BIONEER

Investor Presentation

KOSDAQ:064550



Forward-looking statements

This presentation includes forward-looking statements. The forward-looking statements in this presentation include projections and outlook of the Bioneer Corporation (the "Company" or "Bioneer") concerning its business status and financial results, and include but are not limited to words, such as 'expectation', 'forecast', 'plan', 'anticipation' or '(E)'. The forward-looking statements are subject to changes in business environment and involve inherent risks and uncertainties.

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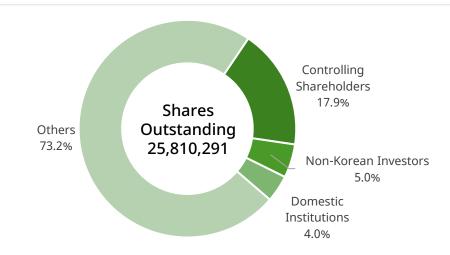
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BIONEER Corporation

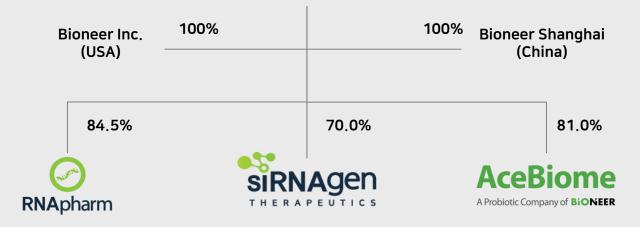
Shareholder Structure (as of 22.9.30)

CEO	Han-Oh Park
Established on	Aug. 28 th , 1992 (Listed on KOSDAQ : Dec. 29 th , 2005)
No. of Employees	619 (as of 2022.12)
No. of R&D Employees	225 (as of 2022.12)



BIONEER Innovation • Value • Discovery

BIONEER Family



Core Competencies – What Makes Us Different?

Irreplaceable cash cow businesses based on our own technologies

Cosmeceuticals

Anti Hair Loss Cosmetics



CosmeRNA

- v siRNA-based Hair Loss Cosmetics
- Received Safety Report and Efficacy Study
 Report from Dermatest
- A Game Changer that replaces all the competitions which come with adverse reactions

Probiotics



BNR17

- Achieved semi-annual sales of 109 Million USD and continued to expand market share
- South Korea's first functional raw material for reducing body fat approved by the Ministry of Food and Drug Safety (Korea) (individually approved type)
- More than 820 citations in BNR17® international journal articles and SCIE-level papers



Anti Hair Loss Cosmeceutical: CosmeRNA

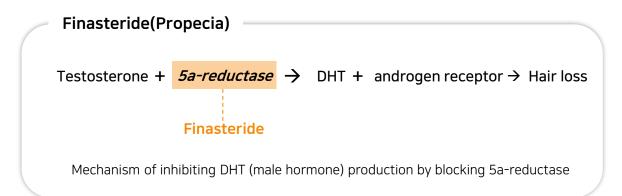
Cosmeceutical + RNA

The main causes of androgenetic alopecia

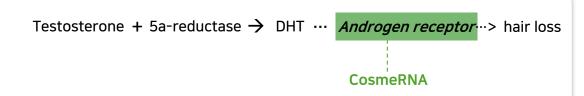
- √ 5a-reductase An enzyme in the body that combines with testosterone to produce DHT
- ✓ DHT (dihydrotestosterone), the male hormone that causes hair loss
- ✓ A protein that binds to the androgen receptor DHT to produce a substance that induces hair follicle cell suicide

Finasteride CosmeRNA Hair root Hair follicle cells cell suicide inducer 모근 세포 모근 세포 자살 유도 물질 CosmeRNA 안드로겐 수용체 생성 억제 Inhibition 5a-환원효소 Finasteride 5a-환원효소 androgen testosterone testosterone receptor production reductase reductase blocking blocking

Mechanism







Mechanisms that inhibit androgen receptor production without direct hormonal effects



Dermatest's Safety Report





Study number: 2207259870

Safety Report

Product/Project CosmeRNA ARI

Formula: -
CPNP-Notification-No.: None

Product type: Serum

Manufacturer: Bioneer Co.

Bottler: Distributor:

Version: 1

Cosmetic Safety Report (CPSR, Cosmetic Product Safety Report)

Part A. Cosmetics Safety Report (Bioneer Submission)

- Quantitative/qualitative compositional information
- Physical and chemical properties
- microbiological testing
- information on residual impurities
- foreseeable expected usage
- cosmetic exposure information
- substance exposure/toxicity information
- · information on serious side effects

Part B. Cosmetic Safety Assessment (Evaluator)

- Safety Assessment
- User Instructions
- Warnings and precautions from the evaluator
- Approval status from the safety assessor (name, qualifications, signature, etc.)

CosmeRNA Safety Report

Evaluator: Dermatest

- The review was conducted in compliance with the European Cosmetics Regulation (EC)
 1223/2009 and its guidelines
- The safety assessment concluded that the main ingredient and each individual ingredient have been reviewed and deemed safe
- All three safety assessors signed (Unconditional approval)

Confirmed that our product will be launched, as we have secured a safety report which is the most critical requirement for registration on the European Cosmetic Product Notification

Portal (CPNP).



Dermatest Efficacy Evaluation Report





Sponsor

Study Number 2106259429

BIONEER CORPORATION 8-11, Munpyeongse-ro, Daedeok-gu, Daejeon 34302 REPUBLIC OF KOREA

Muenster, December 16th 2022

Expert report by dermatological specialists about a randomised double-blind placebo controlled clinical-dermatological application study

on 120 subjects with application once per week resp. once per 2 weeks resp. once per 4 weeks on scalp over a period of 6 months

> Examination of dermal tolerability Quantification and differentiation of hair Quantification of hair loss

> > CosmeRNA ARI product A, product B

CosmeRNA efficacy evaluation report and photos of participants





Category	Dermatest (n=120)	Domestic human application test (n=60)			
# of Participants	120 persons	60 persons			
Female Participants	40 Females	24 Females			
Evaluation Methods	Once per week, Once per 2 weeks	Once per week			
Efficacy (per 1 cm²)	Once a week, once every 2 weeks in the test group (Data came out better than domestic tests)	Response Rate: 91%After 4 months: +7.6 (+1.9/month)			
Thesis Published	Preparing the paper to be published in an international journal such as Nature	Scientific Report (22.01) Weekly treatment with SAMIRNA targeting the androgen receptor ameliorates andregenetic alopecia			
Adverse Reaction	None	None			



Technologies that are protected by patents

SAMiRNA™ technology validated successfully through publications international academic journals and patent portfolio

CosmeRNA

Our Study has been published in Scientific Reports in January 2022

scientific reports

Weekly treatment with SAMiRNA targeting the androgen receptor ameliorates androgenetic alopecia

Sung-Il Yun^{1,5}, Sang-Kyu Lee^{2,5}, Eun-Ah Goh², Oh Seung Kwon², Woorim Choi², Jangseon Kim², Mi Sun Lee², Soon Ja Choi², Seung Sik Lim¹, Tae Kee Moon³, Sin Hae Kim³, Keeyeol Kyong⁴, Gaewon Nam^{4™} & Han-Oh Park^{1,2™}

Androgenetic alopecia (AGA) is the most common type of hair loss in men and women. Dihydrotestosterone (DHT) and androgen receptor (AR) levels are increased in patients with AGA. and DHT-AR signaling correlates strongly with AGA pathogenesis. In this study, treatment with self-assembled micelle inhibitory RNA (SAMiRNA) nanoparticle-type siRNA selectively suppressed AR expression in vitro. Clinical studies with application of SAMiRNA to the scalp and massaging to deliver it to the hair follicle confirmed its efficacy in AGA. For identification of a potent SAMiRNA for AR silencing, 547 SAMiRNA candidates were synthesized and screened. SAMiRNA-AR68 (AR68) was the most potent and could be efficiently delivered to human follicle dermal papilla cells (HFDPCs) and hair follicles, and this treatment decreased the AR mRNA and protein levels. We confirmed that $10\,\mu\text{M}$ AR68 elicits no innate immune response in human PBMCs and no cytotoxicity up to $20\,\mu\text{M}$ with HFDP and HaCaT cells. Clinical studies were performed in a randomized and double-blind manner with two different doses and frequencies. In the low-dose (0.5 mg/ml) clinical study, AR68 was applied three times per week for 24 weeks, and through quantitative analysis using a phototrichogram, we confirmed increases in total hair counts. In the 24-week long high-dose (5 mg/ml) clinical study, AR68 showed average additional hair growth of 1.3-1.9 hairs/cm² per month, which is comparable to finasteride. No side effects were observed. Therefore, SAMiRNA targeting AR mRNA is a potential novel topical treatment for AGA.

✓ Publication of a clinical trial paper on human application of CosmeRNA in Scientific Reports, demonstrating our technological capabilities.

The research paper has been published in multiple SCIindexed academic journals.

Scientific Reports, International Journal of Toxicology, Drug and Chemical Toxicologyl, etc.

The SAMiRNA™ platform is protected by more than 150 patents both domestically and internationally

CosmeRNA has filed and registered patents related to its technology in 12 countries, including Korea, the United States, and Europe



CosmeRNA safety and efficacy data

CosmeRNA cosmetics clinical trial data

4	WEEK 0	Category	CosmeRNA (n=60)	Best in class (n=3177)
Placebo CosmeRNA		Active Principle	CosmeRNA	Finasteride (Propecia)
		Classification	Cosmetics	Prescription Drug
		Drug Administration Route	Topical	Oral
	WEEK 0.4	Target	Androgen Receptor	5α-Reductase
Placebo CosmeRNA	WEEK 24	Battery Duty Cy	Once per week	Daily Oral Administration
		Efficacy (per 1 cm²)	Response Rate: 91% Efficacy after 4 months: +7.6 hair (+1.9 hair/month)	Response Rate: 87.1% Efficacy after 6 months: +9.3 hair (+1.6 hair/month)
		Adverse Reactions	None	ED, Decreased of sexual desire, etc

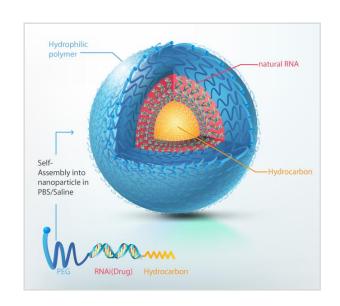
No drug-related adverse reactions were identified in the human application test (n=60) and safety test (n=35) for domestic functional cosmetic registration procedures.

Yun, Sl., Lee, SK., Goh, EA. et al. Weekly treatment with SAMiRNA targeting the androgen receptor ameliorates androgenetic alopecia. Sci Rep 12, 1607 (2022). https://doi.org/10.1038/s41598-022-05544-w



SAMiRNA™ (Self-Assembled-Micelle inhibitory RNA)





The world's only native siRNA/miRNA single-molecule nanoparticle RNAi drug platform

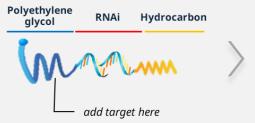
A modular platform that doublelinks RNAi sequences with hydrophilic and hydrophobic materials

Wide range of RNAi sequences applicable to the platform, ensuring universality as a drug

With a particle size (60-100nm) and charge optimized for absorption of various tissues in the body, it is specifically absorbed through the EPR effect.

Absorbed selectively to target inflammatory/cancer tissues

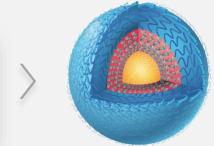
SAMiRNA™ platform formation



Double bonds of PEG (hydrophilic) and Hydrocarbon (hydrophobic) substances to both ends of RNAi



RNAi double binding process



Self-assembled nanoparticles in a spherical shape based on double bond modules_

SAMIRNA-AREG preclinical thesis





Thesis demonstrating the efficacy and safety of SAMiRNA-AREG material in suppressing Amphiregulin (AREG) expression



SAMIRNA

www.nature.com/scientificreports

scientific reports



OPEN In vivo silencing of amphiregulin by a novel effective Self-Assembled-Micelle inhibitory RNA ameliorates renal fibrosis via inhibition of EGFR signals

Seung Seob Son¹,7, Soohyun Hwang¹,7, Jun Hong Park¹, Youngho Ko¹, Sung-Il Yun², Ji-Hye Lee³, Beomseok Son¹, Tae Rim Kim¹, Han-Oh Park¹,2²³³ & Eun Young Lee^{4,5,633}

Amphiregulin (AREG) is a transmembrane glycoprotein recently implicated in kidney fibrosis Previously, we reported that the AREG-targeting Self-Assembled-Micelle inhibitory RNA (SAMIRNA-AREG) alleviated fibrosis by stably silencing the AREG gene, and reduced the side effects of conventional siRNA treatment of pulmonary fibrosis. However, the therapeutic effect of SAMiRNA AREG in renal fibrosis has not been studied until now. We used two animal models of renal fibrosis generated by a unilateral ureteral obstruction (UUO) and an adenine diet (AD) to investigate whethe SAMIRNA-AREG inhibited renal fibrosis. To investigate the delivery of SAMIRNA-AREG to the kidney, Cy5-labeled SAMiRNA-AREG was injected into UUO- and AD-induced renal fibrosis model In both kidney disease models, SAMiRNA-AREG was delivered primarily to the damaged kidney. We also confirmed the protective effect of SAMIRNA-AREG in renal fibrosis models. SAMIRNA-AREG markedly decreased the UUO- and AD-induced AREG mRNA expression. Furthermore, the mRNA expression of fibrosis markers, including α -smooth muscle actin, fibronectin, $\alpha 1(1)$ collagen and α1(III) collagen in the UUO and AD-induced kidneys, was diminished in the SAMiRNA-AREG treated mice. The transcription of inflammatory markers (tumor necrosis factor- α and monocyte chemoattractant protein-1) and adhesion markers (vascular cell adhesion molecule 1 and intercellul adhesion molecule 1) was attenuated. The hematoxylin and eosin, Masson's trichrome, and immunohistochemical staining results showed that SAMIRNA-AREG decreased renal fibrosis AREG expression, and epidermal growth factor receptor (EGFR) phosphorylation in the UUO- and AD-induced models. Moreover, we studied the effects of SAMiRNA-AREG in response to TGF-β1 in mouse and human proximal tubule cells, and mouse fibroblasts. TGF-β1-induced extracellular matrix production and myofibroblast differentiation were attenuated by SAMiRNA-AREG. Finally, we confirmed that upregulated AREG in the UUO or AD models was mainly localized in the distal tubules. In conclusion, SAMIRNA-AREG represents a novel siRNA therapeutic for renal fibrosis by suppressing Published a paper proving the deliverability and stability of the SAMiRNA platform

Proved that our platform overcomes the limitations of existing RNAi-based treatments

(Proven advantages over existing siRNA for pulmonary fibrosis treatment in terms of delivery and side-effects)

✓ Proved the mechanism, stability, and efficacy of the platform by publishing in Scientific Reports, a sister journal of Nature

Through animal experiments, SAMiRNA-AREG confirmed the effect of reducing fibrosis and inflammation indicators delivered to the kidneys

Demonstrated anti-fibrotic and anti-inflammatory effects by reducing phosphorylation of epidermal growth factor receptor (EGFR)



siRNA Challenges

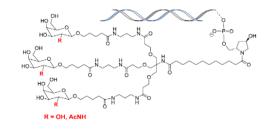


siRNA challenges are conjugation and encapsulation methods, which come with significant downsides

Naked (Conjugated) RNAi

SAMIRNATM

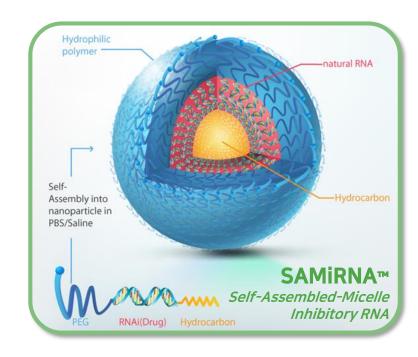
Nanoparticle Encapsulation







GalNAc-siRNA conjugates, Cholesterol, RGD, DPC2.0, PNP



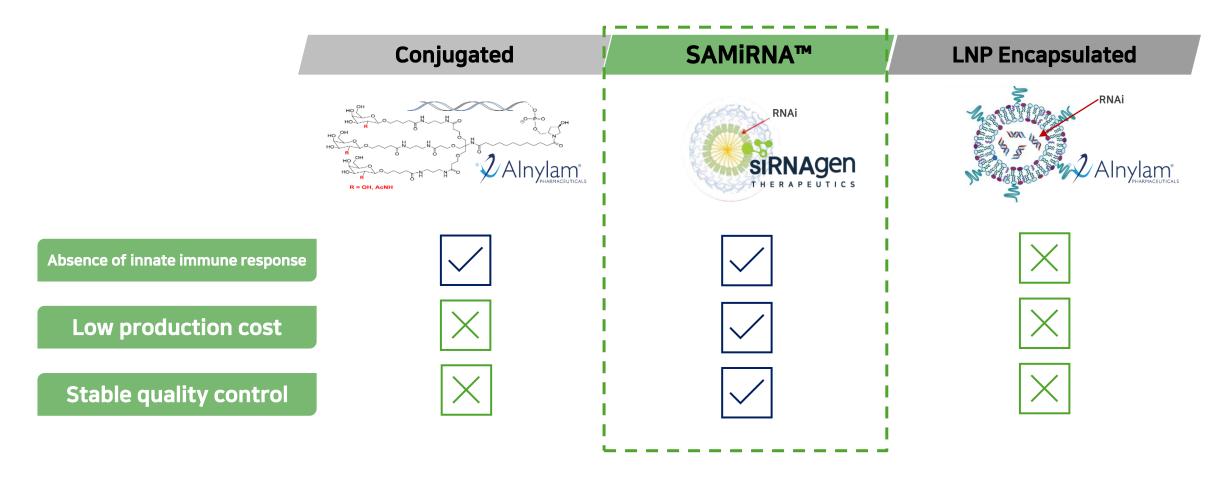


Dlin-DMA, Dlin-MC3-DMA Lipid Nanoparticles (LNPs), Inorganic nanoparticles, Exosomes





SAMiRNA™ complements the unmet needs of Naked Oligo and LNP methods

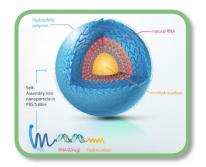


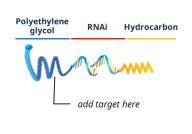
Approval of SRN-001 phase 1 clinical trial for pulmonary fibrosis treatment using SAMiRNA™



IPF (Idiopathic Pulmonary Fibrosis) Phase 1 clinical trial plan (CTN) in Australia

SAMiRNA™





Administration method and mechanism of action

- ✓ Designed for delivery to inflammatory, fibrotic and solid tumors with the realization of EPR effect* by intravenous (IV) administration
- ✓ This makes it possible to target organs regardless of organs so that a single injection can reach various organs and cells.

EPR (Enhanced Permeability and Retention) Effect: Appears in incomplete blood vessels around cancer and inflammatory tissues, and can penetrate the body of high-molecular drugs

Australian HREC Phase 1 Clinical Trial Plan (CTN) Application Details

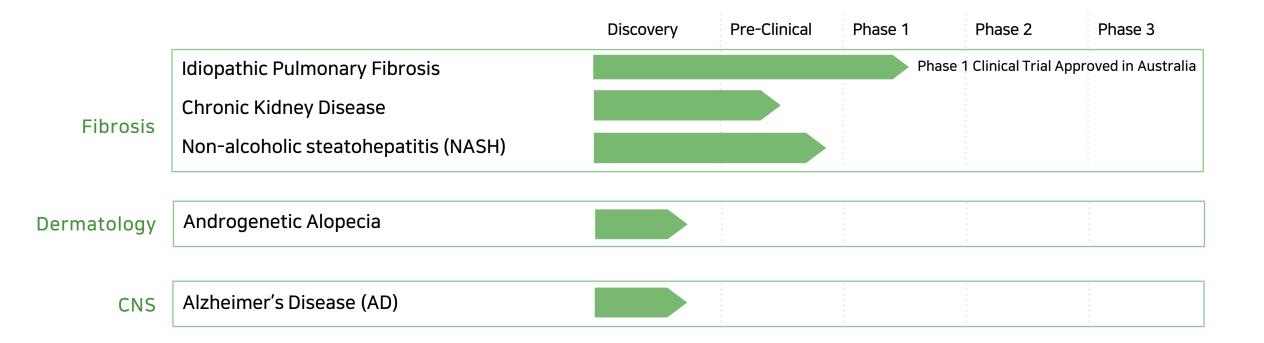
Clinical Trial Name	A randomized, double-blind, placebo-controlled, single-dose escalation clinical trial to evaluate the safety, tolerability and pharmacokinetics of SRN-001 in healthy persons
Approval Institution	Australia HREC (Human Research Ethics Committee)
Indication	Idiopathic Pulmonary Fibrosis
Purpose of the Clinical Trial	Evaluation of safety, tolerability, and pharmacokinetic characteristics of step-by-step, single intravenous administration of SRN-001 in healthy adults
Methods	Randomization, double-blind, placebo-controlled, escalated dose, single dose
Period	From the clinical trial plan approval date to December 26, 2023



SAMiRNA™ Pipelines



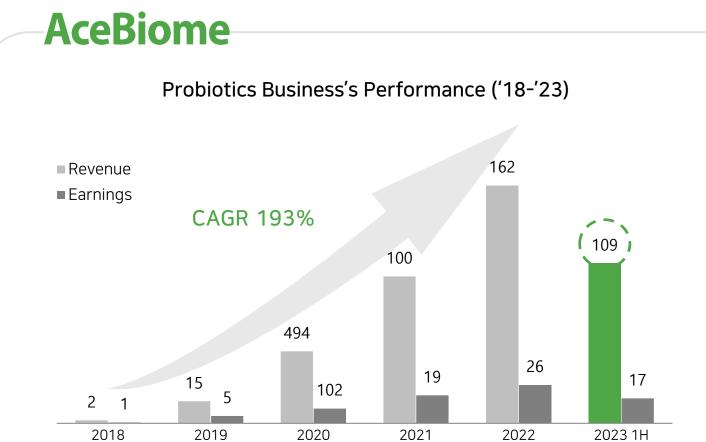
Pipeline Research Status by Indication



Sales growth based on proven product quality

160 Million US in sales in 5 years with only 30 employees, record-high sales in the 2nd quarter of 2023

Unit: Million \$ US



✓ Record-high Sales in first Half of 2023 (109 Million \$)

Recorded sales of 109 Million in the first half of this year

Expected to exceed 200 Million in annual sales by 2023

Achieved sales in the first half of the year that are higher than annual sales in 2021 in just two years

AceBiome always verifies with thesis





Experience in supplying raw materials to global big pharma with body fat reduction efficacy proven in thesis

AceBiome



J. Microbiol. Biotechnol. 2021. 31(9): 1281–1287 https://doi.org/10.4014/jmb.2105.05032

The Effect of Lactobacillus gasseri BNR17 on Postmenopausal Symptoms in Ovariectomized Rats

Sol Lee 1.2, Dong Hoon Jung 1.2, Miri Park 1.2, Seung-Woo Yeon 1.2, Sang-Hyuk Jung 3, Sung-Il Yun 3, Han-Oh Park 1.2,3.4, and Wonbeak Yoo 1.2*

AceBiome Inc., Seoul 06164, Republic of Korea

²R&D Center, AceBiome Inc., Daejeon 34013, Republic of Korea

Clinical and preclinical studies have reported that Lactobacillus gasseri BNR17, a probiotic bacterial strain isolated from human breast milk, reduces body weight and white adipose tissue volume. In order to further explore the actions of L. gasseri BNR17, we investigated the anti-menopausal effects of L. gasseri BNR17 in an ovariectomized (OVX) rat model. The serum alanine aminotransferase levels of the rats in the OVX-BNR17 group were lower than those of the rats in the OVX-vehicle only (OVX-Veh) group. Upon administration of L. gasseri BNR17 after ovariectomy, calcitonin and Serotonin 2A levels increased significantly, whereas serum osteocalcin levels showed a decreasing tendency. Compared to the rats in the OVX-Vehy group, those in the OVX-BNR17 group showed lower urine deoxypyridinoline levels, lower pain sensitivity, and improved vaginal cornification. Furthermore, L. gasseri BNR17 administration increased bone mineral density in the rats with OVX-induced femoral bone loss. These results suggest that L. gasseri BNR17 administration could alleviate menopausal symptoms, indicating that this bacterium could be a good functional probiotic for managing the health of older women.

Keywords: Lactobacillus gasseri BNR17, menopause symptoms, ovariectomized rats

✓ The excellent efficacy and strains of our original technology-based products are protected by patents

The first raw material in Korea to obtain individual recognition certification from the Ministry of Food and Drug Safety (Lactobacillus gasseri BNR17®)

Patents registered in 10 countries including the US, Europe, China, Japan, and Korea Trademark registration completed in over 20 countries

 Efficacy proven worldwide by publication of a paper in a SCIE-level academic journal

Cited more than 820 times in papers published in BNR17® international academic journals and SCIE-level papers

The Journal of Microbiology (JM), Food Science and Biotechnology, etc



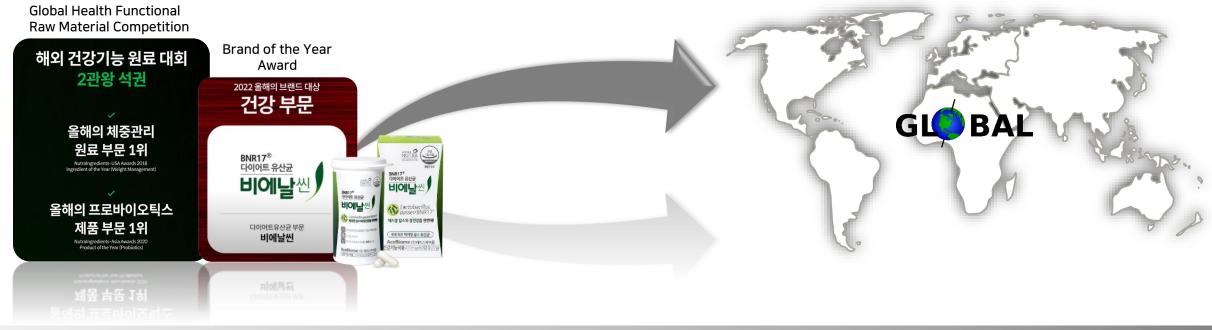
³siRNAgen Therapeutics, Daejeon 34302, Republic of Korea ⁴Bioneer Corporation, Daejeon 34302, Republic of Korea

AceBiome's expansion to the global markets





Plan to enter the global markets with its brand power proved in domestic markets



Probiotics that reduces body fat BNR Series

* Expanding target customer base through product line diversification

Joint Health Supplements Annaparactin (ParActin)

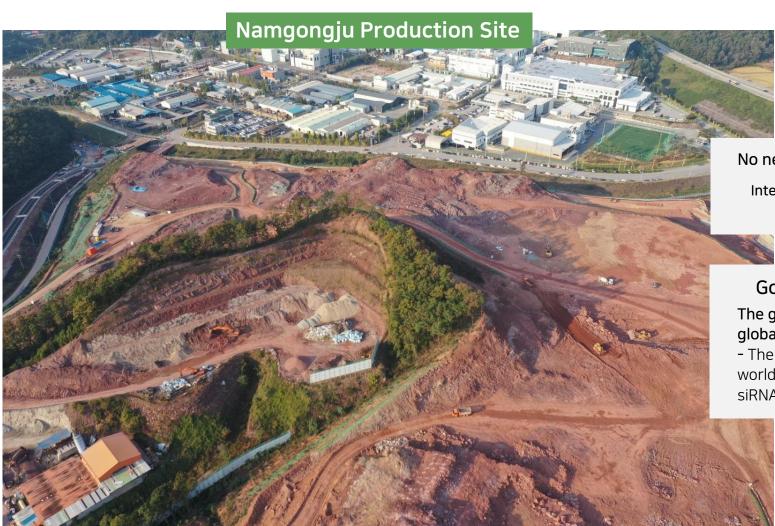






Core Competencies - Our Capability to Have Super-Low Production Cost

Mass production and ultra-low cost ratios possible through in-house production



No need for import due to ownership of the entire value chain

Internalization of the value chain from oligonucleotide production, synthesis, material and raw material production

Goal to become the world's No.1 siRNA producer The goal is to produce 10 tons of siRNA, exceeding the current global production of 6 tons.

- The goal is to secure 640497.15 Sq.ft of land and build the world's largest siRNA production facility, producing 10 cans of siRNA, exceeding 6 tons of global production.

CosmeRNA® raw materials' production system and current production capacity

Laid the foundation for mass production by developing a large-capacity oligo synthesizer and successfully producing CosmeRNA® using it

CosmeRNA® Production Synthesizer



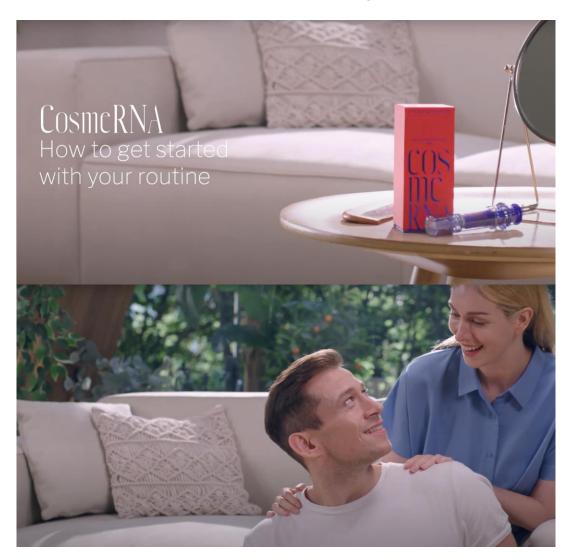
Current Production Capacity

Monthly Production Scale of siRNA	2kg/m ~3kg/m
In Annual Conversion	24kg ~ 36kg
When Converted to mg	24,000,000mg ~ 32,000,000mg
siRNA Required per 1 Unit	30mg
Units	800,000 ~ 1,200,000
Maximum Possible Revenue	320M \$ ~ 480M \$ (US) (300 EUR per unit)



CosmeRNA® Marketing Strategy

Plan to enter the European market through Amazon and partnership with CosmeRNA® Business to Business (B2B) partners





CosmeRNA® flagship website covers domestic markets

Covers the B2C market across Europe with Amazon



Now available in the UK, Italy, Germany, France and Spain Plan to enter Amazon Japan, Singapore and Australia

Aim to enter major markets in the world through B2B

Plan to enter major markets through Business to Business (B2B)

- Negotiating and reviewing proposals with partner companies in various countries, weighing business-to-business (B2B) contracts in India and Japan for products of different formulations



Meaning of CosmeRNA Release

Plan to diversify product portfolio based on proven SAMiRNA™ technology



The first human application of our platform SAMiRNA™

World's first siRNA-based functional cosmetics for hair loss

siRNA-based hair loss relief product with no innate immune response (Three domestic human application tests and European human application tests conducted)

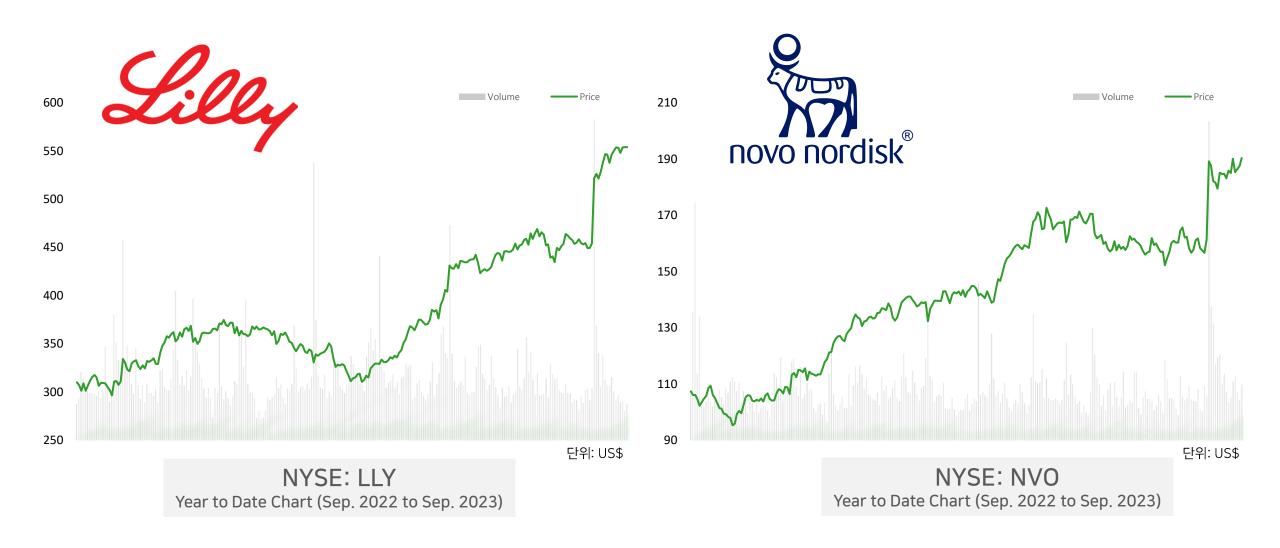
Proving the excellent stability of our platform (can be stored for 30 months at room temperature)

Plan to release various products based on a proven platform (Considering skin and beauty fields first)



Quality of Life Drug Comparable Companies

Quality of Life healthcare firms receive high valuations as they address the unmet needs in Healthcare



Major Catalyst - *Quality of Life* Company in Korea



Transforming from a gene synthesis company to a QOL (Quality of Life) healthcare company that contributes to improving the human quality of life





A Company that cares about improving the quality of life

Owns businesses closely related to improving the quality of human life, such as weight loss and hair loss

- Operating related businesses such as CosmeRNA and BNR17, and planning to continuously advance into these areas

A company that follows the paradigm of the global pharmaceutical/biotech industry

The investment paradigm has recently changed with the approval of weight loss medicine in the pharmaceutical/biotech industry

- As companies with the relevant businesses are gaining attention in Korea, Bioneer, which owns the QOL business, is also engaging public attention.

Bioneer is gaining attention along with companies engaged in related businesses in Korea

- As global pharmaceutical companies are gaining momentum recently with their entry into the obesity market and approval of new drugs related to obesity, Bioneer is recognized as a representative of the same momentum in Korea.



2023 2Q Financial Highlights

69.3 Billion KRW Revenue (QoQ +23.8% YoY 31.9%)

- v 8.6 Billion KRW Revenue recorded by Bioneer Corporation (QoQ 19.5% YoY -16.5%)
- AceBiome Inc, recorded quarterly sales of KRW 60.7 Billion
 (QoQ +24.4%, YoY +43.8%)
- Monthly average revenue recorded as 20 Billion KRW for Acebiome, and for the first half revenue recorded all-time high of 109.5 Billion KRW

5.1 Billion KRW Operating Profit (QoQ Turned to Profit, YoY 11834.9%)

- Bioneer continues operating loss -4.3 Billion KRW
 (QoQ 9.8%, YoY 11.4%)
- , Acebiome Inc, recorded quarterly operating profit 10 Billion (QoQ 27.1%, YoY 67.0%)
- Acebiome Operating Margin recorded 16.4%



Financial Summary (K-IFRS Consolidated)

Balance Sheet

[KRW 100 Million]

	[ratter ree ranner					
	FY'23 Q2	FY'22	FY'21			
Assets	1,449	1,341	1,364			
└ Cash Equ.	699	668	494			
Non-Current Assets	1,679	1,647	1,260			
└ Tangible Assets	1,533	1,515	1,135			
Total Assets	3,128	2,987	2,624			
Current Liabilities	424	508	351			
Non-Current Liabilities	204	40	81			
Total Liabilities	628	548	432			
Equity	129	129	126			
Retained Earnings	(122)	(281)	(523)			
Total Capital	2,499	2,436	2,076			

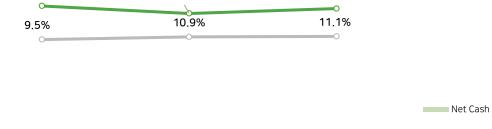
Net Cash & Financial Ratio

[KRW: 100 Million, %]

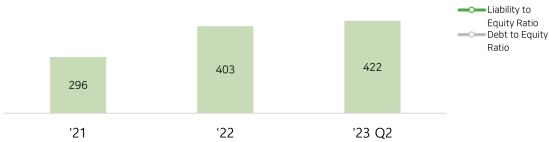
25.1%

*Net Cash = Cash Equivalents - Borrowings

26.4%



22.7%



Income Statement (K-IFRS Consolidated)

[KRW:	Mil	lion]
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	FY'22				FY'23		QoQ	YoY	
	Q1	Q2	Q3	Q4	SUM	Q1	Q2	QUQ	101
Revenue	61,455	52,560	54,591	49,789	218,394	56,030	69,349	23.8%	31.9%
└ Bioneer Corp.	25,055	10,304	10,938	9,878	56,175	7,198	8,604	19.5%	-16.5%
^L AceBiome	36,395	42,256	43,653	39,911	162,215	48,832	60,744	24.4%	43.8%
^L Others	5	-	-	-	5	5	-		
Gross Income	47,924	40,459	41,724	36,541	166,648	43,394	55,456	27.8%	37.1%
(%)	78.0%	77.0%	76.4%	73.4%	76.2%	77.4%	80.0%		
SG&A Expenses	37,525	40,416	37,555	39,613	155,109	43,497	50,323	15.7%	24.5%
Operating Income	10,399	43	4,169	-3,072	11,540	-103	5,132	Turned to Profit	11834.9%
(%)	16.9%	0.1%	7.6%	-6.2%	4.6%	-0.2%	7.4%		
└ Bioneer Corp.	6,869	-4,906	-5,737	-8,269	-12,043	-7,267	-4,345	40.2%	11.4%
^L Acebiome	4,065	6,030	10,817	5,748	26,660	7,925	10,073	27.1%	67.0%
└ Others	-535	-1,080	-911	-552	-3,078	-761	-594	21.9%	45.0%
Non-Operating Income	1,256	1,716	2,318	-3,924	1,366	2,038	1,249	-38.7%	-27.2%
Income Before Tax	11,655	1,791	6,487	-6,029	13,904	1,935	6,381	229.8%	256.3%
Net Profit	9,137	1,242	10,010	-5,267	15,122	400	3,476	1252.5%	335.6%
Net Margin (%)	14.9%	2.4%	18.3%	-11.0%	6.1%	0.0%	5.0%		



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