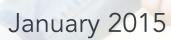


Investor Presentation





Safe Harbor Statement

This presentation and other matters discussed today or answers that may be given to questions asked include forward-looking statements within the meaning of the federal securities laws. These statements, among other things, relate to Supernus' business strategy, goals and expectations concerning its product candidates, future operations, prospects, plans and objectives of management. The words "anticipate", "believe", "could", "estimate", "expect", "intend", "may", "plan", "predict", "project", "will" and similar terms and phrases are used to identify forward-looking statements in this presentation. Supernus' operations involve risks and uncertainties, many of which are outside its control, and any one of which, or a combination of which, could materially affect its results of operations and whether the forward-looking statements ultimately prove to be correct. Supernus assumes no obligation to update any forward-looking statements except as required by applicable law.

Supernus has filed with the U.S. Securities and Exchange Commission (SEC) reports and other documents required by Section 13 or 15(d) of the Securities Exchange Act of 1934, as amended. Before you purchase any Supernus securities, you should read such reports and other documents to obtain more complete information about the company's operations and business and the risks and uncertainties that it faces in implementing its business plan. You may get these documents for free by visiting EDGAR on the SEC website at http://www.sec.gov.



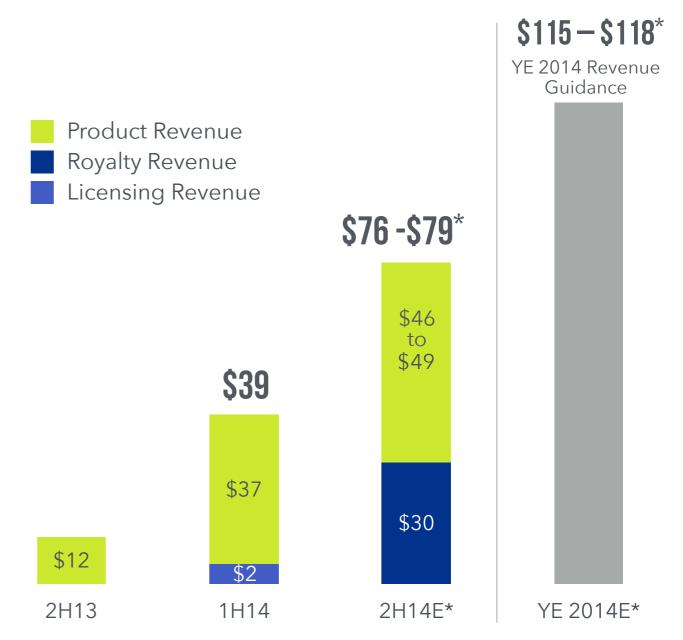
Commercial Stage CNS Specialty Pharma

Strong Execution Since 2012 IPO

(dollars in millions)

2 CNS PRODUCTS LAUNCHED IN 2013

- 25-year track record of bringing products to market
- Successful launch of leading anti-epileptic drugs
- Strong operating performance within 2 years of commercial operations
- Achieving profitability in 2014

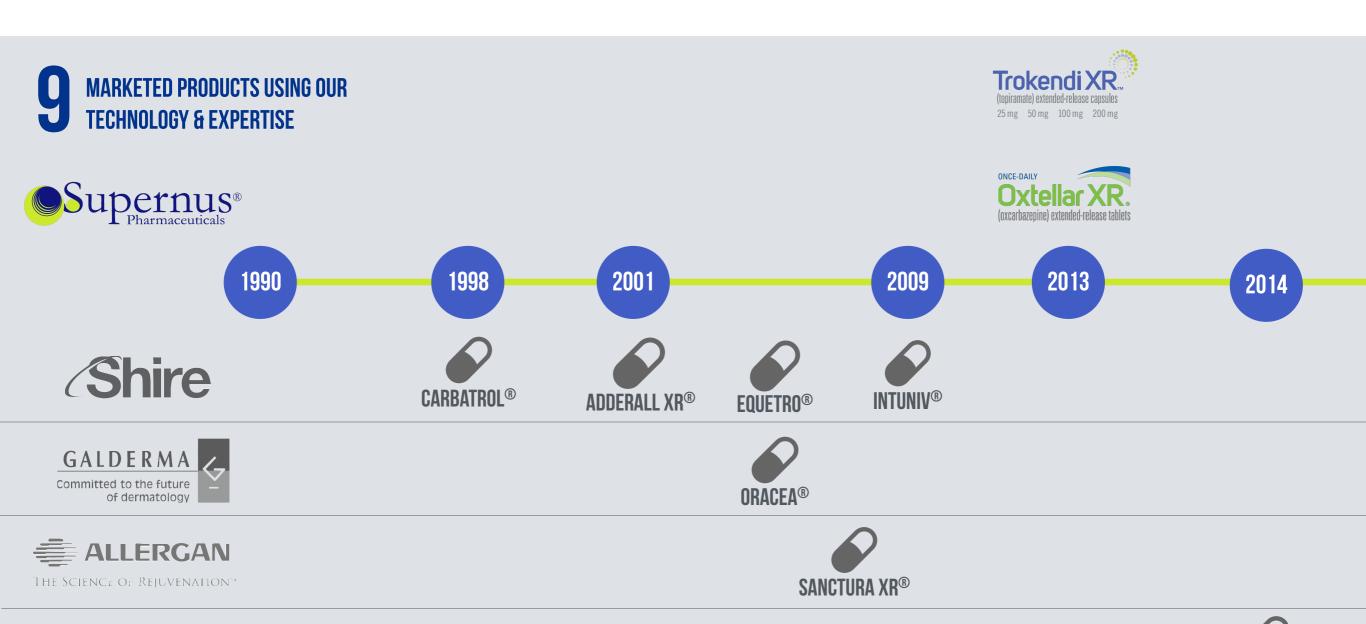


- Reported as of 3Q14
- * Based on guidance provided as of November 12, 2014, which has not been updated since that date.



Proven Execution

25 Years of Successful Product Development & Commercialization









ORENITRAM®

Strong Portfolio of CNS Products

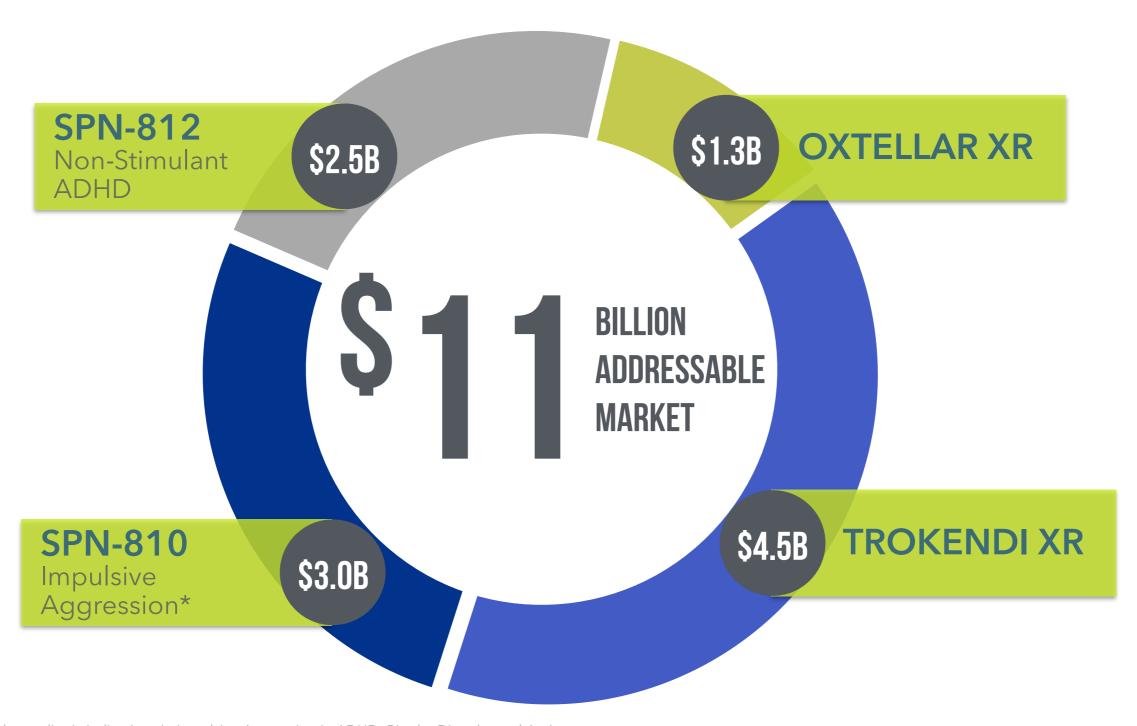
Addresses \$11 Billion Market Opportunity

Product	Indication	Development	NDA	Launch	
Oxtellar XR®	Epilepsy				2013 FEBRUARY
Trokendi XR®	Epilepsy		•		2013 AUGUST
SPN-810	Impulsive Agression in ADHD		Phase III Trial 2H2015		
SPN-812	ADHD		Phase IIb Trial 2H2015		
SPN-809	Depression		IND		



Product Portfolio Opportunity

Penetrate & Gain Share of a Large Market Opportunity



^{*} Includes pediatric indications in Impulsive Aggression in ADHD, Bipolar Disorder and Autism Source: SHA, Global Data, Company Estimates





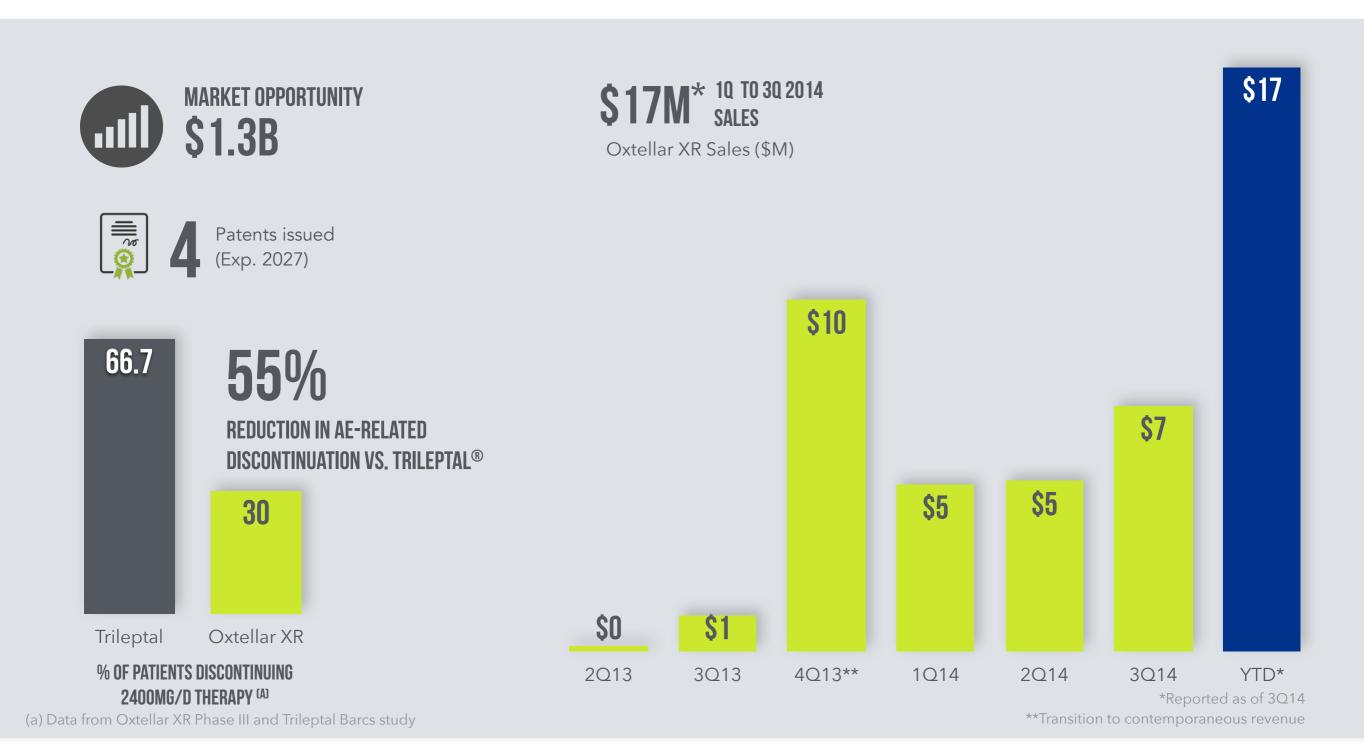
Commercial Success

Oxtellar XR® & Trokendi XR® to Drive Future Growth



Oxtellar XR®

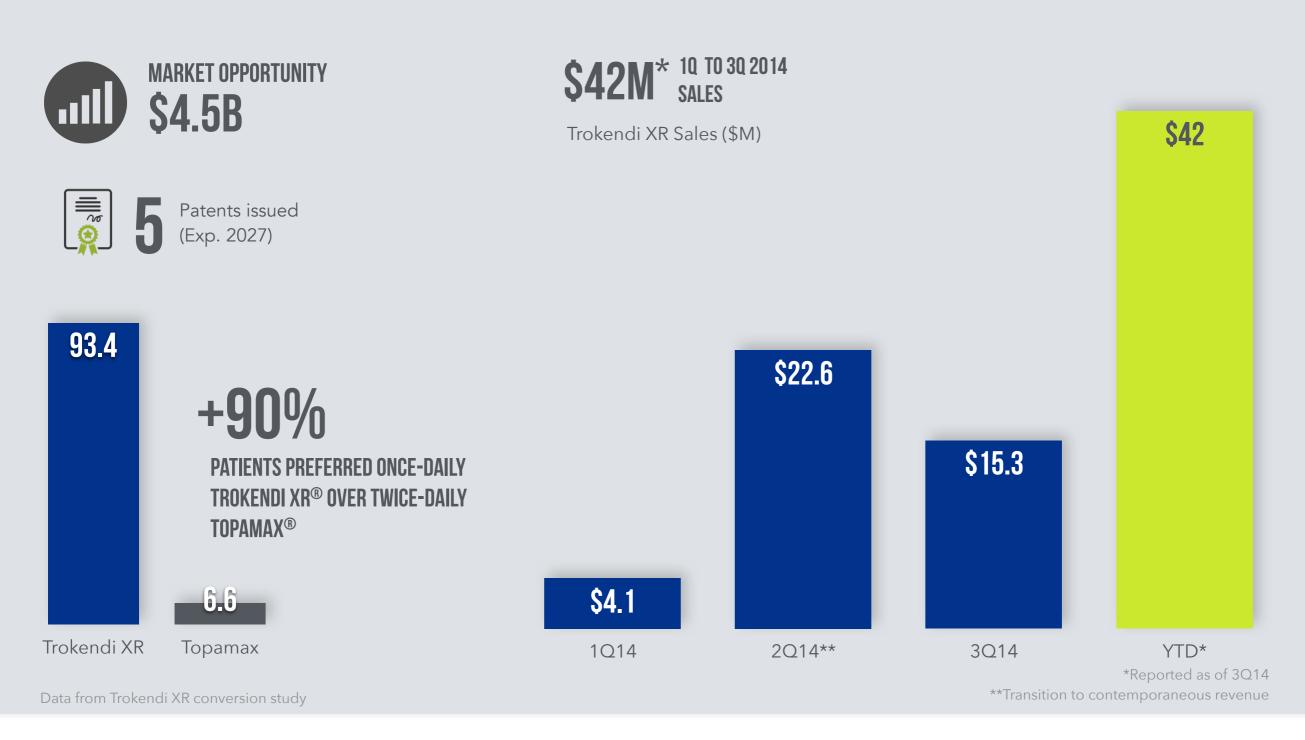
The Only Once-Daily Oxcarbazepine Product in the U.S.





Trokendi XR®

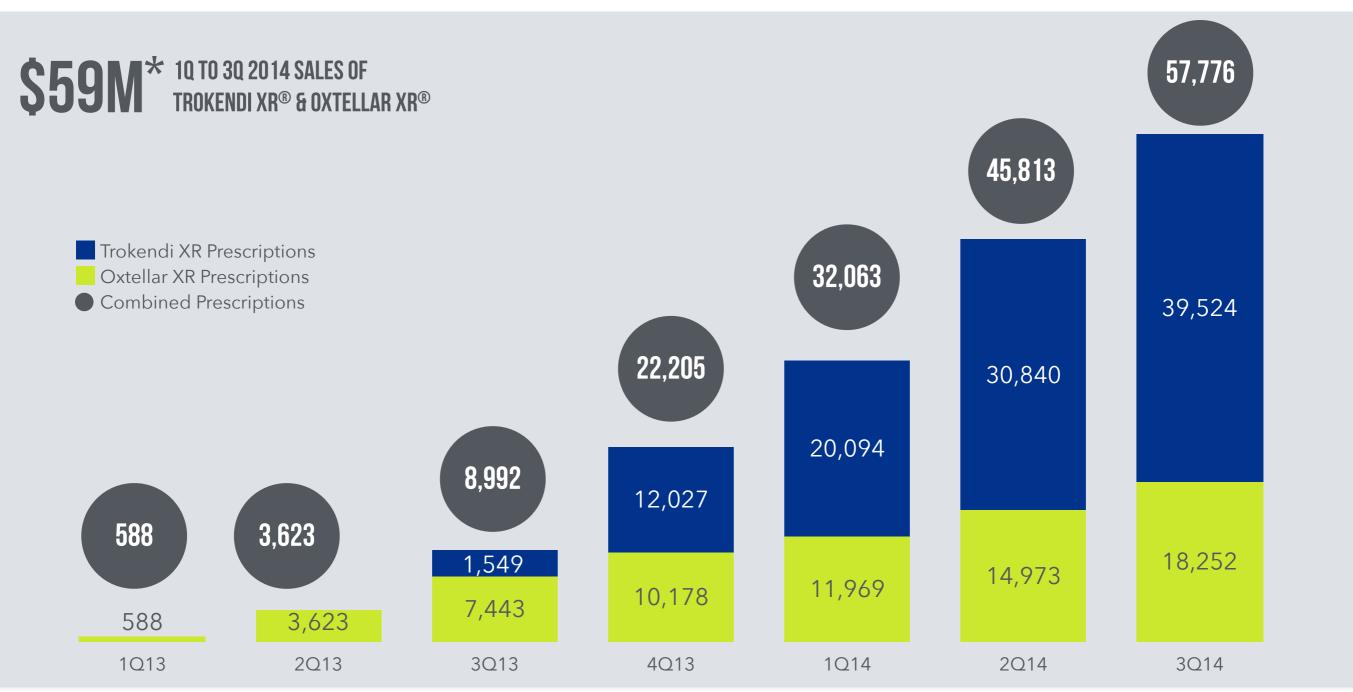
The First Once-Daily Topiramate Product in the U.S.





Strong Prescription Growth

Two Successful Product Launches



Source: SHA Monthly Prescriptions *Reported as of 3Q14



Robust, Late-Stage Pipeline

Fueling Sustainable, Long-term Growth



Robust, Late-Stage Pipeline

Addresses \$5.5 Billion Market Opportunity

Product	Indication	Development	NDA	Launch
Oxtellar XR®	Epilepsy			
Trokendi XR®	Epilepsy			
SPN-810	Impulsive Agression in ADHD		Phase III Trial 2H2015	
SPN-812	ADHD		Phase IIb Trial 2H2015	
SPN-809	Depression			



SPN-810

Novel Product for Impulsive Aggression (IA)

+\$3B ADDR

ADDRESSABLE MARKET OPPORTUNITY

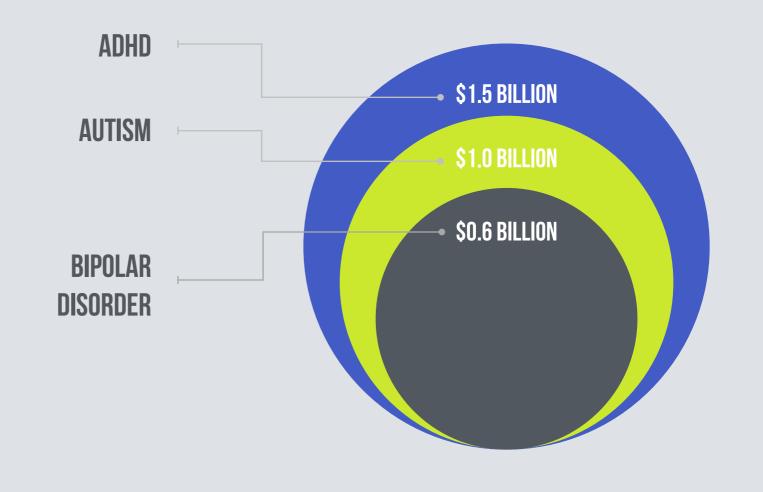


Disorder with impairment of self regulation

Characterized by episodes of unplanned, maladaptive aggression

Currently no FDA-approved treatments

Off-label use of atypical antipsychotics common, which have serious safety and tolerability issues





SPN-810

Novel Product for Impulsive Aggression (IA) in ADHD



Granted Fast Track
Development Designation

1ST

Expected to be first product approved to treat IA



Established safety and efficacy at low and medium doses in Phase IIb trial



Meeting with FDA held in December 2014

2015

Begin Phase III clinical program in second half of 2015

25% CHILDREN WITH ADHD HAVE PERSISTENT CONDUCT PROBLEMS SUCH AS IMPULSIVE AGGRESSION





SPN-812

Novel Non-Stimulant Product for ADHD

\$2.5B

ADDRESSABLE MARKET OPPORTUNITY*



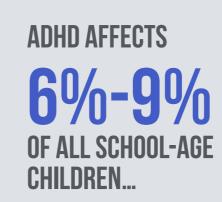
Expected to have a better AE profile than current therapies

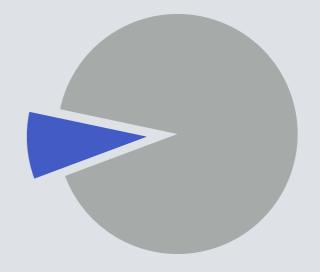


Extended-release formulation selected for the product to be used in the pivotal trials

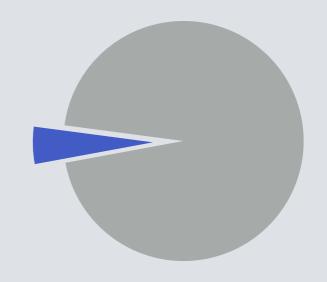
2015

Begin first pivotal trial in second half of 2015









* Represents pediatric and adult, non-stimulant ADHD market



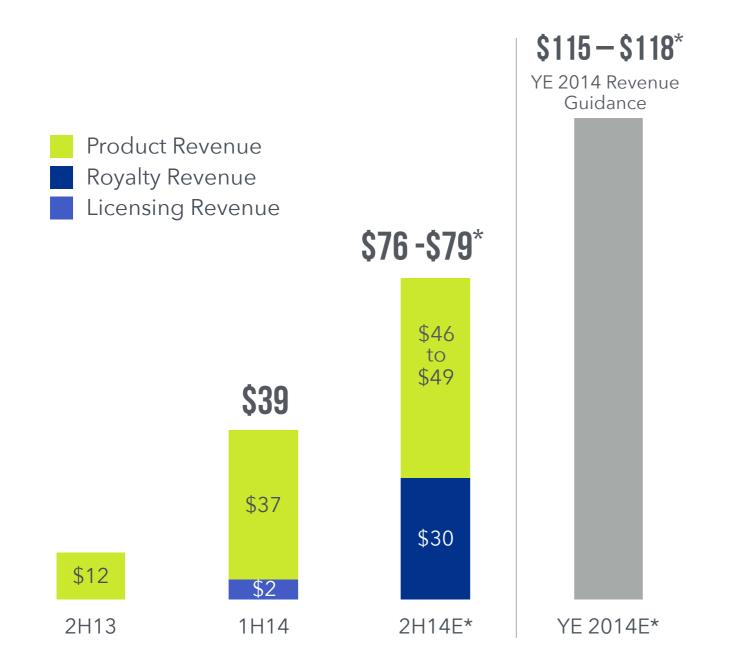
Successful Commercial Business within 2 Years

Financial Overview



Strong Operating Performance

Solid Revenue Growth Since Product Launches (dollars in millions)





which has not been updated since that date.



Strong Operating Performance

Quarterly Improvement in Cash Burn

(dollars in millions)



Note:

Reported as of 3Q14



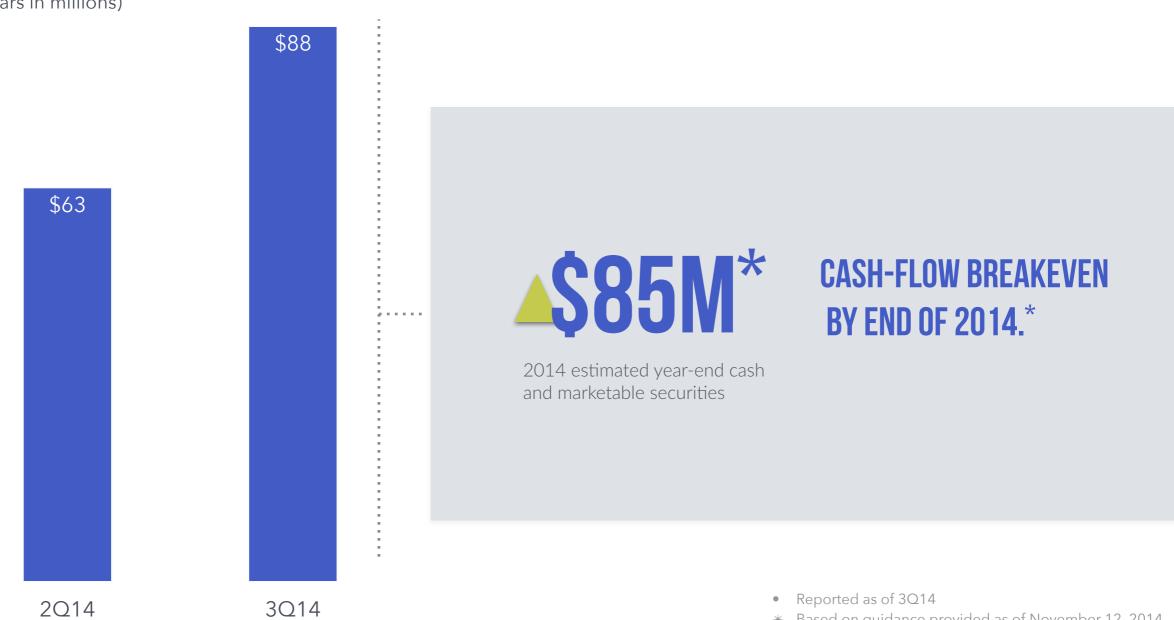
^{*}Based on guidance provided as of November 12, 2014, which has not been updated since that date.

^{**} Excludes \$30M royalty monetization receipt

Strong Balance Sheet

Solid Cash Position Supports Path to Profitability





* Based on guidance provided as of November 12, 2014, which has not been updated since that date.



Going Forward

Positioned for Continued Growth



CONTINUE GROWTH & PROFITABILITY

Trokendi XR and Oxtellar XR peak sales potential of \$400-500 million



ADVANCE PIPELINE TOWARD COMMERCIALIZATION

Advancing both SPN-810 and SPN-812 into pivotal trials



TARGET STRATEGIC BUSINESS DEVELOPMENT OPPORTUNITIES

Execute on strategic near-commercial stage CNS opportunities

