 2014 - $Nil). Receivables increased due to the higher amount of refundable Canadian commodity tax. Payables and prepaid expenses increased due to our increase in operations.

Cash Provided by Investing Activities

During the three months ended June 30, 2015, cash provided by investing activities totaled $378,000 (June 30, 2014 - $173,904). This was made up of the closing of a number of interest-bearing short-term investment accounts held. The previous period's activities included cash acquired from the recapitalization of $23,904 and proceeds from a bridge financing of $150,000.

Cash Provided by Financing Activities

During the three months ended June 30, 2015, financing activities provided cash of $2,394,684(June 30, 2014 - $7,240,597). Financing activities during the three month period ended June 30, 2015, consisted of: issuance of share capital of $2,519,561(June 30, 2014 - $6,607,402) stemming from multiple private placements, share issue costs of $124,877 (June 30, 2014 - $Nil) and proceeds from the debenture of $nil (June 30, 2014 - $633,195).

Off Balance Sheet Arrangements

To the best of management's knowledge, there are no off-balance sheet arrangements that have, or are reasonably likely to have, a current or future effect on our results of operations or financial condition.

Recently Issued Accounting Pronouncements

In April 2015, the FASB issued Accounting Standards Update No. 2015-03 Interest – Imputation of Interest (Subtopic 835-30). This guidance is to simplify the presentation of debt issuance costs by recognizing debt issuance costs in the balance sheet as a direct deduction from that debt liability consistent with the presentation of a debt discount. The amendments in this Update are effective for financial statements issued for fiscal years beginning after December 15, 2015, and interim periods within those fiscal years. The Company is currently evaluating the impact of adoption of this new accounting pronouncement on its financial statements.

Subsequent Events

On July 17, 2015, we closed a private placement to four accredited investors, which included two institutions and two individuals, consisting of an aggregate of 125,756 units at a price of $2.15 per unit for gross proceeds of $270,375. Each unit issued in the private placement consisted of one share of our common stock and one-half of one common share purchase warrant. Each whole warrant entitles the holder to purchase one additional share of our common stock at a purchase price of $3.00 for a period of thirty-six months. We relied on the exemption from registration provided by Section 4(a)(2) of the Securities Act and Rule 506(b) thereunder for the private placements. In connection with the July 17, 2015 private placement, we issued 7,545 warrants to an institutional accredited investor that served as finder for the private placement. The finder's warrants permit the holder to purchase one share of our common stock at a price of $2.15 per share for a period of sixty months. We relied on the exemption from registration provided by Section 4(a)(2) of the Securities Act for the issuance of the finder's warrant.

Critical Accounting Policies and Estimates

Our discussion and analysis of our financial condition and results of operations are based upon our financial statements that have been prepared in accordance with U.S. GAAP. This preparation requires management to make estimates and assumptions that affect the reported amounts of assets, liabilities, revenues and expenses, and the disclosure of contingent assets and liabilities. U.S. GAAP provides the framework from which to make these estimates, assumption and disclosures. We choose accounting policies within U.S. GAAP that management believes are appropriate to accurately and fairly report our operating results and financial position in a consistent manner. Management regularly assesses these policies in light of current and forecasted economic conditions. Actual results could differ from those estimates made by management. While there are a number of significant accounting policies affecting our financial statements, we believe the critical accounting policies involving the most complex, difficult and subjective estimates and judgments are: valuation of non-monetary transactions, stock compensation for services, valuation of options and valuation of income taxes.

Stock-Based Compensation

We account for all of our stock-based payments and awards under the fair value based method. We recognize our stock-based compensation using the accelerated attribution method.

Stock-based payments to non-employees are measured at the fair value of the consideration received, or the fair value of the equity instruments issued, or liabilities incurred, whichever is more reliably measurable. The fair value of stock-based payments to non-employees is periodically re-measured until the counterparty performance is complete, and any change therein is recognized over the vesting period of the award and in the same manner as if we had paid cash instead of paying with or using equity based instruments. The fair value of the stock-based payments to non-employees that is fully vested and non-forfeitable as at the grant date is measured and recognized at that date.

We account for the granting of share purchase options to employees using the fair value method whereby all awards to employees will be recorded at fair value on the date of the grant. The fair value of all share purchase options are expensed over their vesting period with a corresponding increase to additional capital surplus. Upon exercise of share purchase options, the consideration paid by the option holder, together with the amount previously recognized in additional paid-in capital is recorded as an increase to share capital. Share purchase options granted to employees are accounted for as liabilities when they contain conditions or other features that are indexed to other than a market, performance or service condition.

We use the Black-Scholes option pricing model to calculate the fair value of our share purchase options. Option pricing models require the input of highly subjective assumptions, including the expected price volatility. Changes in these assumptions can materially affect the fair value estimate.

Derivative Liabilities

We evaluate our financial instruments and other contracts to determine if those contracts or embedded components of those contracts qualify as derivatives to be separately accounted for in accordance with ASC 815. The result of this accounting treatment is that the fair value of the embedded derivative is marked-to-market at each balance sheet date and recorded as a liability and the change in fair value is recorded in the consolidated statements of loss. Upon conversion or exercise of a derivative instrument, the instrument is marked to fair value at the conversion date and then that fair value is reclassified to equity.

The classification of derivative instruments, including whether such instruments should be recorded as liabilities or as equity, is re-assessed at the end of each reporting period. Derivative instruments that become subject to reclassification are reclassified at the fair value of the instrument on the reclassification date. Derivative instrument liabilities will be classified in the balance sheet as current or non-current based on whether or not settlement of the derivative instrument is expected within 12 months of the balance sheet date.

We use the Black-Scholes option valuation model to value derivative liabilities. This model uses Level 3 inputs in the fair value hierarchy established by ASC 820 Fair Value Measurement

Item 3. Quantitative and Qualitative Disclosures About Market Risk

Financial Instruments and Other Risks

We are exposed to credit risks and market risks related to changes to interest rates and foreign currency exchange rates, each of which could affect the value of our current assets and liabilities. We invest our cash equivalents in fixed rate, highly liquid and highly rated financial instruments such as guaranteed investment contracts, or GICs. At June 30, 2015, our cash and cash equivalents were held as GICs, the majority of which were denominated in U.S. dollars. We do not believe that the results of operations or cash flows would be affected to any significant degree by a sudden change in market interest rates relative to our investment portfolio, due to the relative short-term nature of the investments. We have not entered into any forward currency contracts or other financial derivatives to hedge foreign exchange risk. We are subject to foreign exchange rate fluctuations that could have a material effect on our total net assets or net loss. We are exposed to interest rate cash flow risk on our cash and cash equivalents as these instruments bear interest on current market rates

Item 4. Controls And Procedures

In connection with this quarterly report on Form 10-Q, under the direction of our Chief Executive Officer and our Chief Financial Officer, our management has evaluated our disclosure controls and procedures, as defined in Rules 13a-15(e) and 15d-15(e) of the Exchange Act, and has concluded that our disclosure controls and procedures were ineffective as of June 30, 2015. As of the date of this filing, we are still in the process of remediating the material weakness that caused our disclosure controls and procedures to not be effective.

Disclosure controls and procedures include, without limitation, controls and procedures designed to ensure that information we are required to disclose in the reports we file or submit under the Exchange Act is accumulated and communicated to our management, including our Chief Executive Officer and our Chief Financial Officer, or persons performing similar functions, as appropriate, to allow timely decisions regarding required disclosure. In connection with the preparation of the consolidated financial statements for the year ended March 31, 2015, our management determined that our accounting staff does not have sufficient technical accounting knowledge relating to accounting for income taxes and complex U.S. GAAP matters, which our management determined has caused our disclosure controls and procedures to be ineffective.

We intend to take appropriate and reasonable steps to make the necessary improvements to our accounting staff to remediate this material weakness in our disclosure controls and procedures as resources to do so become available. There are inherent limitations to the effectiveness of any system of disclosure controls and procedures, including the possibility of human error and the circumvention or overriding of the controls and procedures. Accordingly, even effective disclosure controls and procedures can only provide reasonable assurance of achieving their control objective.

PART II — OTHER INFORMATION

Item 1. Legal Proceedings

On January 27, 2015 we received a demand letter containing allegations that we had entered into a consulting arrangement with the complainants and breached certain of its terms, and used certain intellectual property in the form of business and marketing plans allegedly prepared by the complainants, and seeking damages. On May 7, 2015, Mr. Rainier Maas and Dr. Jochen Scheld filed a complaint in the U.S. District Court for the Eastern District of Pennsylvania seeking monetary damages in excess of $225,000. We intend to conduct a vigorous defense of this matter. At this point management is unable to determine the outcome of this matter.

Except as described above, we are not aware of any legal proceedings contemplated by any governmental authority or any other party involving us or our properties. As of August 14, 2015, no director, officer or affiliate is (i) a party adverse to us in any legal proceeding, or (ii) has an adverse interest to us in any legal proceedings. We are not aware of any other legal proceedings pending or that have been threatened against us or our properties.

Item 1A. Risk Factors

There have been no material changes to the risk factors previously disclosed in our annual report on Form 10-K for the year ended March 31, 2015. You should carefully consider the risk factors discussed in Part I, “Item 1A. Risk Factors” in our annual report on Form 10-K for the year ended March 31, 2015, which could materially affect our business, financial condition and/or operating results. The risks described in our annual report on Form 10-K are not the only risks facing our Company. Additional risks and uncertainties not currently known to us or that we currently deem to be immaterial also may materially and adversely affect our business, financial condition and/or operating results.

Item 2. Unregistered Sales of Equity Securities and Use of Proceeds

RECENT SALES OF UNREGISTERED SECURITIES

On April 30, 2015, we closed a private placement to 12 accredited investors, which included one institution and 11 individuals, consisting of an aggregate of 849,273 units at a price of $2.15 per unit for gross proceeds of approximately $1,825,937. Each unit consisted of one share of our common stock and one-half of one common share purchase warrant. Each whole warrant entitles the holder to purchase one additional share of our common stock at a price of $3.00 per share until April 30, 2018. We relied on the exemption from registration provided by Section 4(a)(2) of the Securities Act and Rule 506(b) thereunder for the private placement. In connection with the private placement, we issued 27,396 warrants to one institutional accredited investor that served as a finder for the private placement. The finder’s warrant permits the holder to purchase one share of our common stock at a price of $3.00 per share until April 30, 2018. We relied on the exemption from registration provided by Section 4(a)(2) of the Securities Act for the issuance of the finder’s warrant.

On June 26, 2015, we closed a private placement to seven accredited investors, which included one institution and six individuals, consisting of an aggregate of 335,463 units at a price of $2.15 per unit for gross proceeds of $721,243. Each unit issued in the private placements consisted of one share of our common stock and one-half of one common share purchase warrant. Each whole warrant entitles the holder to purchase one additional share of our common stock at a purchase price of $3.00 for a period of thirty-six months. We relied on the exemption from registration provided by Section 4(a)(2) of the Securities Act and Rule 506(b) thereunder for the private placements. In connection with the June 26, 2015 private placement, we issued 18,978 warrants to an institutional accredited investor that served as finder for the private placement. The finder’s warrants permit the holder to purchase one share of our common stock at a price of $2.15 per share for a period of sixty months. We relied on the exemption from registration provided by Section 4(a)(2) of the Securities Act for the issuance of the finder’s warrant.

Item 3. Defaults Upon Senior Securities

None.

Item 4. Mine Safety Disclosures

Not applicable.

Item 5. Other Information

None.

Item 6. Exhibits

EXHIBIT INDEX

Exhibit No.	Description of Exhibit
2.1	Agreement and Plan of Merger among Helius Medical Technologies, Inc., HMT Mergersub, Inc. and NeuroHabilitation Corporation, dated June 6, 2014 (incorporated by reference to Exhibit 10.6 to the Form S-1 filed with the Securities and Exchange Commission on July 14, 2014)
3.1	Articles of Continuation (incorporated by reference to Exhibit 3.1 to the Form S-1 filed with the Securities and Exchange Commission on July 14, 2014)
3.2	Articles of Amendment filed with the Wyoming Secretary of State on July 3, 2014 (incorporated by reference to Exhibit 3.2 to the Form S-1 filed with the Securities and Exchange Commission on July 14, 2014)
3.3	Articles of Amendment filed with the Wyoming Secretary of State on April 27, 2015 (incorporated by reference to Exhibit 3.3 to amendment no. 1 to the Form 10 filed on May 4, 2015)
3.4	Bylaws (incorporated by reference to Exhibit 3.3 to the Form S-1 filed with the Securities and Exchange Commission on July 14, 2014)
3.5	First Amendment to the Bylaws (incorporated by reference to Exhibit 3.4 to the Amendment to Form S-1 filed with the Securities and Exchange Commission on September 23, 2014)
3.6	Second Amendment to the Bylaws (incorporated by reference to Exhibit 3.3 to amendment no. 1 to the Form 10 filed on May 4, 2015)
31.1	Certification of Chief Executive Officer pursuant to Rule 13a-14(a) and Rule 15d-14(a), as adopted pursuant to Section 302 of the Sarbanes-Oxley Act of 2002 *
31.2	Certification of Chief Financial Officer pursuant to Rule 13a-14(a) and Rule 15d-14(a), as adopted pursuant to Section 302 of the Sarbanes-Oxley Act of 2002 *
32.1	Certification 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 *
101.INS	XBRL Instance Document *
101.SCH	XBRL Taxonomy Extension Schema Document *
101.CAL	XBRL Taxonomy Extension Calculation Linkbase Document *
101.LAB	XBRL Taxonomy Extension Label Linkbase Document *
101.PRE	XBRL Taxonomy Extension Presentation Linkbase Document *
101.DEF	XBRL Taxonomy Extension Definition Linkbase Document *

* filed herewith

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.

HELIUS MEDICAL TECHNOLOGIES, INC.

Date: August 14, 2015

By /s/ Philippe Deschamps
Philippe Deschamps
President, Chief Executive Officer and a Director

Date: August 14, 2015

By /s/ Amanda Tseng
Amanda Tseng
Chief Financial Officer (Principal Accounting Officer), and Corporate Secretary

Exhibit 31.1

Certification of Chief Executive Officer of Period Report Pursuant to Rule 13a-14(a) and Rule 15d-14(a)

I, Philippe Deschamps, certify that:

1. I have reviewed this quarterly report on Form 10-Q of Helius Medical 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(s) 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(s) 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 reasonably 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 control over financial reporting.

Date: August 14, 2015

/s/ Philippe Deschamps

Philippe Deschamps

President, Chief Executive Officer and Director

Exhibit 31.2

Certification of Chief Financial Officer of Periodic Report Pursuant to Rule 13a-14(a) and Rule 15d-14(a)

I, Amanda Tseng, certify that:

1. I have reviewed this quarterly report on Form 10-Q of Helius Medical 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(s) 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(s) 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 reasonably 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 control over financial reporting.

Date: August 14, 2015

/s/ Amanda Tseng

Amanda Tseng
Chief Financial Officer

Exhibit 32.1

Certification of Chief Executive Officer and Chief Financial Officer
Pursuant to
18 U.S.C Section 1350

In connection with the quarterly report on Form 10-Q of Helius Medical Technologies, Inc. (the "Company") for the quarter ended June 30, 2015 as filed with the Securities and Exchange Commission on the date hereof (the "Report"), Philippe Deschamps, as Chief Executive Officer of the Company, and Amanda Tseng, as Chief Financial Officer of the Company, each hereby certifies, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that, to the best of his and her knowledge:

1. The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and

2. The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: August 14, 2015

/s/ Philippe Deschamps
Philippe Deschamps
President, Chief Executive Officer and a

/s/ Amanda Tseng
Amanda Tseng
Chief Financial Officer

A signed original of this written statement required by Section 906, or other document authenticating, acknowledging, or otherwise adopting the signature that appears in typed form within the electronic version of this written statement required by Section 906, has been provided to Helius Medical Technologies, Inc. and will be retained by Helius Medical Technologies, Inc. and furnished to the Securities and Exchange Commission or its staff upon request.