

rehabilitation

Martin Jamieson | CEO & Chairman of the Board | Nexstim



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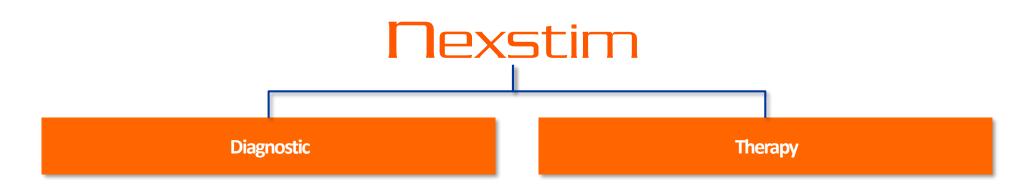
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Nexstim's NBS System is cleared by the FDA for assessment of the motor and speech cortices for pre-procedural planning. The NBT System is not cleared for commercial distribution in the United States.

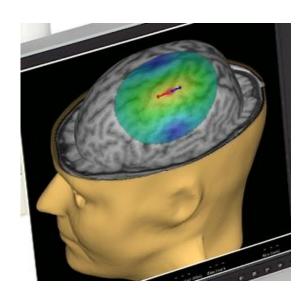
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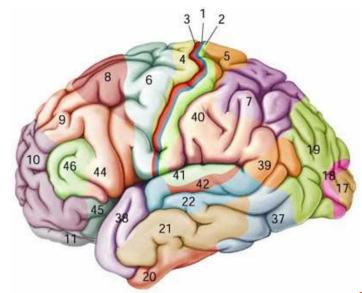
Nexstim Transcranial Magnetic Stimulation (TMS) platform



Navigated Brain Stimulation (NBS)



Navigated Brain Therapy (NBT®)



Platform protected with 17 patent families

- 63 granted patents, 35 pending
- Right to software: Nexstim owns rights to its NBT® and NBS Systems' software developed in-house.
- Core algorithms kept as trade secrets





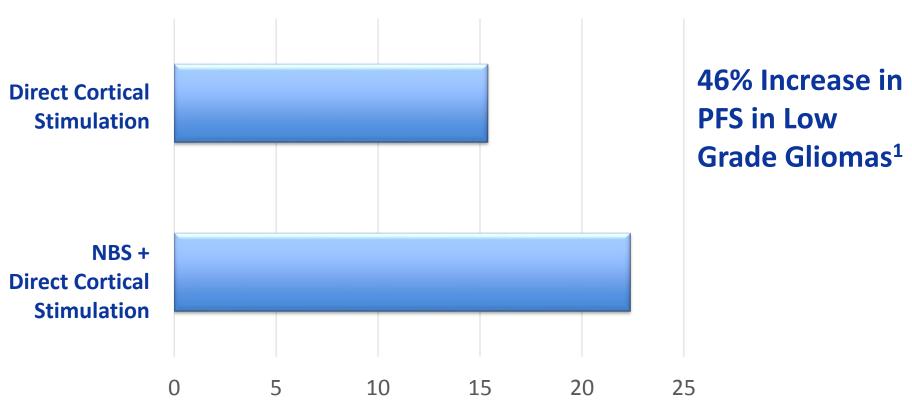
Dexstim

Navigated Brain Stimulation® for pre-surgical mapping of motor and speech cortex



NBS Pre-Surgical Mapping makes the difference to clinical outcome





¹ Dietmar Frey, Peter Vajkoczy, and Thomas Picht Navigated transcranial magnetic stimulation improves the treatment outcome in patients with brain tumors in motor eloquent locations Neuro Oncology 2014: nou110v1-nou110

NBS Pre-Surgical Mapping Major Centers



























BERN UNIVERSITY HOSPITAL





























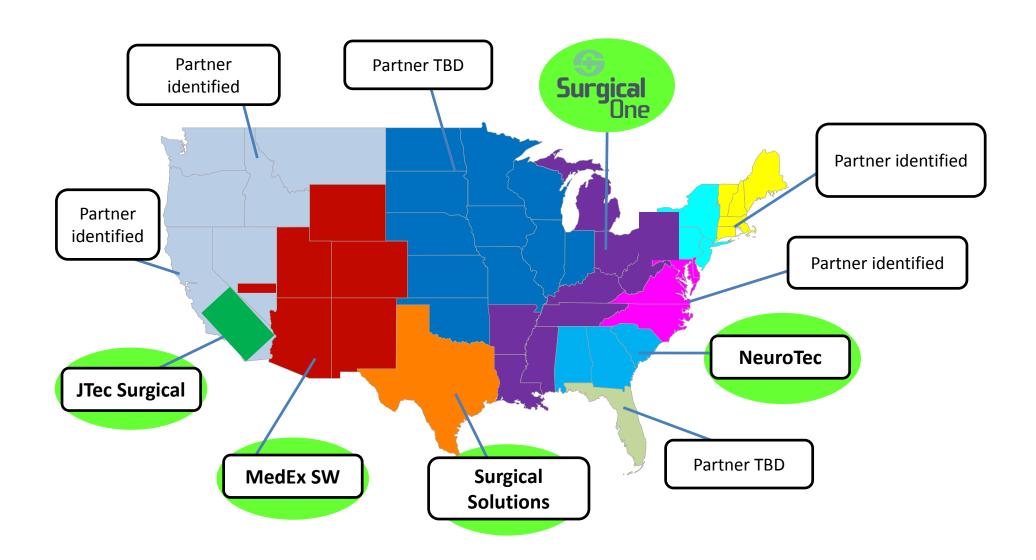








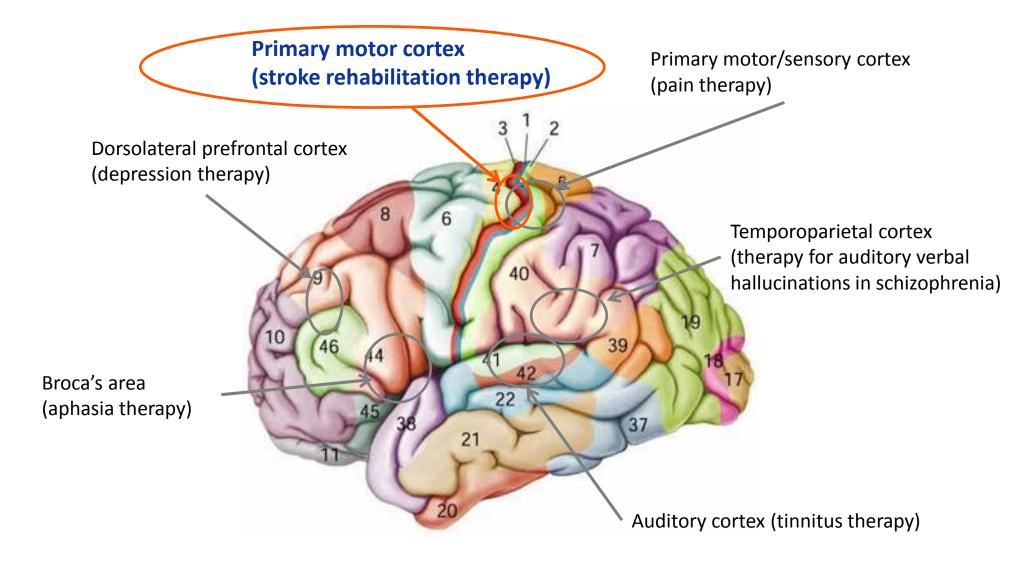
NBS Pre-Surgical Mapping US Distribution Partners







NBT® (Navigated Brain Therapy) has potential for multiple therapeutic applications due to precise navigation

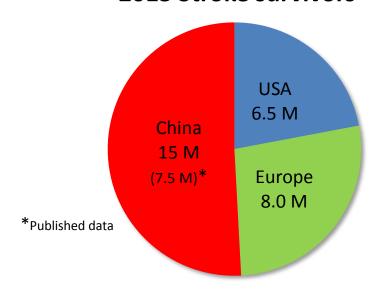


Market opportunity in stroke rehabilitation



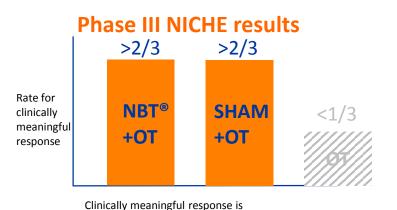
USA 0.8 M China Europe 2.5 M 1.2 M

2015 Stroke survivors

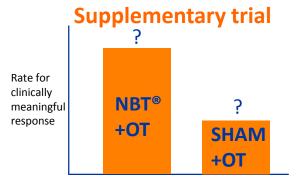


- Large and rapidly growing market
- 50% of stroke survivors will have upper limb paralysis with few available treatment options
- Focus on the period over 3 months post stroke

Clinical data comparing NBT® + occupational therapy (OT) with OT



- 199 patients 12 leading US centres
- Excellent results in active group
- Sham coil in control group proved active
- Safety endorsed by DSMB



>5points in UEFM score

Clinically meaningful response is >5points in UEFM score

- 60 patients 5 leading US centres
- FDA approved trial protocol December 2016
- Study commenced February 2017 with expected completion by March 2018
- Active group data from Phase III and supplementary trial to be pooled

Key performance indicators

EUR in thousands	H2 2016	H2 2015	FY 2016	FY 2015	Full year expenses
	6 months	6 months	12 months	12 months	EUR 2,546.1
Net sales	1,590.1	1,884.7	2,482.6	2,527.9	thousand lower related to Phase III multi-centre
Personnel expenses	-2,038.7	-2,063.8	-4,253.2	-3,969.8	trial ending in H1
Depreciation and amortisation	-203.5	-217.3	-372.3	-386.0	2016.
Other operating expenses	-1,496.8	-4,420.3	-3,907.6	-7,843.0	Achieved H2 2016
Profit/ -Loss for the period	-2,506.2	-5,272.0	-6,732.7	-9,827.0	targeted lower cash burn
Earnings per share (EUR)	-0.16	-0.73	-0.56	-1.37	
Cash flows from operating activities	-1,984.6	-4,333.3	-7,225.3	-9,608.6	Cash plus cash generated from
Cash in hand and at banks	8,156.4	6,874.7	8,156.3	6,874.7	sales, and the
Total equity	4,478.5	3,545.1	4,478.5	3,545.1	remaining financing
Equity ratio (%)	55.12	44.16	55.12	44.16	arrangements with Bracknor and
Number of shares in the end of the period (pcs)	47,087,749	8,010,758	47,087,749	8,010,758	Sitra are now estimated to
Average number of shares during the period (pcs)	15,826,991	7,178,584	11,950,664	7,154,868	finance the
Diluted number of shares in the end of the period (pcs)	78,830,446	8,797,698	78,830,446	8,797,698	Company until Q4
Diluted average number of shares during the period (pcs)	22,521,296	7,965,524	15,762,543	7,941,808	2018.

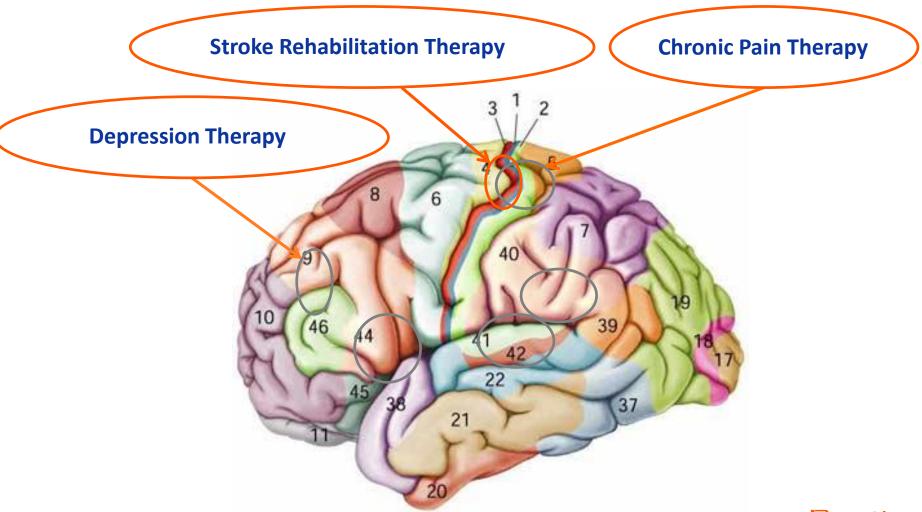
Nexstim Funding

- Accelerated pace in the two year funding arrangement with Bracknor Investment and Finnish Innovation Fund Sitra
- Funding arrangement is a combination of:
 - EUR 5 million of convertible bonds 100% funded
 - EUR 6.5 million stand-by equity facilities ~92% funded
 - EUR 0.5 million direct share issue 100% funded
 - EUR 15.0 million warrants strike range EUR 0.14 0.85
- Open elements in the funding are the Sitra standby-equity facility of EUR 0.5 million and the EUR 15.0 million in warrants



Nexstim Strategic Actions

 Focus on therapeutic applications – Primarily Stroke, **Depression and Pain**



Nexstim Strategic Actions

Timeline of US Regulatory Development for Stroke

December 2016

FDA approval of **Trial Protocol**

February 2017

Trial Commenced March 2018

Expected Trial Completion **April** 2018

Expected FDA 510(k) de novo filing











Rehabilitation Institute of Chicago (central site)

Spaulding Rehabilitation Hospital (Boston)

Rancho Los Amigos National Rehabilitation Center

Burke Rehabilitation Hospital (White Plains, NY)

University of Cincinnati

Nexstim Strategic Actions

• File 510(k) for depression during H1 2017

 Exploit CE marked approval in Stroke – EU sales to begin in H1 2017

 Partner agreements to include major Asian territories

 Prepare funding plan for US therapy commercialisation





Thank you

