

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

FORM 8-K

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

March 26, 2018

Date of Report (Date of earliest event reported)



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

WYOMING
(State or other jurisdiction of
incorporation or organization)

000-55364
(Commission
File Number)

36-4787690
(I.R.S. Employer
Identification No.)

(Exact name of registrant as specified in charter)

642 Newtown Yardley Road Suite 100
Newtown, Pennsylvania, 18940
(Address of principal executive offices) (Zip Code)

Registrant's telephone number, including area code: (215) 944-6100

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a12 under the Exchange Act (17 CFR 240.14a12)
- Precommencement communications pursuant to Rule 14d2(b) under the Exchange Act (17 CFR 240.14d2(b))
- Precommencement communications pursuant to Rule 13e4(c) under the Exchange Act (17 CFR 240.13e4(c))

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

Emerging growth company

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

Item 7.01 Regulation FD Disclosure

On March 26, 2018, Helius Medical Technologies, Inc. (the “**Company**”) posted an updated corporate fact sheet to its website at <http://heliusmedical.com/index.php/investor-relations/overview>, which the Company may use from time to time in communications or conferences. A copy of the corporate fact sheet is attached as Exhibit 99.1 to this Current Report on Form 8-K (this “**Report**”).

The information in this Report, including Exhibit 99.1 hereto, is furnished pursuant to Item 7.01 and shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liabilities of that Section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing. The Company’s submission of this Report shall not be deemed an admission as to the materiality of any information required to be disclosed solely to satisfy the requirements of Regulation FD.

This Report and Exhibit 99.1 hereto contain forward-looking statements within the meaning of the federal securities laws. These forward looking statements are based on current expectations and are not guarantees of future performance. Further, the forward-looking statements are subject to the limitations listed in Exhibit 99.1 and in the other reports of the Company filed with the Securities and Exchange Commission, including that actual events or results may differ materially from those in the forward-looking statements.

Item 9.01 Financial Statements and Exhibits

(d) Exhibits

Exhibit Number	Exhibit Description
99.1	Corporate Fact Sheet, dated March 26, 2018.

SIGNATURE

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 hereunto duly authorized.

HELIUS MEDICAL TECHNOLOGIES, INC.

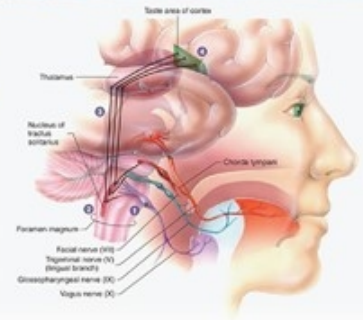
Dated: March 26, 2018

By: /s/ Joyce LaViscount
Joyce LaViscount, Chief Financial Officer



Developer of the Investigational Portable Neuromodulation Stimulator (PoNS®) Treatment

Helius Medical Technologies is a United States-based medical device company developing and commercializing the PoNS® Treatment platform, a specially patterned stimulation technology with the clinical evidence to treat balance and gait disorders in patients with traumatic brain injury (TBI) and chronic neurological disease. The first targeted application of the PoNS® is for chronic balance deficit due to mild to moderate TBI.



CAP TABLE

Price (March 23/18)	CDN\$13.17 / US\$10.10
52 Week H/L	CDN\$25.45 / \$8.60
Market Cap	CDN\$274 M / US\$210 M
Shares Outstanding	20.8 M

TICKERS	HSM: TSX	HSDT: OTCQB
Insider Ownership		25%
Ave. Daily Volume (1 yr)		166,156 (HSM) 26,532 (HSDT)
Cash & Equivalents (Dec. 31/17)		US\$5.6 M
Debt		Nil

COMPANY HIGHLIGHTS

- ✓ Potential first in class, investigational non-invasive PoNS® technology addressing symptoms of neurological disease or trauma
- ✓ Positive clinical data in large registrational trial for first indication
- ✓ Clear regulatory pathway (de novo to 510k in US), contemporaneous submissions to US, Canada, Australia, Europe and China
- ✓ Multiple large, underserved markets the first of which is Traumatic Brain Injury (TBI) - a \$5billion+ indication in the US alone, PoNS would be first and only product indicated for balance disorder related to TBI
- ✓ Well characterized commercialization model through initial workers compensation and disability focus expanding to full commercial coverage
- ✓ Significant barriers to competitor entry with comprehensive IP portfolio
- ✓ Potential to generate revenue in 2018

PoNS® CLINICAL PROGRESS AND OPPORTUNITIES

Indication and Target Patient Population - US*

- Traumatic Brain Injury (5M patients)
- Multiple Sclerosis (1M patients)
- Cerebral Palsy (764K patients)
- Stroke (7M patients; 795K new each year)
- PTSD (25M patients)
- Facial Nerve Palsy (75K patients)
- Depression (15.7M patients)
- Parkinson's Disease (1M patients; 60K new each year)

Completed Registrational and Long Term Clinical Trials



Pilot Studies Completed

Potential for Future Development



Risk Factors Our business is subject to a number of risks of which you should be aware before making a decision to invest in our common stock. These risks are discussed more fully in the "Risk Factors" section of our Annual Report on Form 10-K for the year ended December 31, 2017, which we filed with the SEC on March 12, 2018. These risks include the following: risks related to the Company's limited operating history; the Company being dependent on the ability and expertise of its Chief Executive Officer, Chief Medical Officer and a very limited number of employees; the Company having incurred losses since its inception and its anticipation that it will continue to incur substantial net losses for the foreseeable future and may never achieve or sustain profitability; risks relating to the Company requiring additional financing to carry out its plan of operations; the Company's independent registered public accounting firm having included an explanatory paragraph relating to the Company's ability to continue as a going concern in its report on the Company's audited financial statements for the year ended December 31, 2017; the Company's failure to maintain effective internal controls over financial reporting; risks related to the Company raising additional capital by issuing securities or through debt financing or licensing arrangements that may cause dilution to existing shareholders, restrict its operations or require the Company to relinquish proprietary rights; risks concerning the Company only having one product candidate, which is still in development, and the Company not having obtained clearance from the United States Food and Drug Administration (the "FDA"), CE Mark, TGA (Australia) or Health Canada with respect thereto; the Company's dependence on outside scientists and third-party research institutions for its research and development in order to be able to commercialize its product candidates; the risk that if the Company fails to obtain FDA clearance for commercialization of or otherwise fails to ensure that the PoNS™ device is available for purchase by the U.S. Government by December 31, 2021, the Company will be subject to significant risk of loss of data and proprietary rights and a US\$2,000,000 contract by December 31, 2018 penalty payable to A&I pursuant to the Strategic Agreement with A&I; the risk that the Strategic Agreement with A&I may be terminated; risks related to the limited market awareness of the Company and its product; risks related to the neuromodulation market being new and growing but undefined; the Company's PoNS™ technology being a new "untested" form of neurostimulation therapy and the medical community tending to be very conservative in adopting new therapies; risks related to the Company needing to expand its products beyond its single product by commercializing new product candidates, and the Company not being able to do so in a timely fashion and at expected costs, or at all; and development by others of new or improved devices or products that may result in the Company's present and future products from becoming obsolete.

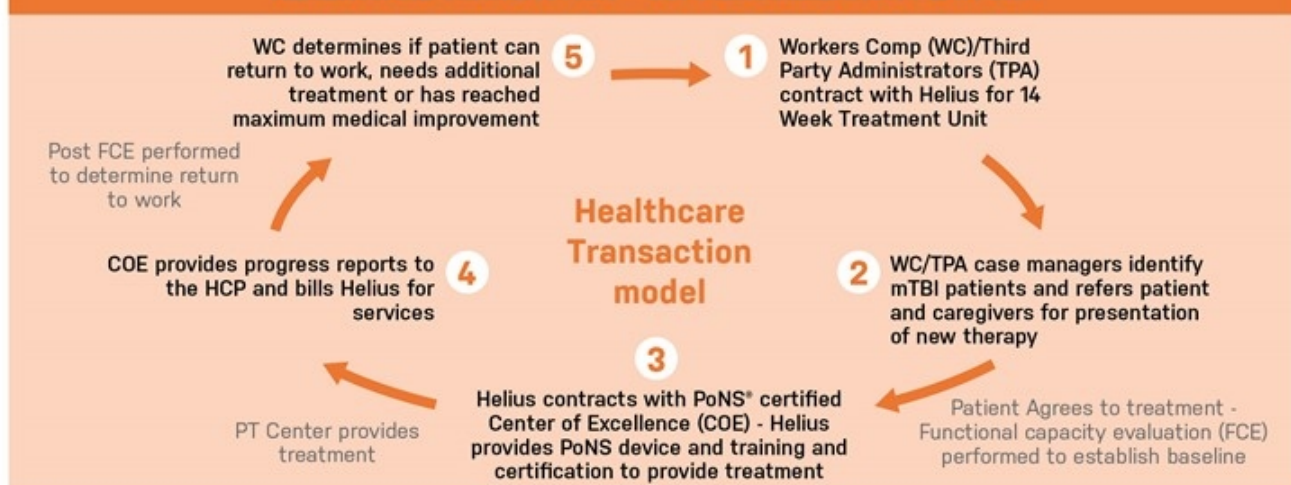
Features of the Portable Neuromodulation Stimulator (PoNS®) Treatment Platform

- ✓ Proprietary non-invasive platform for treating symptoms of neurological disease or trauma;
- ✓ Direct impulses delivered to the brain and brainstem via disposable mouthpiece placed on the tongue (razor blade), powered by pulse generator;
- ✓ In addition to positive registrational trial results in TBI, positive pilot studies support balance and gait disturbance correction, across a range of neurological conditions, including, Multiple Sclerosis (MS) and Stroke and;
- ✓ Protected by over 50 US and international patents including proprietary medical methods as well as design and utility of therapeutic delivery mechanisms.

MILESTONES AND CATALYSTS

Q2 2018	Contemporaneous submissions to the FDA, Health Canada, TGA in Australia, CE Mark in Europe
Q3 2018	Clinical Experience Programs begins to potentially drive commercial infrastructure and reimbursement
H2 2018	Clearance of PoNS device in US, Canada, Australia, Europe
2018	Potential to generate revenue in 2018

HEALTHCARE TRANSACTION – COMMERCIALIZATION PLATFORM



LEADERSHIP TEAM PROFILE



Philippe Deschamps
President and CEO; Chairman and Director

- 30+ years in the health sciences industry
- Healthcare Commercialization expertise
- Former CEO at MediMedia Health Marketing
- Former President and CEO at GSW Worldwide (Division of inVentiv Health)
- Former Director of Neuroscience Marketing at Bristol-Myers Squibb



Joyce LaViscount
CFO and COO

- 29 years in the health sciences industry
- Accomplished pharmaceutical/healthcare public company CAO
- Former Executive Director/group controller at Aptalis Pharmaceuticals
- Former Chief Operating Officer and CFO at MM Pharmaceutical Solutions



Jonathan Sackier
Chief Medical Officer

- 30+ years in the health sciences industry
- Trained surgeon and pioneer of new medical technologies
- Has helped build several companies including medical technology, research and product-design and medical contract sales organizations

Forward-Looking Statements This fact sheet contains forward-looking statements and forward-looking information as such terms are defined under applicable U.S. federal securities and Canadian securities legislation. All statements other than statements of historical fact contained in this fact sheet constitute forward-looking statements and forward-looking information, including, without limitation, statements containing the words "believe", "may", "plan", "should", "predict", "potential", "will", "estimate", "continue", "anticipate", "intend", "expect", "seek", "mission", "goal" and similar words, variations, expressions or the negative thereof. Forward-looking statements are necessarily based on estimates and assumptions made by management of the Helius Medical Technologies, Inc. ("Helius", "we" or the "Company") in light of experience and perception of historical trends, current conditions and expected future developments, as well as the factors management of the Company believes are appropriate. Forward-looking statements and information in this fact sheet include but are not limited to statements relating to: the timing of regulatory submissions; the Company's estimate of the size of the potential markets for its products; the estimated clinical, regulatory and commercial milestones and the timelines for achieving such milestones; the therapeutic benefits, effectiveness; whether the Company will receive, and the timing and costs of obtaining, regulatory clearances in the U.S., Canada, EU and Australia; regulatory developments and the regulatory environments in which the Company operates; and anticipated trends and challenges in the Company's business and the markets in which it operates.

Such statements reflect management of the Company's current views with respect to future events and are subject to risks and uncertainties and are necessarily based upon a number of estimates and assumptions that, while considered reasonable by the Company, are inherently subject to significant business, economic, competitive, political and social uncertainties and contingencies. Many factors could cause our actual results, performance or achievements to be materially different from any future results, performance, or achievements that may be expressed or implied by such forward-looking statements, including, among others, those risks described in "Risk Factors" above and the risks described in our SEC filings, including our Annual Report on Form 10-K for the year ended December 31, 2017 and our other SEC filings from time to time. This fact sheet speaks as of its date and we, our advisors, and our and their affiliates and representatives undertake no obligation to update, revise or correct the information contained herein, except as required by law.

Industry and Market Data This fact sheet contains industry and market data that we obtained from independent industry publications. We have not independently confirmed this data and, although we believe it is generally reliable, it involves a number of assumptions and limitations. It is inherently imprecise, and you are cautioned not to place undue reliance on it.

