



Targeting unmet needs.

Creating solutions.

Improving healthcare.



Rodman & Renshaw 18th Annual Global Investment Conference

September 12, 2016

NASDAQ: CAPN

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Highlights

Multiple Commercial Products

Revenue Generating

CoSense® ETCO Monitor

Serenz® Allergy Relief

Infant pulmonary resuscitation solutions

World-class Partners

Direct Sales plus Global Distributors

Active partnerships in place in the U.S., China, India, Turkey, Italy, Canada & Saudi Arabia

Planning to engage in further geographies in 2016

Clinical Stage Pipeline

Portfolio of Therapeutic and Diagnostic Candidates

Two Phase 2 trials ongoing in cluster headache and trigeminal neuralgia

Top-line data readouts expected in 2016

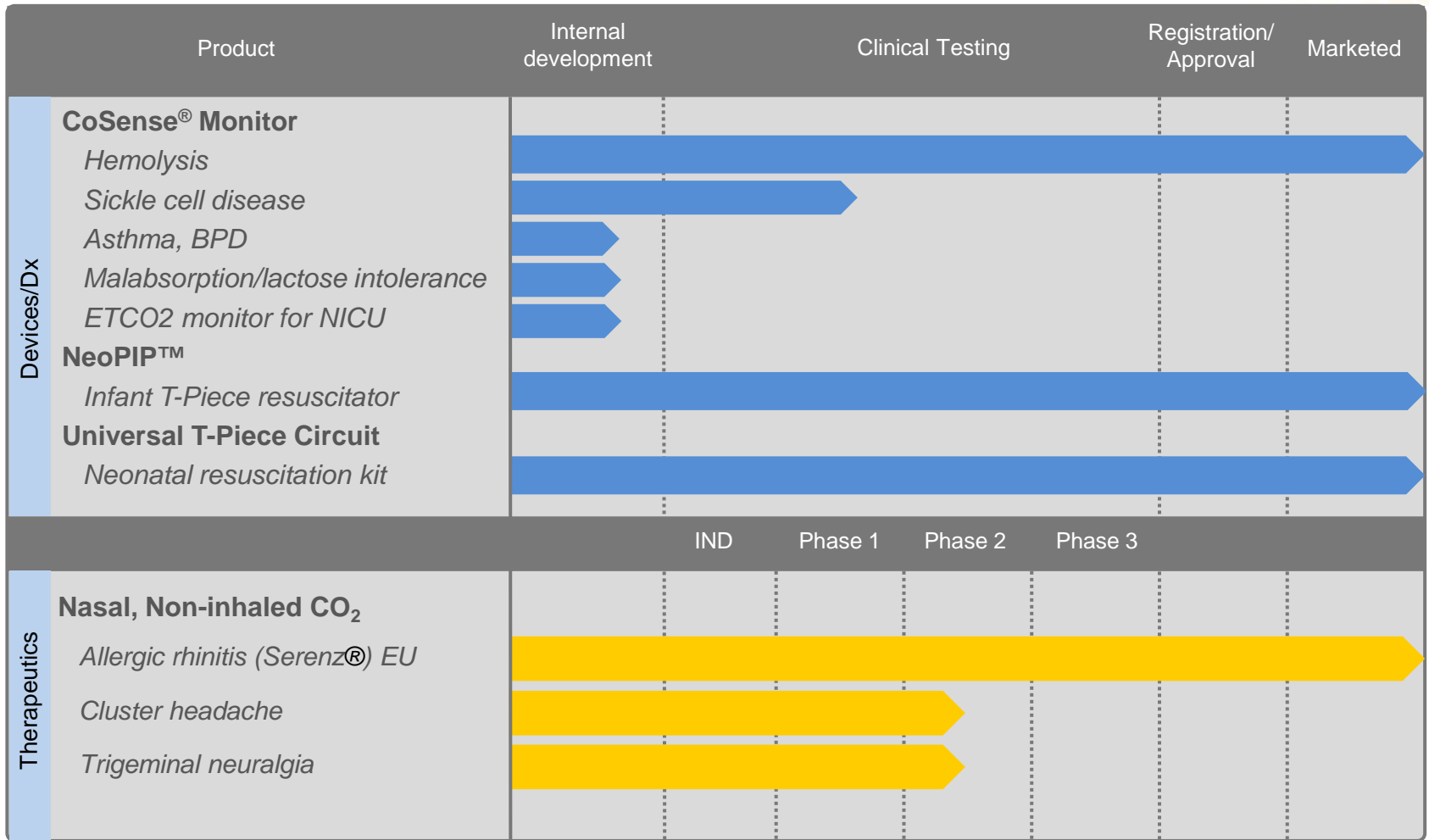
Novel Platform Technology

Cutting edge device for detecting breath analytes

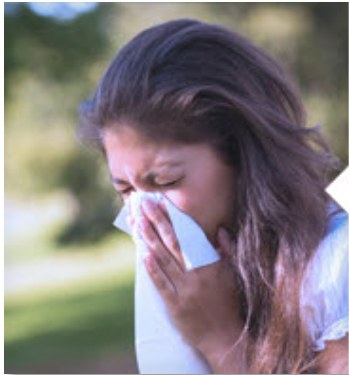
Patent portfolio (issued and pending) with expirations out to 2030s

Applicable across a broad range of therapeutic areas

Pipeline



Diversified, Revenue Generating Product Portfolio



Serenz® Allergy Relief

Serenz is a hand-held device that is designed to provide rapid relief from the symptoms related to allergies.



CoSense® ETCO Monitor

The *CoSense* ETCO Monitor measures carbon monoxide levels in exhaled breath, which may be used to detect the rate of hemolysis.



Infant Solutions

NeoForce is a portfolio of innovative pulmonary resuscitation solutions in the neonatal market.





Serenz[®]

ALLERGY RELIEF



Serenz® **Allergy Relief**

Serenz is the first allergy treatment to use carbon dioxide to relieve nasal allergy symptoms

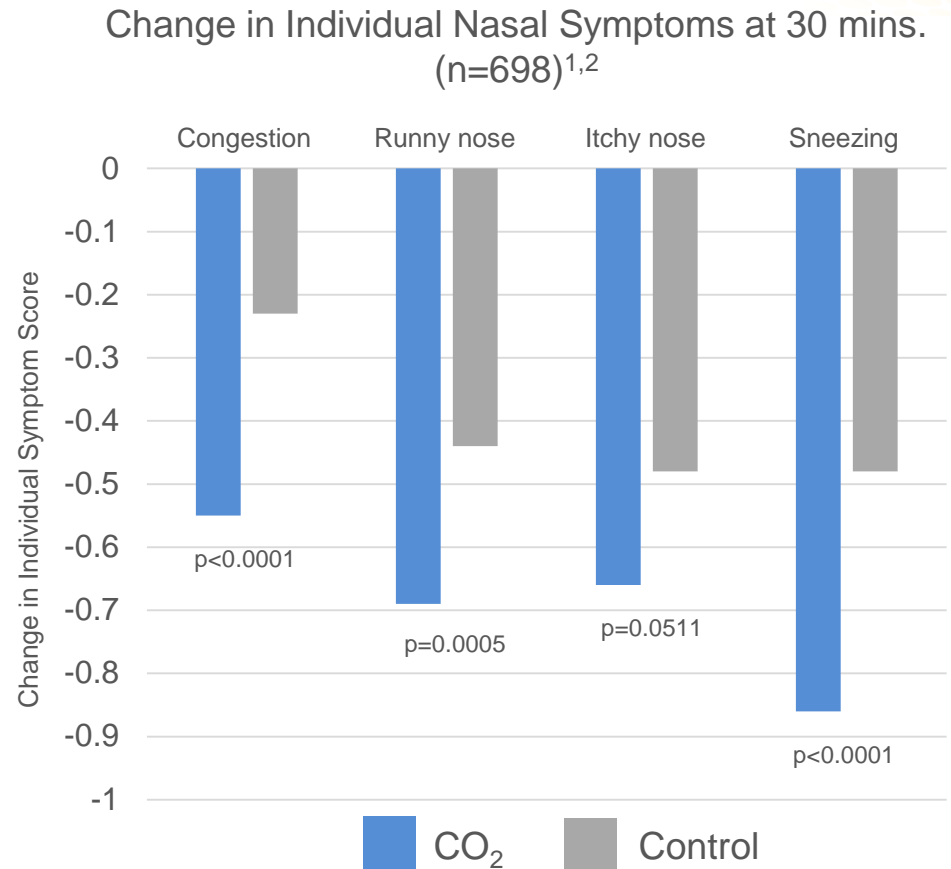
- *Fast acting*
- *Small, portable, disposable*
- *Gentle cleansing of nasal mucosa*
- *Use as needed*
- *Contains 20 cleanses*

Clinically proven to work quickly without causing drowsiness¹

¹While cleansing with Serenz, a temporary nasal discomfort may be felt or eyes may water; these will go away when cleansing is done.

Clinically Proven for Rapid Relief

- ▶ Supported by robust clinical data
- ▶ Studied in a total of 975 patients across six randomized, controlled clinical trials
- ▶ Well-tolerated and statistically significant improvements in standard Total Nasal Symptom Score (TNSS) endpoints



¹Episodes with pre-Use TNSS ≥ 8

²Casale TB et al., Congress of EAACI, 2010; Allergy 65 (S92): 35.

Global Pilot Launches

- ▶ European pilot launch coincided with 2016 allergy season
- ▶ Now available OTC at 150+ retail pharmacy locations and online in the UK and Ireland
- ▶ Full European commercial launch planned for 2017



NEW

Love Spring

Introducing a gentle and effective way to treat nasal allergy symptoms

Serenz™
Allergy Relief

To relieve allergy symptoms such as nasal congestion, sneezing, nasal itching and runny nose.

✓ RAPID RELIEF
✓ USE ONLY WHEN NEEDED
✓ CLINICALLY PROVEN
✓ NASAL CLEANS

Serenz™
Allergy Relief

Net Contents 7.5g (0.26 oz)

Serenz™ gently cleanses the nasal passages with carbon dioxide to relieve allergy symptoms of nasal congestion, itchy, runny noses and sneezing. In clinical studies, the most common side effects were temporary nasal discomfort and watery eyes while cleansing with Serenz™.

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Addressing Unmet Needs: Validated by Physician and Customer Feedback



"It's really reassuring that the clinical trials and the patient feedback in day-to-day practice have been so overwhelmingly positive for Serenz."

*-- Dr. Mariette Grant,
General Practitioner*



"I've tried every single nasal spray, tablet, and there's not really much of an effect from any of them. Serenz has definitely made a big difference in my life. I'd highly recommend it."

*-- Jordana Wallis,
Allergy sufferer*



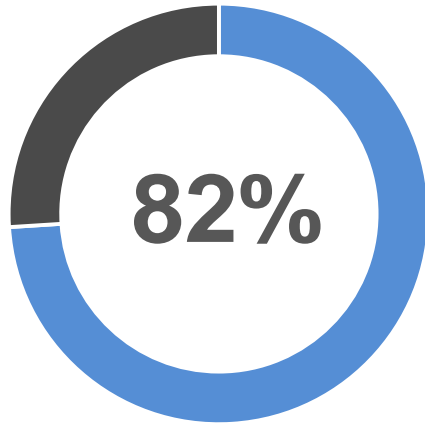
"It's been massively helpful compared to the other products I've used in the past."

*-- Ifetkhar Hussain,
Allergy sufferer*

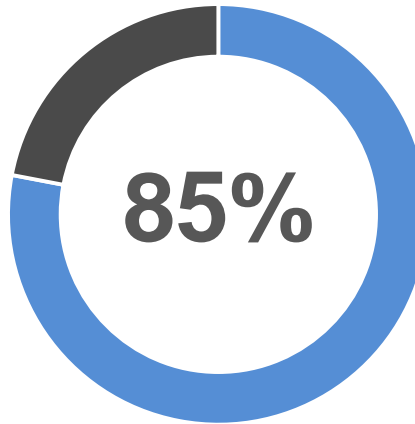
www.serenz.co.uk



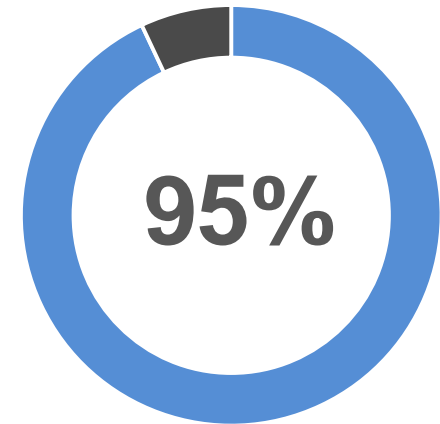
Early Commercial Traction^{1,2}



Very likely or likely
to purchase again



Very likely or likely to
recommend to family/friend



Serenz[®] met or
exceeded expectations

“Rapid relief” most important
purchase consideration

Average reported use:
2.1x / day
6.3x / week

EU Market Opportunity

EU Market Potential^{1,2}

\$621M



	Allergic rhinitis prevalence (M)	# of patients on treatment (M)	Estimated market potential (M; \$USD) ^{3,4}
UK	14.1	7.5	\$133
Germany	15.1	9.0	160
Spain	9.0	3.7	89
France	13.3	6.2	110
Italy	9.3	5.0	120
Ireland ⁵	1.0	0.5	9
Total	60.8	31.4	\$621

¹Charles River Associates (CRA) Market Research and Financial Model 2012

²Capnia has only initiated a pilot launch in the UK and Ireland. This is a competitive market and accomplishing any meaningful market share will be difficult.

³Assumes an average of 27% of patients on treatment across these territories are willing to pay Serenz price point

⁴Assumes an average seasonal revenue value across these territories of \$72

⁵Distributor estimates

Commercial Strategy

- ▶ Pursue additional pilot sales launches in select geographies
- ▶ Select leading local and regional distributors/retailers
- ▶ Engage new consumers with geography-focused, direct-to-consumer marketing efforts utilizing digital and traditional media
- ▶ Establish brand as first CO₂ allergy treatment
- ▶ Patent protection to 2020 or beyond





CoSense® ETCO Monitor

The *CoSense* ETCO Monitor is indicated for the monitoring of carbon monoxide from endogenous sources (including hemolysis) and exogenous sources (including CO poisoning and smoke inhalation) in exhaled breath. The end-tidal carbon monoxide level can be used for the monitoring of carbon monoxide in medical conditions in which the rate of hemolysis may be relevant. It is also for use in smoking cessation programs and can be used for the screening of CO poisoning and smoke inhalation.


Clinical Management of Jaundiced Infants: A Significant Unmet Need



140 million births
worldwide

9 million births in
US + Europe



 **60-80%**

develop JAUNDICE

Condition caused by elevated levels of
bilirubin or hyperbilirubinemia

In some cases

Hyperbilirubinemia + JAUNDICE

are caused by a **hemolytic condition**
(red blood cells degrade rapidly)



***Increased
risk of
neurological
damage***

Hemolysis: A Key Risk Factor for Neurological Damage

▶ CO and Bilirubin are produced **1:1** during hemolysis



▶ CO is elevated in the newborn with a hemolytic condition



▶ CO is eliminated through lungs (ETCO)



CoSense® ETCO Monitor

- Measures ETCO
- May improve treatment decisions

*“ETCOc levels can confirm the presence or absence of hemolysis, and measurement of ETCOc is the **only clinical test** that provides a direct measurement of the rate of bilirubin production...”¹*

The *CoSense* End Tidal Carbon Monoxide Monitor is not a diagnostic or treatment device for any specific disease or condition. Rather, it is a detection tool to be used for the monitoring of carbon monoxide from endogenous sources (including hemolysis) and exogenous sources (including CO poisoning and smoke inhalation) in exhaled breath.

¹ Management of hyperbilirubinemia in the newborn infant 35 or more weeks of gestation. Pediatrics 2004;114:297-316.

Current Standard of Care: Adverse Outcomes and Higher Costs

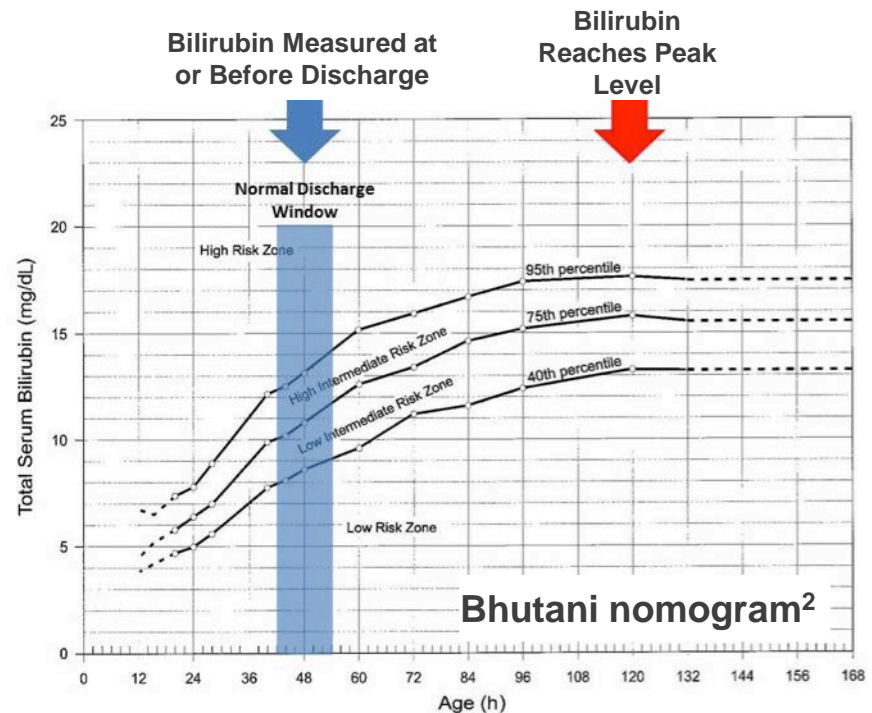
CHALLENGES

- Bilirubin typically peaks after discharge
- Current diagnosis of hemolysis is invasive and inadequate



CONSEQUENCES

- Hemolysis is under-diagnosed
- Treatment is delayed
- Adverse neurological outcomes
- Jaundice is #1 cause of infant readmits to hospital¹



AAP guidelines recommend ETCO for identification of hemolysis but no tool exists

¹ Agency for Healthcare Research and Quality (AHRQ) Healthcare Cost and Utilization Project Fact Book 4: Care of Children and Adolescents in U.S. Hospitals. <http://archive.ahrq.gov/data/hcup/factbk4/factbk4.htm> . April 27, 2014.

² Bhutani VK, Johnson L, Sivieri EM. Predictive ability of a predischarge hour-specific serum bilirubin for subsequent significant hyperbilirubinemia in healthy term and near-term newborns. Pediatrics.1999;103: 6– 14

Capnia Solution: *CoSense*® ETCO Monitor



- ▶ Portable, non-invasive device that measures ETCO, which detects the rate of hemolysis
- ▶ Provides physician information to
 - Assess risk of serious disability and supports early action if needed
 - Identify lower risk newborns needing less intervention, less hospital time
- ▶ FDA 510(k) clearance, CE certification
 - Reusable device + Precision Sampling Set (single use)
- ▶ Quick bedside results, no lab delays



Easy to
use



Point-of-care
testing



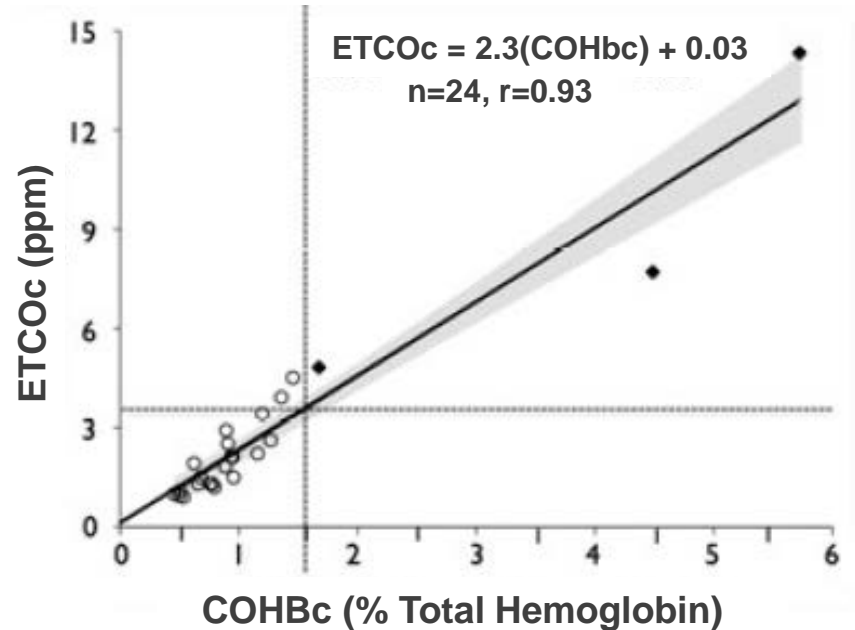
Rapid
results



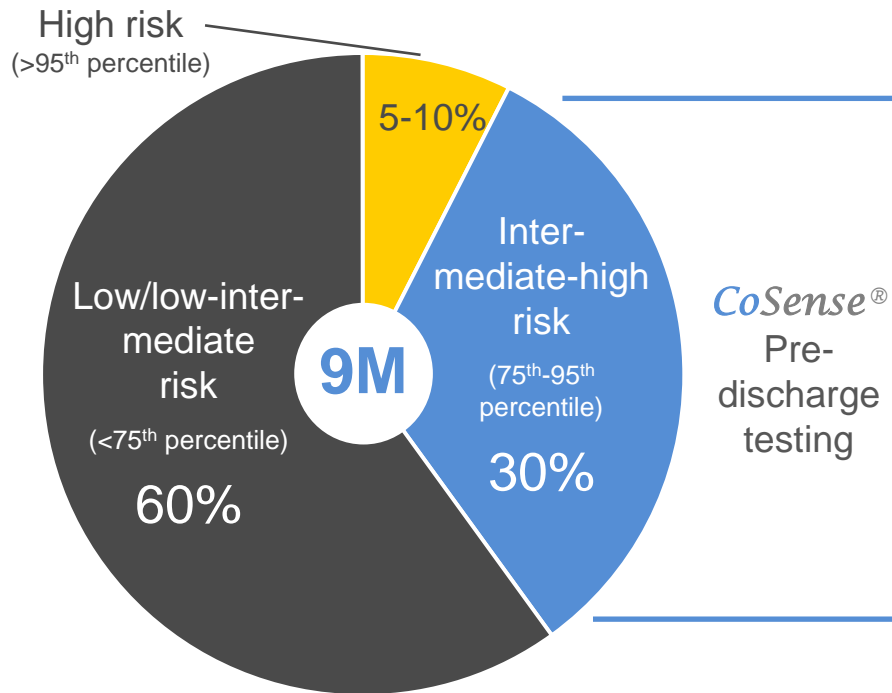
No
calibration

CoSense® Effectively Detects Hemolysis in Infants

- ▶ *CoSense*® ETCO values from 73 newborns (102 separate measurements) at Stanford University
- ▶ Carboxyhemoglobin (COHb) levels in blood are used for experimental measurements of hemolysis
- ▶ Linear correlation between *CoSense*® ETCO and COHb indicates accurate detection of hemolysis by *CoSense*®
- ▶ $r = 0.93$



Neonatal PredischARGE Management: A New Approach



9M births annually in the
US and EU

ETCOc	Approach	# of babies
≥ 1.5 ppm TB-ROR < 0.2	Observe	2.7 million
1.6 – 2.5 ppm TB-ROR > 0.2	Consider for phototherapy	
> 2.5 ppm TB-ROR > 0.2	Urgent phototherapy	

Earlier identification results in:

- Earlier discharge for low risk neonates¹
- More targeted use of phototherapy¹
- More aggressive treatment for high risk neonates¹
- Reduced hospital re-admissions²

Changing the Neonatal Care Landscape

California



Neonatal care pioneer has made *CoSense*® a routine part of newborn care with more than 25% of births being tested with CoSense

California



Moving *CoSense*® from NICU to the well-baby unit to eliminate use of the Direct Coombs Test

Colorado



During a two-week evaluation period, 2 of 18 (11%) *CoSense*®-tested newborns being discharged were found to be high-risk

Florida



Has implemented a Six Sigma quality initiative designed to reduce newborn length of stay

CoSense® is being used as an important tool to identify which newborns can be discharged with confidence

Economics for an Example Hospital



**1,000 births
per year**



One-time initial
CoSense[®]
purchase



\$5-20K¹



Recurring purchase of
Precision Sampling
Sets (PSS)



1,000 births per year
x 30% at intermediate-high risk
300 babies
x 1.5 tests per baby
450 PSS per year
x \$135 retail price
\$60,750 per year

Commercial Strategy: US

- ▶ Nationwide distribution agreement.
 - 50 states; 44 neonatology-focused sales reps
 - Extensive network of sub-distributors

Influential early adopters



Target Market: ~400 Hospitals

Tier 1

- >5k births per year
- 91 hospitals
- Northside, Sharp Memorial, University of New Mexico

Tier 2

- >3k births per year
- 193 hospitals
- Intermountain Healthcare, University of Cincinnati, Lucile Packard Children's Hospital, Vanderbilt

Tier 3

- <3k births per year
- 113 hospitals
- Dartmouth Medical Center, Swedish Covenant, UCSD

Commercial Strategy: ROW

▶ Active distribution partnerships

- China
- India
- Turkey
- Italy
- Canada
- Saudi Arabia
- Qatar



▶ Regulatory submissions in process where needed

▶ Ongoing discussions in several geographies

▶ New agreements expected in 2H 2016



NeoForce Infant Solutions

Highlights

- ▶ Revenue-generating product line
- ▶ Significant potential for growth with Capnia resources
- ▶ Leveraging existing hospital relationships/contracts; NICU services
- ▶ Multiple new product development opportunities
- ▶ Similar call point as CoSense



NeoPIP™ Infant T-Piece Resuscitator



Consumables (e.g., resuscitator circuits, masks)



Other Therapeutic and Diagnostic Development Programs

Nasal, Non-Inhaled CO₂

Our nasal, non-inhaled CO₂ technology is currently undergoing clinical investigations in the U.S.



Trigeminal Neuralgia

A clinical condition characterized by debilitating pain in regions of the face innervated by one or more divisions of the trigeminal nerve.



- ▶ Designated an Orphan Drug by the FDA
- ▶ Phase 2 clinical data (n≈50) expected by year end 2016



Cluster Headaches

Characterized by recurring bouts of excruciating pain in one side of the head.



- ▶ No effective treatment currently available
- ▶ Phase 2 clinical data (n≈25) expected by year end 2016

Sickle Cell Initiative

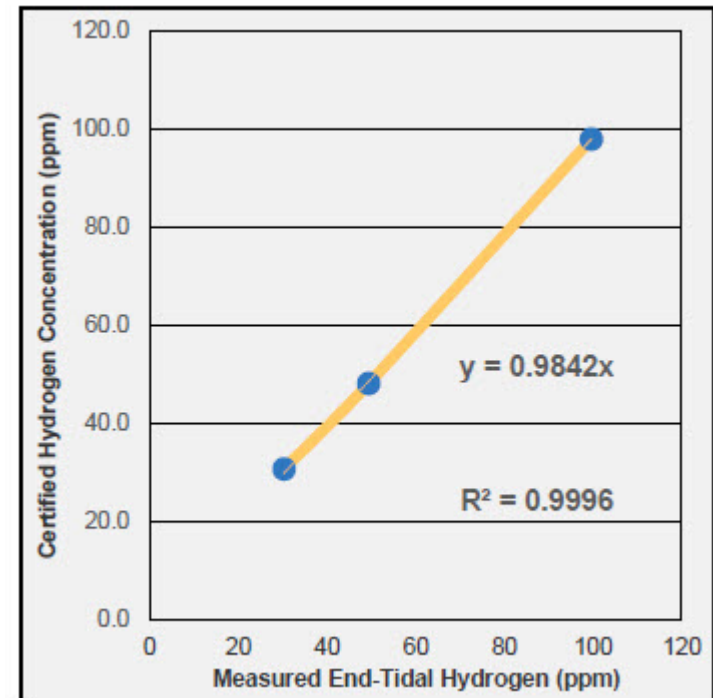
- ▶ Exploring potential of using *CoSense*® Monitor to screen for SCD
- ▶ Phase I SBIR grant received from NIH
 - Developing modified device that accurately and consistently measures ETCO in more extreme temperature conditions in developing countries (e.g., sub-Saharan Africa)
 - Where 90% of the SCD births occur
- ▶ Significant potential for use as:
 - Non-invasive screening device
 - Aid in development of new therapeutics
 - Assist in dose determination
- ▶ Ongoing conversations with governmental and other agencies

Neonatal/Pediatric Hydrogen Monitor

- ▶ Hydrogen breath tests routinely used to diagnose GI disorders, but limited to older patients
 - Lactose/fructose/sucrose intolerance
 - Irritable bowel syndrome
 - Small intestinal bacterial overgrowth
- ▶ Current tests use controlled forced exhalation through a mask or mouthpiece-not feasible for neonates

Sensalyze Hydrogen breath test feasibility data presented at PAS 2016

ETH₂ Linearity (Accuracy vs Hydrogen Level)



GR Gourley, et al. Poster Presentation 1511.499. PAS 2016

Capnia Leadership Team



MANAGEMENT

Anish Bhatnagar, M.D., *President & CEO*

- Senior Capnia executive since 2006
- Product approval experience at Coulter (Bexxar), Capnia (Serenz, *CoSense*), management at Titan

David O'Toole, *SVP & Chief Financial Officer*

- Former CFO for Codexis (CDXS), Response Genetics (RGDX), Abraxis (ABBI)
- Partner at Deloitte focused on biotech and life science industry

Otho Boone, *VP & General Manager of Neonatology*

- 25 years of neonatology medical device sales, operations and product development experience
- Founded NeoForce, which was acquired by Capnia in September 2015

Anthony Wondka, *SVP of R&D*

- Former VP R&D, VP Technology and Clinical Affairs for Breathe Technologies
- Several approvals of devices in the respiratory space in the US and globally

Kristen Yen, *VP of Clinical and Regulatory Affairs*

- Head of Clinical Operations at Capnia since 2006
- Program management of multiple U.S. and global clinical studies

Ann Rich, *VP of Marketing*

- Former Senior Marketing Director at Arthrocare, Apnicure and Johnson & Johnson
- Managed marketing and global development of numerous products

Key Financial Information

- ▶ Revenue being generated from three product lines
 - CoSense® ETCO Monitor w/consumable Precision Sampling Sets
 - Serenz® Allergy Relief
 - Infant Solutions (neonatal resuscitation products)
- ▶ Common stock outstanding¹: 15.4 million
- ▶ Fully-diluted share count¹: 29.4 million
- ▶ Market cap¹: \$ 16 million
- ▶ Cash balance¹: \$2.5 million
- ▶ No Debt

¹ As of June 30, 2016

2016 Priorities

- ▶ Execute on commercial strategy for CoSense
- ▶ Prepare for Serenz full commercial launch in Europe
- ▶ Support recently announced distribution partnerships
- ▶ Report top-line results from Phase 2 trials in cluster headache and trigeminal neuralgia by year end

Key Achievements since IPO

Commercial Products	Product Pipeline	Corporate/Financial
<ul style="list-style-type: none">✓ Launched CoSense® globally✓ Acquired NeoForce assets✓ Secured distribution partnerships in key territories✓ Serenz CE Mark re-activated✓ Commenced Serenz® pilot launch in Europe	<ul style="list-style-type: none">✓ Awarded NIH grant to develop device to detect sickle cell disease✓ Initiated Phase 2 clinical trials in cluster headache and trigeminal neuralgia✓ Received orphan designation for nasal CO₂ in TN	<ul style="list-style-type: none">✓ Remaining Series B warrants expired✓ Secured \$26M in funding through private placements and at-the-market stock purchase agreements

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