Jefferies 2015 Healthcare Conference

June 2, 2015



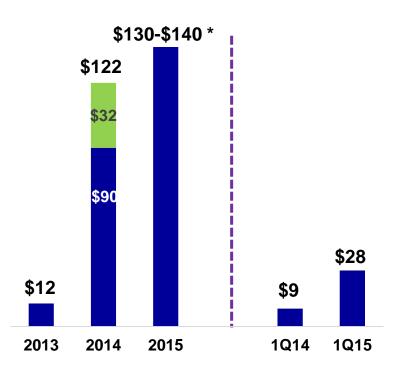
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This presentation and other matters discussed today or answers that may be given to questions asked include forward-looking statements within the meaning of the federal securities laws. These statements, among other things, relate to Supernus' business 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.

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- 25-year track record of bringing products to market
- IPO in 2012
- Successful launch of two epilepsy products in 2013
- Profitable in 2014



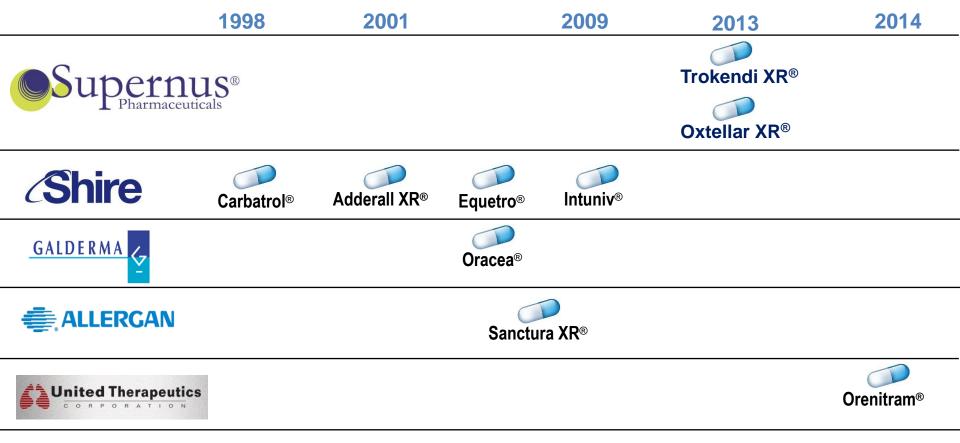
Revenue (\$ Millions)

Product Royalty/Licensing



* Based on guidance provided as of May 5, 2015, which has not been updated.

Proven Execution Nine Marketed Products Using Our Technologies



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Strong Portfolio of CNS Products

Product	Indication	Development	NDA	Launch
Oxtellar XR®	Epilepsy			February 2013
Trokendi XR®	Epilepsy			August 2013
SPN-810	Impulsive Aggression in ADHD		Phase III Trial 4Q201	5
SPN-812	ADHD		Phase IIb Trial 4Q2015	
SPN-809	Depression		IND	



Product Portfolio Targets Large Addressable Markets



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



Trokendi XR® and Oxtellar XR® Drive Future Growth



Non-Compliance – A Serious Problem in Epilepsy

71% of patients miss a dose at least once/month 45% <u>reporting seizures</u> after a missed dose

Serious Quality of Life Issues



Non-compliance leads to breakthrough seizures that cost annually >\$26,000 per patient

Increased Healthcare Costs

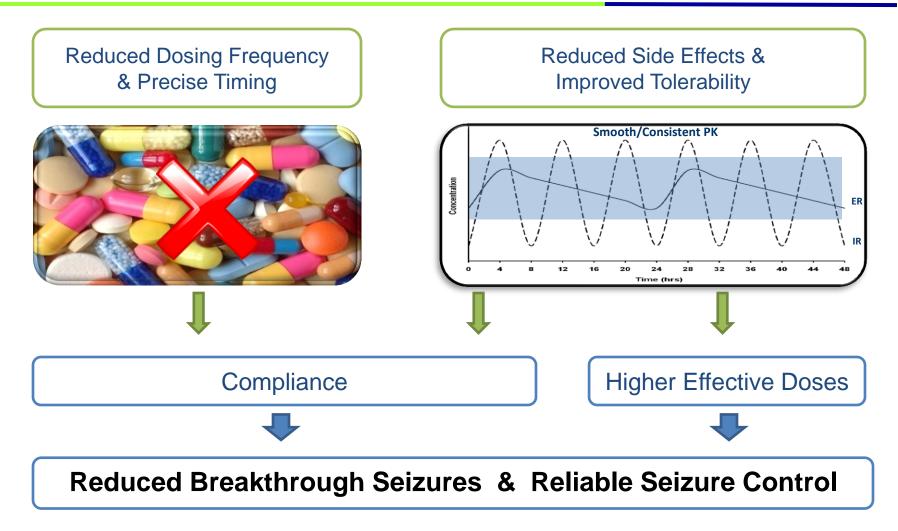


Worsening of Condition





Extended-Release AEDs = Significant Patient Benefits

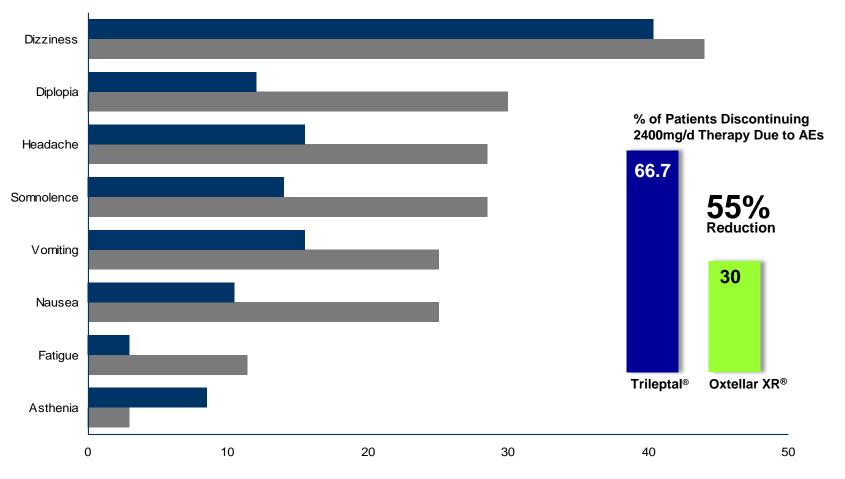


Extended Release ("ER"), Immediate Release ("IR"), Anti-Epileptic Drugs ("AEDs") and Pharmacokinetics ("PK").



Oxtellar XR[®]

Improved AE Profile at Double the Dose of Trileptal®



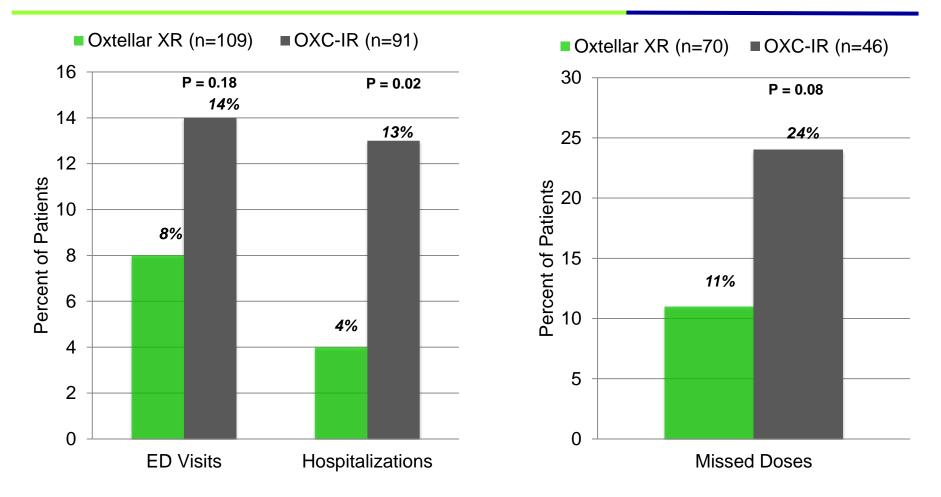
■ Trileptal 1200 mg/d (n=171) ■ SPN-804O 2400 mg/d (n=123)

Based on comparison of Oxtellar XR (SPN-804O) Phase III vs. Trileptal PI (adjunctive therapy study in adults); differences in trial design exist between the two studies. Dizziness includes vertigo in Trileptal group because of change in the MedDRA system



Oxtellar XR®

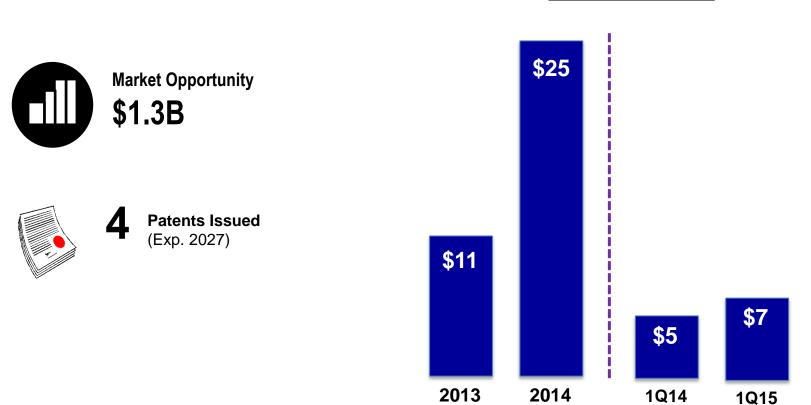
More Favorable Clinical Outcomes & Greater Adherence Compared to OXC-IR¹



¹O'Neal W, et al., Adherence and Resource Utilization with Extended-Release Oxtellar XR® or Immediate-Release Oxcarbazepine (OXC-IR) Treatment in Clinical Practice: A Standardized Case Record Review. Neurology 2015;84 (P1.244)



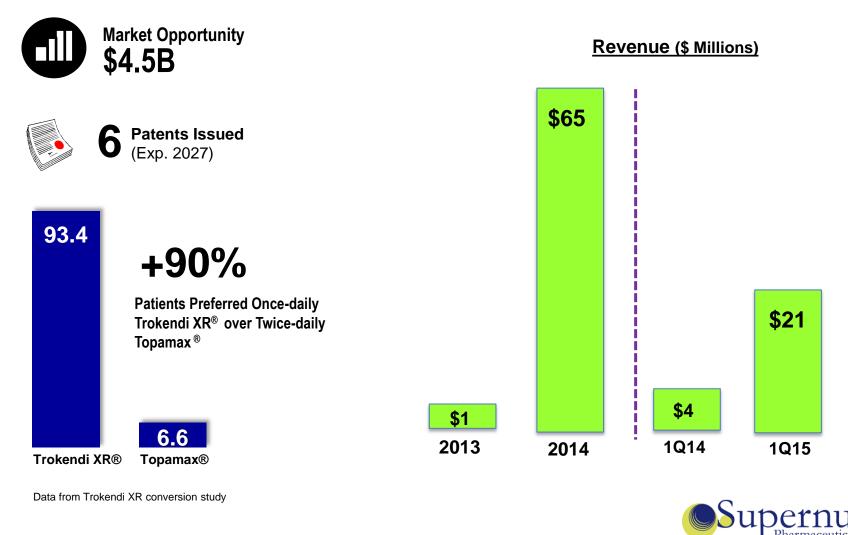
Oxtellar XR[®] Only Once-Daily U.S. Oxcarbazepine Epilepsy Product



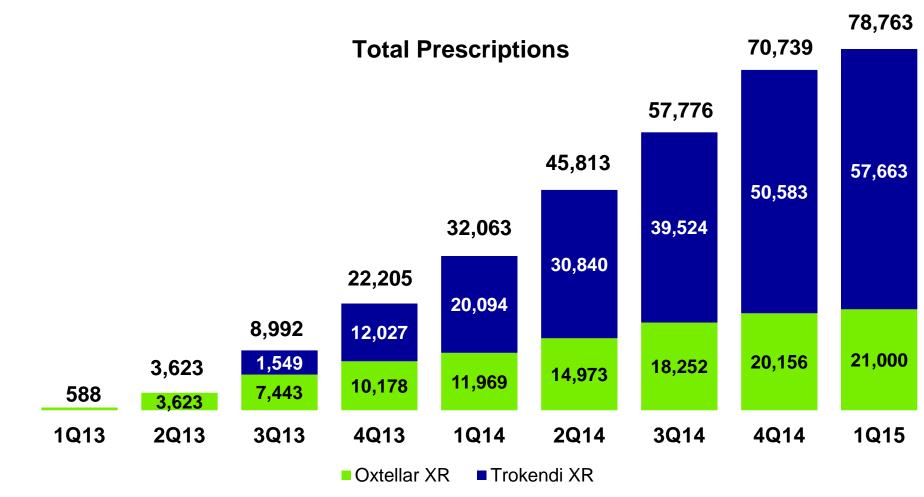
Revenue (\$ Millions)



Trokendi XR[®] First Once-Daily U.S. Topiramate Epilepsy Product



Two Successful Product Launches





Source: SHA Monthly Prescriptions

Robust, Late-Stage Pipeline Fuels 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 Aggression in ADHD	Phase III Trial 4Q2015		
SPN-812	ADHD	Phase IIb Trial 4Q2015		
SPN-809	Depression			



SPN-810: Novel Product for Impulsive Aggression (IA)



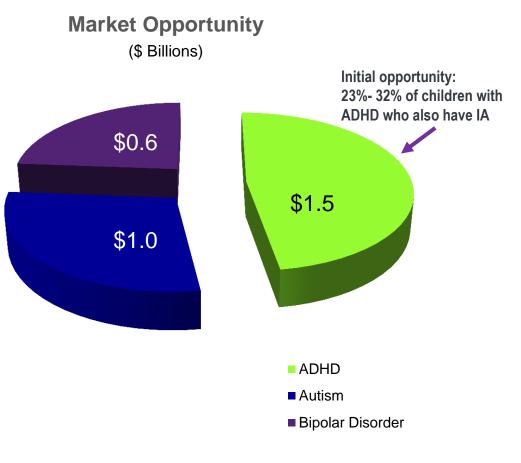
Market Opportunity +\$3B



Disorder with impairment of self regulation, characterized by episodes of unplanned, maladaptive aggression

No FDA-approved treatments

Off-label use of atypical antipsychotics common but with serious safety and tolerability issues





SPN-810: Initial IA Program in ADHD Patients

80%

60%

40%

20%

0%

Placebo

(n=30)



Granted Fast Track **Development Designation**



Expected to be first product approved to treat IA



Phase IIb trial established safety and efficacy at low and medium doses



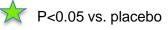
Special protocol assessment to FDA in 2Q 2015

2015

Phase III in 4Q 2015

¹Primary endpoint based on FDA input: Change from baseline at visit 10 in R-MOAS score. LOCF, ITT population

Medium dose High dose Low dose (n=27) (n=30) (n=31)



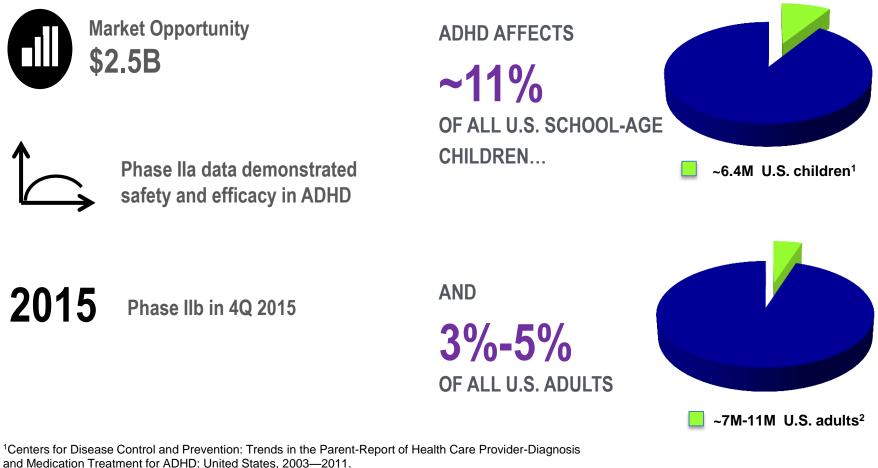


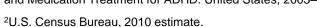
R-MOAS = Retrospective Modified Overt Aggression Scale

Phase IIb Demonstrated Greater Improvement from Baseline¹

Improvement vs. Baseline

SPN-812: Novel Non-Stimulant ADHD Product





Successful Commercial Business Financial Overview



Financial Summary and Guidance

First Quarter 2015 Financial Results

- Net product sales of \$28.1 million, up from \$9.0 million in prior year
- Operating income of \$3.4 million, compared to operating loss of \$(13.4) million in prior year
- Cash flow positive on operating basis
- March 31, 2015 cash balance of \$92.2 million

Full Year 2015 Financial Guidance Provided as of May 5, 2015

- Net products sales: \$130 million \$140 million, up ~50% from prior year
- Operating income: \$6 million \$10 million



Positioned for Continued Success



CONTINUE GROWTH & PROFITABILITY

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



ADVANCE PIPELINE TOWARD COMMERCIALIZATION

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



TARGET STRATEGIC OPPORTUNITIES

Execute on strategic near-commercial stage CNS opportunities

