UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 8-K

CURRENT REPORT
PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

Date of Report (Date of earliest event reported): August 14, 2025

BioRestorative Therapies, Inc.

(Exact name of registrant as specified in its charter)

Nevada	**001-37603**	**30-1341024**
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

40 Marcus Drive Melville, New York	**11747**
(Address of principal executive offices)	(Zip code)

Registrant's telephone number, including area code (631) 760-8100

Not Applicable
(Former Name or Former Address, if Changed Since Last Report)

Securities registered pursuant to Section 12(b) of the Securities Exchange Act of 1934:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, $0.0001 par value	BRTX	NASDAQ Capital Market

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter):

Emerging growth company ☐

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Item 7.01 **Regulation FD Disclosure**.

On August 14, 2025, BioRestorative Therapies, Inc. (the "Company") made available an updated corporate presentation (the "Presentation") that may be used by the Company in connection with presentations at conferences and investor meetings. The Presentation can be found on the Company's website, www.biorestorative.com. The Presentation is furnished as Exhibit 99.1 hereto.

The information in the Presentation is being furnished, not filed, pursuant to this Item 7.01. Accordingly, the information in the Presentation will not be deemed filed for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference into any other filing under the Securities Act of 1933, as amended (the "Securities Act"), or the Exchange Act, except as expressly set forth by specific reference in such a filing. The furnishing of the information in this Current Report on Form 8-K with respect to the Presentation is not intended to, and does not, constitute a determination or admission by the Company that the information in this Current Report on Form 8-K with respect to the Presentation is material or complete, or that investors should consider this information before making an investment decision with respect to any security of the Company.

Item 9.01 **Financial Statements and Exhibits**.

(d) Exhibits.

Number	Description
99.1	Presentation Materials
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

BIORESTORATIVE THERAPIES, INC.

Dated: August 14, 2025

By: /s/ Robert Kristal
Robert Kristal
Chief Financial Officer

Exhibit 99.1


biorestorative
therapies
Regenerative Biology
for Healthier Lives
Investor Presentation
August 2025

Forward-Looking Statements

Statements in this presentation, including the information set forth as to the future financial or operating performance of Biorestorative Therapies, Inc. (the "Company") that are not current or historical factual statements may constitute "forward-looking" information within the meaning of the U.S. federal and state securities laws. When used in this presentation, such statements may include, among other terms, such words as "may," "will," "expect," "believe," "plan," "anticipate," "intend," "estimate," "project," "target" and other similar terminology. These statements reflect current expectations, estimates and projections regarding future events and operating performance and speak only as to the date of this presentation. Readers should not place undue importance on forward-looking statements and should not rely upon this information as of any other date.

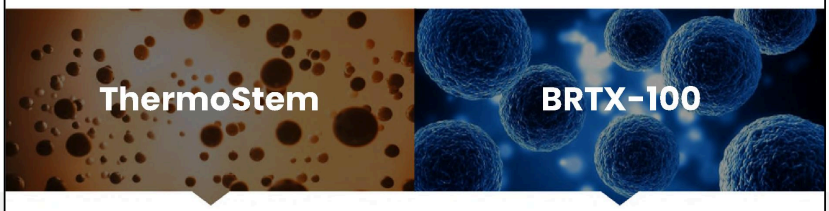
Forward-looking statements involve known and unknown risks, uncertainties and other important factors that could cause our actual results, performance or achievements, business plan or industry results, to differ materially from our expectations of future results, performance or achievements expressed or implied by these forward-looking statements. These forward looking statements may not be realized due to a variety of factors, including without limitation: (i) our limited operating history, lack of significant revenues, and substantial losses since inception;(ii) our ability to obtain sufficient financing to initiate and complete our clinical trials and fund our operations; (iii) our ability to timely and successfully develop and commercialize BRTX-100, our lead product candidate for the treatment of chronic lumbar disc disease; (iv) delays in enrolling patients in our clinical trials; (v) disruption to our access to the media (including cell culture media) and reagents the Company is using in the clinical development of our cell therapy product candidates; (vi) failure of our clinical trials to demonstrate adequately the safety and efficacy of our product candidates; (vii) our lack of manufacturing capabilities to produce our product candidates at commercial scale quantities and lack of an alternative manufacturing supply; (viii) a loss of our exclusive license rights with regard to our disc/spine technology; (ix) safety problems encountered by us or others developing new stem cell-based therapies; (x) ethical and other concerns surrounding the use of stem cell therapy which negatively impact the public perception of our stem cell products and/or services; (xi) our limited experience in the development and marketing of cell therapies; (xii) our reliance on novel technologies that are inherently expensive and risky; (xiii) significant product liability claims and litigation to which the company may be subject, including potential exposure from the use of our product candidates in human subjects; (xiv) our inability to obtain reimbursement for our products and services from private and governmental insurers; (xv) our inability to protect our proprietary rights; and (xvi) compliance with applicable federal, state, local, and international requirements. See also "management's discussion and analysis of financial condition and results of operations – factors that may affect future results and financial condition" set forth in the Company's most recent annual report filed with the SEC.

Many of these issues can affect the Company's actual results and could cause the actual results to differ materially from those expressed or implied in any forward-looking statements made by, or on behalf of, the Company. You are cautioned that forward-looking statements are not guarantees of future performance, and you should not place reliance on them. In formulating the forward-looking statements contained in this presentation, it has been assumed that business and economic conditions affecting the Company and the economy generally will continue substantially in the ordinary course. These assumptions, although considered reasonable at the time of preparation, may prove to be incorrect.

The description of the Company and its business in this presentation does not purport to be complete and is subject to the more detailed description of the Company and its business in the Company's annual, quarterly and current reports filed with the SEC.

Fully Integrated Regenerative Medicine Company

DEVELOPMENT



Brown adipose derived stem cells

PRECLINICAL

Bone marrow derived mesenchymal stem cells

MID-STAGE CLINICAL

COMMERCIAL

Secretome (novel exosome, growth factor/cytokine) biologics based technology for cosmetic applications

Experienced Leadership

Lance Alstodt
Chairman & CEO

- 30+ years leading, advising and operating companies within the Healthcare sector
- Founder of MedVest Capital, a Healthcare fund created in 2013
- Prior to that led the Medical Technology investment banking group at Bank of America Merrill Lynch and Leerink Partners

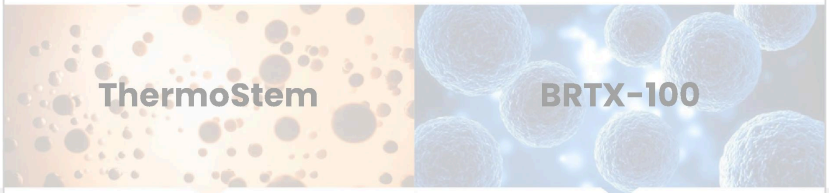
Robert Kristal
Chief Financial Officer

- 25+ years on Bay Street and Wall Street
- Most recently was the DOR for a Healthcare focused Investment Bank
- Career has spanned Trading, Sales, Investment Banking and Research

Francisco Silva
Vice President of R&D

- 20+ years in the R&D of cell-based and off-the-shelf therapeutics
- As BRTX's Vice President R&D, established high throughput Stem Cell Research Program based on his academic and industrial research experience
- Has obtained several patents in cell therapy, and has manuscripts published with regards to translational stem cell research
- Recently appointed section editor of the newly launched Regenerative Medicine section of the peer-reviewed *Journal of Translational Medicine*

DEVELOPMENT



Brown adipose derived stem cells	Bone marrow derived mesenchymal stem cells
PRECLINICAL	MID-STAGE CLINICAL

COMMERCIAL

Secretome (novel exosome, growth factor/cytokine) biologics based technology for cosmetic applications

Going Commercial: Biologics Based Cosmetic Products

Key Features of Cosmetics & Hair Growth Biologics Products

The Market

- Global Cosmetic market $63 B and growing
- Global injectable market 12% CAGR 2021-2026 Est $11.9 B
- Derms need differentiated product
- Bundle with current procedures

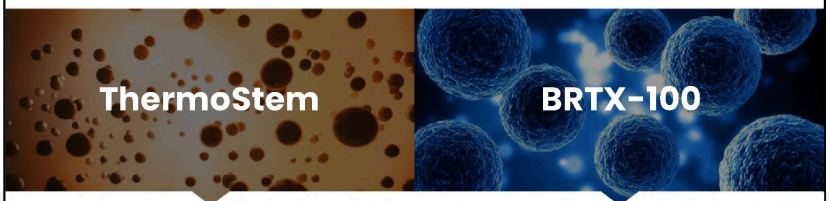
Manufacturing

- cGMP ISO 7 Certified Facility
- Cellular Biology Engineering Expertise
- Multi use facility highlights versatility

Products

- Cell based biologics engineered and targeted for both clinical and aesthetic use
- Exclusive 5-year commercial agreement with Cartessa Aesthetics

DEVELOPMENT



Brown adipose derived stem cells

Bone marrow derived mesenchymal stem cells

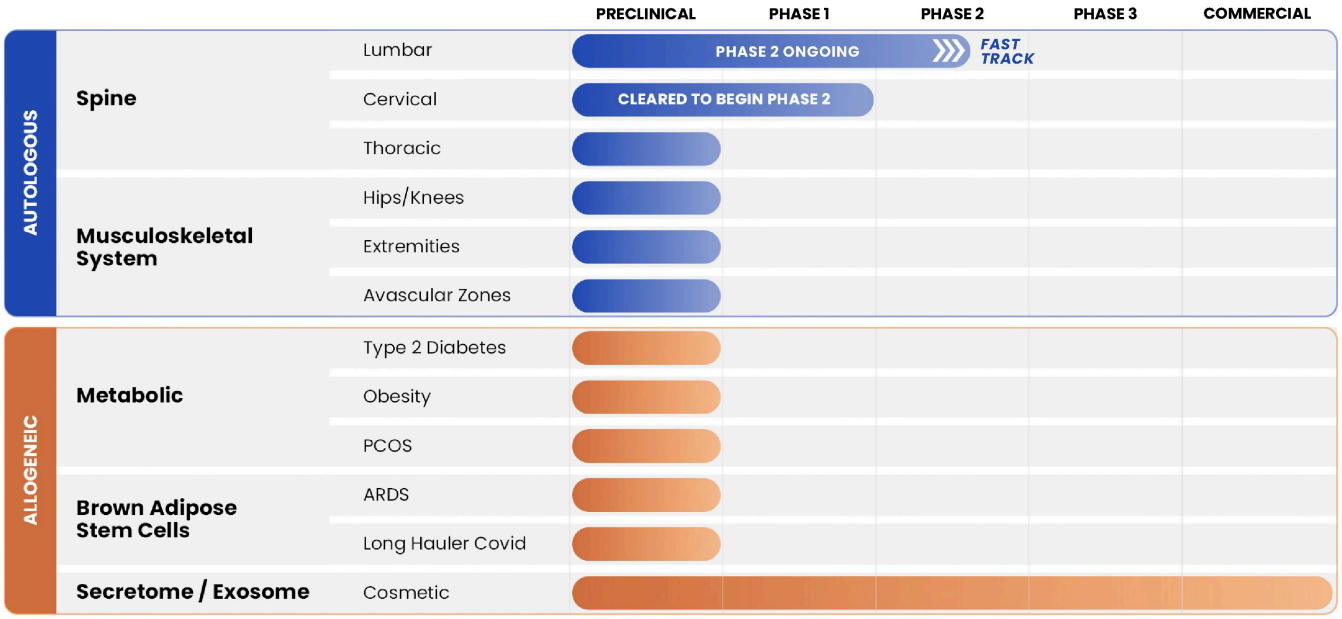
PRECLINICAL

CLINICAL

COMMERCIAL

Secretome (novel exosome, growth factor/cytokine) biologics based technology for cosmetic applications

Robust Preclinical & Clinical Pipeline



* Polycystic ovarian syndrome

Chronic Lumbar Disc Disease (cLDD)

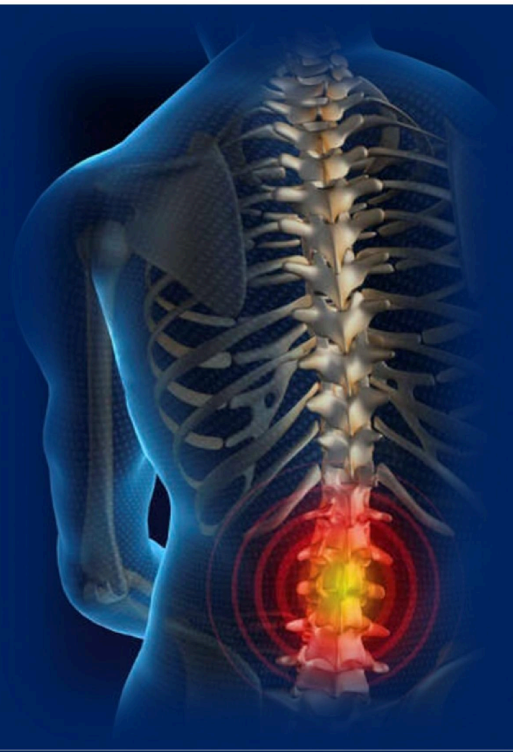
258 M U.S. adult population

64.5 M American adults with chronic lower back pain prevalence

32 M American adults with diagnosed and treated disc degeneration

15 M Americans suffering pain caused by a protruding or injured disc

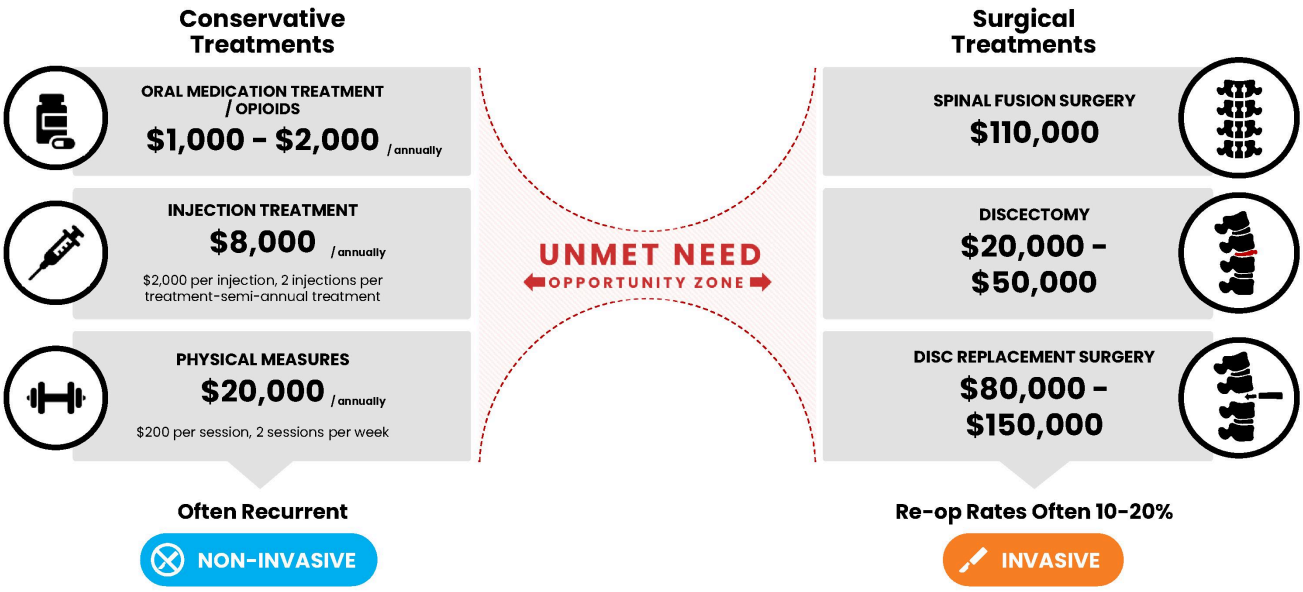
2.5 M Invasive Surgical Procedures per year **$40 billion** in surgeries



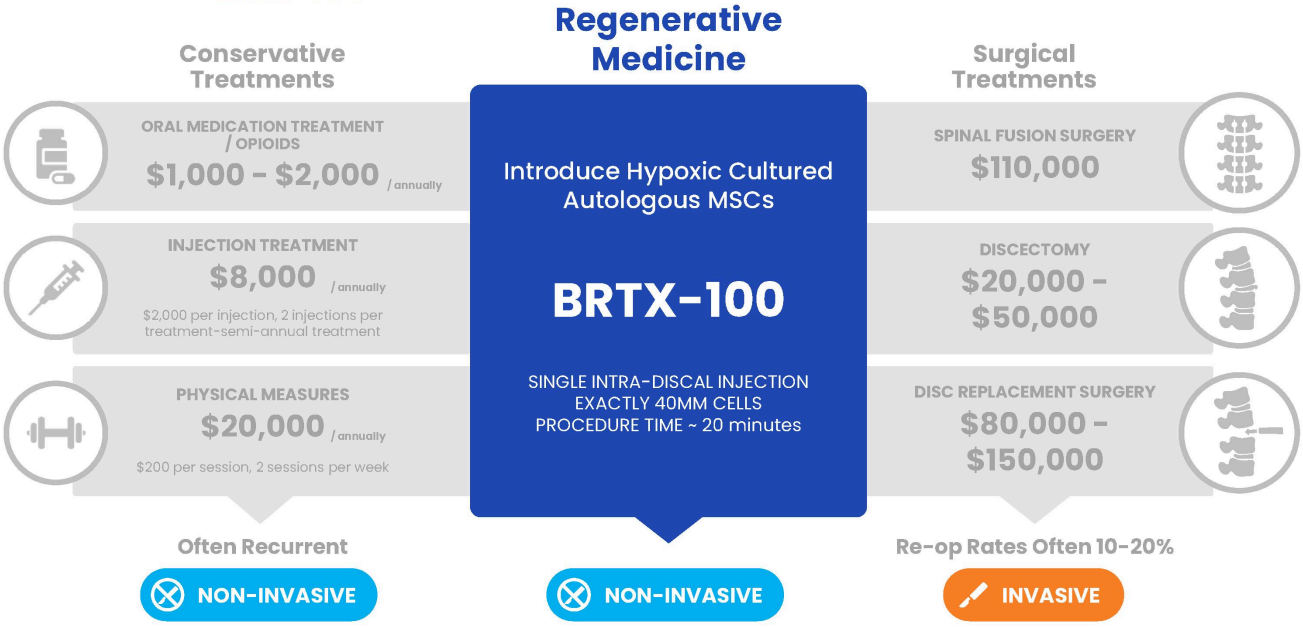
* Trinity Partners Report, "Degenerative Disc Disease US Market Assessment (Phase I)," Feb. 2016, Slide 9 of report.

biorestorative therapies

The Problem: Clinical & Economic



Our Solution: BTRX-100



biorestorative therapies

BRTX-100: Clinical Snapshot

Lead investigational therapeutic product

Autologous (patient's own) cell-based biologic

Hypoxic (low oxygen) cultured, bone marrow-derived

Single intradiscal injection – anticipated 30 minute in-office procedure

Prior human data provides insight into the potential safety and efficacy of BRTX-100

Ongoing FDA authorized Phase 2 clinical trial

Large growing market with few comparable autologous therapies

1) Trinity Partners Report, "Degenerative Disc Disease US Market Assessment (Phase I)," Feb. 2016, Slide 9 of report.

BRTX-100: Key Differentiating Factors

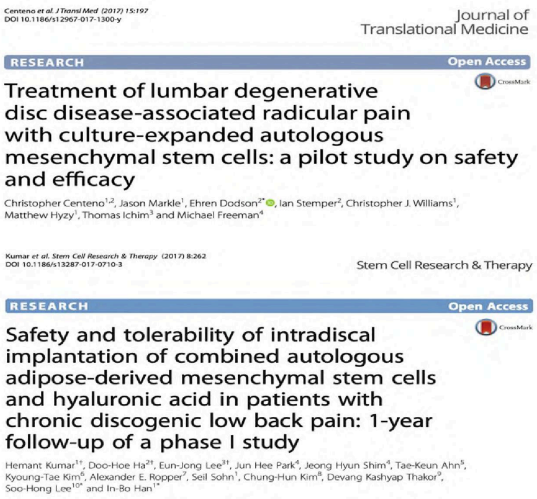
	mesoblast	biorestorative therapies
SOURCE	**Allogeneic** uses human derived stem cells (not from patient) - 6 million	**Autologous** uses patients own stem cells - 40 million
CULTURING	**Normoxic cultured** with normal oxygen environment (~20%)	**Hypoxic cultured** in low oxygen environment (5%)
CARRIER	**Hyaluronic Acid Carrier**	**Autologous Platelet Lysate Carrier & Adjuvant**
MANUFACTURING	**Animal Products Used**	**100% Animal-Free**

BRTX-100 Advantages

- Autologous cells means low to no risk of rejection, greater safety profile (introduction of viral/genetic), potentially streamlined regulatory path
- Hypoxic culturing creates increased cell proliferation, greater plasticity, increased paracrine effect and increased cell survival after application
- Autologous platelet lysate provides growth factors that interact with the cells, allowing for better cell survival
- Low to no risk of safety concerns related to immunological and zoonotic (animal to people) transmission
- Strong runway for value creation with successful clinical results

BRTX-100: Positive Human Data

Human data from studies of therapies similar to BRTX-100 show reduced pain, increased function, and an absence of significant safety issues with a durable response



Centeno et al. J Transl Med (2017) 15:197
DOI 10.1186/s12967-017-1300-y

Journal of Translational Medicine

RESEARCH — Open Access

Treatment of lumbar degenerative disc disease-associated radicular pain with culture-expanded autologous mesenchymal stem cells: a pilot study on safety and efficacy

Christopher Centeno[1,2], Jason Markle[1], Ehren Dodson[2*], Ian Stemper[2], Christopher J. Williams[1], Matthew Hyzy[1], Thomas Ichim[3] and Michael Freeman[4]

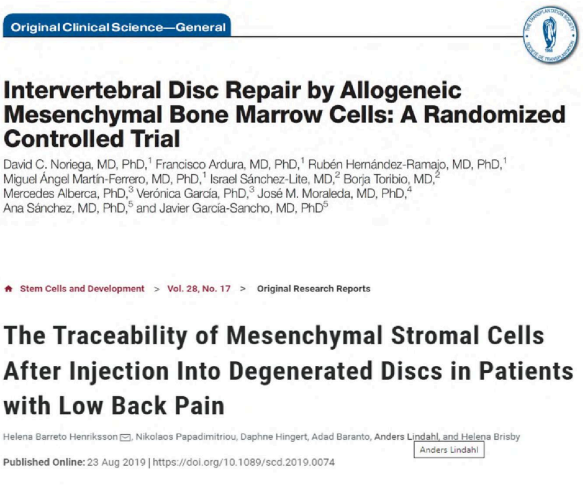
Kumar et al. Stem Cell Research & Therapy (2017) 8:262
DOI 10.1186/s13287-017-0710-3

Stem Cell Research & Therapy

RESEARCH — Open Access

Safety and tolerability of intradiscal implantation of combined autologous adipose-derived mesenchymal stem cells and hyaluronic acid in patients with chronic discogenic low back pain: 1-year follow-up of a phase I study

Hemant Kumar[1†], Doo-Hoe Ha[2†], Eun-Jong Lee[3†], Jun Hee Park[4], Jeong Hyun Shim[4], Tae-Keun Ahn[5], Kyoung-Tae Kim[6], Alexander E. Ropper[7], Seil Sohn[1], Chung-Hun Kim[8], Devang Kashyap Thakor[9], Soo-Hong Lee[10*] and In-Bo Han[1*]



Original Clinical Science—General

Intervertebral Disc Repair by Allogeneic Mesenchymal Bone Marrow Cells: A Randomized Controlled Trial

David C. Noriega, MD, PhD,[1] Francisco Ardura, MD, PhD,[1] Rubén Hernández-Ramajo, MD, PhD,[1] Miguel Ángel Martín-Ferrero, MD, PhD,[1] Israel Sánchez-Lite, MD,[2] Borja Toribio, MD,[2] Mercedes Alberca, PhD,[3] Verónica García, PhD,[3] José M. Moraleda, MD, PhD,[4] Ana Sánchez, MD, PhD,[5] and Javier García-Sancho, MD, PhD[5]

Stem Cells and Development > Vol. 28, No. 17 > Original Research Reports

The Traceability of Mesenchymal Stromal Cells After Injection Into Degenerated Discs in Patients with Low Back Pain

Helena Barreto Henriksson, Nikolaos Papadimitriou, Daphne Hingert, Adad Baranto, Anders Lindahl, and Helena Brisby

Anders Lindahl

Published Online: 23 Aug 2019 | https://doi.org/10.1089/scd.2019.0074

biorestorative therapies

BRTX-100 in cLDD: Phase 2 Trial Design

FDA Cleared IND 17275:
Phase 2 Randomized, Controlled Study Design in Patients with cLDD

Design

- Study includes 99 subjects (2:1 product to placebo)
- 40.000.000 cells/dose
- Included subjects will have only one symptomatic diseased disc

Primary Efficacy Endpoint

12 m, F/U at 24 m

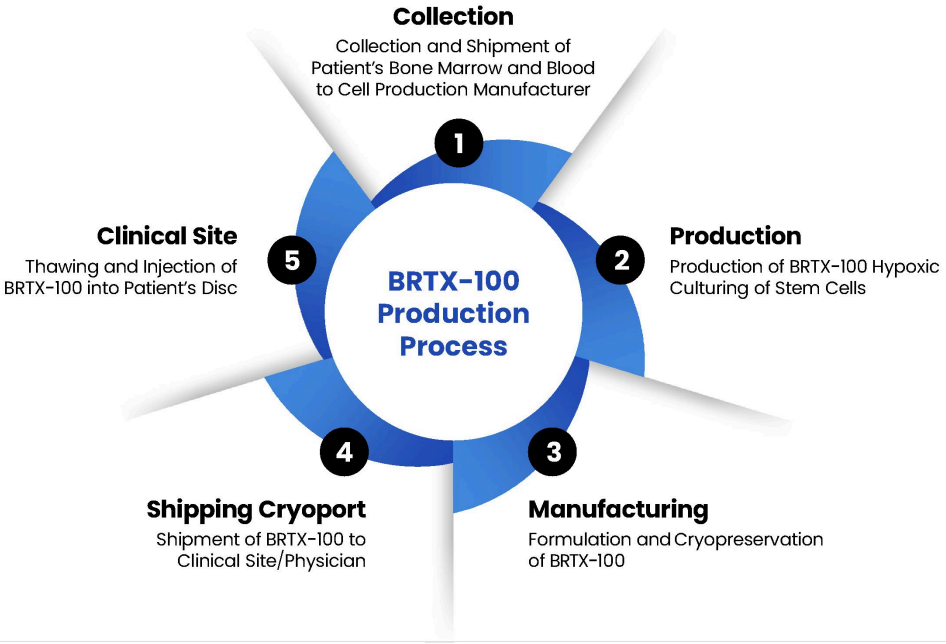
Improvement in function:
at least 30% increase in function based on Oswestry Disability Index questionnaires (ODI)

Reduction of pain:
at least 30% decreased in pain as measured using a Visual Analogue Scale (VAS)

Patient Population

- Subjects must have current diagnosis of cLDD, typical pain with degeneration of a single disc confirmed by history, exam, radiography, or other acceptable means
- Subjects will have exhausted previous conservative non-operative therapies

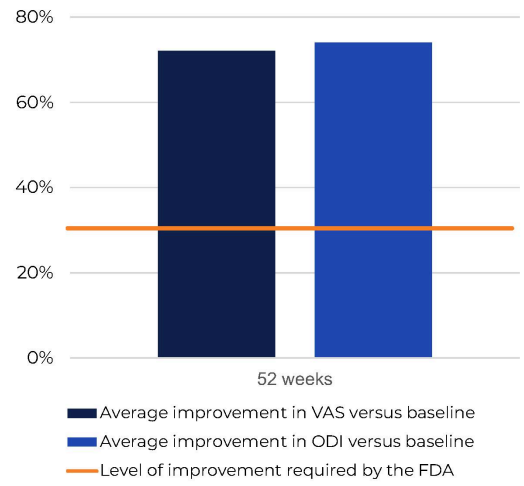
BRTX-100: Logistical /Clinical Process



BRTX-100: Cleared DSMB June 2023

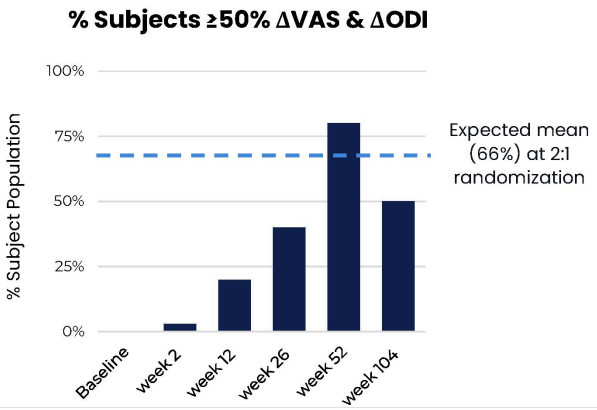
- Unanimous approval by the DSMB to continue trial without changes
- BRTX-100 is safe and well tolerated
- All 4 subjects successfully dosed with either 40 mil hMSCs or placebo
- First time 40 million cells injected in a human subject
- 3:1 randomization
- No Significant Adverse Events
- VAS, ODI, SF-12, RMDQ, and FRI scores to measure pain and function were collected
- Opportunity to leverage this data and clinical package

Compelling Preliminary Phase 2 Clinical Data (36 Subjects): Meaningful Signals



Presented at ISSCR 2025

FDA is requiring at least a greater than 30% improvement in function (ODI) and a greater than 30% reduction in pain (VAS) in determining whether the clinical trial will be allowed to proceed and ultimately gain Biologics License Application approval



biorestorative therapies

ThermoStem Program: Allogeneic Cell-based Therapy

Target Conditions: Obesity, Type 2 diabetes, and Metabolic disorders

Cell Type: Brown Fat

- Has been shown to regulate metabolic homeostasis in the body

Components of Library:

- Human Brown Adipose Tissue (BAT)
- White Adipose Tissue (WAT)
- Brown Adipose-derived Stem Cells (BADSC)

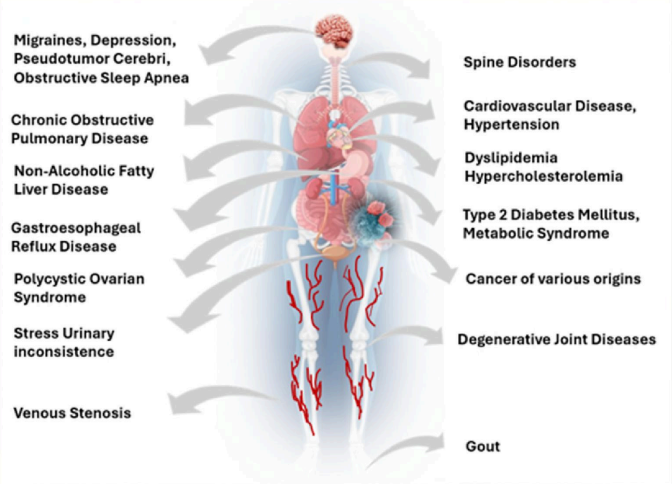
Initial Proof of Concept:

✓ Completed in small animal model

Patent Portfolio: Related BAT patent portfolio, including issued patents in the U.S., Australia and Japan

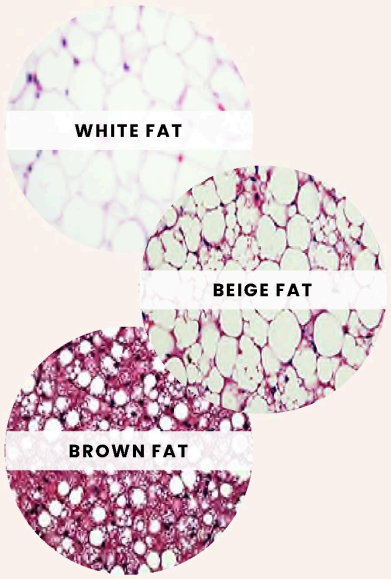
Platform Program:

For the development of cell & small molecule therapies



biorestorative therapies

Metabolic Program Highlights



- First human stem cell derived BAT transfer
- Creation of first human 3D engineered artificial brown adipose tissue construct (aBAT)
- Successful delivery of 3D aBAT construct in mouse model
- Transplantation of aBAT lowered blood glucose levels
- Transplantation of aBAT decreased weight in obese mice
- Published initial proof of concept completed

Metabolic Program Clinical Pathway

File DMF with FDA

Expect filing a Drug Master File ("DMF") with the FDA to facilitate licensing opportunities around ThermoStem.

Pre-IND Meeting with FDA

Scheduling Pre-IND meeting with FDA to discuss first-in-man fast-track regulatory pathways.

Initiate Phase 1/2 Clinical Trial

Upon FDA approval commence Phase 1/2 clinical trial.

Our Opportunities are Well-Protected

PROGRAM	brtdiscx	ThermoStem
INDICATION	Disc / Spine	Metabolic
PATENT TITLES	• Methods and Compositions to facilitate repair of avascular tissue • Surgical Methods and Compositions to facilitate repair of avascular tissue • Therapeutic Delivery Device	• Brown Fat Compositions and Methods • Human Brown Adipose Derived Stem Cells and Uses • Non-naturally occurring three-dimensional (3D) Brown Adipose-Derived Stem Cell aggregates and methods of generating and using the same
STATUS	4	26
NO. OF APPLICATIONS	2 Issued 2 Pending	23 Issued 3 Pending

Scientific Advisory Board

Wayne Marasco, MD, PDt
Chairman of SAB

- Principal Faculty Member of Harvard Stem Cell Institute
- Professor in the Department of Cancer Immunology & AIDS at the Dana-Farber Cancer Institute
- Professor of Medicine at Harvard Medical School

Jason Lipetz, MD
Chairman of SAB Sub Committee
Disc Advisory Board

- Chief of Spine Medicine for the Northwell Health Spine Center
- Founder of Long Island Spine Rehabilitation Medicine

Harvinder Sandhu, MD
Member Disc Advisory Board

- Orthopedic Spine Surgeon at the Hospital for Special Surgery
- Specializes in minimally invasive spine surgery, endoscopic spine surgery, microsurgery, computer-assisted surgery, and the study and use of spinal biologics

Wayne Olan, MD Clinical
Director of Regenerative Disc / Spine Program

- Board-certified Interventional Neuroradiologist
- Director of Endovascular and Minimally Invasive Neurosurgery at the George Washington University Medical Center

Christopher Plastaras, MD
Member Disc Advisory Board

- MossRehabs' Clinical Director of Musculoskeletal Spine & Sports Rehabilitation Medicine

Joy Cavagnaro, PhD
Member

- President and Founder of Access BIO, L.C.
- Previously positions with the FDA Center for Biologics Evaluation and Research (CBER), for a decade

Financial Summary

CURRENT CAPITALIZATION	SHARES
Common Shares Outstanding	8.0 Million*
Common Shares Including Abeyance	9.1 Million**
Preferred Series B Shares Outstanding	Convertible to 1.4 Million Common**
Cash	$ 7.4 Million**
Debt	$0

* As of 08/11/2025
** As of 06/30/2025

Recent Accomplishments

- ✓ Announced FDA clearance of important BRTX-100 clinical study protocol amendment (replaces saline injection with sham injection in control arm)
- ✓ Reported positive preliminary Phase 2 BRTX-100 study data (36 patient) in cLDD
- ✓ Received FDA Fast Track Designation for BRTX-100 in cLDD
- ✓ FDA cleared IND for Phase 2 BRTX-100 tiral in cCDP
- ✓ Developed novel exosome-based biologic for obesity
- ✓ Expanded ThermoStem patents in U.S., Europe, Israel, and Japan
- ✓ Initiated discussions with an undisclosed commercial stage regenerative medicine company on potential ThermoStem IP licensing
- ✓ Announced commercial biocosmeceutical agreement with Cartessa
- ✓ Received expanded tissue license from New York State Dept. of Health

In Conclusion

- cGMP ISO-7 Certified Clean room
- Disruptive Platform Technologies in Cellular Therapy
- Strong Preliminary Data Indicative of Positive Trial Outcomes
- Active Phase 2 Trial in Spine
- Addressing Multi-Billion Dollar Markets with Unmet Needs
- Opportunity for Key Strategic Partnerships in Cosmetic Space
- Strong Intellectual Property Protection
- Experienced Management Team & Scientific Advisory Board

biorestorative
therapies

40 Marcus Drive, Suite 1
Melville, NY 11747
(631) 760-8100
biorestorative.com