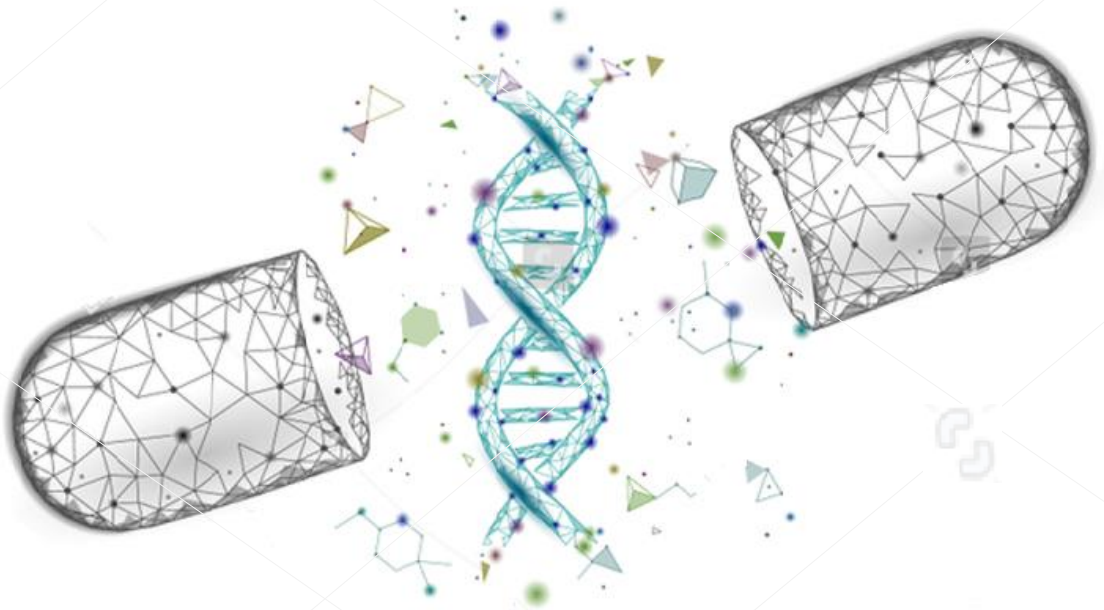
Saving Lives Through Innovation

Genexine

Corporate Overview May 2022



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Genexine: A Global Biopharmaceutical Company

"Focused on the Development of Innovative biologics and Saving the Lives of Patients."

Focused R&D	• Immuno-oncology • Orphan drugs
Proprietary platform technologies	• hyFc™ fusion technology • DNA vaccine technology
Advanced Therapeutic Candidates	• GX-I7 Long-acting Interleukin 7 • GX188E for Cervical Cancer and SCHNC • GX-H9 Long-acting Growth Hormone
Key milestones	• Expected first commercial launch 2023/24 • Key data releases in 2022
Location	• Head office: Seoul, Korea • Expansion to US expected in 2022
Employees	• Total 130 employees • R&D 60.7%
Market Cap	• USD $1.0 B (Q1 2022)

Key Executive Members

Professional leadership from discovery through clinical development, CMC and commercialization

CEO
Neil K. Warma

US GM, I-Mab Biopharma
CEO, Opexa Therapeutics
CEO, Viron Therapeutics
Founder, President, MedExact
Global Mktg, Policy, Novartis Pharma,
Univ. of Toronto HBSc, neuroscience
York Univ. MBA

COO/CEO
Jungwon Woo

SNU MS Pharmacology
Cornell Univ. PhD. Microbiology
Harvard Medical School post doc
Research Professor, Seoul St. Mary's Hospital

CFO
Sung-June Hong

Yonsei Univ. MBA
Vanderbilt Law School LLM
Daewoo Motors, Philip Morris, Nike Korea, Handok CFO
President of Rokit Healthcare

CMO
Jong-Sub Park

Professor, MD
Catholic Univ. College of Medicine
Catholic University
Director of Gynecological Cancer Center
Johns Hopkins
Exchange Professor

VP, Corp Dev
Hyun-Jin Park

SKKU Pharmacy
Aalto Univ. MBA
GSK Korea sales/marketing
Daewoong Pharma
Executive director, Development division

VP, Clinical Dev
Min-Kyu Heo

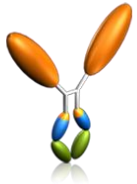
Yonsei Univ. MMed
JW Creagen Clinical Team
JW Pharma Clinical Team

VP, CMC
Ki-Yong Kim

Yonsei University
Ph.D. Biochemistry
NIH Postdoctoral Fellow
Mokam Biotechnology Res. Center
Green Cross Res. Inst.

Two Proprietary Platforms yield late-stage candidates

hyFc™ (Long-acting protein drugs)



Long-acting platform technology which can be applied to various APIs to maximize protein sustainability.

DNA vaccine (Cancer therapeutic/ Infectious disease)

Therapeutic + Preventive DNA vaccine platform with strong T cell immunity.

A **Clinical Stage** biotech company developing **First-in-Class & Best-in-Class** products derived from two **proprietary platforms**

Two on-going Phase 3 trials, targeting **multiple BLA submissions** in the next 2-4 years

Head office in **Seoul, Korea**, **expansion into U.S.** in next 12 months; ability to conduct **global drug development**

Access to multiple **novel drugs** and platforms through **extensive strategic partnerships**

Pipeline Overview: Multiple Clinical Assets, On-going Ph 3 trials

pipeline	indication	Clinical development phase			sponsor	collaborator
		Phase 1	Phase 2	Phase 3		
GX-E4 (long-acting EPO)	**CKD-induced Anemia**	**Phase 2 in KR**		**Phase 3 in Asia**	KG Bio	Genexine
GX-H9 (long-acting growth hormone)	**Growth Hormone Deficiency (**Pediatric)	**Phase 2 in KR/EU**		**Phase 3 in CN**	Handok, I-MAB	Genexine
	Growth Hormone Deficiency (Adult)	**Phase 2 in KR/EU**			Genexine, Handok	
GX-I7	**TNBC, GBM, High risk Skin cancer**	**Phase 1b/2 in KR/US/CN**			Genexine, NIT, I-Mab	MERCK, Roche, Bristol Myers Squibb™
	COVID-19 treatment	**Phase 2 in Indonesia**			KG Bio, Genexine	
GX-188E	**Cervical cancer**	**Phase 2 in KR**			Genexine	

Completed On-going

GX-I7 Long-acting Interleukin 7
(efineptakin alfa)

First-in-Class T-cell amplifier

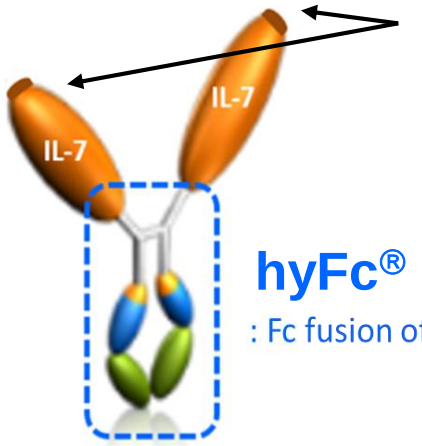
GX-I7: The Only Solution for Lymphopenia

- Only clinical-stage long-acting human IL-7 drug candidate
- No lymphopenia correction drugs have been approved, to date
- IL-7 fundamental cytokine for naïve and memory T-cell development and sustaining immune response
- Favorable PK/PD and safety profile, for combination with Checkpoint Inhibitors or CAR-T therapies.
- Multiple ongoing trials in solid tumors, and in planning for hematologic malignancies

Erythrocyte	Platelet	Neutrophil	Lymphocyte
EPO (Erythropoietin) Anemia	**TPO (Thrombopoietin) Thrombocytopenia**	**G-CSFs Neutropenia**	**GX-I7 (Efineptakin alfa) Lymphopenia**
Epogen: Amgen Mircera: Roche	Nplate: Amgen Promacta: Novartis	Neupogen: Amgen Neulasta: Amgen	GX-I7 (NT-I7 / TJ-107)
Global Market **$ 10.7bn** in 2025 from $ 7.3bn in 2018	**Global Market** **$3.3bn** in 2025 from $ 3.0bn in 2018	**Global Market** **$15.4bn** in 2025 from $ 11.3bn in 2018	

GX-I7: The First Long-acting Lymphopenia Correction Drug Candidate

Hybrid Fc-fused long-acting recombinant human IL-7 which plays an essential role in the development and homeostasis of T-cells



N-terminal engineering

hyFc®

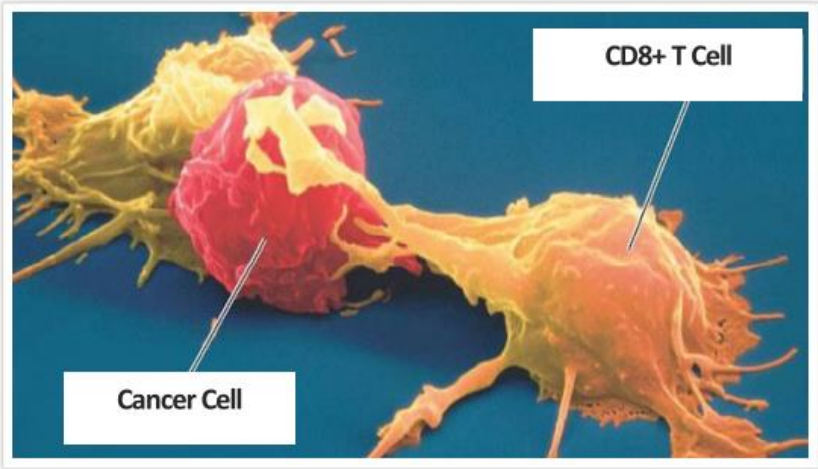
: Fc fusion of IgD + IgG4

No cytotoxicity on target cells
(No ADCC of IgD & No CDC of IgG4)

- **Higher protein stability**
- **Higher productivity**
- **Longer *in vivo* half-life**

T cell Amplifier

Long-acting Interleukin-7



T cell Activator

- Cancer vaccine
- IL-2, IL-15, IFN-alpha
- CD137 L, OX40 L, ICOSL
- TLR agonists, etc

T cell Suppressor blockade

- anti-PD-1, anti-PD-L1
- anti-CTLA4
- anti-TIM-3, anti-TIGIT
- anti-TGF-beta

GX-I7(NT-I7/TJ-107): Clinical Development

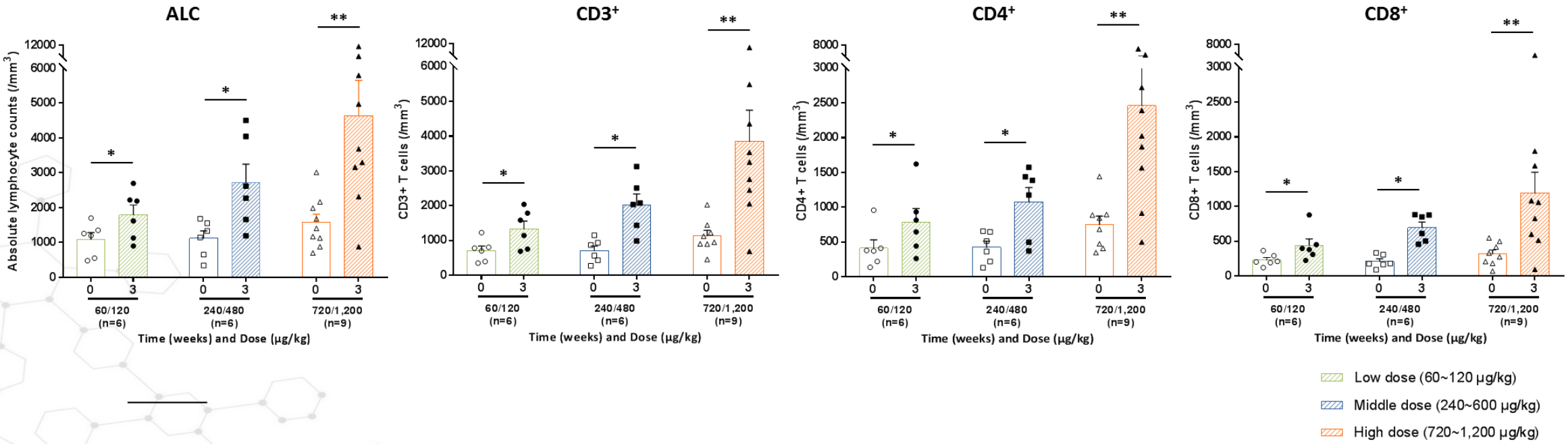
GX IMAB NIT

Field	Type	Treatment	Indication	Preclinical	Phase 1	Phase 2	Phase 3	Partner
Oncology	COM	KEYTRUDA®	TNBC		Phase 1b/2, KEYNOTE-899			MERCK, NeoImmuneTech
	COM	Avastin	Recurrent GBM		Phase 2			Roche
	COM	GX-188E/Opdivo	Recurrent HNSCC (HPV16)		Phase 2a			
	Mono	-	Solid Tumor		Phase 1/2a (STM101)			Genexine, NeoImmuneTech
	COM	Temozolomide	GBM		Phase 2 (GBM201)			
	COM	Pembrolizumab	TNBC, HNSCC		Phase 2 (STM202)			
	Mono	-	GBM (n=75)		Phase 1 (NIT-104, Single dose)			Johns Hopkins University
	COM	Temozolomide	GBM (n=46)		Phase 1/2 (NIT-107)			Washington University in St. Louis School of Medicine
	COM	Tecentriq®	High risk skin cancer		Phase 1b/2a (NIT-106)			Roche, cancer immunotherapy trials network
	COM	KEYTRUDA®	TNBC, Lung, Pancreatic, Colorectal cancer		Phase 1b/2a (NIT-110), KEYNOTE-A60			MERCK
	COM	Opdivo®	Gastric, GEJ, and Esophageal Adenocarcinomas		Phase 2 (NIT-109)			Bristol Myers Squibb
	COM	Tecentriq®	NSCLC, Non-Small Cell Lung Cancer		Phase 2 (NIT-119)			Roche
	Mono	-	Recurrent GBM		Phase 1 (NIT-115)			UCSF
	Mono	-	Kaposi's sarcoma		Phase 1 (NIT-108)			Fred Hutch Cures Start Here
	COM	Kymriah®	Diffuse large B-cell lymphoma		Phase 1b (NIT-112)			
Infectious Disease	Mono	-	Idiopathic $CD4^+$ T Lymphopenia	Under IND submission (NIT-114)				NIH National Institute of Allergy and Infectious Diseases
	COM	Vaccine	Preventative vaccine (Elderly cancer survivors)		Phase 1a/1b (NIT-105)			NIH National Cancer Institute
	Mono	standard	Covid19 treatment		Phase 1b			
	Mono	standard	Covid19 treatment		Phase 2			KGbio
	Mono	standard	Covid19 treatment		Phase 1 (NIT-116)			NIH National Institute of Allergy and Infectious Diseases
	Mono	-	Progressive polysomal leukoencephalitis		Phase 1 (NIT-113)			NIH National Institute of Allergy and Infectious Diseases

GX-I7: the only stable and long-acting IL-7 agent

- **GX-I7 amplifies T cell count which is key to the cancer immunotherapy**
- **GX-I7 injection to solid tumor patients, ALC, CD3+, CD4+, and CD8+ T cell all increased.**

(2019, Nov. SITC)



Source: Poster Presentation SITC 2019

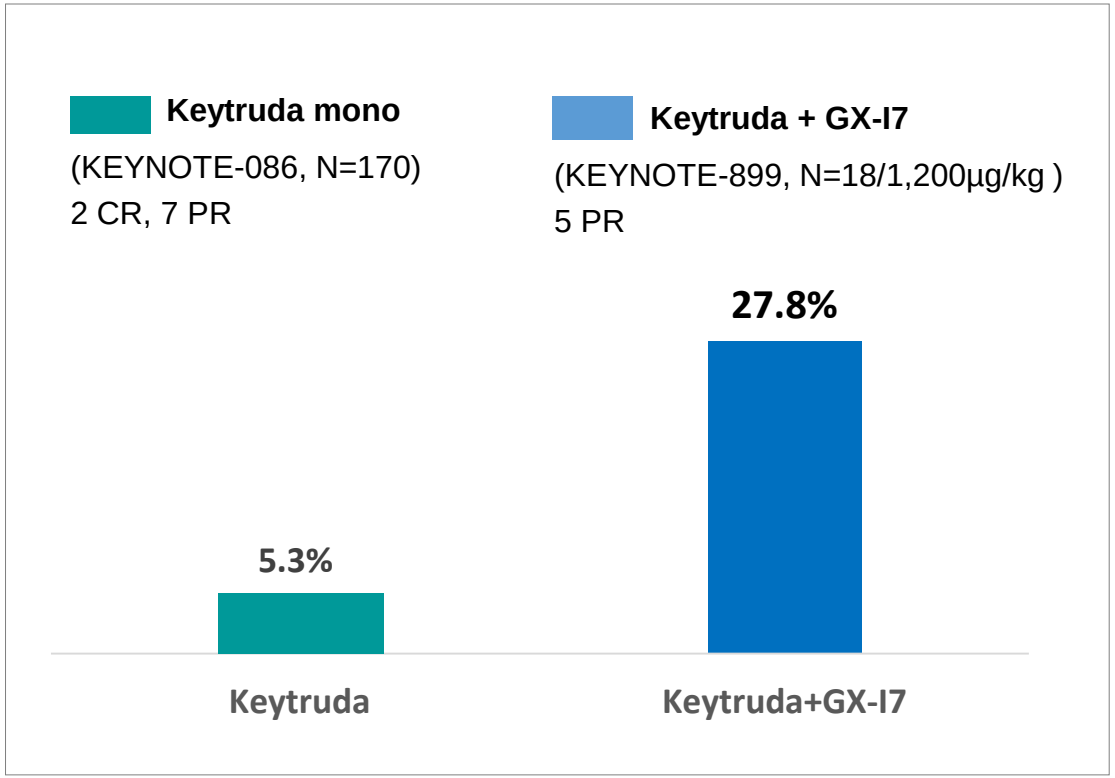
GX-I7 (NT-I7) : Promising Efficacy and Safety in mTNBC

- **Top-line results** of phase 1b/2 study of GX-I7 plus pembrolizumab in patients with R/R mTNBC to be presented at **ASCO June 2022**
  - Interim analysis(n=36) reported (KEYNOTE-899) at SITC in November 2020; 1,200ug/kg dose of GX-I7 combo with pembro showed ORR 27.8 %
- SITC 2021 Interim analysis: GBM Ph1b Combo with Chemo & colorectal cancer Ph2a Keytruda Combo. (NT-I7)

Ph2 Interim Result in mTNBC

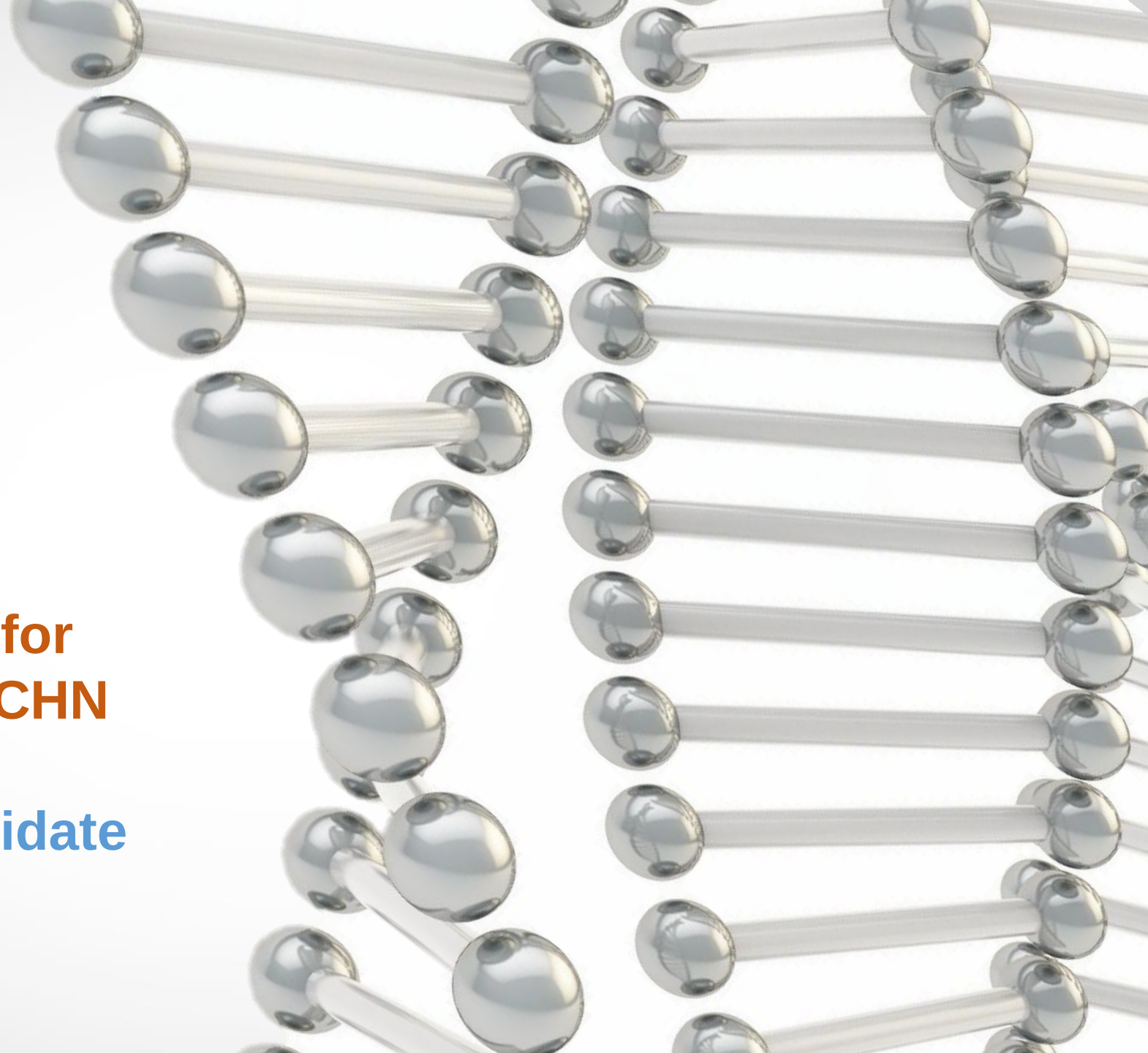
Response (RECIST v1.1) N (%)	GX-I7/Pembrolizumab Simultaneous Treatment (N=36)			
	720 µg/kg	960 µg/kg	**1,200 µg/kg**	1,440 µg/kg
ORR	**1 (16.7)**	**1 (16.7)**	**5 (27.8)**	**-**
CR	-	-	**-**	-
PR	1 (16.7)	1 (16.7)	5 (27.8)	-
SD	1 (16.7)	2 (33.3)	3 (16.7)	1 (16.7)
PD	4 (66.7)	3 (50.0)	10 (55.6)	5 (83.3)
DCR	**3 (50.0)**	**4 (66.7)**	**8 (44.4)**	**1 (16.7)**

Source: SITC 2020 Poster Presentation



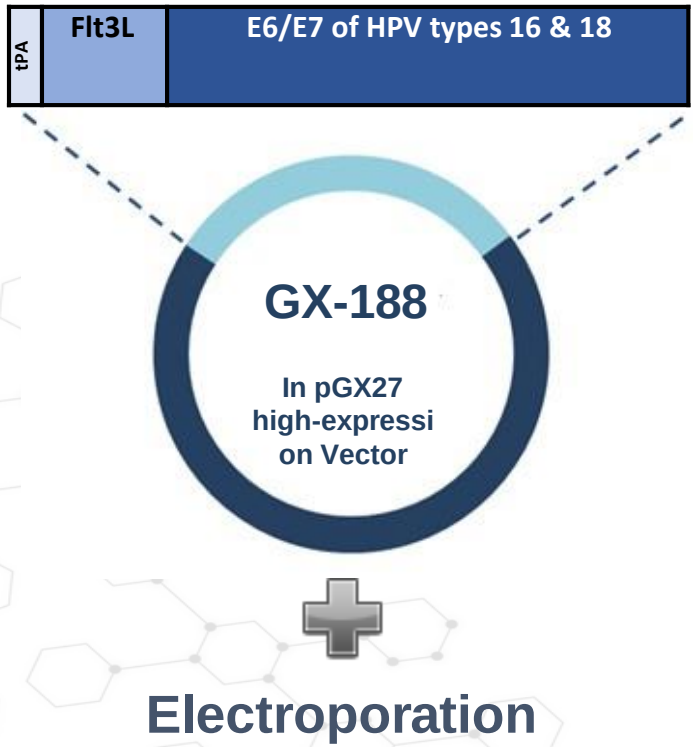
GX-188E DNA Vaccine for Cervical Cancer and SCCHN

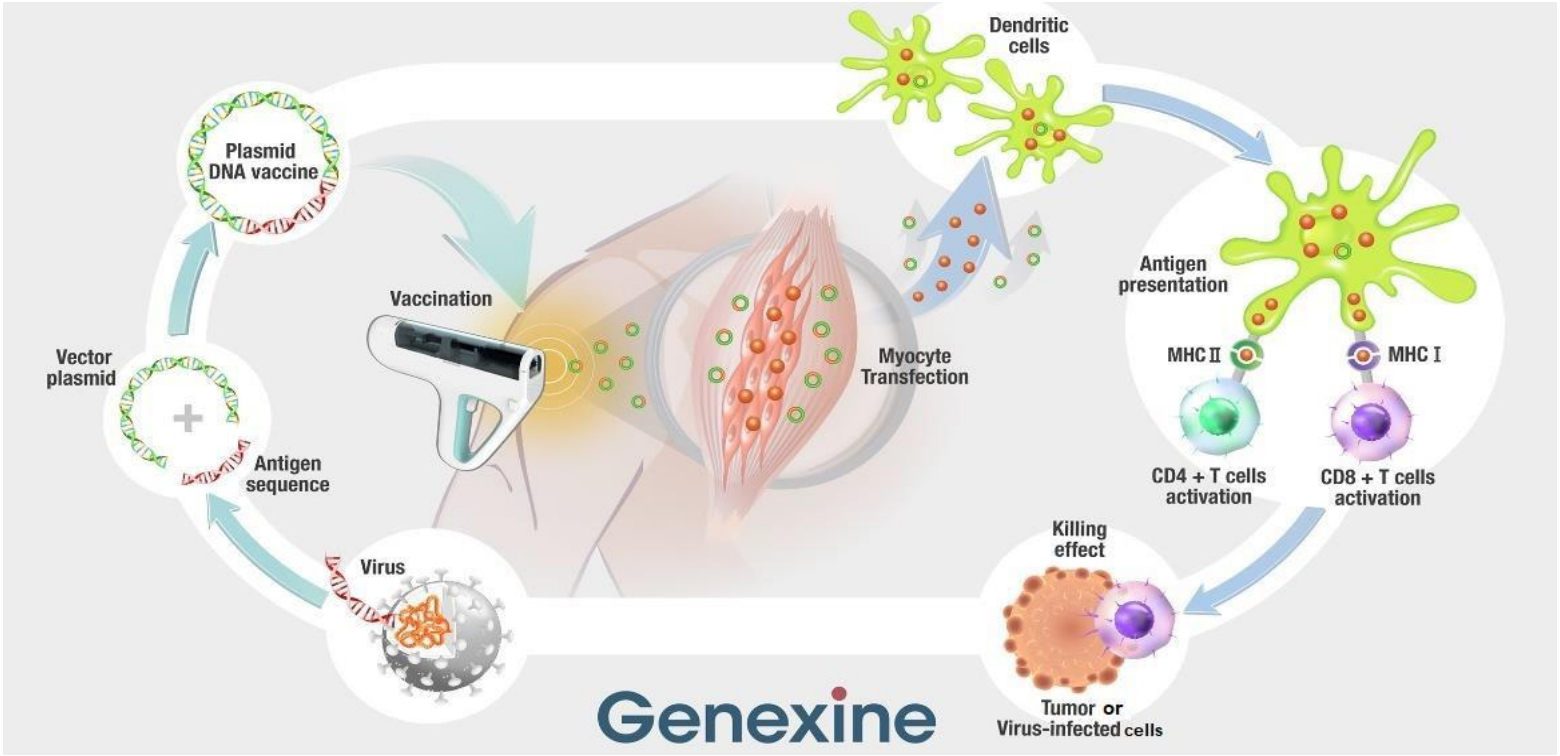
First-in-Class Drug Candidate



GX-188E: Therapeutic DNA Vaccine against HPV-based diseases

- Therapeutic DNA Vaccine for HPV genotypes 16 & 18 (responsible for 70% of cervical cancer)
- Rationally designed DNA vaccine to enhance HPV 16/18, E6- and E7-specific CD8 + T cell responses
- Electroporation improves delivery and uptake
- Dramatic increase in efficacy as combination therapy with Check Point Inhibitors (CPI)





Source; TJ Kim et al, Nat Commun.; YJ Choi et al, Clin Cancer Res., 2020; JW Youn et. al, Lancet Oncol, 2020

GX-188E + Keytruda: Overall Timeline & Current Status (Cervical Cancer)

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Phase 2 clinical timeline

Product/Indication		1	2	3	4	5	6	7	8	9	10	11	12
GX-188E + KEYTRUDA® Cervical Cancer, >2L (n=60)				▲ ORR at W14	▲ ORR_W24 by Investigator			▲ ORR_W24 by BICR	▲ DB lock	Top line results ▲ ESMO 2022	▲ TLF		▲ CSR ▲ CA Sub.

Interim analysis

- (n=48/60 total) reported at ASCO 2021, results from ph2 for combination with Keytruda (KEYNOTE-567)
- **33 % ORR (16/48) with 6 CR and 10 PR.**
- 48.0% (12/25) ORR under condition of PD-L1 + patients, HPV 16 and SCC (squamous cell carcinoma).
- High safety and tolerability: similar AE level compared to Keytruda mono.

Expected Milestones

- Presentation of **top-line results** in 2H ’22 (possibly ESMO)
- Submission of conditional approval in Korea, early ’23
- Commercial plans:
  - On-going discussions with potential partners for Phase 3 and commercial
  - Evaluating commercial launch in Korean market by Genexine

GX-188E + CPI + GX-I7: Overall Timeline & Current Status (Head & Neck Cancer)

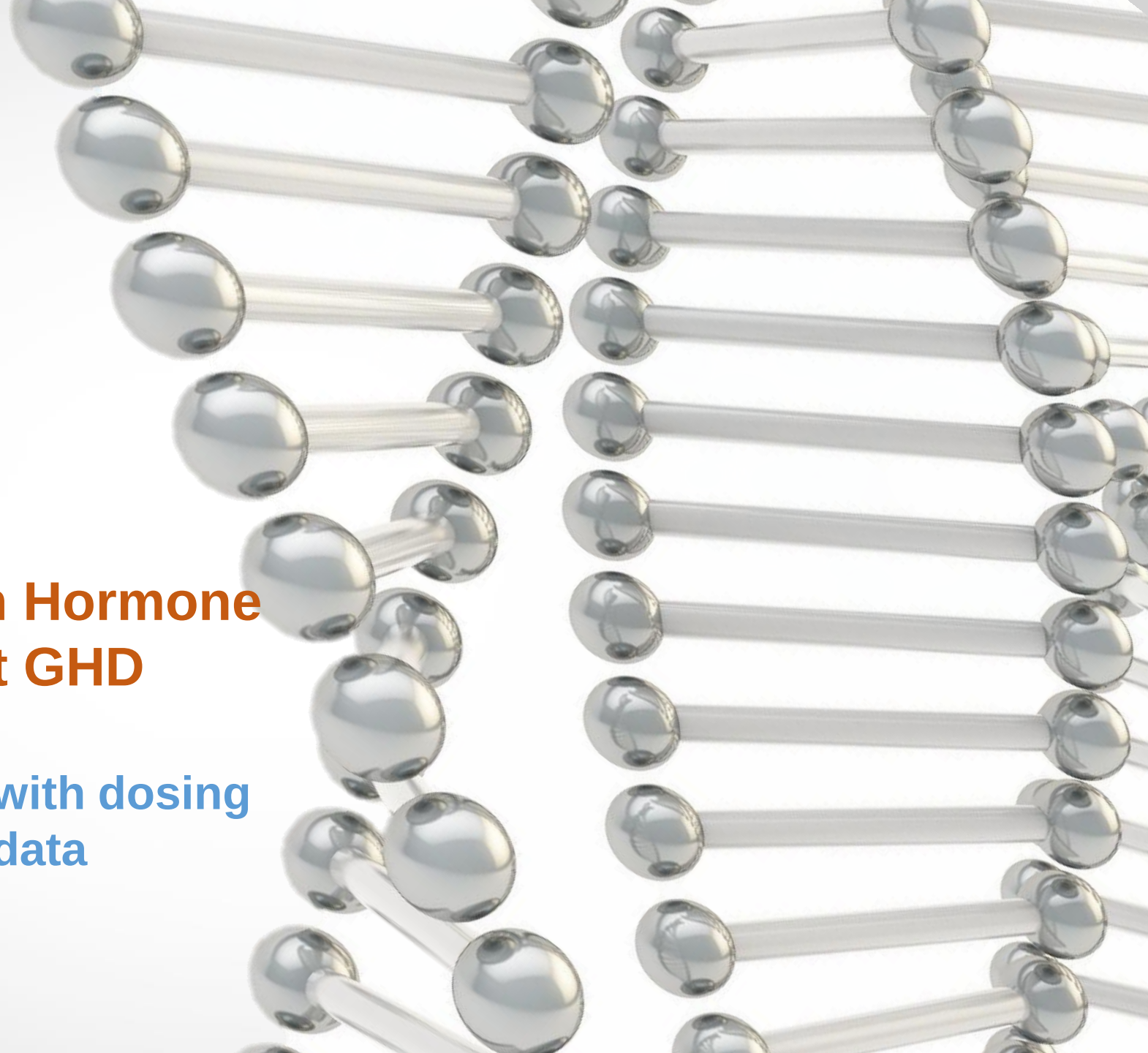
Product/Indication	Objectives	1	2	3	4	5	6	7	8	9	10	11	12
GX-188E+ GX-I7 + OPDIVO® r/m HNSCC, ≥2L (n=21)	▪ Anti-tumor PoC and safety of triple combo in HPV+ r/m HNSCC	IND App./Contract		IND Approval	Opdivo Register in site	Enroll & Tx.	FPI	Lead-in safety (n=3)			ORR at W9 (n=6)	LPI	
GX-188E+GX-I7 + KEYTRUDA® LA HNSCC (n=11)	▪ Safety of triple combo in LA HNSCC as neoadjuvant therapy ▪ Histological PoC of triple combo in tumor microenvironment	IND App./Contract			IND Approval	FPI	Enroll & Tx.			LPI			MPR data

Expected 2022 Milestones

- IND submissions Q1/2 ’22
- Last patient dosed Q4 ’22 for both studies

GX-H9 Long-acting Growth Hormone for Pediatric and Adult GHD

Best-in-Class Drug Candidate with dosing advantage and growth data



Phase 3 : GX-H9 (Long-Acting Growth Hormone Deficiency Treatment)

AGHD and PGHD

Growth Hormone Deficiency Treatment

Daily Injection	Weekly/Bi-weekly injection
312 ~ 365 shots/year	Weekly : 52 shots/year Bi-weekly : 26 shots/year

Clinical Unmet Needs
Long-Acting & Convenience

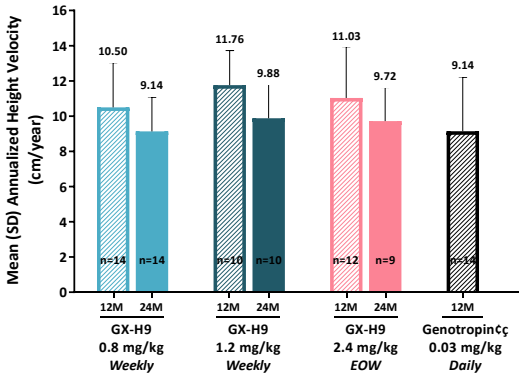
Clinical Trials

- **Ph3 Trial in China by I-Mab**
  - Ph3 approval: 2020 Oct.
  - FPI: 2021 Feb.
  - BLA: 2024 first half
  - Patients: n=165 (expected completion of recruitment 2H 2022)

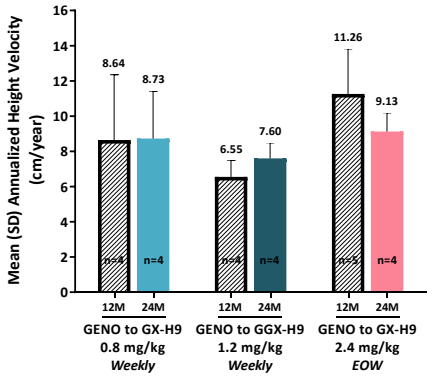
Growth data comparison, 1st yr vs 2nd average growth

- **Weekly or bi-weekly based injection** clinical data for 5 to 12 yrs-old growth hormone deficiency patients.
- Growth rate continued in 2nd yr compared to 1st yr. (both weekly and bi-weekly)
- Growth rate continued even after switching to H9 2nd yr after injection of other growth hormone product during 1st yr. (both weekly and bi-weekly)
- Developing pen type injector for better convenience.

aHV at 1st and 2nd year by doses



Switching to GX-H9 at 1 year



* EOW : Every Other Week

GX-H9 Global Development Status

Early stage development	Global development	Global market

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HANDOK

GX-H9

Growth hormone deficiency-Pediatric
Ph1 in EU, Ph2 multi-national trial completed

Growth hormone deficiency-adult
Ph1 in EU, Ph2 multi-national trial completed

Co-developing with Handok
CMC improvement completed
- Purity
- Getting ready for mass-production end of 2021

L/O
(Oct. '13)

I-MAB BIOPHARMA

Ph3 in China (on going)

- Total deal $ 100mn (GX-H9, G6, G3 combined)
- **Sales milestone**
- **BLA submission expected 2024**
- **Expected Launch 2025**

- **China** (excl. HK, Macau, Taiwan)
- **Strategic alliance with Jumpcan Pharmaceutical for fast commercialization in China** (Nov. '21)

Genexine

- **Evaluating ROW Ph3 Pediatric Growth hormone deficiency**

L/O

America, EU, Korea, Japan ASEAN, MENA, Oceania.

GX-H9 Market & Competitive Landscape

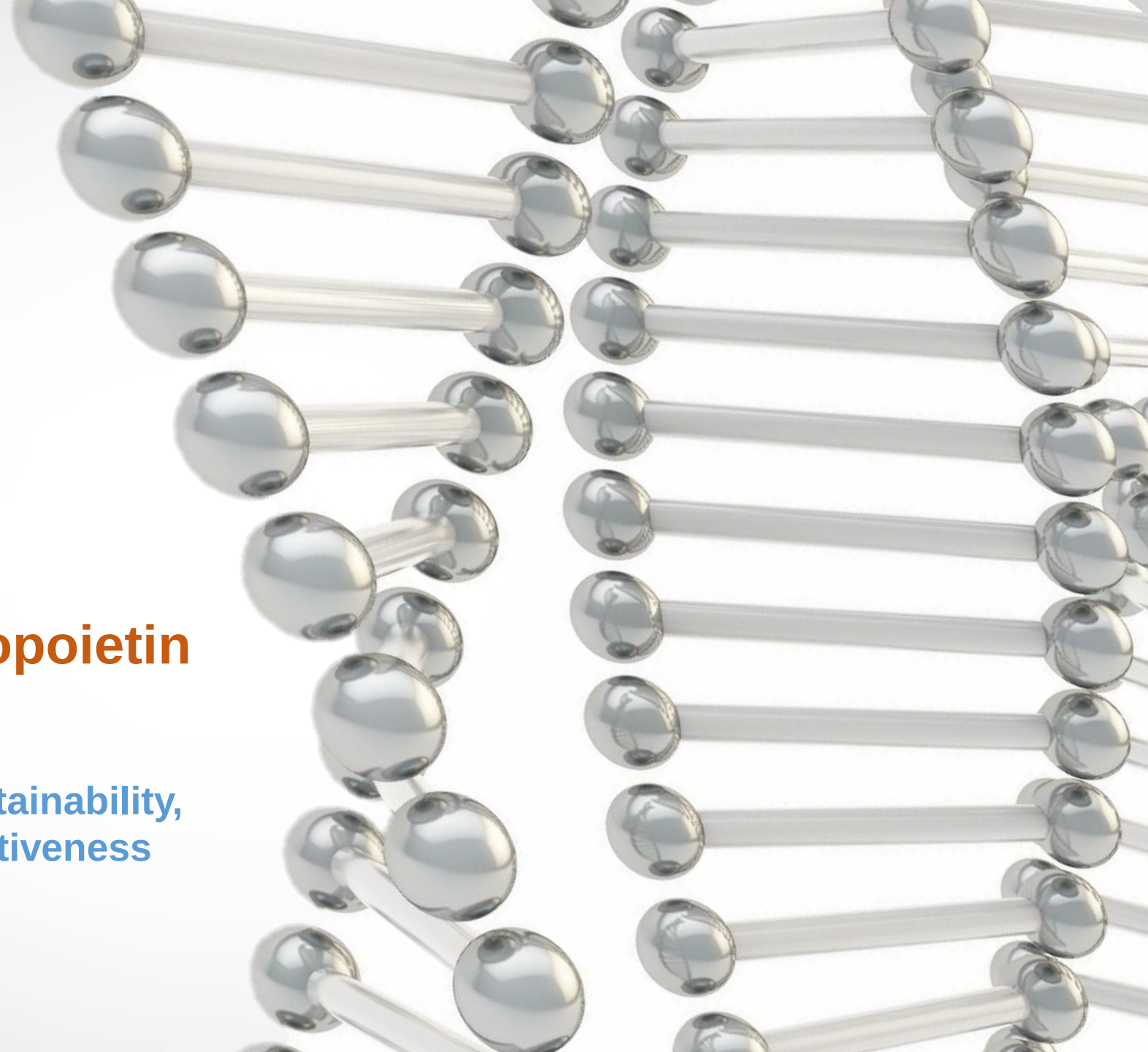
- Pediatric growth hormone deficiency treatment, global market expected $6.0 B (CAGR6.4%) in 2030.
- Chinese pediatric growth hormone deficiency treatment market expected $3.2B (CAGR 15.7%) in 2030, 60% share of global market

Reference : F&S, CMBIS

Company	Genexine HANDOK	ascendis pharma	OPKO Pfizer	novo nordisk	GenSci 金赛药业	特宝生物 AMOYTOP BIOTECH
Product	GX-H9	TransCon hGH	somatrogon	somapacitan	Jintrolong®	Xiamen Amoytop Biotech
Frequency	Weekly/**Twice-monthly**	Weekly	Weekly	Weekly	Weekly	Weekly
Stage	I-MAB BIOPHARMA CN, Phase 3	US, Marketed (‘21.8) CN, Phase 3	US, BLA submission	US, Phase 3	CN, Marketed	CN, Phase 2/3
Orphan Drug Designation	FDA: Nov. 2016 EMA: July 2021	FDA: Apr. 2020 EMA: Oct. 2019	FDA: Sep. 2010 EMA: Jan.2013	EMA: Aug. 2018	-	-
# of Patients	n=165	n=150	n=224	n=200	n=1500	n=400
Height Velocity	**11.76cm/yr** (1.2mg/kg)	11.2cm/yr (0.24mg/kg)	10.12cm/yr (0.66mg/kg)	11.5cm/yr (0.16mg/kg)	2.26±0.87cm ~ 13.41±3.72cm/yr (0.2mg/kg)	-

GX-E4 Long-acting erythropoietin

Long-acting EPO with superior sustainability, efficacy, safety and price competitiveness



Phase 3 : GX-E4 (Long-acting EPO)

EPO(Erythropoietin)
Anemia treatment

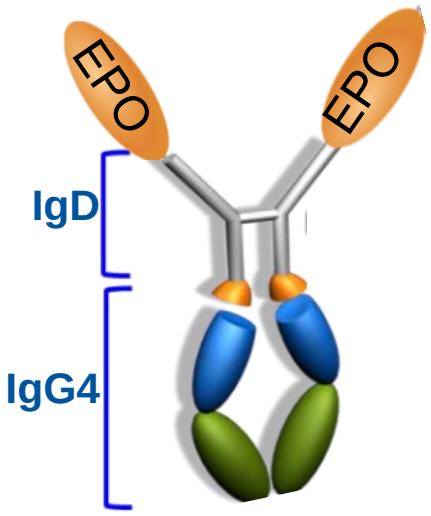
1st Gen	2nd Gen	3rd Gen
3 shots/wk	1 shot/wk	Bi-weekly/monthly
Epogen (Amgen)	Aranesp (Amgen)	Mircera (Roche)
		Unmet needs **Sustainability & safety**

Clinical Trials

- **Phase 3 trial in multi-nations (2020 March~)**
  - CKD-induced Anemia (non-dialysis)
  - 7 countries: Korea, Australia, Taiwan, Indonesia, Malaysia, Philippines, Thailand.
  - Recruited 327 / 386(Dec. '21)
  - Non-inferiority study against Mircera

Competitiveness

Long-acting EPO drug with superior sustainability, efficacy, safety and **price competitiveness**



No mutation

Outstanding efficacy and safety

- High activity and efficacy
- Safely treat patients without hypersensitive immune reaction nor adverse events.

Price competitiveness

- Simplified production process
- Price competitive due to low mfg. cost

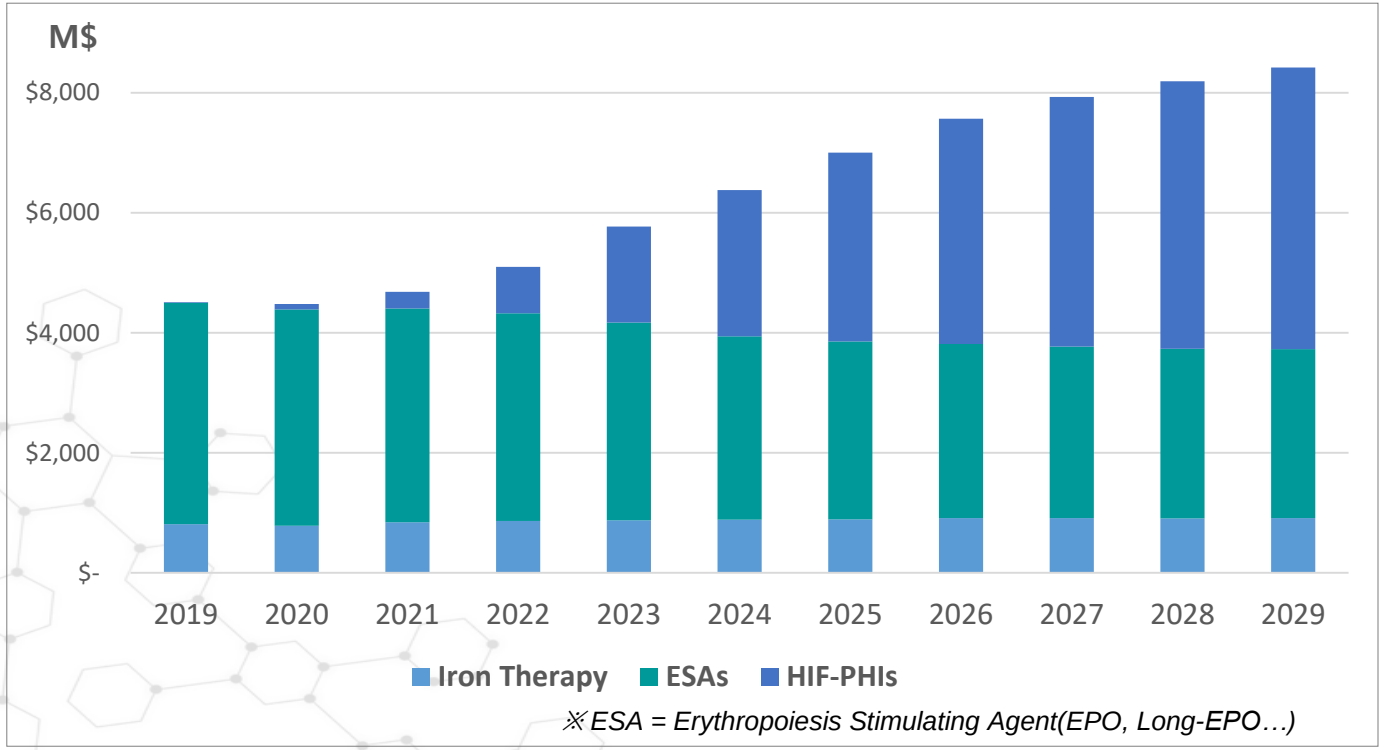
Superior sustainability

- Bi-weekly or monthly dose

GX-E4 (Long-Acting EPO) Market & Competitive Landscape

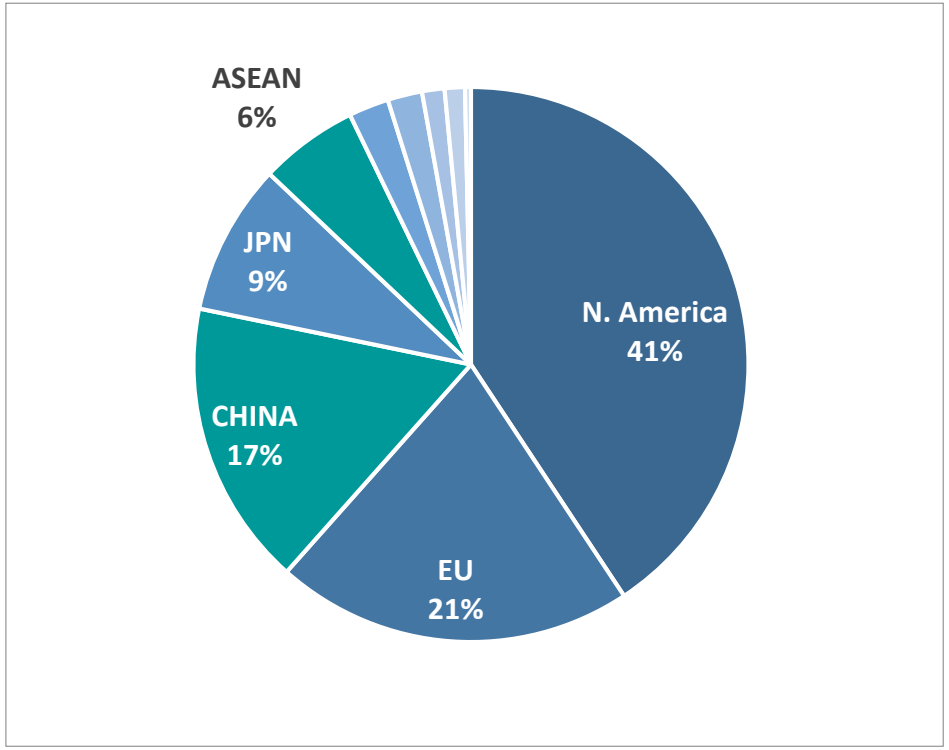
- Over 60% of Global CKD Anemia market consisted of major market such as US, EU.
- ESA (Erythropoiesis Stimulating Agent) market expected $2.9B in 2029

Global CKD Anemia treatment market expectation



※ESA = Erythropoiesis Stimulating Agent(EPO, Long-EPO…)

Global CKD Anemia treatment market 2027



Reference: IMS(~2017) + Global Data(HIF-PHi) + Roche Data Base + VIFOR annual report + Amgen annual report

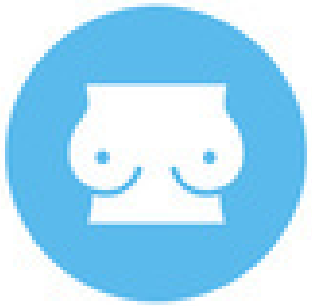
Business Focus and Upcoming Milestones

Genexine leverages global development via strategic partnerships

Contract Date	Partner Company	Product	Territory
2014.07	BSK	GX-188E	China
2015.06	NeoImmuneTech	GX-I7	Europe/USA
2015.10	I-Mab	GX-H9, GX-G6, GX-G3 etc.	China
2016.02	CWB	GX-E4	China
2016.03	KG BIO	GX-E4	ASEAN
2017.12	I-Mab	GX-I7	China
2020.01	GENBIO	GX-P1, GX-P10	Worldwide
2020.12	Turret Capital	GX-P1	Worldwide
2021.02	KG BIO	GX-I7	ASEAN, MENA, Oceania, etc.

Genexine's Commercialization Plan

Multiple BLA submissions over next 5 years

2023	2024-2025			2025-
GX-188E +Keytruda® (conditional approval)	**GX-I7** (NT-I7) +Keytruda® (conditional approval)	**GX-E4**	**GX-H9**	**GX-I7** +Temozolomide
				
Cervical cancer	**CRC/PaC/ TNBC/GBM**	**CKD-induced Anemia**	**Growth Hormone Deficiency**	**GBM**
Phase 2 (Korea)	Phase 2 (Korea/US)	Phase 3 (Asia)	Phase 3 (China)	Phase 3 (China)

Genexine Investment Thesis

Poised for global expansion and value generation

- Genexine pursuing a global path to drug development that will enable it to leverage global markets, technology and capital
- With solid platform technologies and multiple first-in-class products, Genexine now in position to commercialize its first products over the next 2-3 years.
- Uniquely positioned to access novel technology in South Korea, Asia and the U.S. provides competitive advantage
- Globally networked and experienced leadership team for efficient execution of global development plan
- Key near-term clinical data readouts & inflection points will drive shareholder value

THANK YOU

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