



Cells for Therapy and Research

Rodman
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Annual Global
Investment Conference

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New York City, NY 2014



International Stem Cell Corporation
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OTCQB ISCO
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Forward Looking Statement

This presentation includes certain statements, estimates and projections with respect to the anticipated future business and performance of the Company, which are collectively referred to as forward-looking statements.

Forward-looking statements reflect various assumptions of management that may or may not prove to be correct, and are intended solely to convey our expectations or predictions about the future performance of the Company.

All forward-looking statements are inherently uncertain as they are based on our current expectations and assumptions concerning the future performance of our Company. This is not an offer to sell nor a solicitation to buy any security.

Cells for Therapy and Research

Unique stem cell platform

Proprietary

Scalable

Ethical

Commercially
successful human
cell manufacturing
subsidiary

Robust pre-clinical
pipeline addressing
large unmet
medical needs

Partnering Opportunities
Intellectual Property

UniStemCell
Human Cell
Culture

Neural stem cells
Parkinson's disease
Stroke

Liver cells
Crigler-Najjar syndrome
 α_1 -antitrypsin deficiency

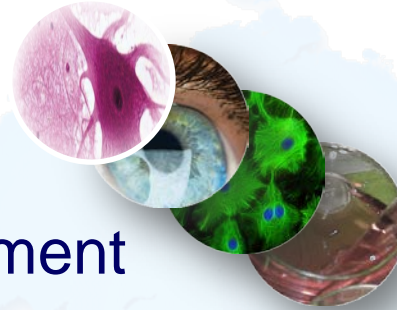
Cornea & Retina cells
Corneal blindness
Age-related Macular Degeneration

\$6 MM revenue
Primary cells & media
Dermatology products

Three Businesses in One



Core stem cell technology and IP
Therapeutic Research and Development



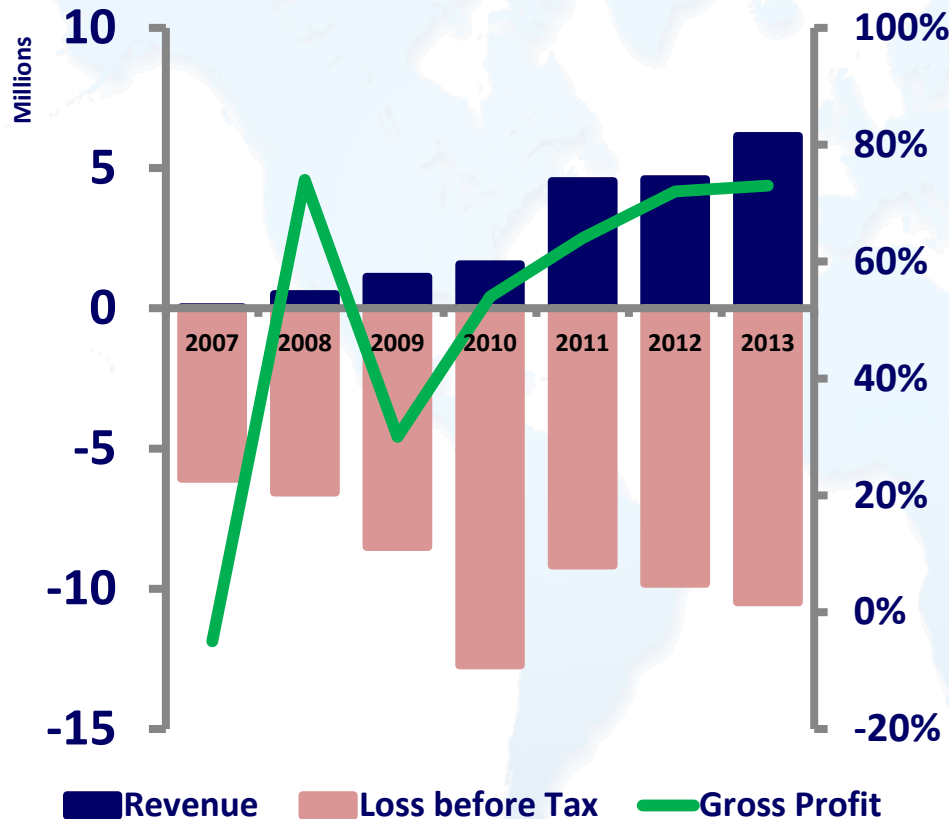
*Biomedical research products
Primary human cells &
optimized cell culture media*

**\$6.2 MM revenue
73% gross margin**



*Branded skin care products
Proprietary stem cell skin care
technology*

Growing Revenues from Subsidiaries



| | |
|---------------------------------------|----------------------|
| Trading symbol | OTCQB ISCO |
| Headquarters | Carlsbad, CA |
| Stock price | \$0.11 |
| 52 week range | \$0.06 - \$0.30 |
| Market cap. | \$25.5 MM |
| 90 day volume | 1.04 MM |
| Long-term debt | \$0 |
| Cash on hand | \$0.75 MM |
| Net cash used in operating activities | (\$3,357k) |
| Full time employees | 38 |
| Audit firm | Mayer Hoffman McCann |

*As of 7/31/14 or last 10-Q unless otherwise states

UniStemCell

Unique stem cell platform technology
Human parthenogenesis (unfertilized oocytes)

| | PSC | ESC | iPS | Adult |
|---|----------|------------------------------------|----------------------------|----------------------------|
| Potential to immune match | Yes | Impractical each line is unique | Individual only | Individual only |
| Fully proliferative/ Economic source | Strong | Strong | Varies | Weak |
| Genes manipulated/ Use of viruses | No | No | Yes | No |
| Genetic diseases/ Carries defective gene | Superior | Superior | Carries damaged gene | Carries damaged gene |

Intellectual Property

Freedom to operate in major jurisdictions
Potential competitive advantages in EU

130 patents and licenses in
14 families

90 patents pending in 5
families

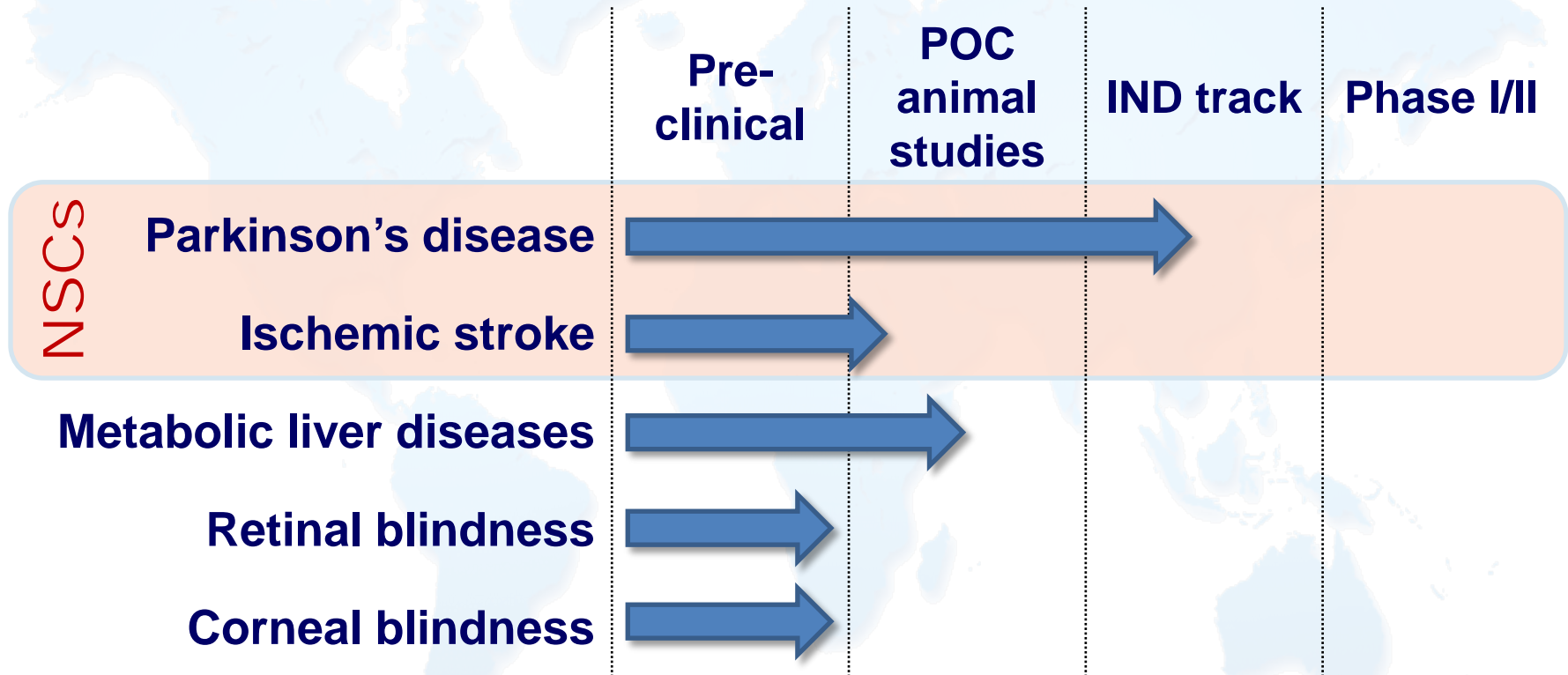
Significant peer-reviewed
publications

Nature's Scientific Reports (2013)
Cell Stem Cell (2011,2012)



Multiple Therapeutic Opportunities

Success unlocks significant value



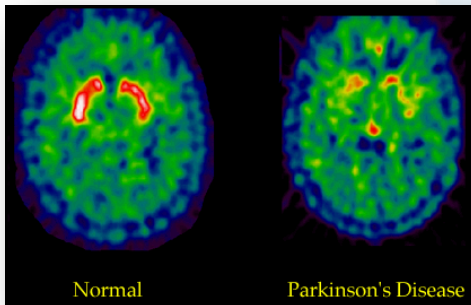
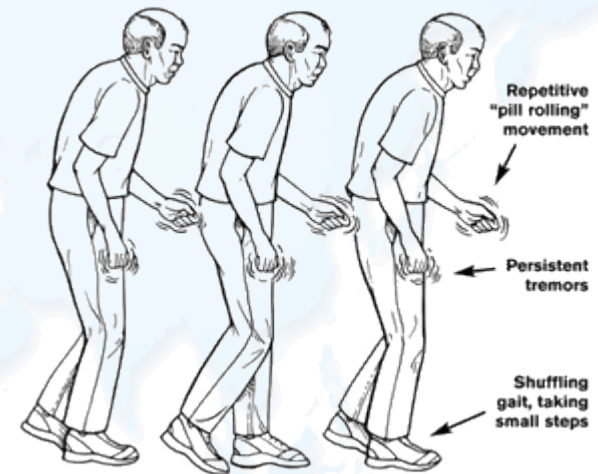
Partner with human data

Parkinson's Disease

2 Neurodegenerative disease

Limited treatment options (symptomatic only)

- About 1 million people in US have PD
- Direct and indirect costs ~\$25 billion
- Rarely diagnosed prior to 50 years of age
- Hallmark motor symptoms
 - Bradykinesia (slow movement)
 - Rigidity
 - Tremor



- Motor symptoms arise as a consequence of the degeneration of dopamine neurons in SNPC
- Greater than 50-70% of SNPC neurons degenerate prior to the appearance of overt symptoms

New Treatment for Parkinson's Disease

One-time MRI-guided injection of NSC

- *Disease modifying*
- Rebuild nigro-striatal pathway
 - Migrates and innervate
 - Produce DA neurons
- Prevent further damage
 - Express neuro-protective factors

Standard
neurosurgical
procedure

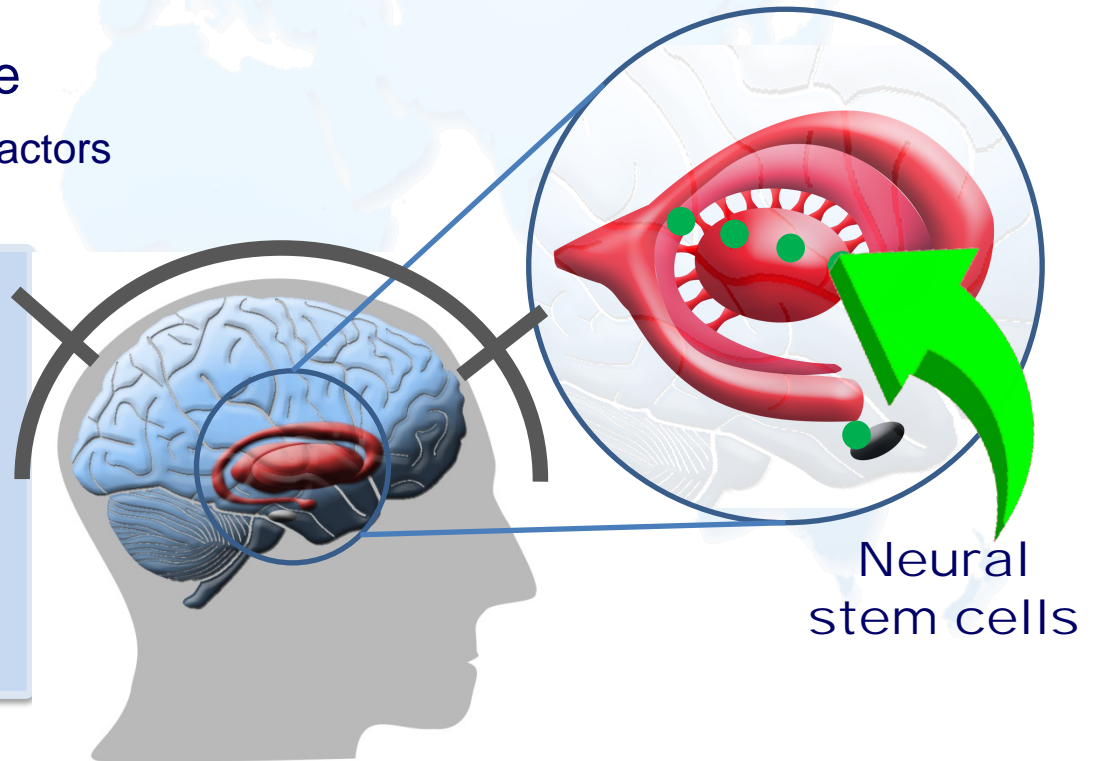
Previous human trials using fetal
tissue with over 400 patients showed:

- Generally good safety and efficacy
- Cells persist > 10 years

Lack of donors

Ethically controversial

Graft induced dyskinesias

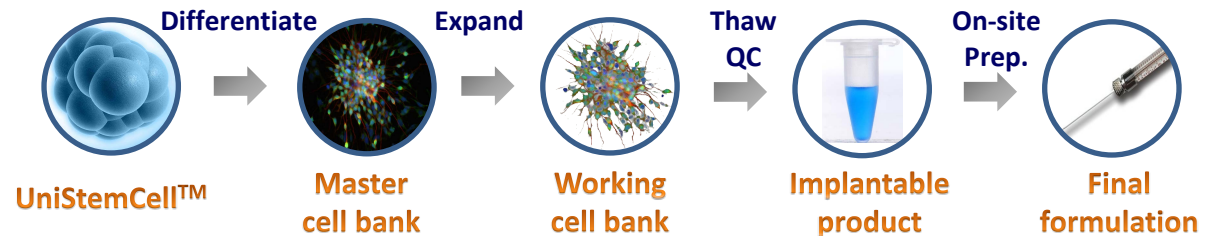


Neural
stem cells

cGMP Manufacturing of NSC

Highly optimized GMP process

- Virtually unlimited quantities of NSC
- Produces extremely pure cell populations
- Patented



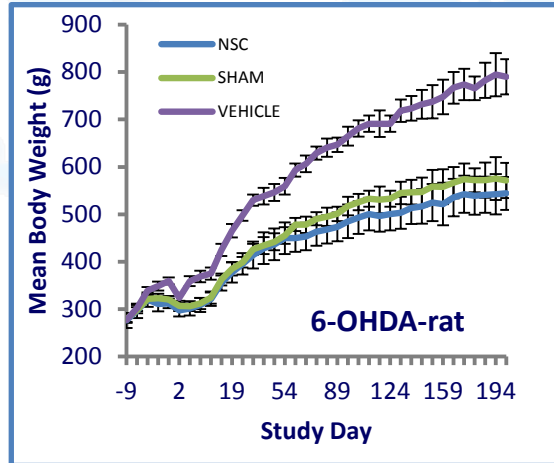
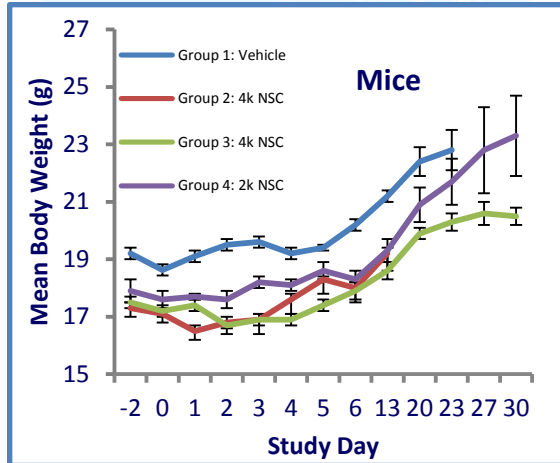
Ideal product characteristics

- Cryopreservable
- Centralized manufacturing
- Easily scalable

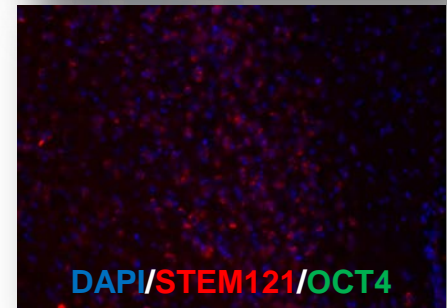
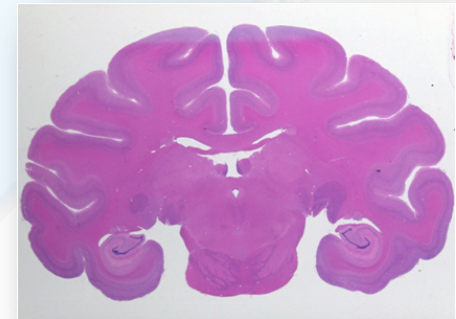


Pre-Clinical Shows Good Safety Profile

Body weights remains within normal ranges



Coronal section of the monkey brain stained with hematoxylin and eosin showing a normal morphology without the presence of ectopic tissue



Implantation site stained for human specific antibody STEM121 (in red) and OCT4 (in green), a marker of undifferentiated pluripotent stem cells. OCT4 is completely absent in the graft site.

No visible tumors or pluripotent cells

Necropsies showed organs to be normal

Biodistribution indicates no human cells in peripheral organs

Pre-clinical Evidence of Efficacy

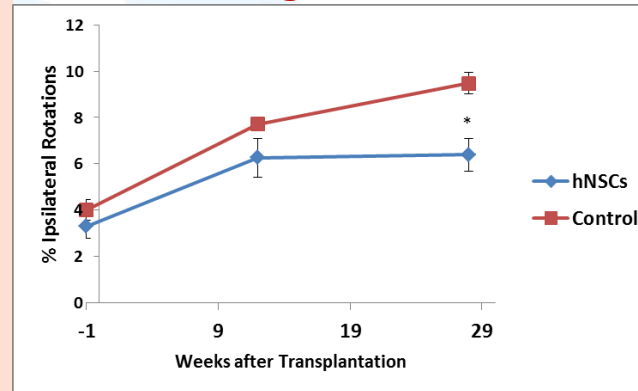
Unilateral implantation of hNSCs into the striatum of 6-OHDA-lesioned rats demonstrates functional activity post-transplantation

hNSC transplanted animals showed significant slow down in the progression of the disease and increased dopamine levels compared to sham control rats

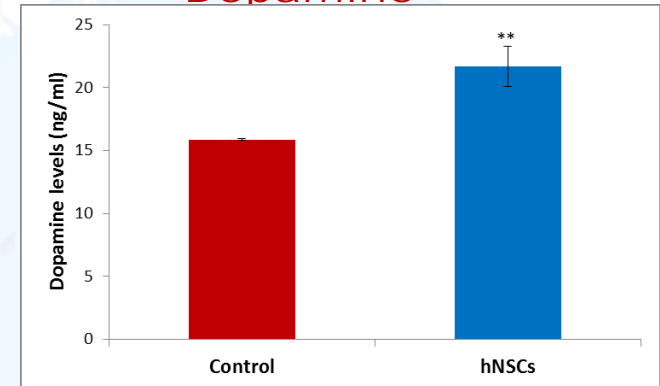
hNSCs successfully survived and engrafted 28 weeks post-transplantation and migrated from site of implantation to the substantia nigra. A small percentage of hNSC differentiated into DA neurons

hNSC therapy was safe and well-tolerated by the animals as determined by necropsy, histology, and biodistribution analysis

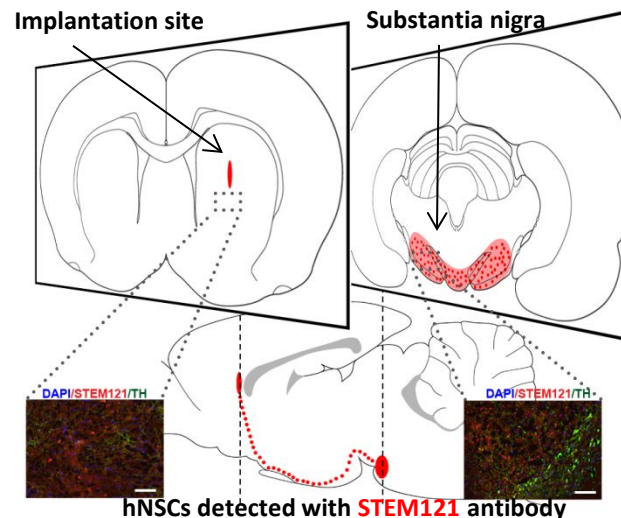
Slows Disease Progression



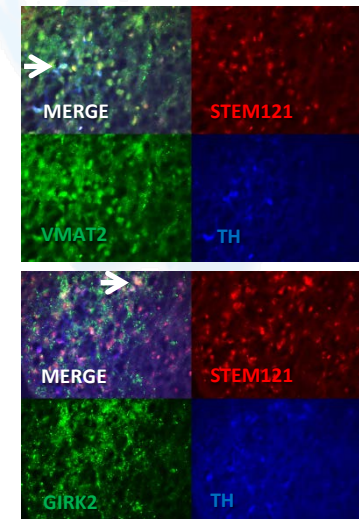
Increases Brain Dopamine



Engraftment and migration

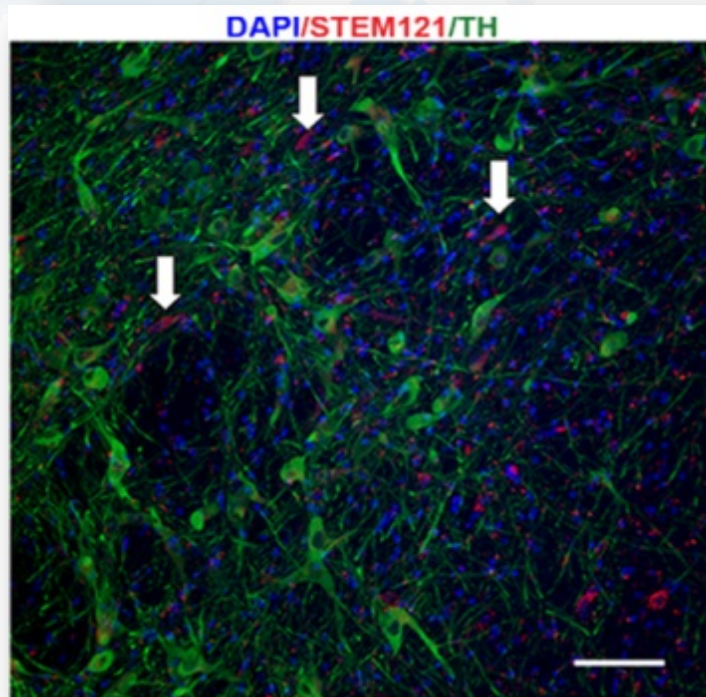


Differentiates into DA neurons *in situ*



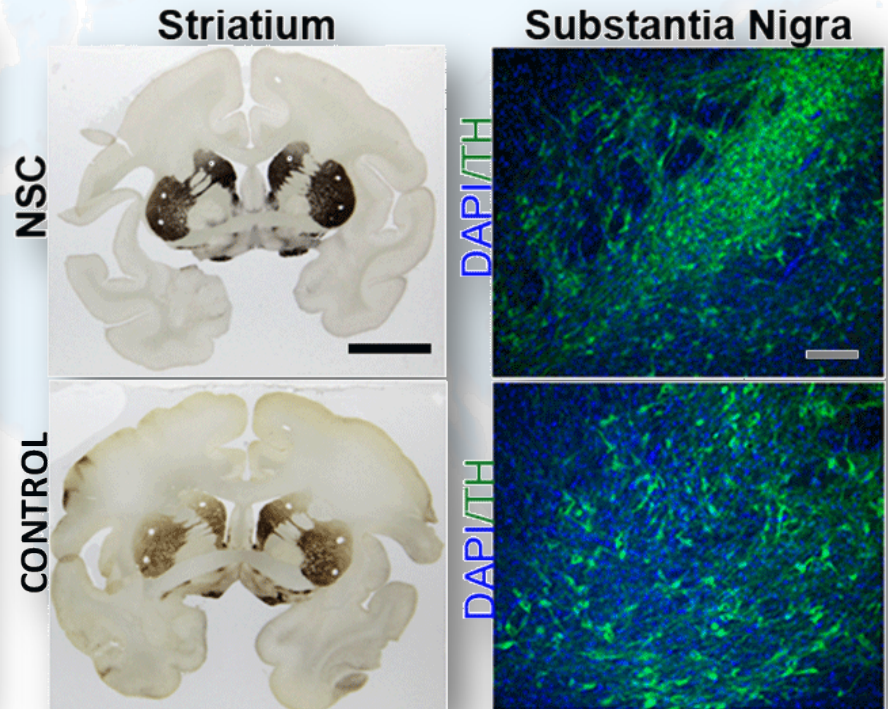
Primate Proof of Concept

NSCs integrate around
host DA neurons



Green = host DA neurons
Red = human NSC

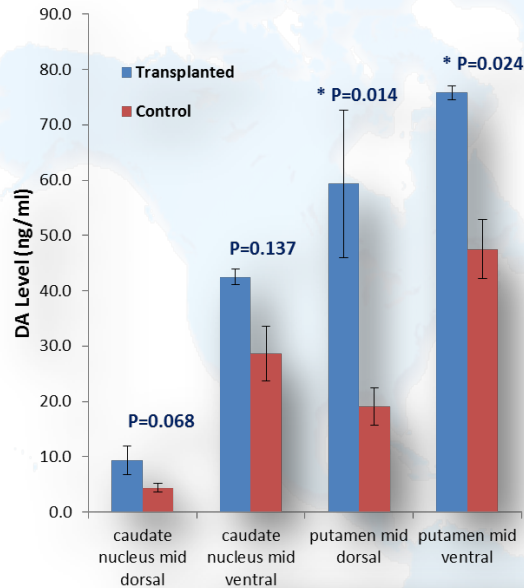
Greater density of DA neurons
and higher DA production



Green = host DA neurons

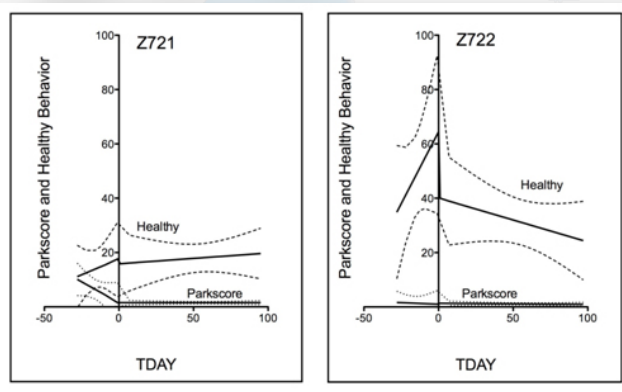
Functional Activity Established

3 Month study



Elevated DA levels *post mortem*

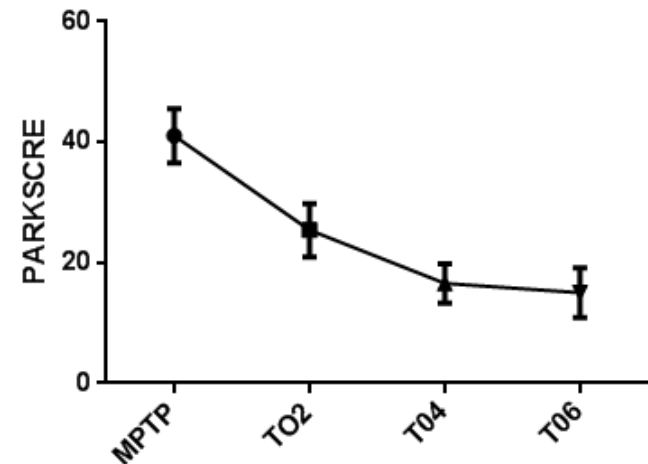
Stable Park and Health scores



(n=6, asymptomatic)

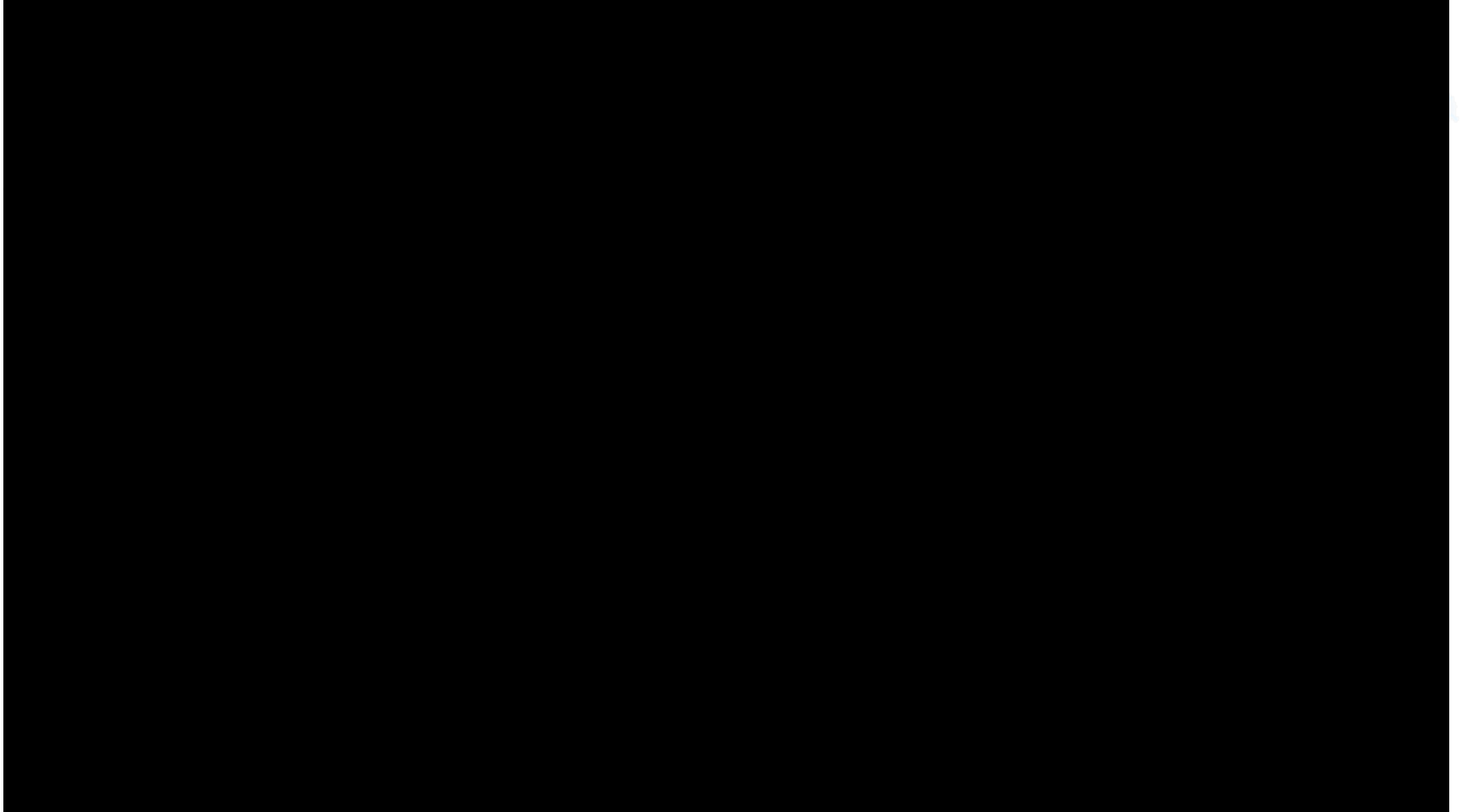
12 Month Study

18 severely symptomatic African green monkeys



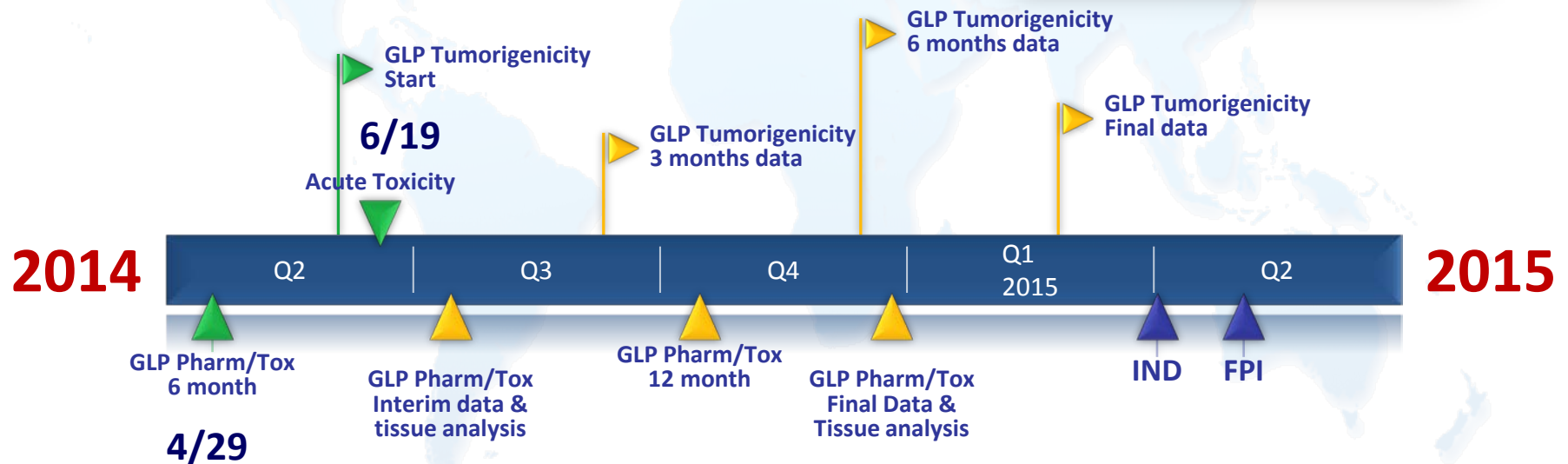
*Prof. Gene Redmond,
Yale School of Medicine

Return of Normal Behaviors



Parkinson's Disease Milestones

- ✓ Manufacturing optimized
- ✓ Good Pre-clinical safety
- ✓ Primate proof of concept



Expanding beyond PD

Pipeline of follow-on indications

NSCs

| Indication | Market Potential | Current approach |
|---------------------------------|--|-------------------------|
| Stroke ¹ | 600,000 strokes / year \$500 million / year | 3 hour treatment window |
| Spinal cord injury ² | 12,000 case / year \$360 million / year | No effective treatment |
| Dry AMD | 15 million in US \$20 billion | None |
| CytoHep | Metabolic liver disease | Transplant |
| CytoRet and CtyoCor | Forms of blindness | None |

1. Global Data 2012

2. CDC data

Strategic Partners

Rohto Pharmaceutical Company Inc.

Ophthalmology indications



DukeMedicine

Duke University

Parkinson's clinical program

Prof. Stacy MD, Vice Dean of Clinical Research

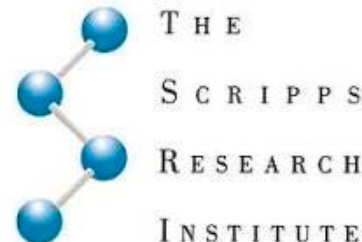
– World-renowned movement disorder expert

Academic

Dr. Evan Y. Snyder, MD PhD, Sanford-Burnham

Dr. D. Eugene Redmond Jr. MD - Yale

Dr. Jeanne Loring, PhD - Scripps



ISCO Growth Opportunities



Fast growing commercial subsidiaries

- Revenues +30% (2012 – 2013)
- Gross Profit 73%

Pipeline addressing large unmet medical needs

- Pre-clinically validated targets
- Upcoming catalysts



Broadly applicable platform

- Almost inexhaustible source
- Ethically superior
- Good IP, distinct advantages in EU



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