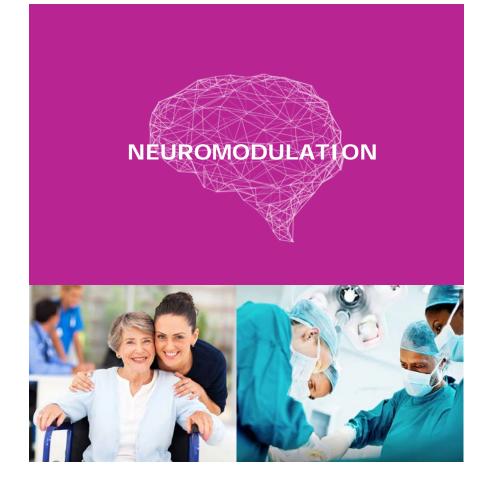


Health innovation that matters

LivaNova Investor Day

Jason Richey General Manager, Neuromodulation & President, North America

September 14, 2017

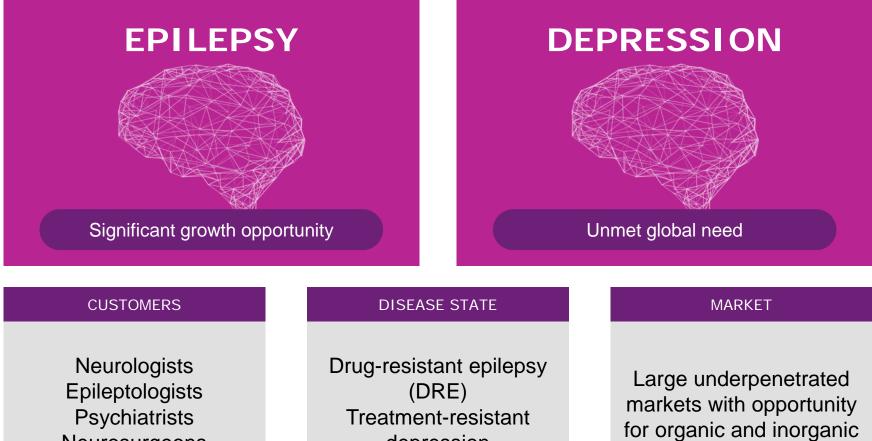


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Certain statements in this presentation, other than purely historical information, are "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, Section 27A of the Securities Act and Section 21E of the Securities Exchange Act of 1934, as amended (the "Exchange Act"). These statements include, but are not limited to, LivaNova's plans, objectives, strategies, financial performance and outlook, trends, the amount and timing of future cash distributions, prospects or future events and involve known and unknown risks that are difficult to predict. As a result, our actual financial results, performance, achievements or prospects may differ materially from those expressed or implied by these forward-looking statements. In some cases, you can identify forward-looking statements by the use of words such as "may," "could," "seek," "guidance," "predict," "potential," "likely," "believe," "will," "should," "expect," "anticipate," "estimate," "plan," "intend," "forecast," "foresee" or variations of these terms and similar expressions, or the negative of these terms or similar expressions. Such forward-looking statements are necessarily based on estimates and assumptions that, while considered reasonable by LivaNova and its management based on their knowledge and understanding of the business and industry, are inherently uncertain. These statements are not guarantees of future performance, and stockholders should not place undue reliance on forward-looking statements. There are a number of risks, uncertainties and other important factors, many of which are beyond our control, that could cause our actual results to differ materially from the forward-looking statements contained in this press release, including those described in the "Risk Factors" section of Annual Reports on Form 10-K, Quarterly Reports on Form 10-Q, Current Reports on Form 8-K, the Registration Statement on Form S-4 and other documents filed from time to time with the United States Securities and Exchange Commission by LivaNova.

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We are the market leader in Vagus Nerve Stimulation (VNS) Therapy®



Neurosurgeons **Patients**

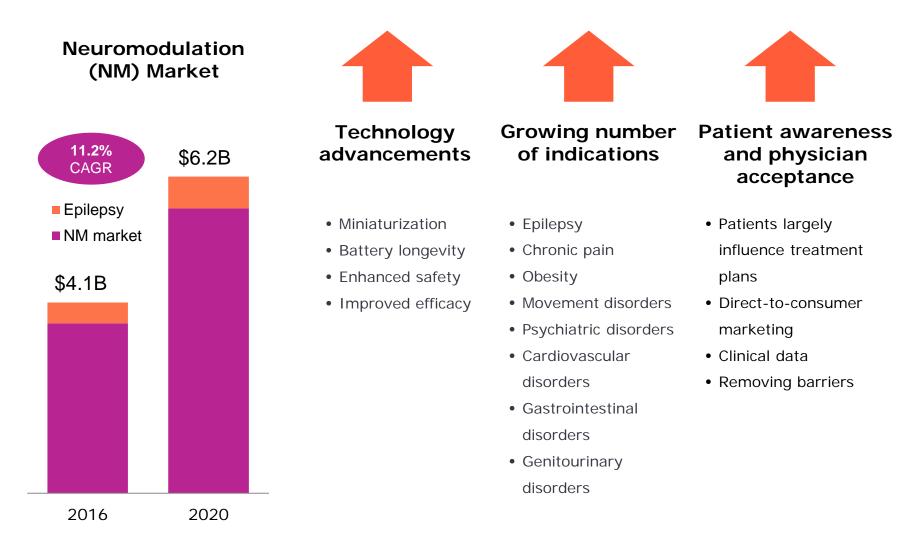
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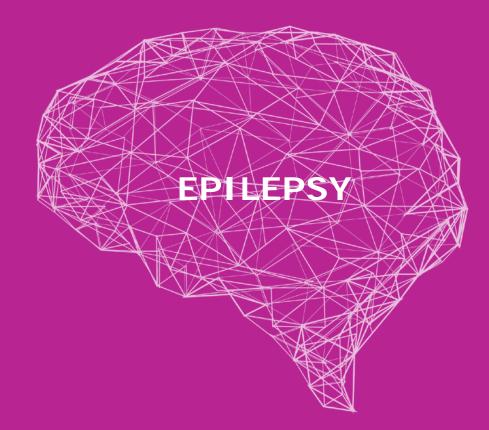
depression (TRD)

3

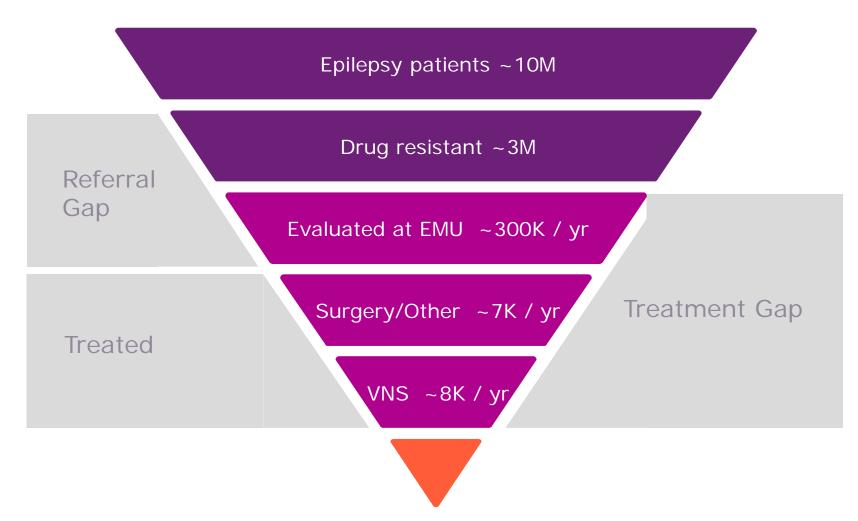
growth

Neuromodulation is a large and growing market





~35% of patients need treatment beyond medication Significant growth opportunity remains



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Source: Prevalence data and approximations based on CDC; EMU = Epilepsy Monitoring Unit – contained inside a comprehensive epilepsy center, which is a specialized extensive center that focuses on treatments for epilepsy

Growing through innovation and patient awareness

Remove barriers	MRI (U.S. and OUS) and pediatric labeling expansions (U.S.)
Innovation	Maintain new product cadence every 12-24 months
International growth opportunities	Strategic targeting of key markets
Patient awareness	Direct-to-consumer marketing and clinical studies
Universal platform	Supports cost reduction and inorganic growth through M&A

Unlocking potential via labeling expansions

	FDA expanded VNS Therapy access to patients as young as age 4 (2017)
	OUS CE Mark approval with no age restrictions (1994)
Pediatric	Clinical data proves early adjunctive use yields better results
	Patients having seizures for <10 years respond better to VNS Therapy
	VNS Therapy has shown to help children reach important developmental milestones
	FDA approval and CE Mark for expanded MRI labeling (2017)
	VNS Therapy continues to be the only implantable epilepsy device approved by FDA for MRI scans
MRI	Patients now have access to 90% of MRI scans routinely performed on people with epilepsy
	Reduces average distance to MRI centers significantly

Liva Nova Source: Adapted from Englot DJ et al. NeurosurgClin N Am. 2011: 22:443-448; Orosz I et al. Epilepsia. 2014; 55(10): 1576-1584

Developing customer-centric innovation Providing ease of use, better patient care, cost effectiveness







SenTiva[™] innovations

- Senses bradycardia and tachycardia
- Monitors patient sleeping position
- Provides scheduled dosing
- Next-generation programmer
- New user interface on tablet
- Wireless wand

Future generations

- Auto gain control minimal device positioning by physician
- Microburst new vagus nerve stimulation method for different area of brain
- Wearable technology
- Patient apps and portals
- Cloud-based capabilities
- Data and analytics tools
- Comprehensive disease management

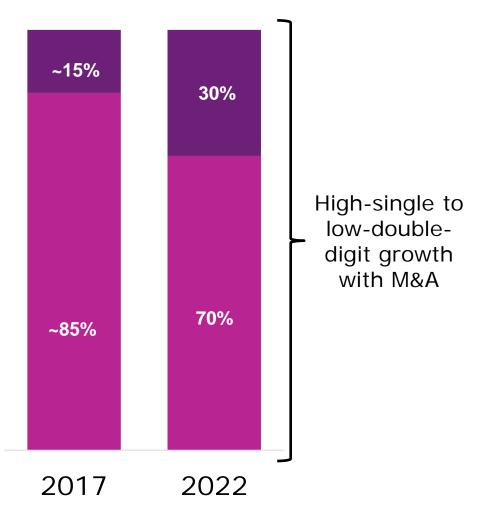
Targeting international markets More than 100K global patients treated



- Focused team with regional marketing
- Key markets: UK, Germany, Nordics, France, Japan



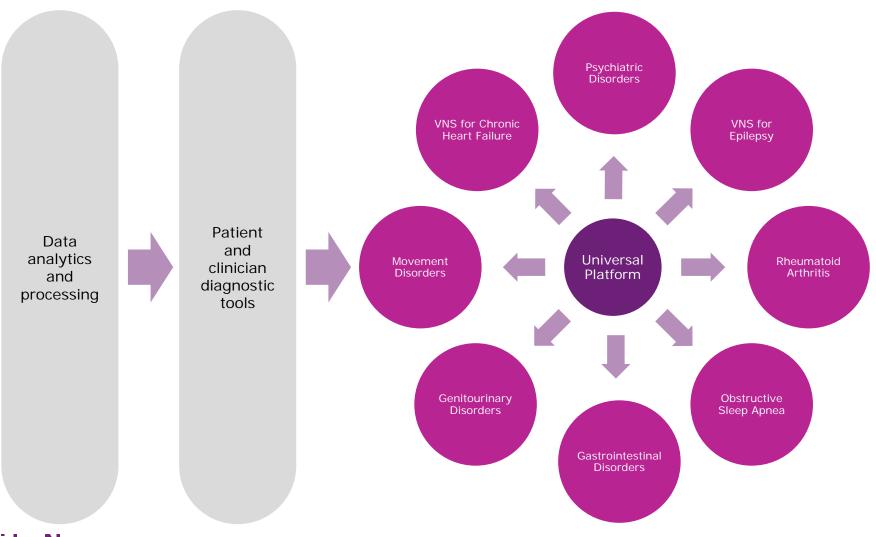
- Increase sales force
- Enhance R&D teams

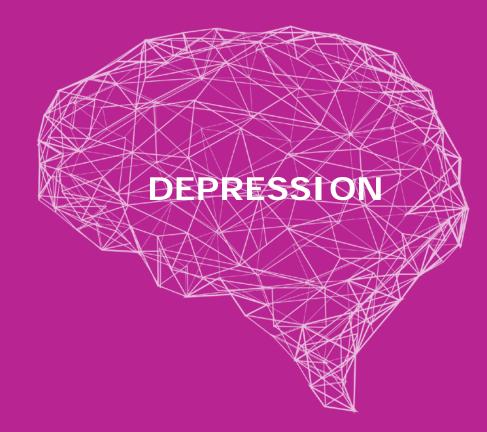


Clinical studies to further support adoption Our customers rely on data

PATIENT REGISTRY	EARLY ADJUNCTIVE USE STUDY	
 Goes live in parallel with SenTiva launch Tracks seizure activity and overall patient quality of life Periodic data reviews Health economics data 	 Prospective multi-center study Patients 4+ years VNS as adjunctive therapy Early use of VNS: Reduces seizures Reduces recovery time Improves cognitive results Enhances performance development 	
Multi-year Registry	Multi-year Study	
Launches 2017	Launches 2018	

Universal platform: foundation for additional indications





Depression affects patients in the prime of their life and burdens families and society

Depression	 300M+ patients globally Median age of onset: 25 years Fewer than half of patients receive treatment Leading cause of disability and major burden
Treatment-Resistant Depression (TRD)	 ~10-30% of patients with depression Median age 40+ after 2+ unsuccessful treatments (medications, psychotherapy, electroconvulsive therapy) More relapse, less remission, more side effects
VNS Therapy as an adjunctive treatment	 800 TRD patients with 4+ unsuccessful prior treatments for 5 years 40.9% responded with treatment as usual 67.9% responded with VNS Therapy

"We are **very encouraged** by the results of this unprecedented study, and hope that VNS becomes more readily available as a viable option for patients who have been through countless interventions for severe, chronic depression."

- Dr. Scott Aaronson, lead investigator, Sheppard Pratt Health System



Sources: Aaronson et al. American Journal of Psychiatry. 2017; 174(7): 640-48.; WHO Depression Fact sheet: http://www.who.int/mediacentre/factsheets/fs369/en/; RC Kessler and EJ Bromet. Annu Rev Public Health. 2013; 34:119–138.; Rush et al. Am J Psychiatry 2006; 163:1905–1917.; Rush et al. Psychiatric Annals. 2008; 38(3):188-193.; Gaynes et al. AHRQ Publication No. 11-EHC056-EF. September 2011.

Exploring options for LivaNova's TRD treatment

Received CE Mark in 2001

Received FDA approval in 2005

Currently have 4,000 devices implanted over ~18-month period

Started pilot trial in Germany in 1Q17

Planning scaled launch in other European countries where we have approval and reimbursement

Ongoing discussions with CMS to gain U.S. reimbursement approval

Summary: a pioneer in Neuromodulation with robust growth opportunities

Epilepsy	 Strong core growth driven by consistent product innovation Label expansion increases patient pool International expansion opportunities Inorganic opportunities drive high-single- to low-double-digit growth trajectory
Depression	 Massive market opportunity with unmet need 800+ TRD patients show positive results with VNS Therapy Piloting in selected European countries In discussions with CMS on reconsideration and coverage with evidence development in the U.S.
Future Technology Enhancements	 Universal platform Enhanced diagnostic capabilities Wireless Wearables Patient/physician apps Improved data and analytics

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