Presentation Material for Business Plan and Growth Potential

StemRIM Inc.（Stock code：TSE4599）

October 30,2026

StemRIM

PURPOSE

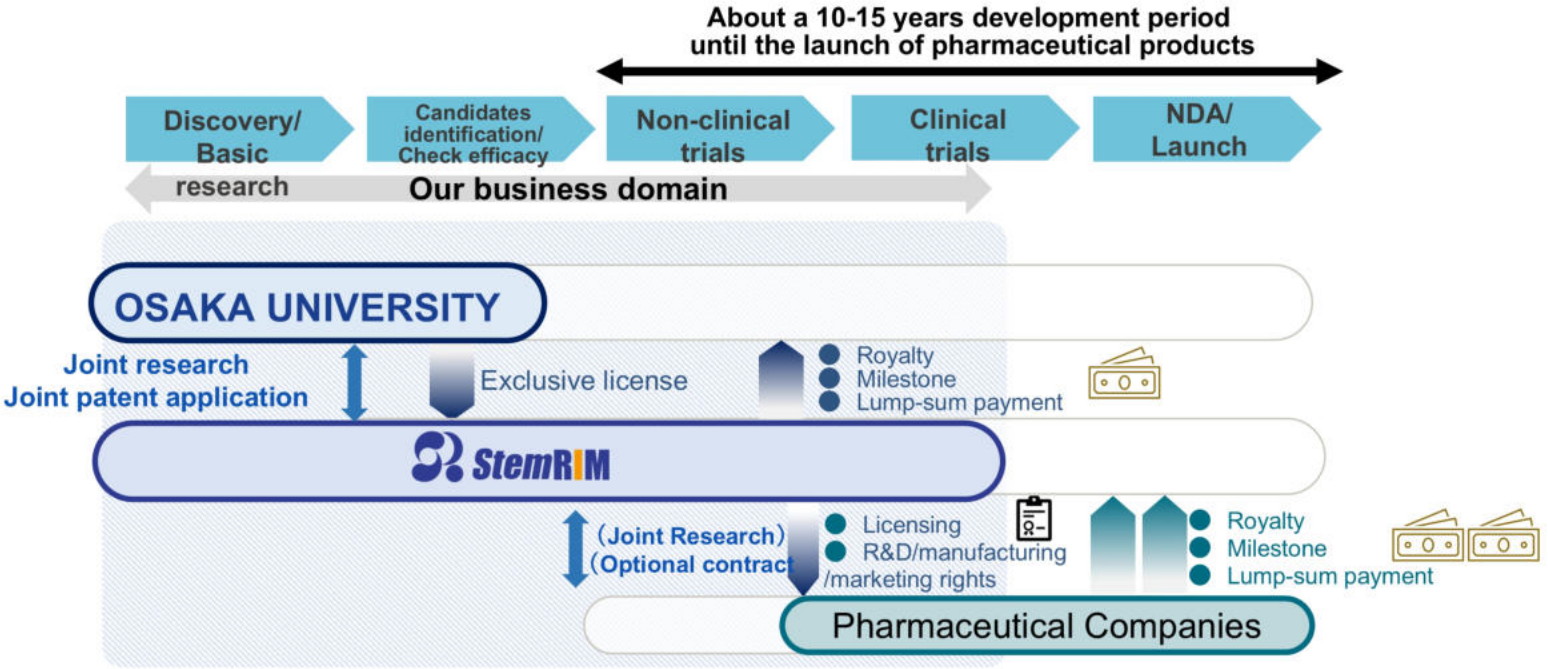
Empower Life
Regeneration.

REGENERATION

1. Business Model and R&D Structure

Business Model

A business model that generates income by licensing out product development, manufacturing, and marketing rights to pharmaceutical companies in Japan and overseas.



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Main Contracts with Pharmaceutical Companies

Contract Name (Date of Agreement)	Company	Contract Details (Excerpt)	Total Contract Amount	Received Contract Amount
Implementation License Agreement(November 2014)	Shionogi & Co. Ltd.	A license is granted for the exclusive development, manufacturing, use, or sale of prior compounds and products for pharmaceutical applications worldwide, based on patents related to the pharmaceutical use of Redasemtide (HMGB1 peptide) or compounds containing it, as well as their methods of manufacture or formulation. As compensation for the license, Stemrim will receive upfront payments, milestone revenues, and royalty income.	Not disclosed*	4,046 million yen
Contract for the accelerated clinical development of the "Regeneration-Inducing Medicine™"candidate Redasemtide for multiple diseases(June 2020)	Shionogi & Co. Ltd.	Utilizing non-clinical research evidence related to the Redasemtide (HMGB1 peptide), physician-led clinical trials will be conducted targeting cardiomyopathy, osteoarthritis of the knee, and chronic liver disease. As compensation for the license, Stemrim will receive a one-time payment based on the achievement of specified conditions.	3,100 million yen	3,100 million yen

*The total contract amount related to this agreement is not disclosed due to confidentiality obligations.

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2026年7月期 決算概要

・For FY 2026, there were no recognition of milestone revenues related to research progress or upfront payments from contracts. As a result, **operating revenue was none.** Since we are a drug discovery bio-venture, we have an unstable revenue structure considering our business model.

・As of the end of FY 2026, we hold **5,377 million yen** in cash and deposits.
The estimated annual expenditure for the FY 2027 is between 1.5 billion yen and 2 billion yen (cash outflows related to R&D: 1.3 billion yen to 1.7 billion yen, cash outflows for general administrative expenses: 230 million to 310 million yen). At present, **we have secured sufficient funds to sustain stable R&D activities until 2028.**

(Millions of yen)

	FY 2022.7	FY 2023.7	FY 2024.7	FY 2025.7	FY 2026.7	Function (FY on FY)
Operating revenue	22	2,350	—	—	—	—
R&D expenses	1,421	1,567	1,453	1,394	1,499	+104
Total operating expenses	2,003	2,207	2,076	1,971	1,998	+27
Operating Income (loss)	(1,980)	142	(2,076)	(1,971)	△1,998	△27
Ordinary Income (loss)	(1,972)	145	(2,077)	(1,970)	△1,879	+90
Net Income (loss)	(1,948)	168	(2,022)	(1,929)	△1,851	+77
Cash and deposit	8,880	10,217	8,410	6,994	5,377	

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Scientific founder : Prof. Katsuto Tamai

Katsuto Tamai, our current Representative Director , Chairman and Chief Scientific Officer (CSO), was inspired to explore the potential of skin regeneration using bone marrow-derived stem cells after encountering a patient with the rare and debilitating disease, Dystrophic Epidermolysis Bullosa. He elucidated the Regeneration-Inducing Mechanism of the HMGB1 protein and subsequently developed “Redasemtide,” a peptide that excludes the inflammatory domain of HMGB1. Through clinical trials, Redasemtide demonstrated both efficacy and safety, leading to the establishment of a new therapeutic concept known as “Regeneration-Inducing Medicine™.”

Academic Society

The Japanese Dermatological Association
The Japanese Society for Matrix Biology and Medicine
The Japanese Cancer Association
The Japanese Society for Regenerative Medicine
Japan Society of Gene and Cell Therapy
The Japanese Society of Inflammation and Regeneration
Japan Organization of Clinical Dermatologists
Society for Skin Structure Research
Society for Investigative Dermatology
American Society of Gene & Cell Therapy

Career

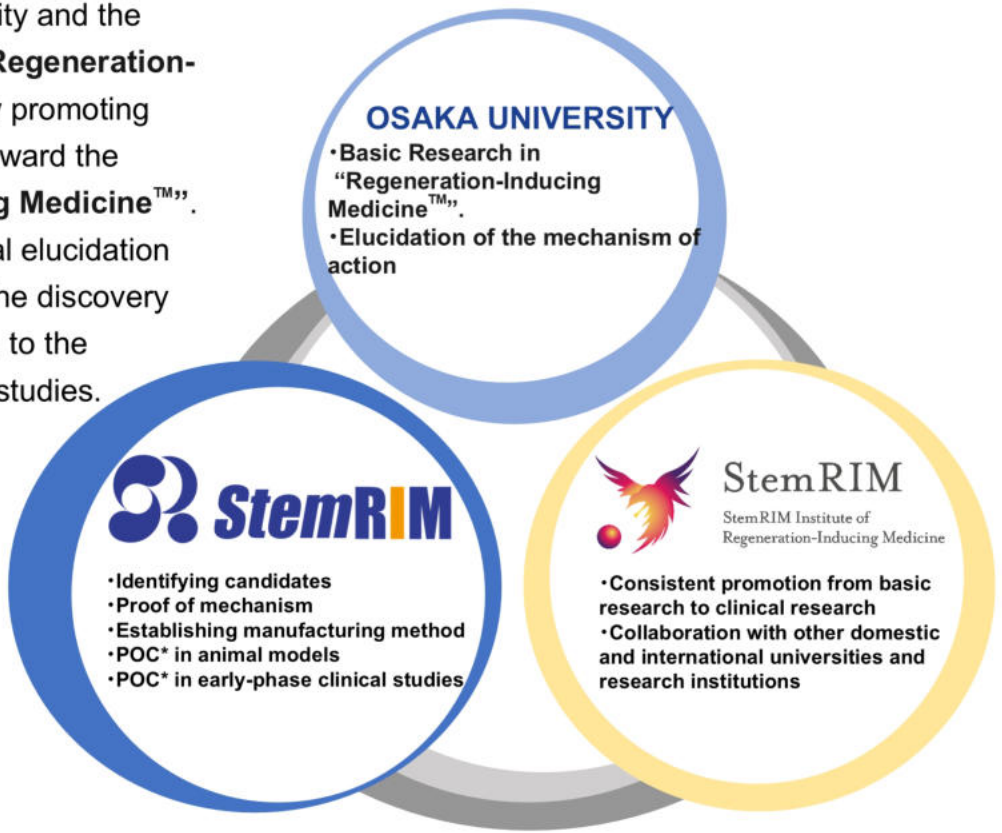
April 1990
Assistant, Department of Dermatology, Hirosaki University Hospital, Faculty of Medicine
October 1990
Postdoctoral Fellow, Jefferson Medical College, United States
December 1992
Assistant, Department of Dermatology, Hirosaki University, Faculty of Medicine
February 1995
Lecturer, Department of Dermatology, Hirosaki University Hospital, Faculty of Medicine
August 1998
Associate Professor, Department of Dermatology, Hirosaki University, Faculty of Medicine
April 2002
Associate Professor, Department of Gene Therapy, Graduate School of Medicine, Osaka University
May 2003
Associate Professor (tenure-track), Department of Gene Therapy, Graduate School of Medicine, Osaka University
February 2007
Director, Genomix Co., Ltd. (now StemRIM Inc.)
October 2009
Professor, Endowed Chair of Regenerative Medicine, Graduate School of Medicine, Osaka University
October 2022
Director and Chief Scientific Officer (CSO), StemRIM Inc.
October 2023
Visiting Professor, Graduate School of Medicine, Osaka University
September 2026
Representative Director , Chairman and Chief Scientific Officer (CSO), StemRIM Inc.



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Research and Development system

At Stemrim, in collaboration with Osaka University and the Institute for Advanced Co-Creation Studies in **"Regeneration-Inducing Medicine™"**, we are comprehensively promoting the entire research and development process toward the practical application of **"Regeneration-Inducing Medicine™"**. This process spans from the molecular biological elucidation of stem cell mobilization mechanisms, through the discovery of next-generation regenerative induction drugs, to the verification of efficacy and safety in non-clinical studies.



*POC: Proof of Concept

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2. Mechanism of Action of "Regeneration-Inducing Medicine™"

Mechanism of Action of "Regeneration-Inducing Medicine™"

Bone marrow mesenchymal stem cells mobilized into the peripheral blood stream induce the tissue regeneration.

Administration of "Regeneration-Inducing Medicine™"

↓

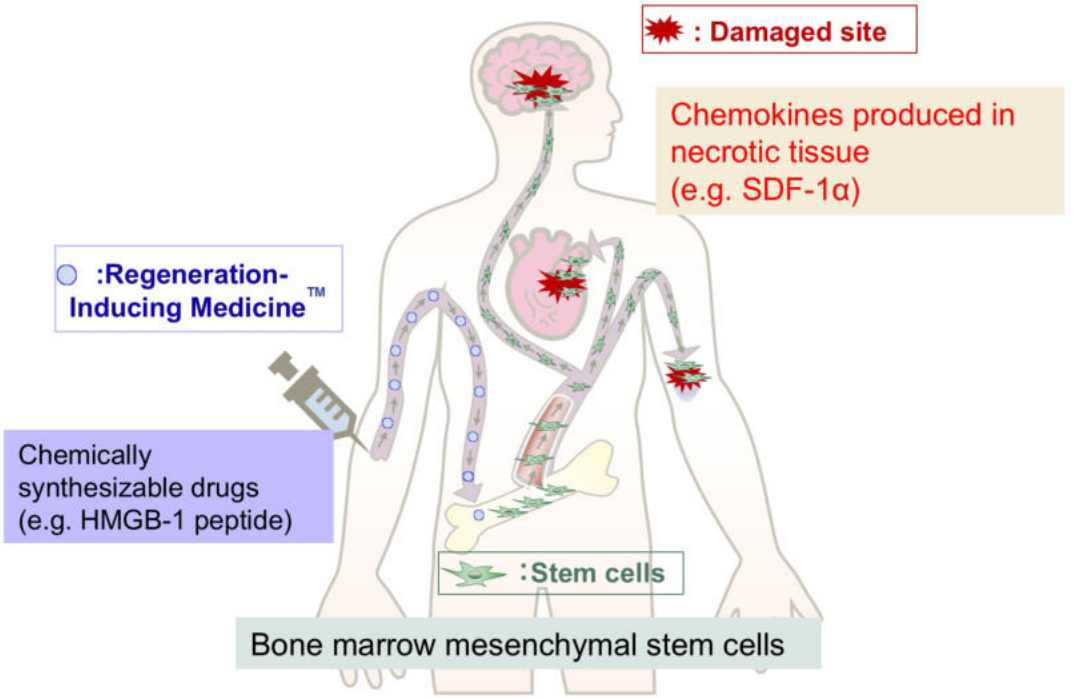
Mobilizing Stem cells from bone marrow into the blood stream

↓

Stem cells accumulation in the damaged tissues

↓

Functional tissue regeneration



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Discovery of in-vivo mechanism inducing tissue regeneration

- **Differences between normal skin and epidermolysis bullosa skin**

In normal skin (Figure A), type 7 collagen functions like an adhesive, bonding the epidermis and dermis, the superficial layers of skin. In epidermolysis bullosa congenita (Figure B), the epidermis and dermis are easily detached with the slightest irritation due to abnormal type 7 collagen.Since epidermal stem cells, which are responsible for supplying epidermal cells, reside in the epidermis, the epidermal stem cells are lost from the skin of patients with epidermolysis bullosa, and the epidermis loses its regenerative capacity.

Fig. A

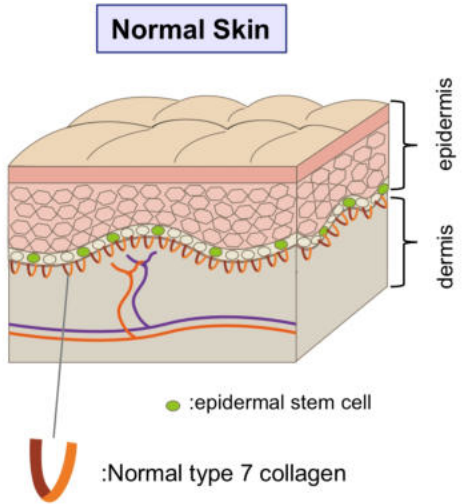
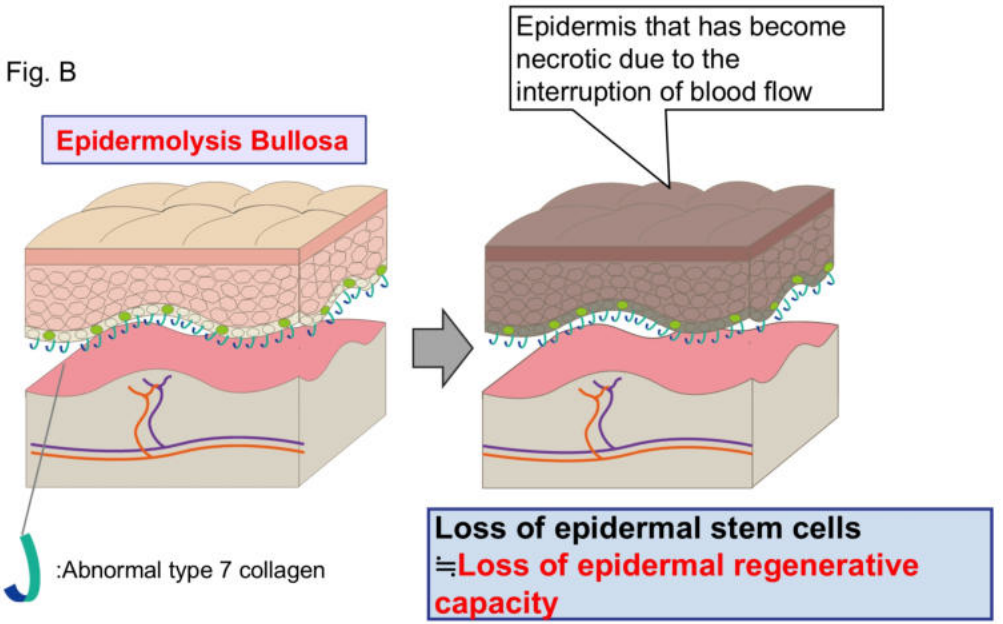


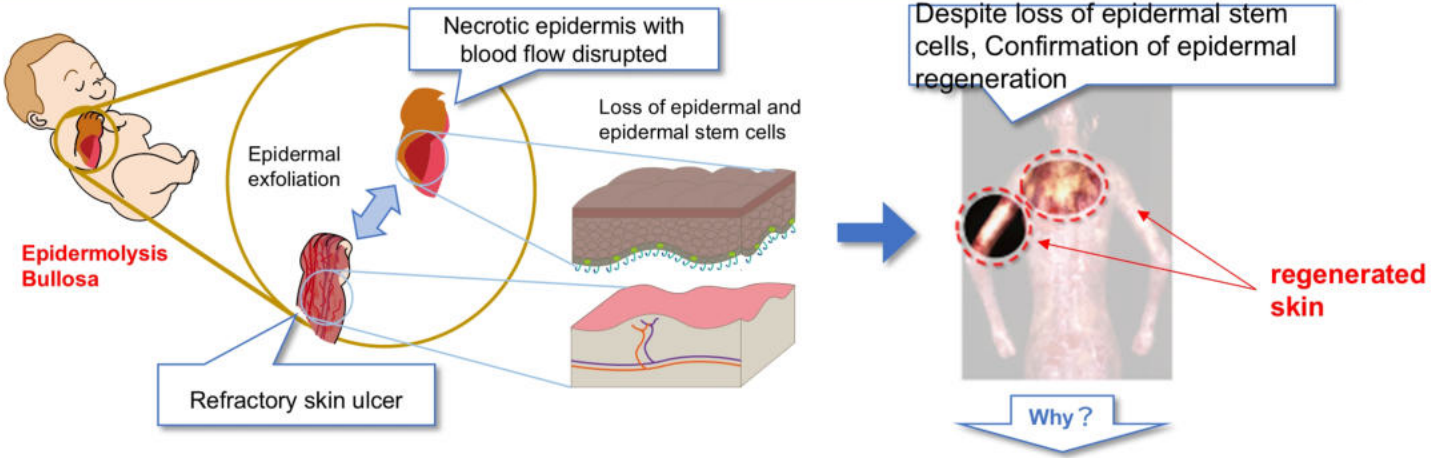
Fig. B

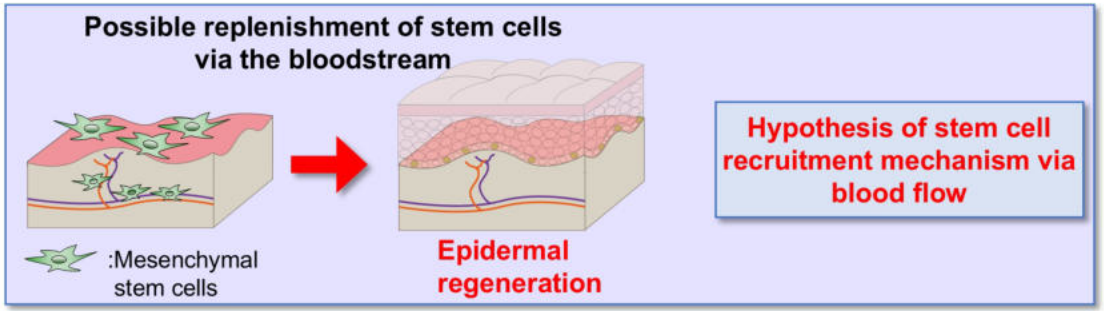


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Discovery of in-vivo mechanism inducing tissue regeneration

The beginning of the research and development on "Regeneration-Inducing Medicine™" : Hypothesis of stem cell recruitment mechanism from bone marrow to damaged skin.



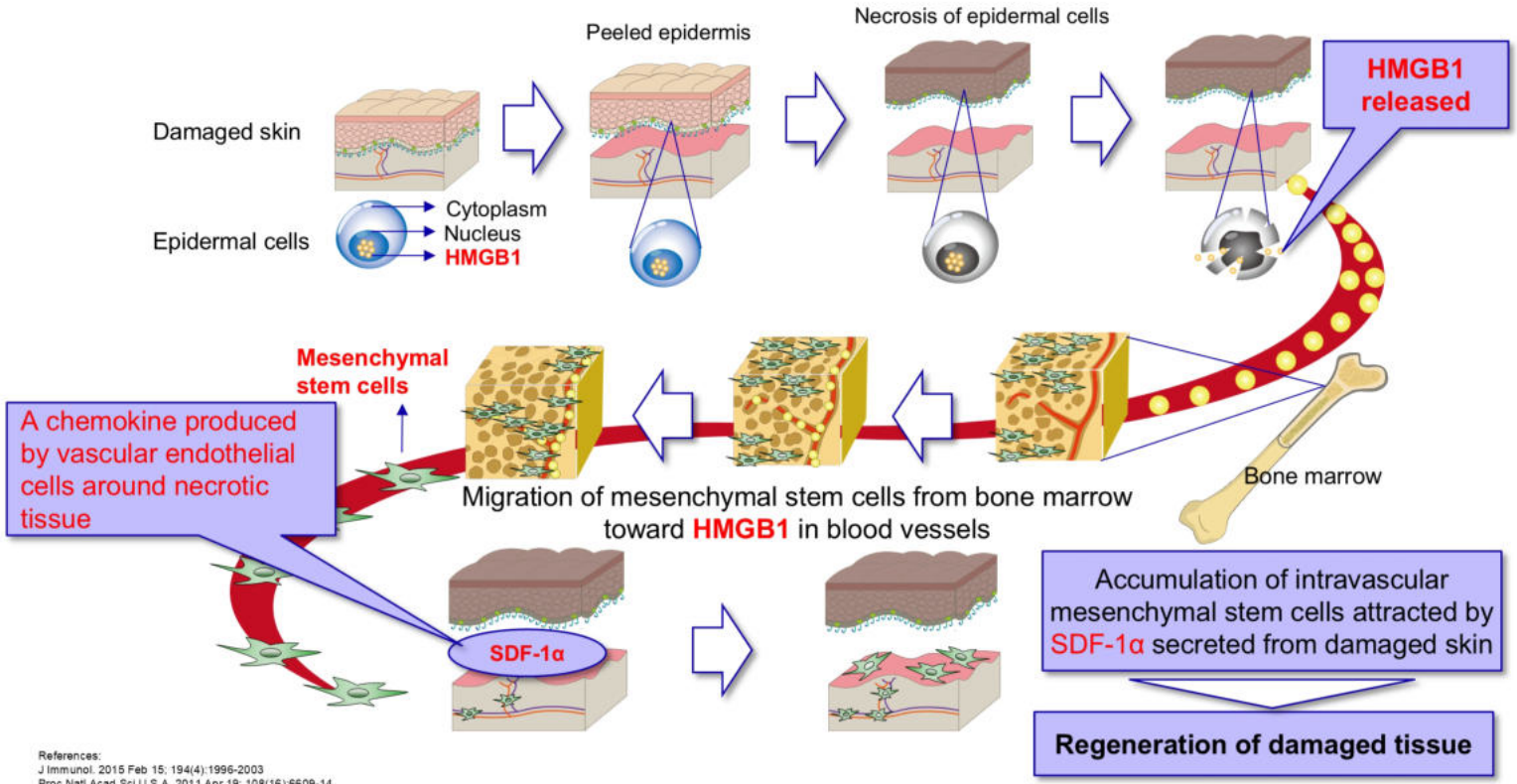


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"Igaku-no-ayumi" Vol.265 No.5 463-468; 2018
Skin Diseases :41(1); 7-12,2019
Photo courtesy of Osaka University

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Discovery of in-vivo mechanism inducing tissue regeneration

Discovery of crosstalk mechanism between damaged skin and bone marrow mesenchymal stem cells via necrotic tissue-derived factor

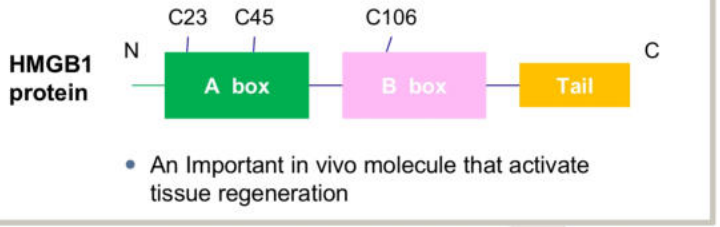


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Proc Natl Acad Sci U S A. 2011 Apr 19; 108(16):6609-14.

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HMGB1 peptide drugs with improved safety

Designing highly safe, chemically synthesized peptide drug from A-Box domain of HMGB1 protein

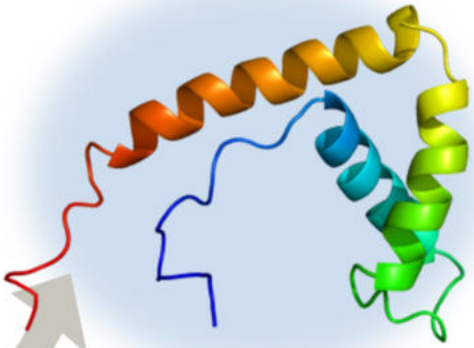


- An Important in vivo molecule that activate tissue regeneration

Katsuto Tamai
Representative Director , Chairman, CSO



Identifying the function of protein domains



A box	Bone marrow mesenchymal stem cell activating domain, named **"KOI2-domain"**
B box	Innate immune response-activating domain that induces inflammation

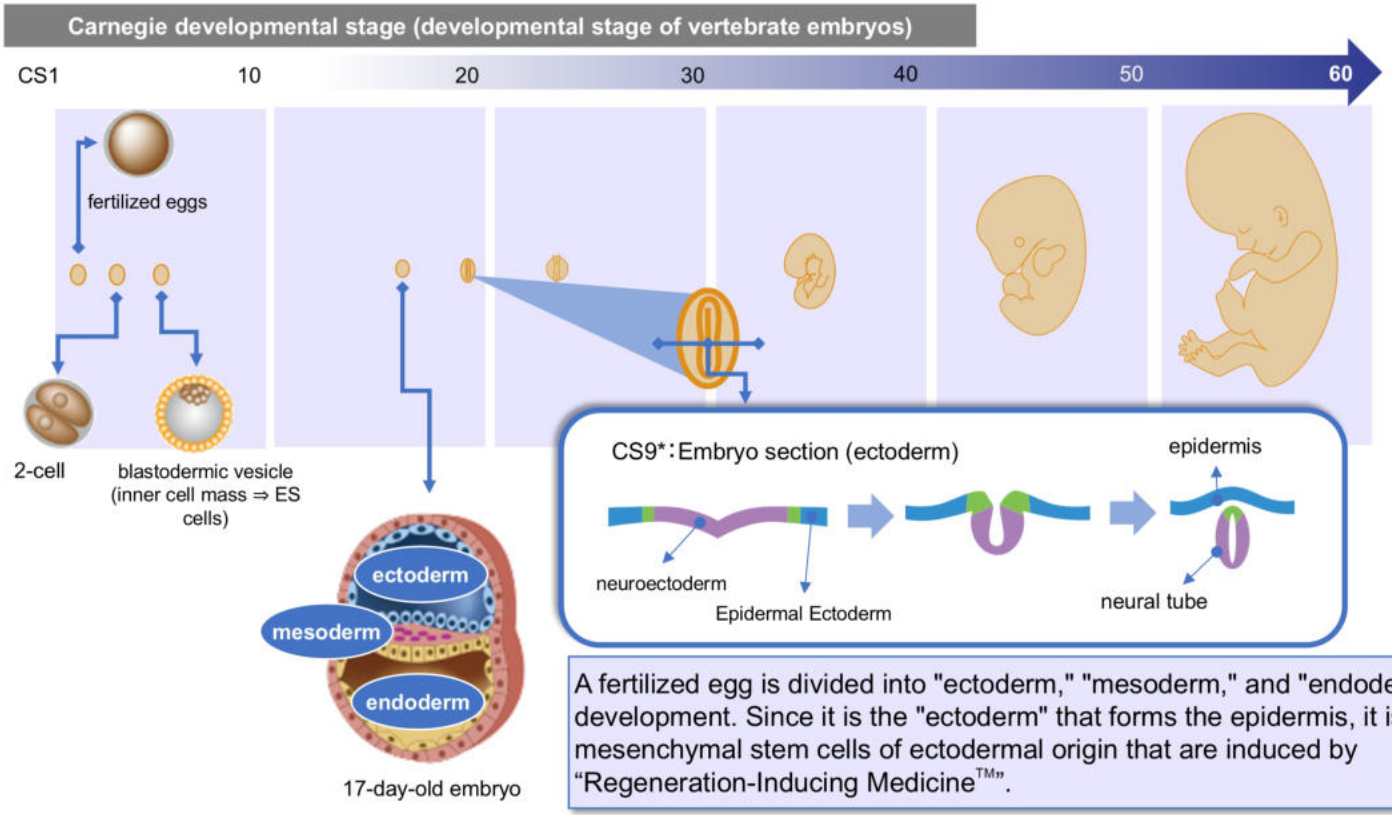
References:
J Intern Med. 2004 Mar ; 255(3):351-66.

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3. Advantages of “Regeneration-Inducing Medicine™”

Advantages of “Regeneration-Inducing Medicine™”

Epidermis formation during human development

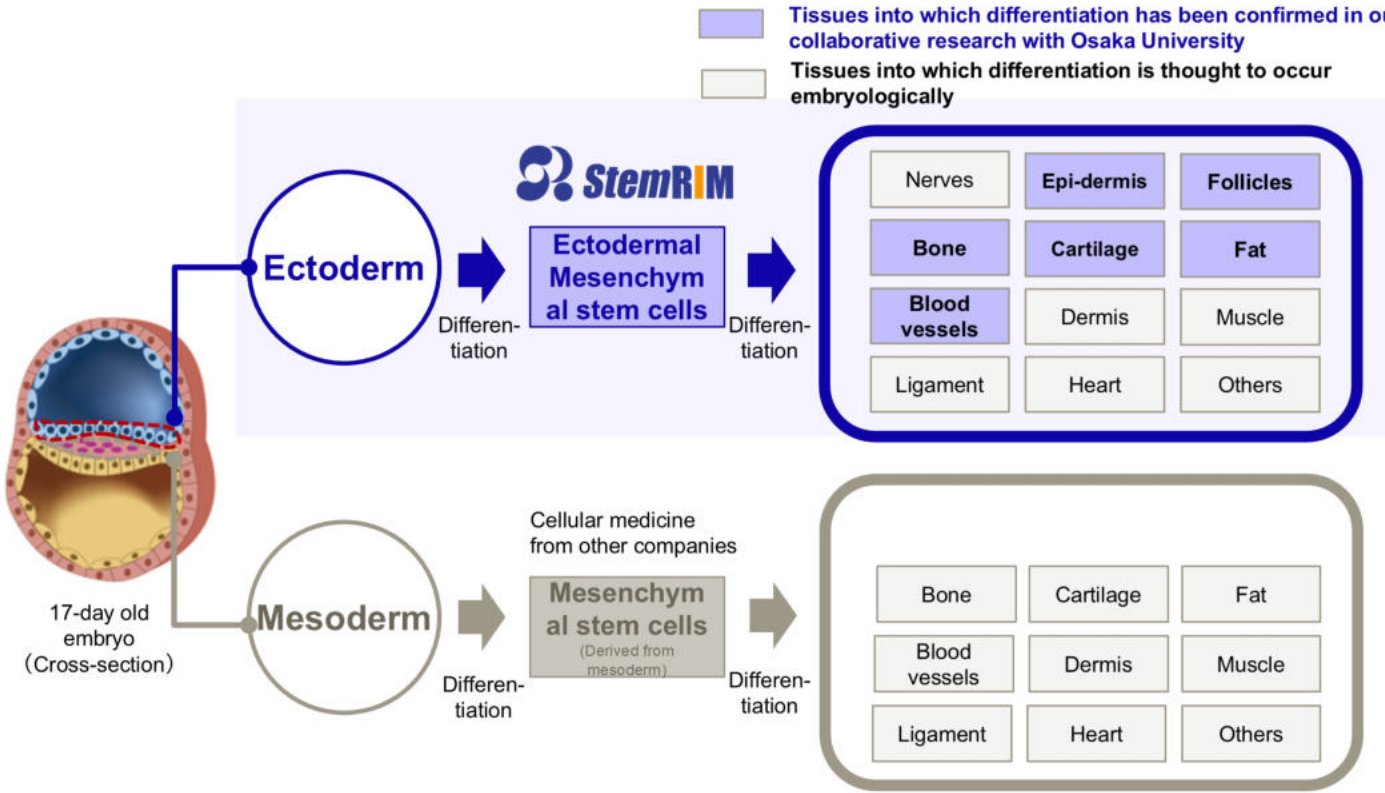


A fertilized egg is divided into "ectoderm," "mesoderm," and "endoderm" during development. Since it is the "ectoderm" that forms the epidermis, it is the mesenchymal stem cells of ectodermal origin that are induced by “Regeneration-Inducing Medicine™”.

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Advantages of "Regeneration-Inducing Medicine™"

Ectodermal mesenchymal stem cells have high pluripotency and differentiation ability to various tissues.

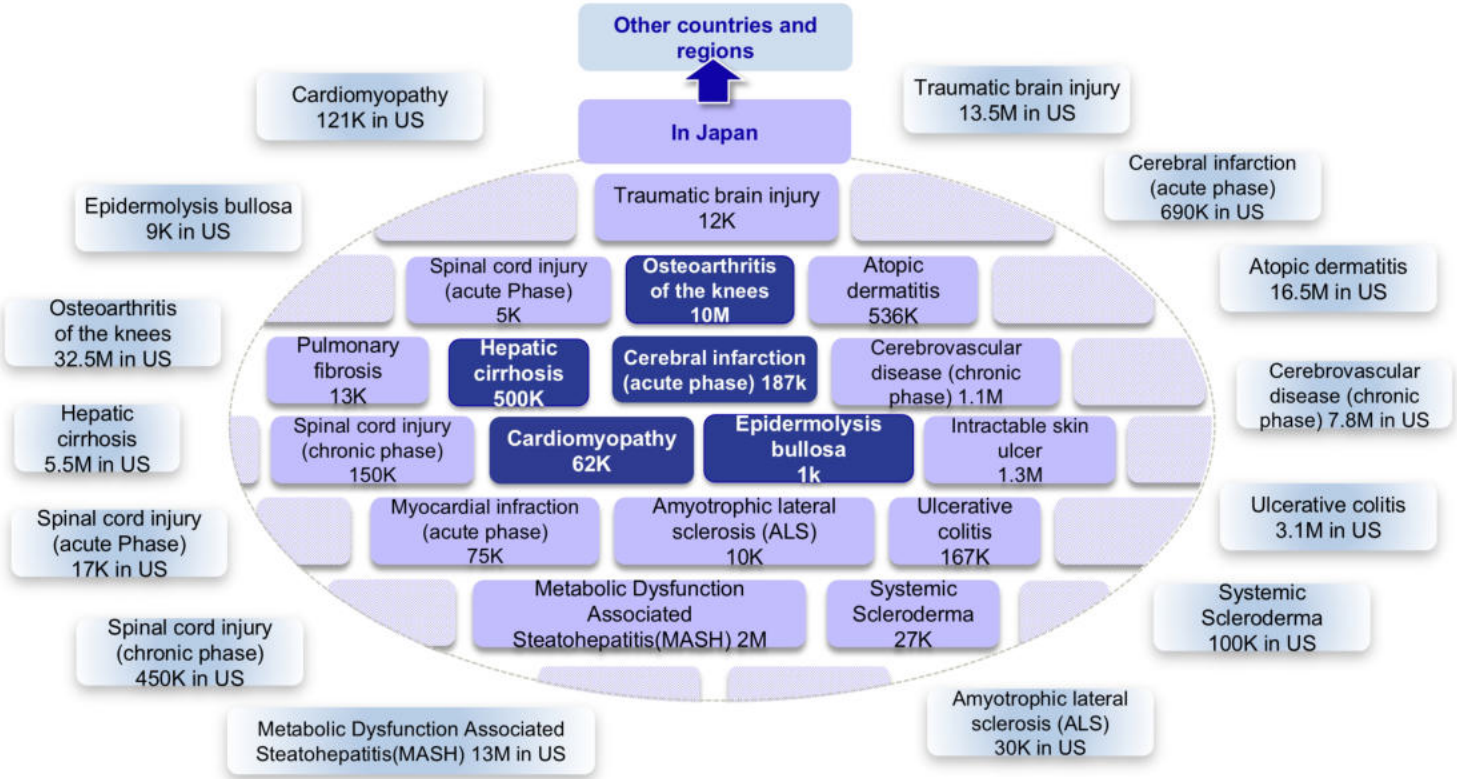


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Expanding Indications and Markets (Number of patients)

Targeting all areas where mesenchymal stem cell therapy can be effective

: Clinical trial on going



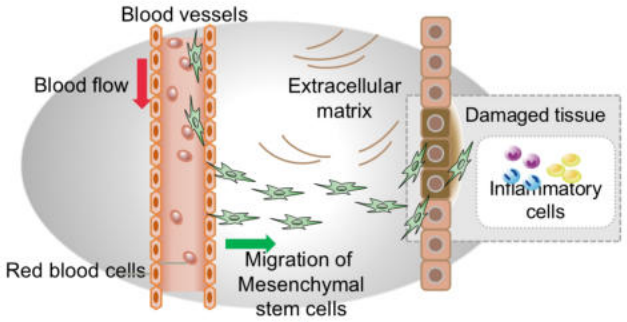
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Functions of mesenchymal stem cells

In-vivo mesenchymal stem cells have 5 distinctive capabilities

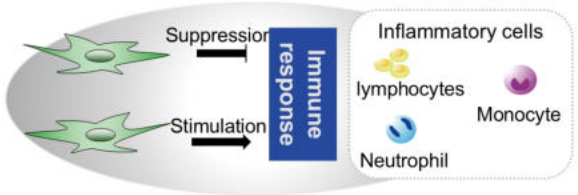
1. Cell migration ability

Mesenchymal stem cells migrate to damaged tissue via the bloodstream



2. Immunomodulatory ability

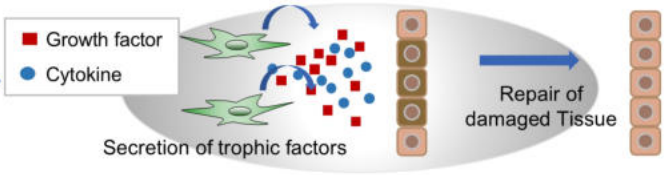
Modulates immune response and inhibits the spread of tissue damage caused by excessive inflammation



* MMP：Matrix metalloproteases

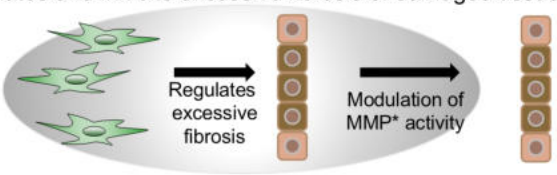
3. Trophic factor secretion ability

Promotes cell proliferation and tissue repair by secreting growth factors and cytokines to cells in damaged tissue

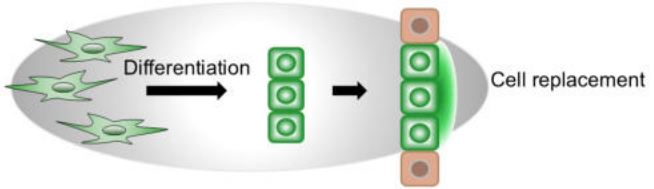


4. Fibrosis regulation ability

Regulates and inhibits excessive fibrosis of damaged tissue



5. Tissue regeneration ability

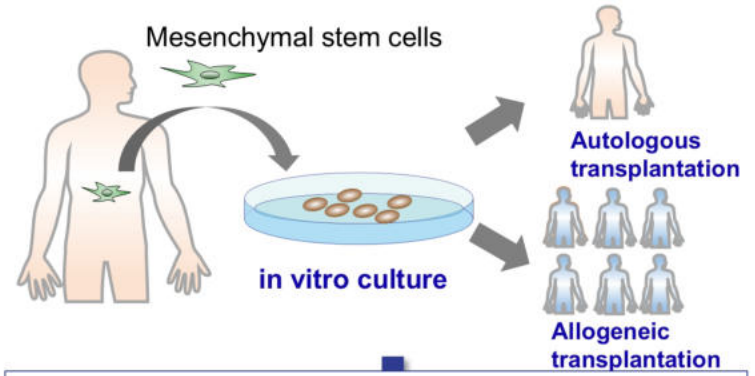


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In vitro culture reduces the functions of MSCs

"Regeneration-Inducing Medicine™" can avoid functional degradation of mesenchymal stem cells due to in vitro culture

Manufacturing process of conventional cellular medicine

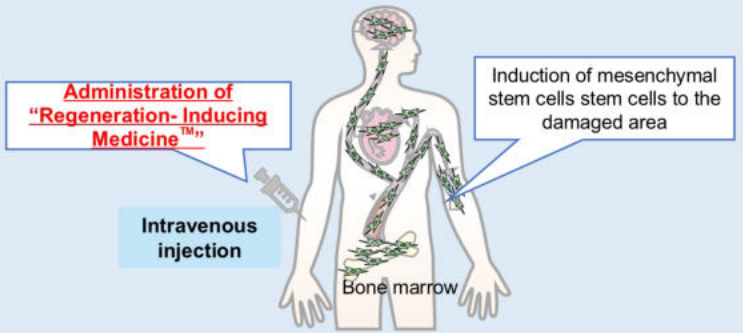


Mesenchymal stem cells lose their functions during in vitro culture

Source: Stem Cell Research & Therapy2018,9:131

Cell Migration ability (×) | Immuno-modulatory ability | Trophic factor Secretion ability | Fibrosis regulation ability (×) | Tissue Regene-ration ability (×)

Induction of MSC in "Regeneration-Inducing Medicine™"



Induction of mesenchymal stem cells into damaged tissues while retaining their native functions

Cell Migration ability | Immuno-modulatory ability | Trophic factor Secretion ability | Fibrosis regulation ability | Tissue Regene-ration ability

"The effects of MSC cell therapy are limited to inflammation suppression and supply of growth factors to the remaining cells", reported by Caplan AI

「Mesenchymal Stem Cells: Time to Change the Name!」 Arnold Caplan June 2017

Source: Stem Cells Transl Med. 2017 Jun; 6(6):1445-1451. doi: 10.1002/sctm.17-0051. Epub 2017 Apr 28.

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Advantages of "Regeneration-Inducing Medicine™"

"Regeneration-Inducing Medicine™" products have an advantage over conventional regenerative medicine products in terms of global expansion

Issues in the Global Development of Regenerative Medicine Products Using Living Cells

Transportation

- Need to be frozen from CPC and other manufacturing facilities and transported with thorough quality control

Manufacturing

- Need for mass culture with uniformity in cell properties
- In the case of Allogeneic transplantation, the need for mass production while keeping costs down

These problems do not occur in "Regeneration-Inducing Medicine™"

→ **Product Transportation Advantages**

→ **Manufacturing Advantages**

Potential for global development "Regeneration-Inducing Medicine™"

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Summary of advantages of "Regeneration-Inducing Medicine™"

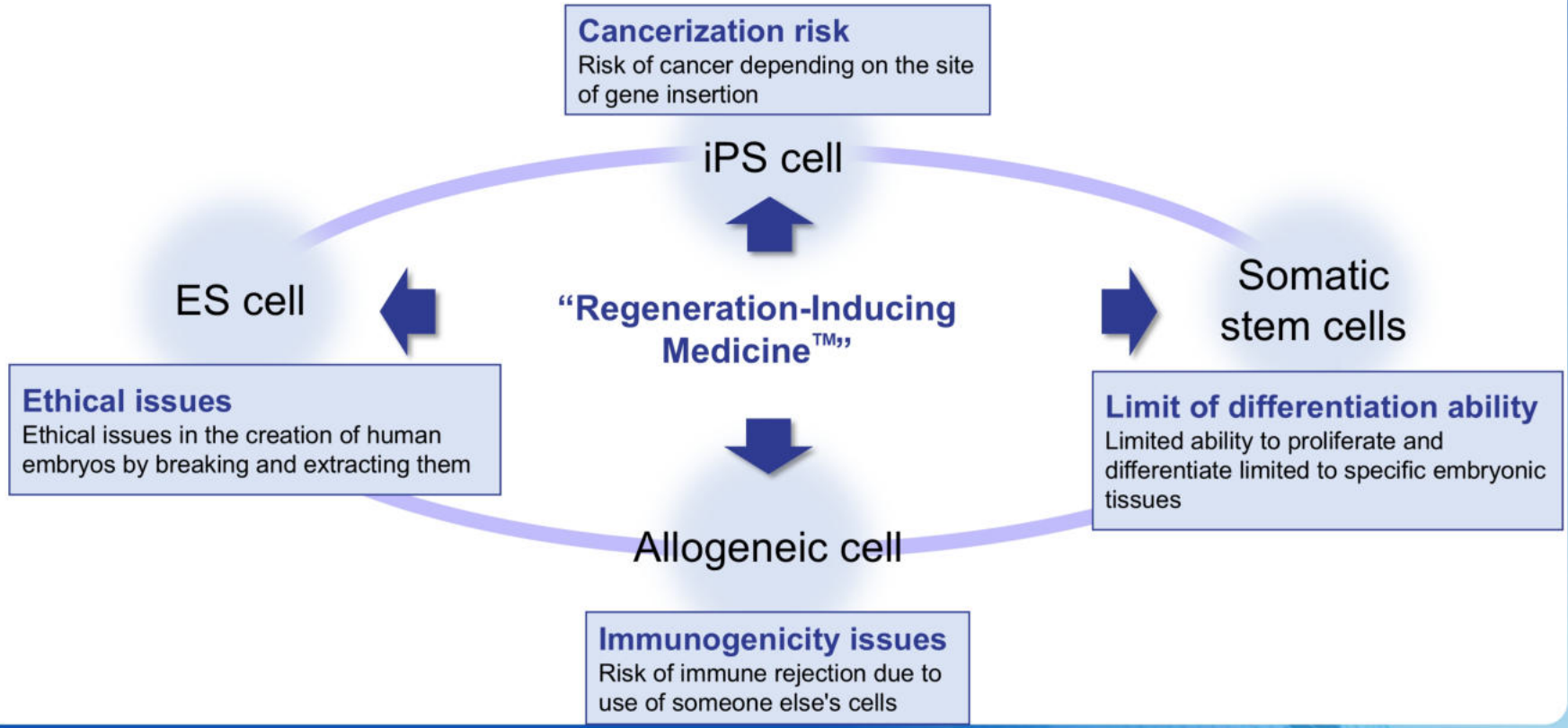
"Regeneration-Inducing Medicine™" includes advantages in both cell therapy and chemicals

		"Regeneration-Inducing Medicine™"	Cell therapy	Chemicals
Efficacy	Tissue regeneration	○ Applicable for large-scale tissue damage	▲ Applicable for large tissue damage with large number of cells	↓ No regeneration
	Mechanism of action	○ Use in vivo native regeneration mechanism	○ Cellular physiological activity	↓ Targeting molecules often including side-effect and off-target
	Indications	○ Same compound can cover a wide range of indications	○ Same platform can cover a wide range of indications	↓ In general, targeting limited indications caused by same mechanism
Safety	Noninvasive	○ Compound mobilizes the patient's cells in vivo and no rejection	↓ Invasive in cell collection Immune-rejection in allogenic case	○ Low noninvasive
Quality	Quality control	○ Easy quality control and stable production	↓ Cell culture includes risk of cellular change	○ Easy quality control and stable production
Other benefit	Cost	○ Normal industrial drug production	↓ CPC and cell collection and transplantation facility is required	○ Affordable and large-scale production
	Regulatory affairs	○ Same as general compound drugs	↓ No standard, and case-by-case regulation is required	○ Standardized regulation

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Summary of advantages of "Regeneration-Inducing Medicine™"

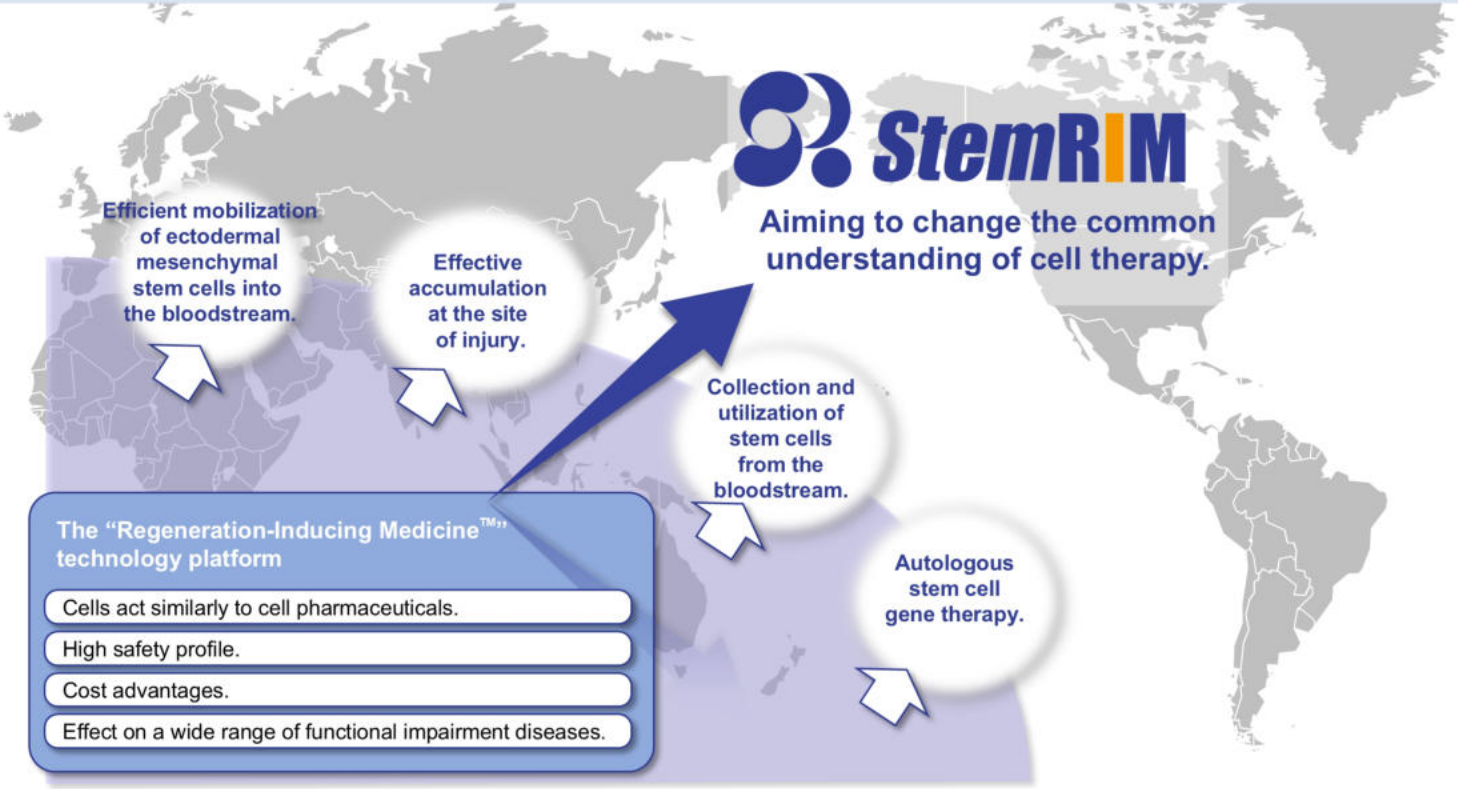
"Regeneration-Inducing Medicine™" can solve the four major problems of conventional cell therapy



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Summary of advantages of "Regeneration-Inducing Medicine™"

The "Regeneration-Inducing Medicine™" technology platform aims to be a game-changer in the field of cell therapy.



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4. Business Areas and Pipeline Progress Status

Overview of Development Pipeline

Project code	Development candidate	Indication	Investigator	Target Area	Research / Pre-clinical / Phase 1 / Phase 2 / Phase 3	Status
Redasemtide (TRIM2)	HMGB1 cell mobilization domain peptides	Epidermolysis bullosa	Shionogi & Co., Ltd.	Japan	Add Phase2	2022.07 Add Phase2 Start 2024.02 Add Phase2 FPI 2025.06 Add Phase2 LPI 2025.09 Add Phase2 Receipt of Study Results
		Acute Ischemic Stroke	Shionogi & Co., Ltd.	Global	Phase2b	2023.04 Global Phase2b start 2025.04 Interim analysis, clinical trial plan change 2025 09 Receipt of Global Phase 2b Study Results
		Ischemic Cardiomyopathy	Osaka University	Japan	Phase2	2024.03 Physician-Initiated Phase2 start 2024.12 Physician-Initiated Phase2 FPI 2026.04 Physician-Initiated Phase2 LPI
		Osteoarthritis of the knee	Hirosaki University	Japan	Phase2	2020.12 Physician-Initiated Phase2 start 2023.04 Physician-Initiated Phase2 completed
		Chronic liver disease	Niigata University	Japan	Phase2	2020.11 Physician-Initiated Phase2 start 2023.05 Physician-Initiated Phase2 completed
TRIM3	Novel Regeneration-Inducing peptide for Systemic administration	Multiple tissue damage diseases	In-house (partnership is planned)	—		Promoting out-licensing activities with multiple domestic and international companies
TRIM4	Novel Regeneration-Inducing peptide for Systemic administration	Multiple tissue damage diseases	In-house (partnership is planned)	—		Promoting out-licensing activities with multiple domestic and international companies
TRIM5	Novel Regeneration-Inducing peptide for Local administration	Multiple tissue damage diseases	In-house (partnership is planned)	—		Expansion of disease model animal data
SR-GT1	Stem cell gene therapy	Epidermolysis bullosa	In-house (partnership is planned)	—		2024.12 Adopted by AMED grant Advance preparations for domestic Phase 1/2 trials

＊Dystrophic epidermolysis bullosa, the target disease, is a rare and intractable disorder affecting approximately 300 patients in Japan. Given the limited availability of approved treatment options and the substantial unmet medical need that remains, the Company plans to submit a marketing authorization application following the evaluation of the results from the additional Phase II clinical trial.

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開発パイプラインにおける1年間の進捗状況

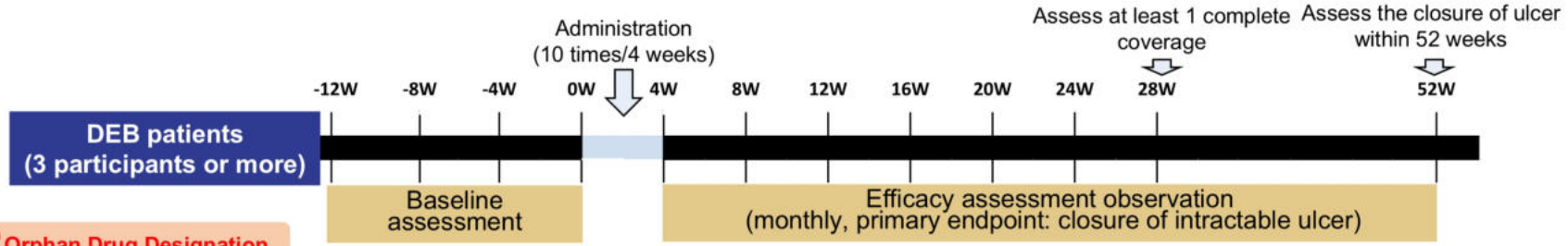
開発コード	内容	対象疾患	開発主体	地域	Progress (as of October 2026)	(Reference) From "Presentation Material for Business Plan and Growth Potential" disclosed on October 31, 2025
Redasemtide (TRIM2)	HMGB1 cell mobilization domain peptides	Epidermolysis bullosa	Shionogi & Co., Ltd.	Japan	Additional Phase 2 Receipt of Study Results	Additional Phase 2 trial currently underway
		Acute Ischemic Stroke	Shionogi & Co., Ltd.	Global	Receipt of Global Phase 2b Study Results	Global Phase 2b clinical trial currently underway
		Ischemic Cardiomyopathy	Osaka University	Japan	Physician-Initiated Phase2 trial currently underway	Physician-Initiated Phase2 trial currently underway
		Osteoarthritis of the knee	Hirosaki University	Japan	Physician-Initiated Phase2 trial completed	Physician-Initiated Phase2 trial completed
		Chronic liver disease	Niigata University	Japan	Physician-Initiated Phase2 trial completed	Physician-Initiated Phase2 trial completed
TRIM3	Novel Regeneration-Inducing peptide for Systemic administration	Multiple tissue damage diseases	In-house (partnership is planned)	–	Non-clinical	Non-clinical
TRIM4	Novel Regeneration-Inducing peptide for Systemic administration	Multiple tissue damage diseases	In-house (partnership is planned)	–	Non-clinical	Non-clinical
TRIM5	(Novel Regeneration-Inducing peptide for Local administration)	Multiple tissue damage diseases	In-house (partnership is planned)	–	Non-clinical	Non-clinical
SR-GT1	SR-GT1 (Stem cell gene therapy)	Epidermolysis Bullosa	In-house (partnership is planned)	–	Preparing for clinical trials	Preparing for clinical trials

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Redasemtide: Additional Phase 2 Trial for DEB

Additional Phase 2 Protocol	
Study objectives	Evaluation of efficacy and safety of Redasemtide in patients with dystrophic epidermolysis bullosa having intractable ulcers
Study design	Single arm, multicenter, open label, uncontrolled
Intervention	Redasemtide (1.0 mg/kg) group: More than 3 participants
Regimen	30-minute intravenous infusion once a day, total 10 times/4 weeks [1st week of administration: 4 times/week, 2nd-4th weeks of administration: twice/week (once every 3-4 days)]
Primary endpoint	**Closure of intractable ulcer**

Clinical Trial Timeline to Date
December 2017: Initiation of Phase 2 investigator-initiated clinical trial
September 2019: Completion of Phase 2 investigator-initiated clinical trial
March 2020: Completion of follow-up study for Phase 2 investigator-initiated clinical trial
July 2022: Initiation of additional Phase 2 clinical trial
March 2023: First patient enrolled in the additional Phase 2 clinical trial
May 2023: Orphan Drug Designation
July 2025: Final patient enrolled in the additional Phase 2 clinical trial
September 2026: Additional Phase 2 Receipt of Study Results



Orphan Drug Designation

In May 2023, the Ministry of Health, Labour and Welfare designated ledasemtid as an orphan drug for hypoparathyroidism-related epidermolysis bullosa. This designation reflects the Ministry's recognition of the development plan's validity for treating hypoparathyroidism-related epidermolysis bullosa. Eligibility for the priority review system is expected to shorten the review period, facilitating earlier approval.

Status

Planned Market Launch by March 2028

* Shionogi & Co. Ltd., 1st Quarter of Fiscal 2025 Financial Results, July 28, 2025, pp.31
** jRCT2031220378

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Receipt of Additional Phase 2 Study Results for Redasemtide in Dystrophic Epidermolysis Bullosa

Primary endpoint
Closure of intractable ulcer

3 of 4 cases

Achieved
Closure of intractable ulcer

Achieved
Primary endpoint

Clinical Findings

Favorable Clinical Outcomes, Including Reduction in Total Ulcer Area, Were Observed in Some Patients

Safety and Tolerability

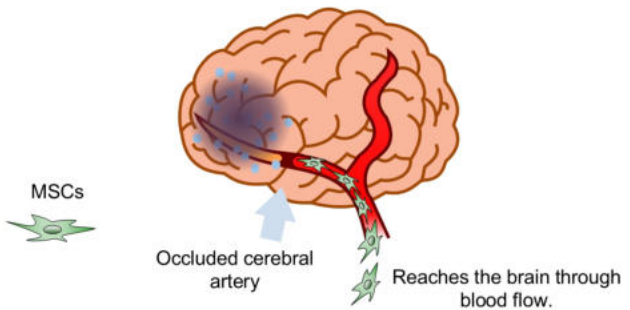
No New Safety Concerns Were Identified for Redasemtide, and Favorable Tolerability Was Confirmed

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Redasemtide: Global Phase 2b Trial for Acute Ischemic Stroke

Developing a stroke treatment that alleviates time constraints compared to conventional therapies.

Acute Ischemic Stroke



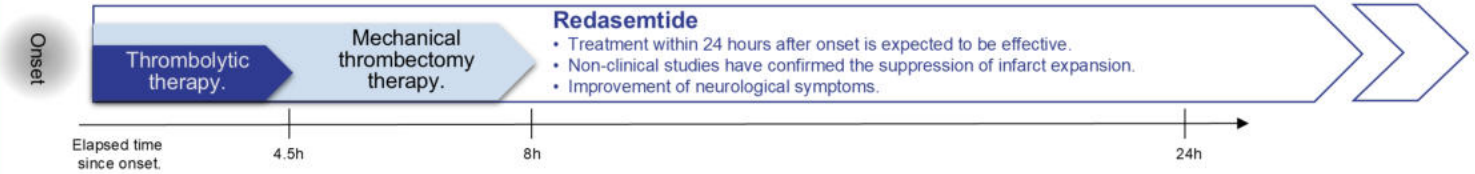
Market Size in Japan

187 thousand people



Mechanism of action

- ✓ Suppression of inflammation
- ✓ Promotion of angiogenesis and secretion of trophic factors to remaining nerve cells
- ✓ Inhibition of fibrosis to suppress the expansion of the infarct area in stroke.



Onset to 4.5 hours

Thrombolytic therapy (t-PA intravenous therapy): A treatment method that dissolves clots and restores blood flow by administering a thrombolytic agent (t-PA).

Onset to 8 hours

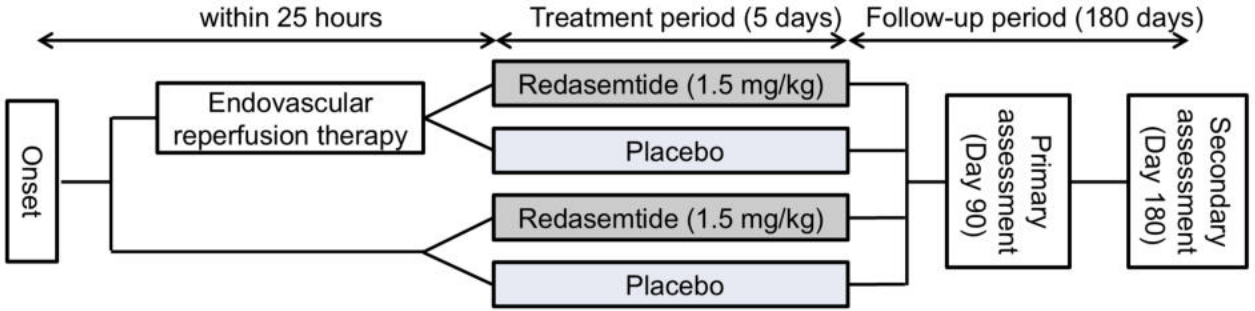
Mechanical thrombectomy therapy: A treatment method that retrieves clots using a catheter and thrombectomy device.

Compared to conventional therapies, Redasemtide offers more relaxed time constraints and is expected to be used in combination with t-PA during the acute phase, as a first-line treatment for t-PA contraindications, and as a standalone treatment in the subacute phase.

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Redasemtide: Global Phase 2b Trial for AIS

Phase 2b Trial Protocol (After interim analysis)	
Study objectives	Evaluation of the Efficacy, Safety, and Tolerability of Redasemtide in Patients with Acute Ischemic Stroke
Subject population	Patients aged 18 years or older who can receive treatment within 25 hours of stroke onset, with a baseline NIHSS* score between 8 and 22.
Study design	Multicenter, Randomized, Placebo-Controlled, Double-Blind
Intervention	Cohort A: Patients Who Did Not Received Thrombolytic Therapy and/or Mechanical Thrombectomy ・Redasemtide (1.5 mg/kg) treatment group ・Placebo group Cohort B: Patients Who Receive Thrombolytic Therapy or Mechanical Thrombectomy ・Redasemtide (1.5 mg/kg) treatment group ・Placebo group　　Total: 679 subjects
Dose	Intravenous Administration Once Daily for 90 Minutes Over 5 Days
Primary End Point	Modified Rankin Scale (mRS) at 90 Days After Initial Dosing
Region	Japan, Europe, North America, China, etc.



*NIHSS(National Institutes of Health Stroke Scale)：Stroke Neurological Severity Rating Scale (42 points in total, the higher the score, the more severe)

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Receipt of Global Phase 2b Study Results for Redasemtide in Acute Ischemic Stroke

Primary endpoint
Modified Rankin Scale (mRS) at 90 Days After Initial Dosing

Compared with Placebo

No Statistically Significant Difference

Cohort Without Endovascular Recanalization Therapy

Primary Endpoint Not Met

Efficacy Signal

Consistent with the Previous Domestic Phase 2 Study, a Trend Toward Improvement in mRS Was Observed in the Redasemtide Group

Exploratory Subgroup Analysis Suggested a Trend Toward Improvement in mRS Among Patients with Greater Baseline Stroke Severity

Safety and Tolerability

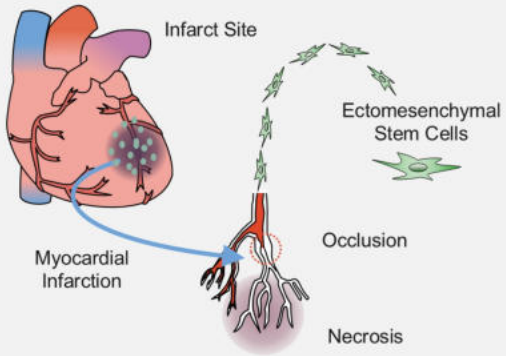
Comparable Rates of Adverse Events in Both Groups.

No New Safety Concerns Identified/Favorable Tolerability Confirmed

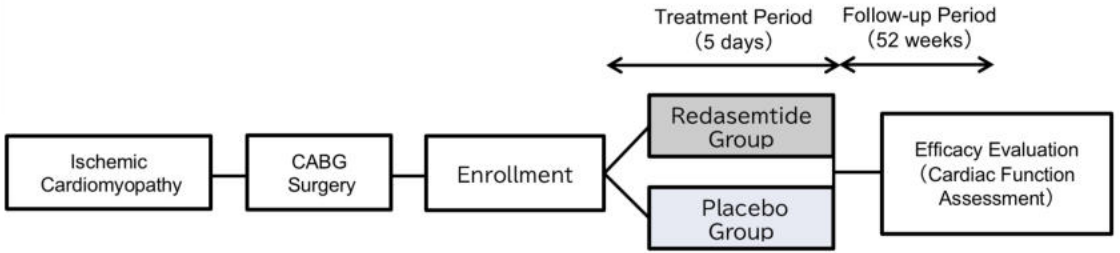
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Redasemtide: Phase II Investigator-Initiated Clinical Trial for Ischemic Cardiomyopathy

Ischemic Cardiomyopathy



Objective	Evaluate the efficacy and safety of Redasemtide
Patients	Patients with ischemic cardiomyopathy who have undergone coronary artery bypass grafting (CABG).
Study Design	Multicenter, randomized, placebo-controlled, double-blind study.
Treatment Arms / Number of Patients	Redasemtide and Placebo Groups: 14 Patients in Total
Administration Method / Duration	Intravenous administration for 5 days.
Efficacy Endpoints	Assessment of cardiac function by echocardiography and other cardiac function tests at Week 52.



Status

April 2026 : Enrollment Completed in Investigator-Initiated Phase Ⅰ/Ⅱ Clinical Trial

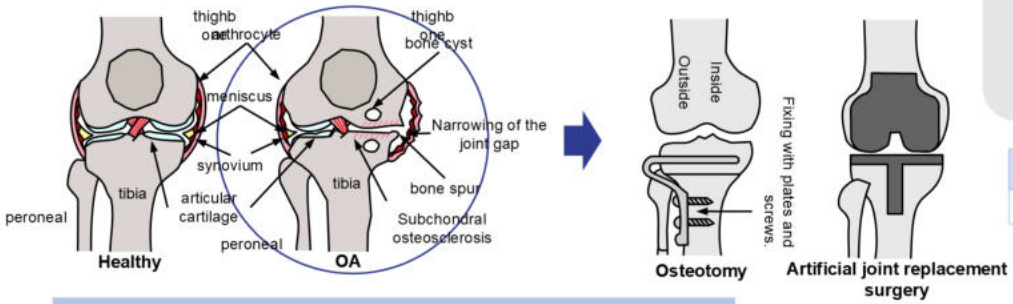
CABG (Coronary Artery Bypass Grafting): A surgical procedure that restores blood flow to the heart by creating a new pathway (bypass) around a narrowed or blocked coronary artery.
Japan Registry of Clinical Trials (jRCT): https://jrct.niph.go.jp/latest-detail/jRCT2051240065

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Redasemtide：Osteoarthritis of the Knee

Developing a treatment for osteoarthritis of the knee that does not rely on surgical interventions.

Osteoarthritis of the Knee（OA）



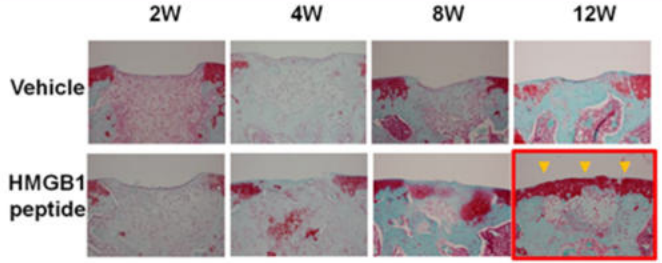
Market size in Japan

・Number of patients with symptoms: approximately 8 million

・Estimated number of potential patients: approximately 25 million

Mechanism of action
✓ Regeneration of articular cartilage tissue.

Cartilage regeneration with Redasemtide in a mouse model



(Note)Joint research between our company and Osaka University

Cartilage regeneration by blood-induced bone marrow-derived mesenchymal stem cells following Redasemtide administration (the red-stained area indicated by ▼)

Conventional treatment methods.

・For mild cases, analgesics (oral or topical) and hyaluronic acid injections into the knee joint are prescribed.

・There is no radical treatment available, and in severe cases, the decline in quality of life (QOL) is significant, necessitating surgical interventions (such as total knee arthroplasty and high tibial osteotomy).

A radical treatment for osteoarthritis of the knee through the administration of pharmaceuticals, without the need for surgical intervention, is anticipated

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Redasemtide：Osteoarthritis of the Knee

Phase 2 Protocol	
Study objectives	Evaluation of efficacy and safety of Redasemtide in patients with Osteoarthritis (OA) of the knee
Subject population	Patients with knee OA who have undergone high tibial osteotomy (HTO) and arthroscopic microfracture
Study design	Multiple arms, Single center, randomized, placebo-controlled, double blinded
Intervention	**Redasemtide (1.5 mg/kg) : 10 participants** **Placebo : 10 participants** **total 20 participants**
Regimen	90-minute intravenous infusion, total 8 times / 4 weeks [once every 3-4 days]
Primary endpoint	**Presence/absence and percentage of adverse events**
Secondary endpoint	**Morphological assessment (based on MRI images) and functional assessment (KOOS)**

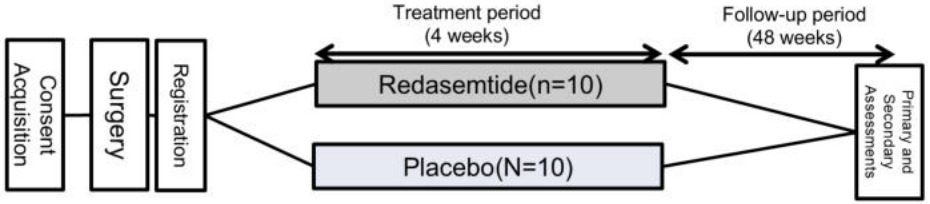
Phase 2 Clinical Trial Results

- Safety Evaluation

  No severe adverse events or side effects related to Redasemtide were identified. The safety of Redasemtide administration for osteoarthritis of the knee was confirmed.

- Morphological and Functional Evaluation

  As a morphological assessment of cartilage damage, a primary cause of osteoarthritis of the knee, MRI imaging was conducted. At 52 weeks after the start of administration, the median change in the defect area of the medial femoral condyle cartilage was -3.5% in the placebo group, compared to -7.5% in the Redasemtide group, showing a tendency toward greater reduction in defect area with Redasemtide. Additionally, endoscopic observation by specialists revealed favorable cartilage regeneration in 5 cases in the Redasemtide group (compared to 2 cases in the placebo group).



Status

We are currently reviewing our future development strategy following favorable results from a physician-initiated Phase 2 clinical trial.

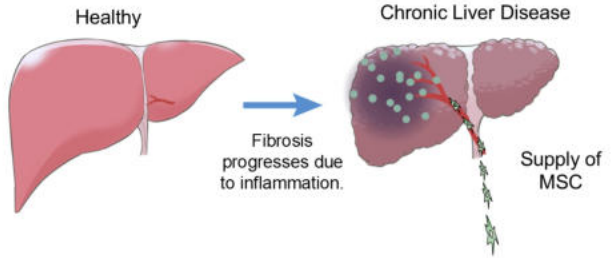
* High tibial osteotomy(HTO): Surgery to reduce knee pain by making an incision on tibia to correct the surface angle of the tibial joint of the O-leg so that the weight is applied to the lateral side of the joint where intact cartilage and meniscus are left.
** Arthroscopic microfracture: A treatment procedure that promotes the recruitment of bone marrow stem cells for tissue repair by making small holes in the subchondral bone at the mother bed of the damaged cartilage to flow out the blood and bone marrow fluid.
*** Knee Injury and Osteoarthritis Outcome Score (KOOS)：One of the scores to measure the outcome of knee injury and osteoarthritis
**** jRCT2021200034

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Redasemtide：Chronic Liver Disease

Developing a treatment for chronic liver disease that works by inhibiting fibrosis.

Chronic Liver Disease (CLD)



Market size in Japan

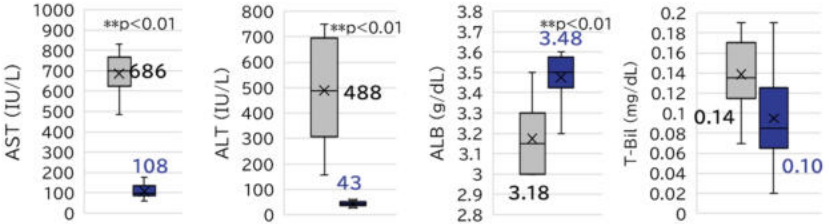
Approximately 400,000 to 500,000 people



Mechanism of action

- ✓ Exhibits strong anti-inflammatory effects
- ✓ Improves liver tissue fibrosis
- ✓ Regenerates liver function through stem cell activation

In a cirrhotic mouse model, improvements in serum liver damage markers (AST, ALT) and liver function indicators (ALB, T-Bil) were confirmed with Redasemtide treatment.



*Nojiri S, Tsuchiya A, Tamai K, Terai S et al. Inflamm Regen. 2021

Conventional treatment methods

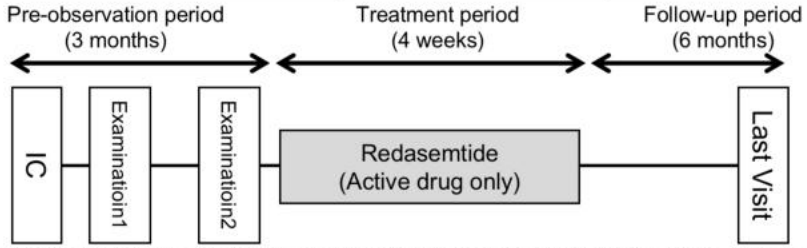
No established treatment can be expected to completely cure the condition. Liver transplantation is effective for advanced cirrhosis with fibrosis, but there is a concern about the shortage of organ donors.

Industrially producible Redasemtide is expected to provide a fundamental treatment for chronic liver disease through drug administration.

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Redasemtide：Chronic Liver Disease

Phase 2 Protocol	
Main purpose	Evaluate the safety and exploratory efficacy in patients with chronic liver disease
Clinical trial design	Single arm study, Open label, Uncontrolled
Target patient	Patients with chronic liver disease with liver hardness test results of 4 kPa or greater by MR elastography.
Administration group/number of cases	1.5 mg/kg (free form), 90minutesintravenous infusion •Cohort A: 4 times / 4 weeks [once a week] •Cohort B: 7 times / 4 weeks [Week 1: 4 days, Week 2-4: once a week (1 dosage/3-4 days)]
Endpoint	**Rate of change in liver stiffness, rate of change in liver stiffness using ultrasound elastography, and rate of change in Child-Pugh score, etc.**
Site	Division of Gastroenterology and Hepatology, Niigata University Medical and Dental Hospital



* MR elastography: Magnetic Resonance Elastography (MRE) is one test that can quantitatively evaluate liver fibrosis.
** Child-Pugh score：Child-Pugh score is an assessment method mainly used to evaluate liver reserve function in patients with chronic liver diseases such as liver cirrhosis. It scores the severity of liver dysfunction using hepatic encephalopathy, ascites, serum bilirubin level, serum albumin level, and prothrombin activity, and classifies it into three stages, A to C.
*** jRCT2031200232

Phase 2 Clinical Trial Results

Safety Evaluation: Presence and Incidence of Adverse Events

2 mild adverse events (voice disorder and fever) with a possible relationship to the investigational drug were observed, both of which resolved. 1 severe adverse event (bleeding during liver biopsy) occurred, but it resolved without treatment and was deemed unrelated to the investigational drug. These findings confirm the tolerability of Redasemtide administration for chronic liver disease.

Efficacy Evaluation: Rate of Change in Liver Stiffness, Change in Liver Stiffness by Ultrasound Elastography, and Change in Child-Pugh Score

In Cohort A (5 cases), an improvement in liver stiffness based on MR elastography was observed at 78 days and 162 days after the start of treatment, showing an average reduction of 12% and 8%, respectively, compared to baseline. In addition to improvements in liver stiffness by MR elastography, several cases showed an accompanying improvement trend in other fibrosis indicators (Fibrosis Index, Fibrosis Markers, and modified HAI Fibrosis Stage values). Based on these various efficacy evaluation indicators, the principal investigator's overall assessment suggested an improvement trend in liver fibrosis in 3 of 5 cases (60%) in Cohort A and 2 of 5 cases (40%) in Cohort B.

Status

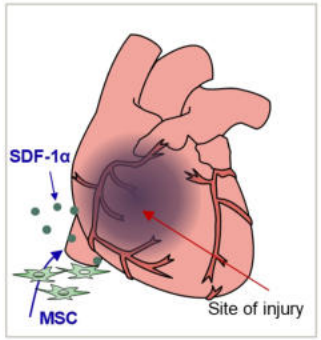
We are currently reviewing our future development strategy following favorable results from a physician-initiated Phase 2 clinical trial.

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TRIM5：Novel Regeneration-Inducing Peptide for Local Administration

Developing protein drugs that accumulate mesenchymal stem cells at the site of injury

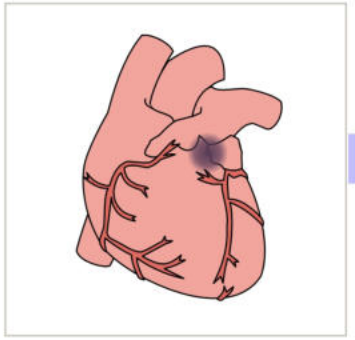
Large injury



- SDF1-a is released, and mesenchymal stem cells mobilized in the blood accumulates at the injury

＝Mechanism of action in Redasemtide, TRIM3, or TRIM4 is effective

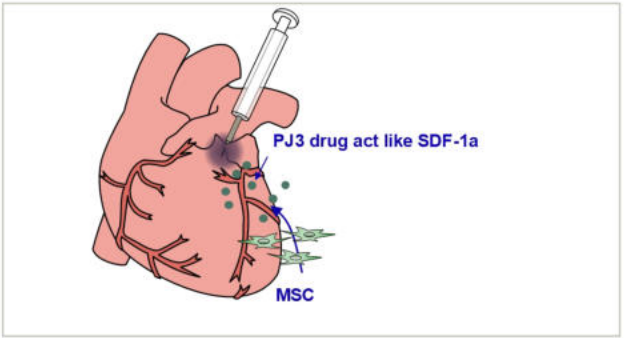
Small injury or chronic phase



- SDF1-a is not released, and mesenchymal stem cells cannot accumulate efficiently

＝Combination therapy that maximizes the effects of "Regeneration-Inducing Medicine™" is effective

Efficient accumulation of mesenchymal stem cells by topical administration ofTRIM5



- Effective accumulation of mesenchymal stem cells at the site of injury by topical administration or intravenous injection

＝Maximize damage repair effect of mesenchymal stem cells

- ✓ Multiple candidate proteins have been identified so far
- ✓ Confirmed good results in animal experiments
- ✓ Currently, the most suitable indication is being selected through multiple animal model experiments

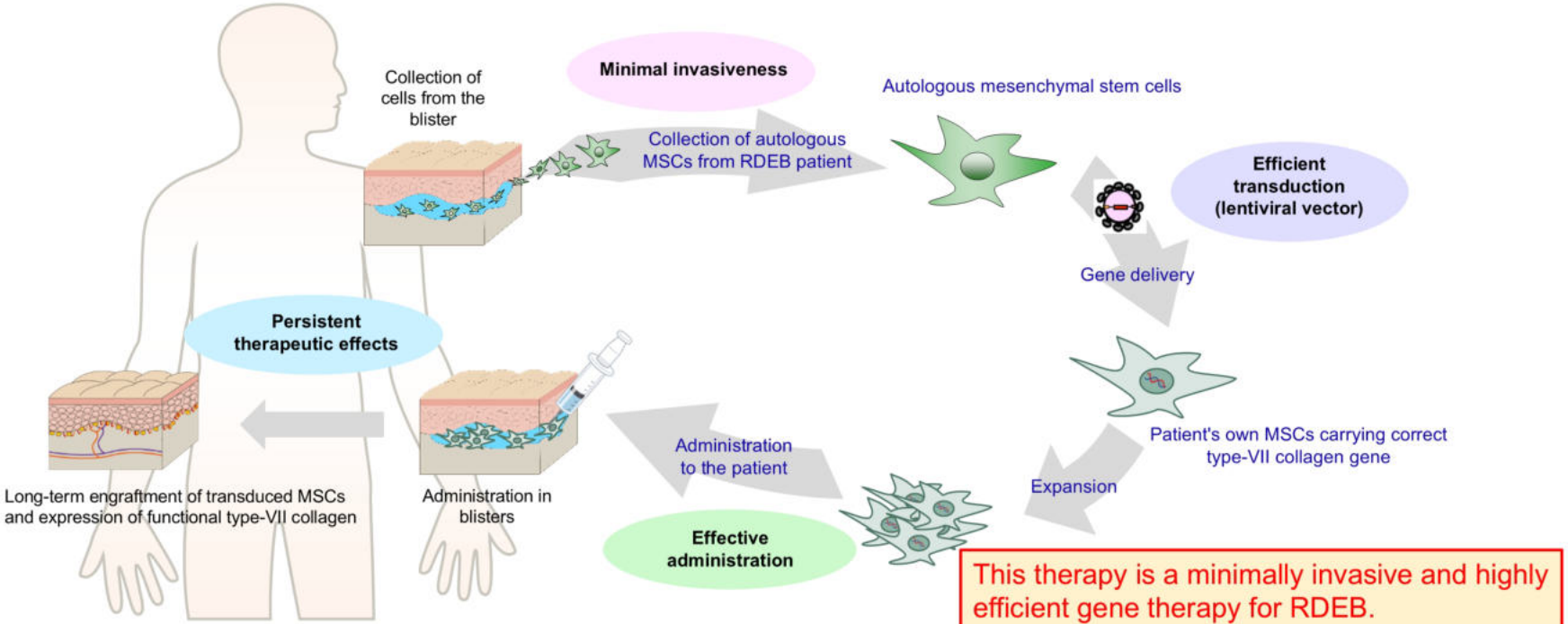
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SR-GT1：Stem cell gene therapy

Aim to cure intractable genetic disease by stem cell gene therapy

Concept

Ex vivo gene therapy involving the introduction of correct type VII collagen gene into autologous mesenchymal stem cells (MSCs) and administration of the cells in the blisters of the patient.



This therapy is a minimally invasive and highly efficient gene therapy for RDEB.

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SR-GT1：Stem cell gene therapy

Ex vivo gene therapy with minimal invasiveness, high efficacy, and persistent effect
This therapy employs a novel method of isolating Bf-MSCs from a patient, efficient delivery of functional type VII collagen gene to the cells, and novel administration method to the patient with minimal invasiveness.

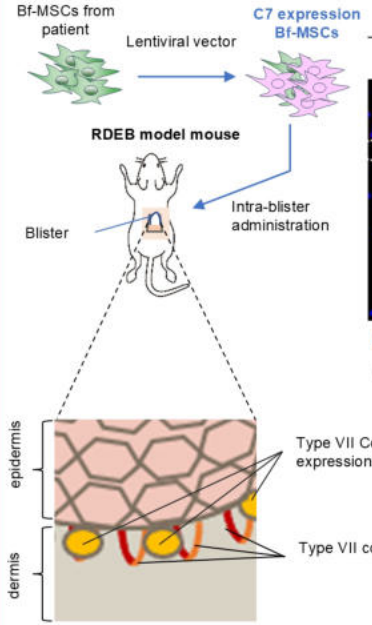
Company	Brand, Generic or Code name	*in/ex vivo*	Target cell	Formulation	Administration route	Comparison with StemRIM			
						Area	Patient's burden	Effective length	Efficacy
StemRIM	**SR-GT1**	*ex vivo*	Mesenchymal stem cells	Cell suspension	Intra-blister administration	**Non-ulcer surface**	Low	Long-term (sustained)	High
Krystal Biotech	Vyjuvek	*in vivo*	-	Virus containing gel	Local application	**Ulcer surface**	Low	Long-term (limited)	High
Abeona Therapeutics	prademagene zamikeracel	*ex vivo*	Skin keratinocytes	Epidermal Sheet	Epidermal sheet transplantation	**Ulcer surface**	High	Long-term (limited)	High
Castle Creek Biosciences	dabocemagene autoficel	*ex vivo*	Dermal fibroblasts	Cell suspension	Intradermal administration	**Ulcer surface**	High	Long-term (limited)	Low
Amryt Pharma	AP-103	*in vivo*	-	Protein solution	Intravenous administration	**Whole body**	Low	Short-term	High

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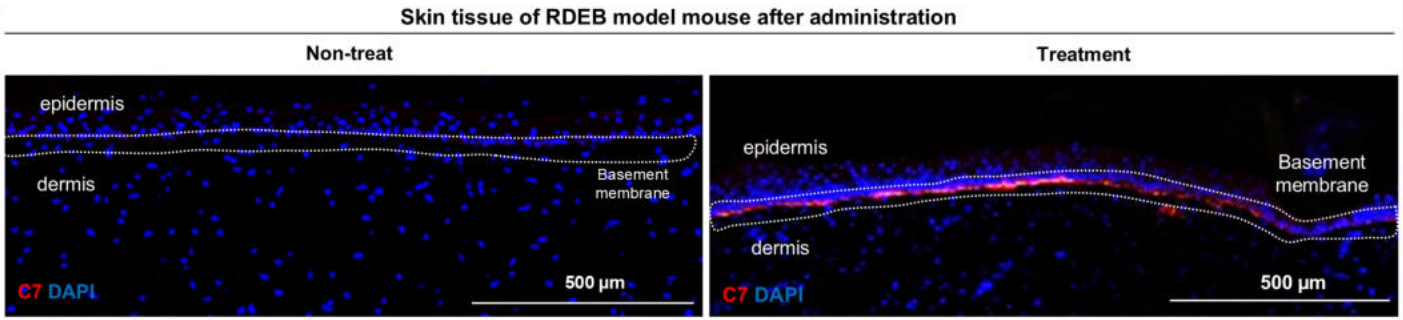
SR-GT1：Stem cell gene therapy

Verification of therapeutic efficacy and duration of drug effect of this treatment using RDEB model mice

■ Restoration of the type-VII collagen protein（C7）at the basement membrane on RDEB model mouse

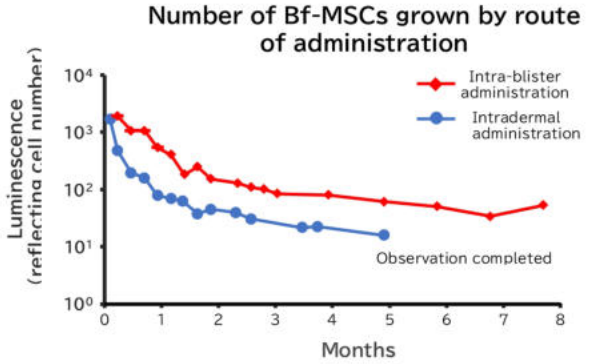


Skin tissue of RDEB model mouse after administration

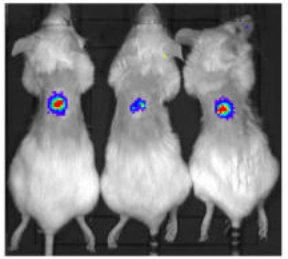


DAPI:

■ Long-lasting strong signal compared to the intradermal administration



Intra-blister (5 months)



*Joint research between and Osaka University

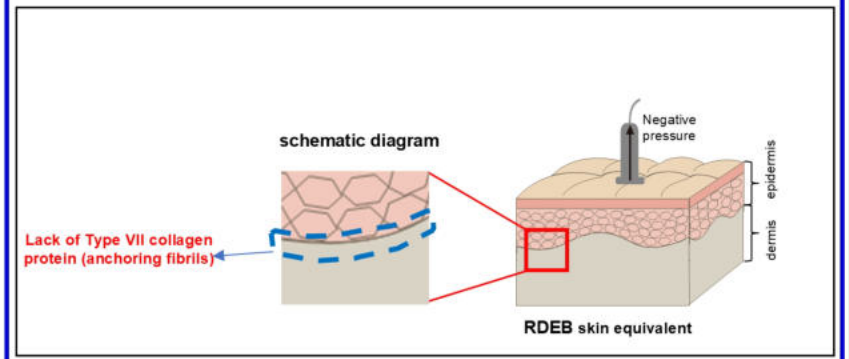
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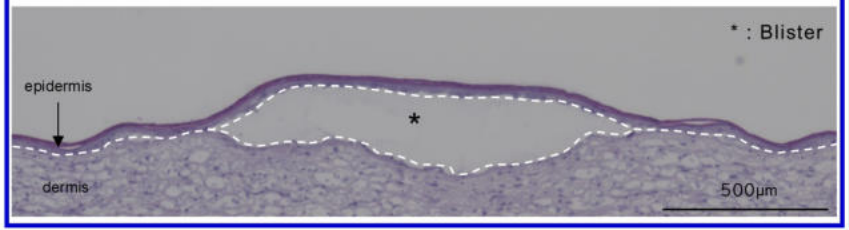
SR-GT1：Stem cell gene therapy

Therapeutic effects on RDEB skin model

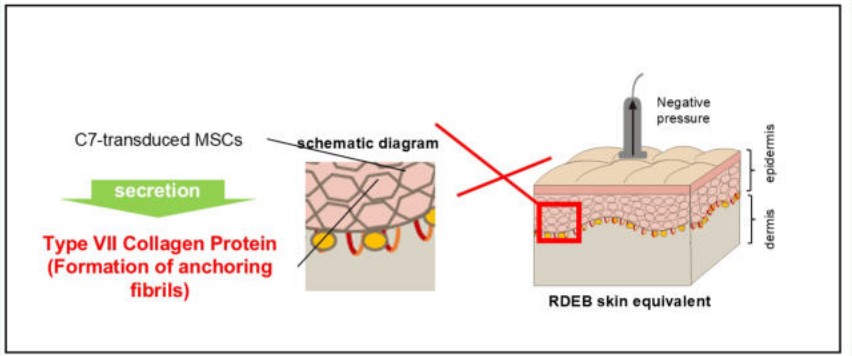
- We confirmed the effect of gene therapy using patient-derived Bf-MSCs by RDEB skin model and artificially forming blisters by suction method.

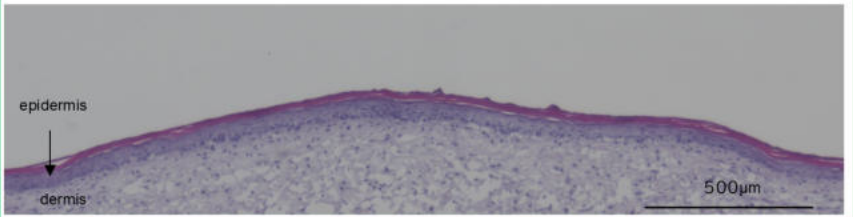
RDEB skin equivalents





RDEB skin equivalents with gene transduced Bf-MSCs





Joint research between and Osaka University

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5. Future Growth Strategy

Future Growth Strategy

Growth Strategy to Maximize the Potential Value of "Regeneration-Inducing Medicine™"

Out-Licensing of Next-Generation Regeneration-Inducing Medicine™

TRIM3, TRIM4, and TRIM5 are next-generation "Regeneration-Inducing Medicine™" candidates discovered through joint research with Osaka University and proprietary screening efforts. TRIM3 and TRIM4 have demonstrated efficacy in non-clinical studies, and we plan to continue early-stage development in-house, followed by out-licensing to pharmaceutical companies. Similarly, for TRIM5, we intend to pursue out-licensing after selecting appropriate target indications. Through these out-licensing activities, we aim to establish diverse revenue streams including upfront payments, milestone payments, and royalties.

Support for the development of Redasemtide.

To support the smooth progress of clinical development by Shionogi & Co., Ltd., we will continue to provide assistance for investigator-initiated clinical trials at universities and other institutions (for ischemic cardiomyopathy, chronic liver diseases, and osteoarthritis). In the case of epidermolysis bullosa, we aim for early approval and commercialization by utilizing the designation as an orphan drug. As these programs advance, we aim to maximize milestone and royalty revenues.

The clinical development of stem cell gene therapy (SR-GT1)

SR-GT1 is a therapeutic product designed for the treatment of severe genetic disorders, utilizing a process in which the patient's own stem cells are genetically edited ex vivo and then reintroduced into the body. Development is currently focused on severe epidermolysis bullosa as the initial indication, with efficacy and safety confirmed in animal studies. Non-clinical studies are ongoing in preparation for the start of clinical trials. Following initial in-house development, we aim to out-license the program and are also considering applications to other genetic disorders.

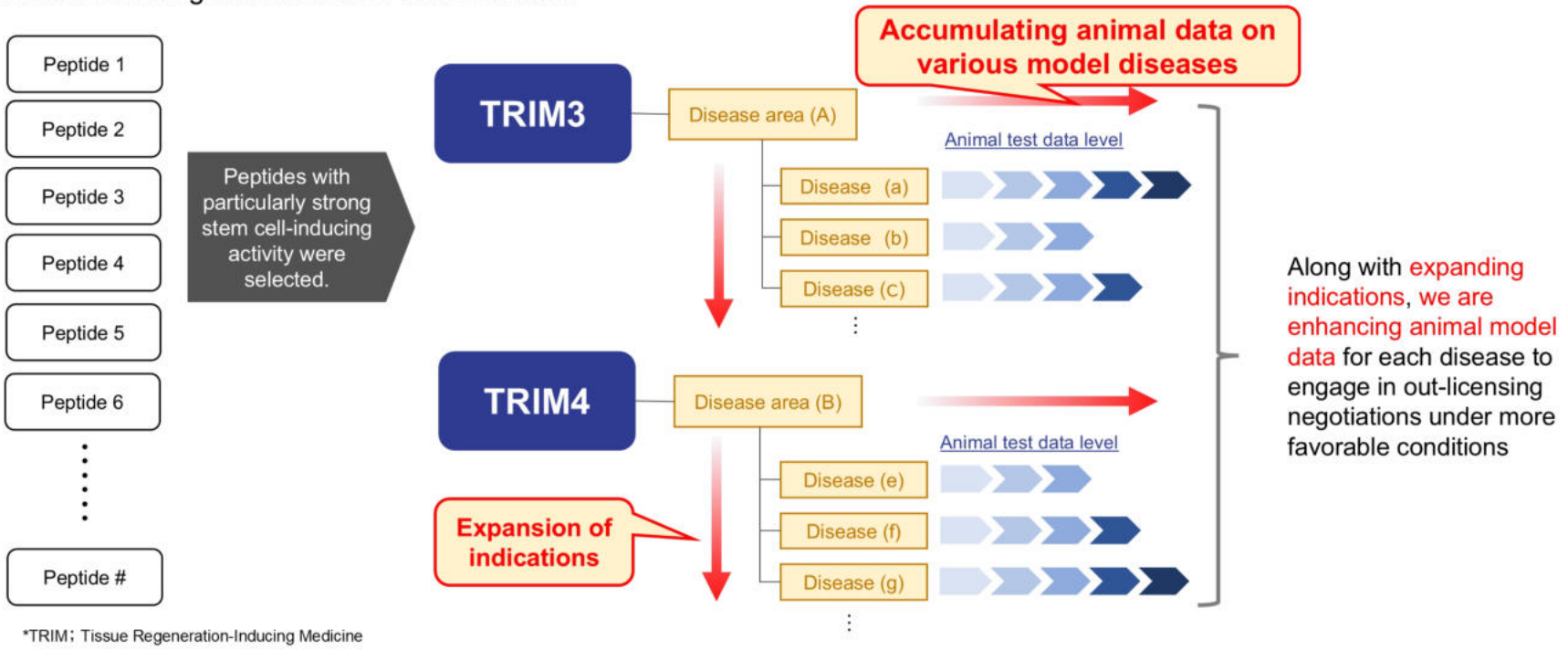
Expansion of the development pipeline.

To complement the existing pipeline, enhance corporate value, and diversify risk, we are actively pursuing the discovery of new "Regeneration-Inducing Medicine™" candidates. Through multifaceted screening in academia-industry collaborations, we are identifying new compounds, filing patent applications, and advancing non-clinical studies. Utilizing single-cell functional evaluation technologies, we are developing high-performance stem cell-based pharmaceuticals. By sharing these technologies with both domestic and international academic institutions and pharmaceutical companies, we aim to establish business partnerships and license-out opportunities, thereby diversifying revenue sources and achieving sustainable growth.

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TRIM3, TRIM4

We have identified several peptides that mobilize mesenchymal stem cells from the bone marrow into the bloodstream, accumulate in damaged tissues, and induce functional regeneration. Among them, two peptides with particularly prominent activity have been selected as candidates for the next-generation “Regeneration-Inducing Medicine™” : TRIM3 and TRIM4, and out-licensing activities have been initiated.



*TRIM: Tissue Regeneration-Inducing Medicine

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事業開発活動

Continuing from last year, out-licensing negotiations were conducted with multiple pharmaceutical companies both domestically and internationally.

BioJapan

2025.10.8~10 @Yokohama, JPN

J.P.Morgan
Healthcare Conference

2026.1.12~15 @San Francisco, CA

Bio International Convention

2026.6.22~25 @Boston, MA



2026.6 Bio International Convention

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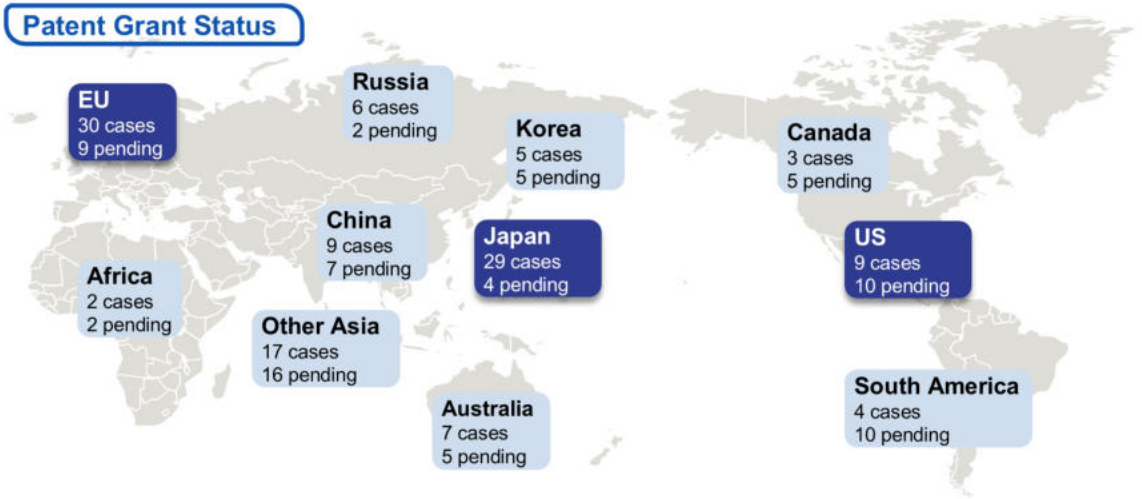
IP Strategy

Patents related to “Regeneration-Inducing Medicine™” have been granted in various countries. We are steadily promoting the intellectual property protection of our research outcomes, paving the way for global expansion.

Total Patents 121

Patent Pending 75

Patent Grant Status



Countries of Grant and Application

JPN USA CHN DEU KOR IND ESP POLE SWD RUS BEL IDN BRA MEX ZAF

ISR HKG GBR FRA AUS CAN ITA IRL NLD CHE DNK FIN TUR SGP TWN

* As of July 2025

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IP Strategy

During the fiscal year ended July 2026, patents related to Regeneration-Inducing Medicine™ were granted in multiple countries. Steady progress was made in securing intellectual property rights for research achievements. (Based on timely disclosures issued between August 2025 and July 2026.)

Pipeline Products	Indication	Countries/Regions Where Patents Have Been Granted(Grant Date)
Redasemtide	**Cartilage Disorders** Traumatic Cartilage Defects, Osteoarthritis, and Osteochondritis Dissecans	Canada Feb.2026 / Australia Jun.2026 / Mexico Jul.2026
Redasemtide	**Cardiomyopathy and Chronic Heart Failure** Cardiomyopathy, Prior Myocardial Infarction, and Associated Chronic Heart Failure	Canada Jul.2026
Redasemtide	**Fatty Liver Disease and NASH** Fatty Liver Disease and Non-Alcoholic Steatohepatitis (NASH)	Japan Dec.2025 / Russia Jun.2026
Redasemtide	**Psoriasis** Plaque Psoriasis and Related Disorders	USA Oct.2025
SR-GT1	**Epidermolysis Bullosa(EB)** Curative Therapy for Dystrophic Epidermolysis Bullosa/Stem Cell Gene Therapy	Japan Oct.2025 / Japan Mar.2026 / South Korea Jul.2026 / South Korea Aug.2026

5 Therapeutic Area — Patented in Cartilage, Cardiac, Hepatic, and Dermatological Fields

11 Patents — Patent Registrations(Since August 2025)

7 Countries and Regions — Global Expansion Across Multiple Fields

2 Pipeline Product(s) — Redasemtide＋SR-GT1

* Prepared by the Company based on its timely disclosures from August 2025 to August 2026. The number of patents is based on patent grant announcements disclosed during the period. The number of countries and regions is an approximate figure excluding duplicates.

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Activities of “StemRIM Institute of Regeneration-Inducing Medicine, Osaka University”

StemRIM

StemRIM Institute of Regeneration-Inducing Medicine

In June 2020, StemRIM Institute of Regeneration-Inducing Medicine, Osaka University (covering an area of 1,540 square meters) was established on the 6th and 7th floors of the Techno Alliance Building at Osaka University's Suita Campus. Professor Masayuki Endo (Department of Children’s and Women’s Health, Graduate School of medicine and Division of Health Sciences, Osaka University) was appointed as the institute's director. The team includes distinguished members such as Specially Appointed Professor Shinya Murakami (Department of Periodontology and Regenerative Dentistry, Osaka University, Graduate School of Dentistry.), Professor Masaru Ishii (Department of Immunology and Cell Biology, Graduate School of medicine and Frontier Biosciences, Osaka University), and Professor Manabu Fujimoto (Department of Integrated Medicine, Graduate School of medicine, Osaka University). Together, they aim to explore and advance the multi-faceted development of “Regeneration-Inducing Medicine™”. To date, several collaborative research projects have made significant progress.

Joint Research Projects

(number of events)

	FY 2022	FY 2023	FY 2024	FY 2025	FY 2026	FY on FY	Notes
Division of Health Sciences	2	3	2	2	1	-1	Neonatal-Associated Diseases
Division of Biofunctional Research	—	—	—	—	—	±0	
Division of Medical Research	1	2	2	3	5	+2	Nervous System Diseases, Orthopedic-Related Diseases
Division of Dentistry	5	5	5	6	6	±0	Periodontitis-Related Diseases
Total	8	10	9	11	12	+1	

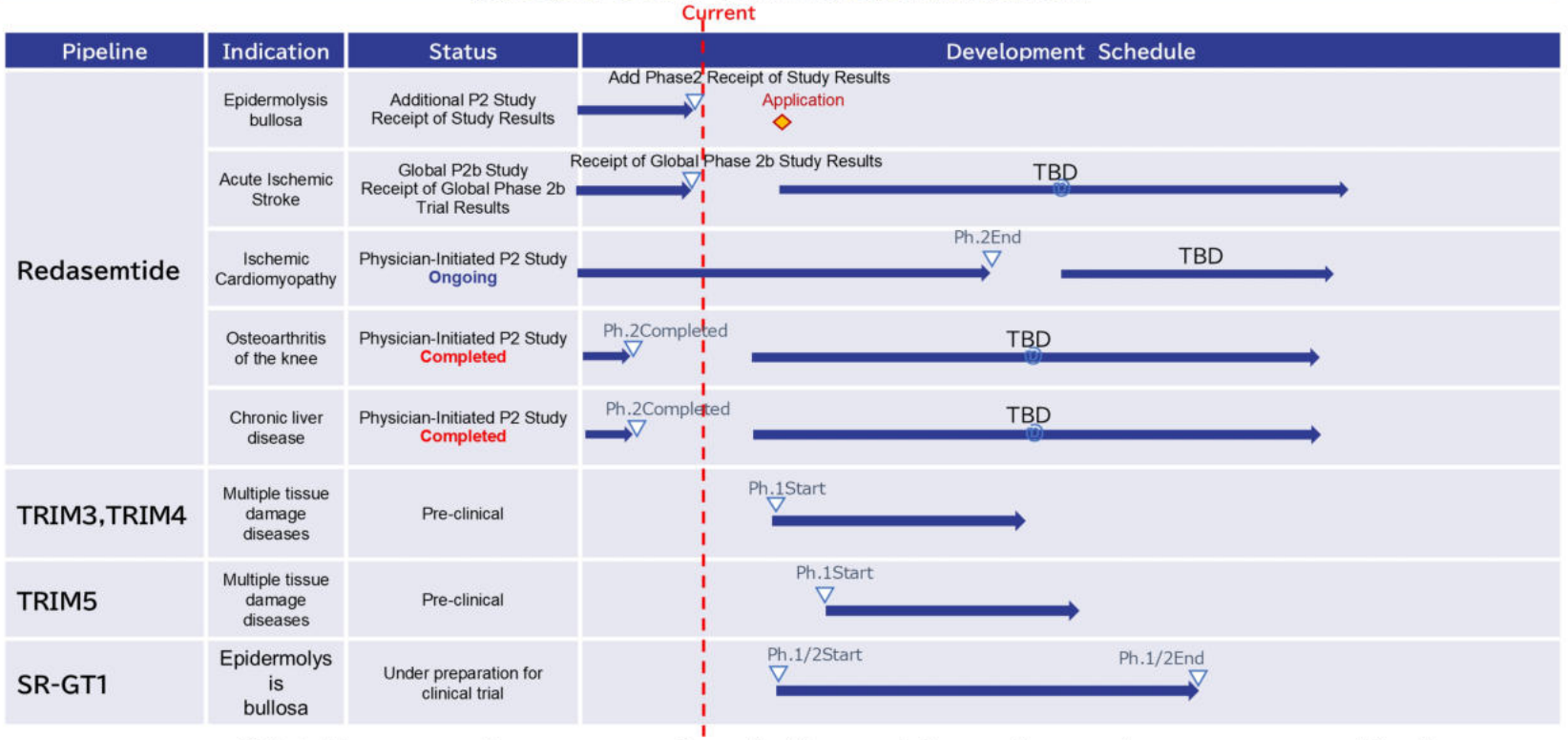


Website (Japanese):
https://stemrim-osaka-u.jp/

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パイプライン開発計画

Given the long development timeline for pharmaceuticals, a pipeline development plan is formulated based on a long-term management vision.



*TBD: to be determined

This table represents our company's projections and does not guarantee progress as outlined.

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Forecast for the fiscal year ending July 2027

Research and Development Progress for the Fiscal Year Ended July 2026

Previous Disclosure (October 31, 2025)

Research Progress Outlook for the Fiscal Year Ending July 2026

Research and development for Redasemtide is progressing in clinical trials and toward expanded indications. Additionally, negotiations for clinical trials and licensing for candidate "Regeneration-Inducing Medicine™" following Redasemtide continue to advance.

Cash Outflow Forecast for the Fiscal Year Ending July 2026

Cash Outflow for R&D	1,300 million to 1,700 million yen
Cash Outflow for General and Administrative Expenses	230 million to 310 million yen

Actual Results

Research Progress Results for the Fiscal Year Ended July 2026

Received Phase 2 trial results for the additional indication of RDEB for Redasemtide. Additionally, Received Global Phase 2b trial results for acute ischemic stroke for Redasemtide. And negotiations for licensing next-generation regenerative induction pharmaceutical candidates were conducted at multiple conferences with domestic and international pharmaceutical companies.

Cash Outflow for the Fiscal Year Ended July 2026

Cash Outflow for R&D	1,349 million yen
Cash Outflow for General and Administrative Expenses	260 million yen

Research and Development Progress Forecast for the Fiscal Year Ending July 2027

Most business revenue relies on development milestones, which are highly dependent on the development strategy and schedule of partners, making it difficult to predict the timing of receipts. Additionally, there is a possibility of receiving upfront payments from new partnerships, but the timing of contract finalization is uncertain. For these reasons, it is challenging to reasonably forecast earnings for the fiscal year ending July 2027, so earnings projections are not disclosed.

Research Progress Outlook for the Fiscal Year Ending July 2027

Research and development for Redasemtide is progressing in clinical trials and toward expanded indications. Additionally, negotiations for clinical trials and licensing for candidate "Regeneration-Inducing Medicine™" following Redasemtide continue to advance.

Cash Outflow Forecast for the Fiscal Year Ending July 2027

Cash Outflow for R&D	1,300 million to 1,700 million yen
Cash Outflow for General and Administrative Expenses	230 million to 310 million yen

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Allocation Status of Raised Funds

(Amount: million yen)

Funding Method	Amount Raised	Use of Funds	Planned Disbursement Amount(Timing)	Actual Disbursement Amount(Timing)	Undisbursed Amount(Timing)
Stock issuance associated with new listing (August 2019)	8,625	Funds for the establishment and maintenance of the "Regeneration-Inducing MedicineTM" research institute and an animal testing facility.	3,050	2,350 [Breakdown] Establishment Costs 940 Maintenance Costs 1,410	803
			(FY 2020 to FY 2030)	(FY 2020 to FY 2026)	(FY 2027 to FY 2030)
		Costs for promoting R&D for the existing pipeline and R&D for new pipeline development.	5,471	4,135 [Breakdown] Outsourcing Costs 724 Research Material Costs 1,296 Collaborative Research Costs 2,115	1,336
			(FY 2020 to FY 2029)	(FY 2020 to FY 2026)	(FY 2027 to FY 2029)

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6. Risk Information

Risk Information

Risk		Possibility and Timing of Materialization	Countermeasures
A) Uncertainty in the development of the pharmaceutical pipeline and associated revenue generation	・Risk of development delays or discontinuation due to the inability to discover beneficial effects in clinical trials. ・If the commercialization of our developed pharmaceutical candidates or those licensed out to other companies is delayed or canceled, there is a significant risk of adversely impacting our performance and financial condition.	Medium / As needed	・We aim to maintain multiple pipelines in clinical development stages. ・We will advance non-clinical pipeline projects swiftly to the clinical development stage.
B) Business plan reliant on specific partnership agreements	・Risk of dependence on limited joint research and licensing agreements with specific pharmaceutical companies. ・Risk of early termination of contracts before the expiration date due to factors beyond our control, such as significant deterioration in the business environment or changes in the management policies of the licensing partner company.	Medium / As needed	・We aim to generate revenue from subsequent pipelines to reduce dependence on income from current partnership agreements. ・We will minimize the impact on our business plan by establishing new partnerships with other pharmaceutical companies.
C) Cash Flow Management	・Research and development-oriented companies like ours require significant R&D funding, and due to the burden of R&D costs, there is a prolonged period of upfront investment. During this period, there is a risk of incurring continuous operating losses and negative cash flow from operations. ・Currently, our company lacks stable revenue sources, such as ongoing royalty income, and future revenue generation heavily depends on the progress of Redasemtide development and the outcomes of licensing negotiations for other pipeline projects. Therefore, we are not yet in a position to consistently generate funds from operating activities.	Low / As needed	・We aim to secure necessary funds by obtaining upfront payments and milestone revenue from existing pipelines, along with planned fundraising activities.

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Appendix

Management Team



Representative Director, Chairman & CSO

Katsuto Tamai

Representative Director, Chairman（Sep. 2026－Present）
Director, StemRIM Inc. (Oct. 2022)
Guest Professor, Endowed course of Regeneration-Inducing Medicine Graduate School of Medicine/ Faculty of Medicine,
The University of Osaka (Oct. 2023 – Present)
Professor, Endowed course of Regeneration-Inducing Medicine Graduate School of Medicine/ Faculty of Medicine,
The University of Osaka (Oct. 2010 – Sep. 2023)
Director, StemRIM Inc. (Feb. 2007 – Aug. 2010)
Associate professor, Department of Gene Therapy, Graduate School of Medicine/ Faculty of Medicine,The University of Osaka (May 2003 – Sep. 2009)



Representative Director, President & CEO

Masatsune Okajima

President and CEO, StemRIM Inc. (Oct. 2023 – Present)
President, StemRIM Inc. (March 2019 – Oct. 2023)
Vice President, Medicinova Inc. (Sep. 2006 – March 2019)
Deputy General Manager, Daiwa Securities SMBC Co., Ltd.(April 2002 – Aug. 2006)
Manager, Daiwa Securities SB Capital Markets Co., Ltd. (currently Daiwa Securities SMBC Co., Ltd.) (April 1999 – March 2002)
Sumitomo Capital Securities Co., Ltd. (Oct. 1996 – April 1999)
Sumitomo Bank, Ltd. (currently Mitsui Sumitomo Bank) (April 1991 – Oct. 1996)

Audit & Supervisory Board Member

Kazuyoshi Okuhira

Audit & Supervisory Board Member

Akihiro Mizukami

Audit & Supervisory Board Member

Yoichiro Shimada



External director

Noriko Sawai

Acting General Manager, New Business Creation Office, Business Development Department, NTT Precision Medicine Co., Ltd., (Feb.2025 – Present)
Social Innovation and Investment Foundation
(Feb. 2020 – July 2022)
External director, StemRIM Inc. (Oct. 2019 – Present)
DeNA Co. (June 2014 – Jan. 2020)
CSK Venture Capital Co. (April 1995 – May 2014)



External director

Hirotada Nagai

President, HyakusanSoken KK (July 2022 - Present)
External directors, StemRIM Inc. (Oct. 2020 - Present)
Auditor, Regional Fish Institute, Ltd.
(May 2020 – Present)
Director, PRDM Co., Ltd. (March 2018 – Present)
Director, PorMedTec Co., Ltd. (Dec. 2017 – Present)
Director, Kyoya KK (Dec. 2017 - Present)
Pharmaceuticals and Medical Devices Agency (PMDA)
(Sep. 2012 – July 2014)
Pharmaceutical and Food Safety Bureau of Ministry of Health, Labour and Welfare (April 2001 – Sep. 2017)



External director

Masaru Iida

External directors, StemRIM Inc. (Oct. 2026 - Present)
Exawizards Inc External Audit & Supervisory Board Member（Jun, 2019 – Present）
Ichigo Hotel REIT Investment Corporation Supervisory Officer（July, 2015 – Present）
Iida Management Law Office Representative Attorney（July, 2011 – Present）
DeNA Co., Ltd External Audit & Supervisory Board Member
Masuda & Partners Law Office（Jan, 2010 – July, 2011）
Sumitomo Bank, Ltd. (currently Mitsui Sumitomo Bank) (April, 1989 – Jan. 2010)

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Corporate Information

Corporate Name	StemRIM Inc.
Chief Executives	Masatsune Okajima (Representative Director)
Established	October 30, 2006
Business Description	Research and Development of “Regeneration Inducing-Medicine™”
Shareholders’ Equity	4,161 million yen
Equity Ratio	70.3 %
Number of Employees	70

Head Office
7-7-15, Saito-Asagi, Ibaraki-City,
Osaka, Japan



StemRIM Institute of Regeneration-Inducing Medicine, Osaka University
Techno-Alliance Building, 2-8,
Yamadaoka, Suita-City, Osaka, Japa



Endowed Chair for Regeneration-Inducing Medicine／Joint Research Course in Stem Cell and Gene Therapy
The Center of Medical Innovation and
Translational Research, 2-2, Yamadaoka,
Suita-City, Osaka, Japan



As of the End of July 2025

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