



GenScript Biotech Corporation Company Presentation

(Stock Code: 1548.HK)

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Business Highlights

Financial Highlights

Future Strategy

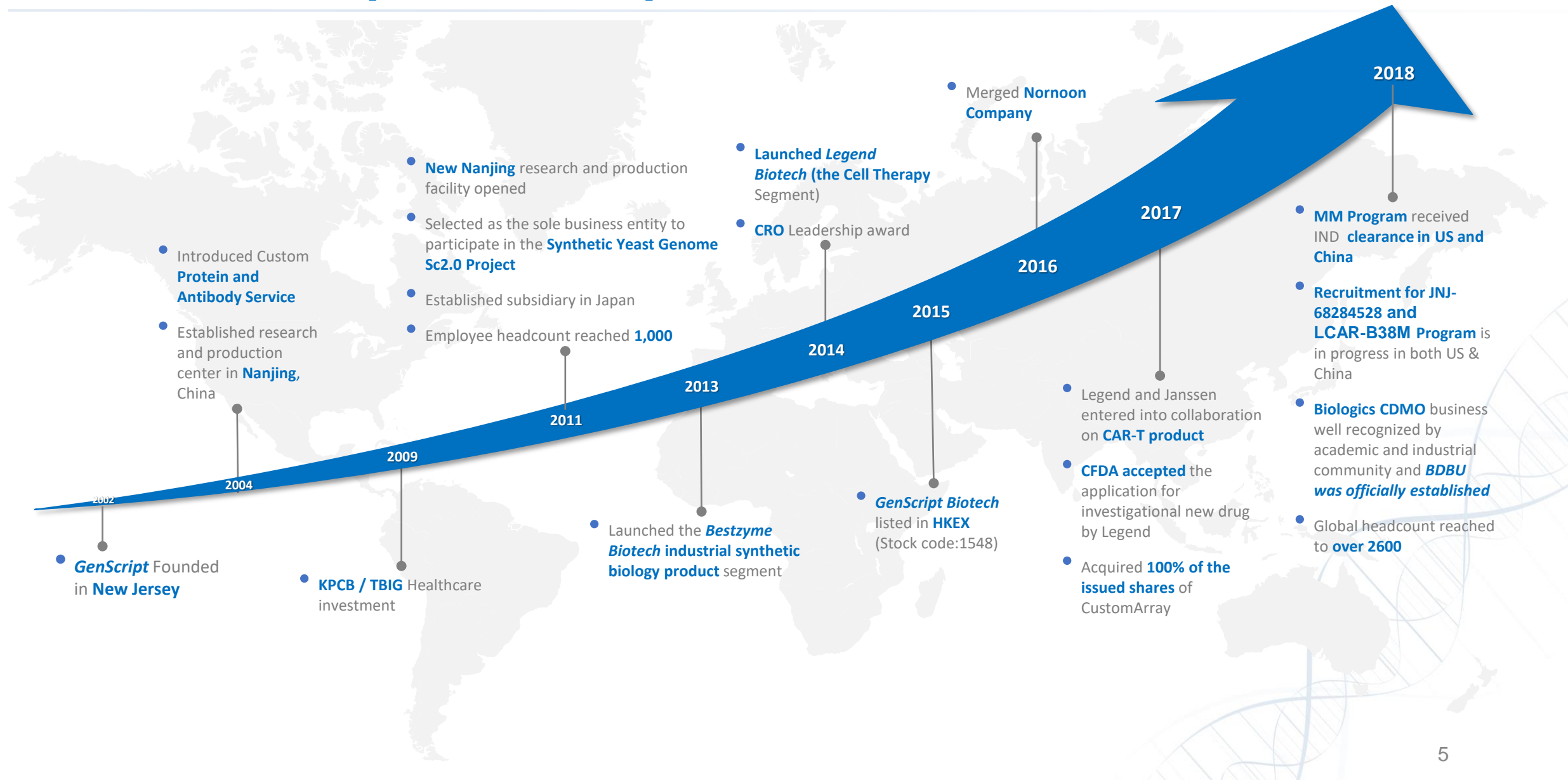
Q&A

Mission

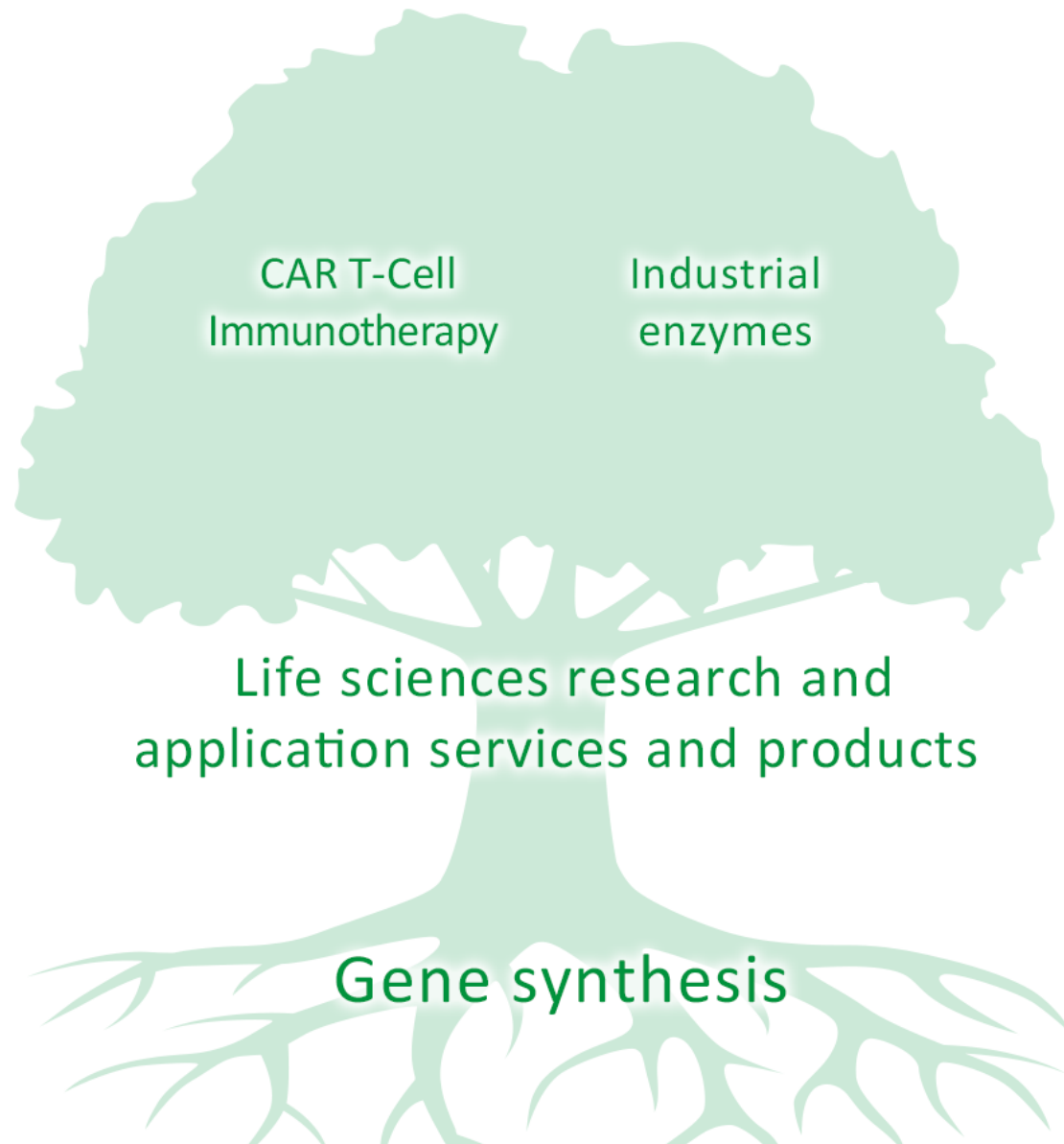
Make Humans and Nature Healthier
through Biotechnology



Global Development Footprint



Business Blueprint – Incubating the Future



New Area of Business Development
Generating Higher Return

Core Business Generating Cash Flow
For Future Development

GenScript Proprietary Technologies

Company Positioning & Major Achievements

Company Positioning

- ✓ Become Global Leader in Gene Synthesis Services and Synthetic Biology Products;
- ✓ Become world class leader in Cell Therapy and Gene Therapy fields

Major Achievements

- ✓ Group Sales Revenue increased over 50%;
- ✓ Expanded global leading position in Gene Synthesis Services;
- ✓ Successfully launched antibody purification devices
- ✓ Received FDA & China FDA clearance to initiate clinical trials with JNJ-68284528 / LCAR-B38M for treatment of Multiple Myeloma;
- ✓ US Phase 1b clinical trial has been running as planned, and entitled to First milestone payment US\$25M;
- ✓ FDA has granted the Orphan Drug Designation to JNJ-68284528 / LCAR-B38M
- ✓ EMA has granted the PRIME designation to JNJ-68284528 / LCAR-B38M
- ✓ Finished Preparation for China Clinical trials
- ✓ Built up world class management team for Legend;
- ✓ Constructing US and PRC GMP facilities as planned;
- ✓ Reached strategic collaboration with Biologists CDMO partners
- ✓ Finished the construction of Industry Enzyme manufacture facility;

Company Overview



Global Presence

- **New Jersey**-USA
- **Nanjing**-China
- **Amsterdam**-Netherland
- **Dublin**-Ireland
- **Tokyo**- Japan;



Strong IP Position

- Strong **IPs and know-how proprietary** technology in the area of synthetic biology;
- **Holds 70+ registered patents and 200+ patent application**¹;



Diversified Customer Base

- **5,000+ customers globally**
- Including global pharmaceutical and biotech companies, colleges and universities, research institutes, government organizations and distributors in over **100 countries**;



Well Trained Employees

- Over **2,600 employees** globally;¹
- Over 75% of employees hold Bachelor and above degrees,
- Over 33% of employees hold Master and PHD degrees

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Strategic Business Positioning



GenScript
Make Research Easy

Life Science CRO

Continuously Being the Global Leader in Gene Synthesis Services Market



GenScript
Make Research Easy

Biologics CDMO

Becoming an Emerging Leader of CDMO Service provider

Gene Synthesis Technology



Bestzyme

Industrial Synthetic Biology Products

Make the best Enzyme products to improve the quality of life and maintain a better nature environment

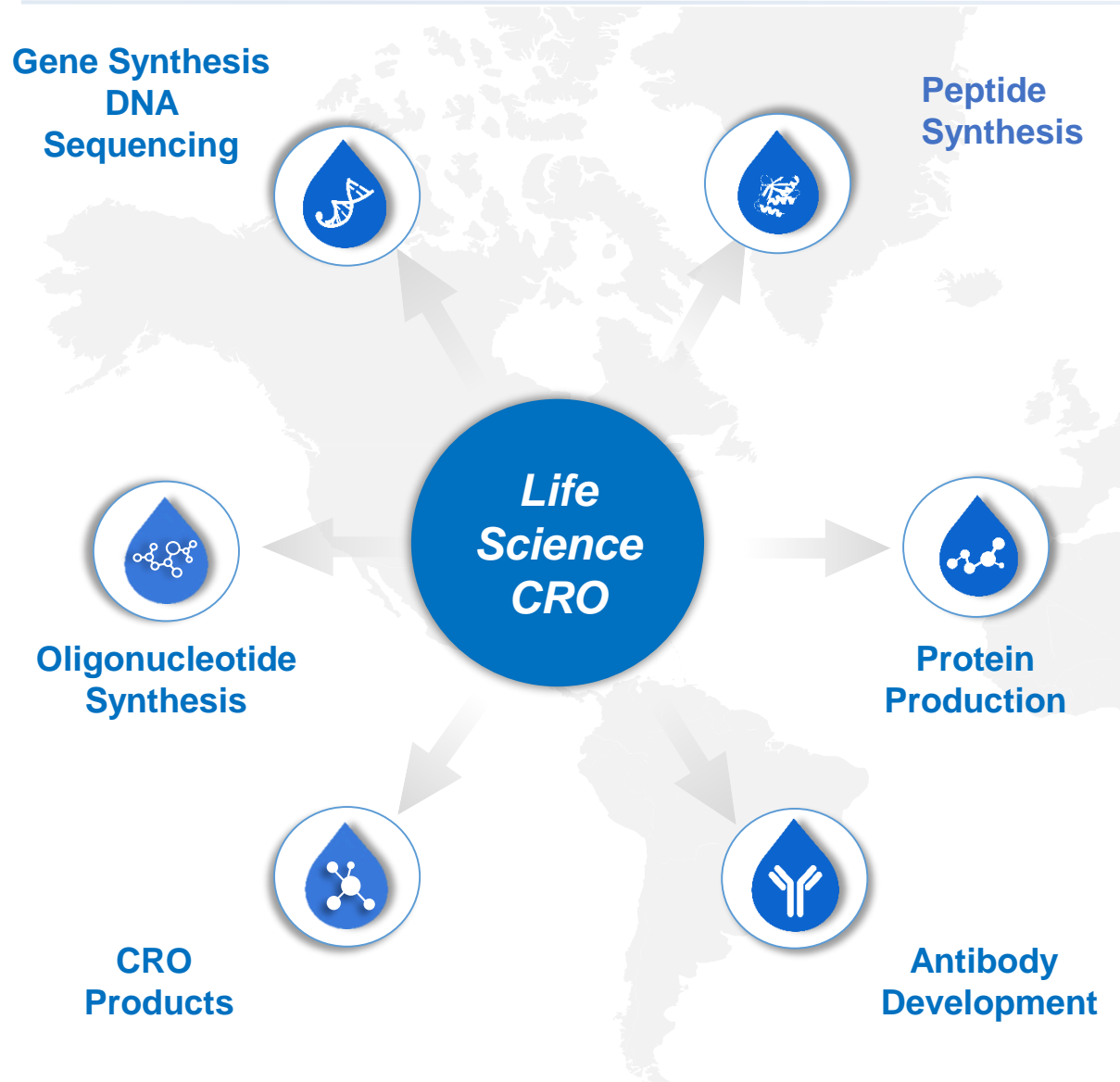


**Legend
Biotech**

Cell Therapy

The Pursuit of A Cure

One Stop Life Science CRO Solution



Global Leader in Gene Synthesis

- Largest gene synthesis provider;
- Successful delivery of more than 1.2M genes;
- Largest capacity at 100 million bp/month;
- Successful rate 99.95% ;
- Over 98.5% on-time delivery;

Becoming the Largest Oligo Provider in Precise Medicine

- Rich experience and solid foundation in oligo synthesis over past 16 years;
- Supply oligo in most promising health territory, precise medicine;
- MDx oligo (US\$ 427.8M addressable market);
- MDx Kits (US\$1,100M addressable market);

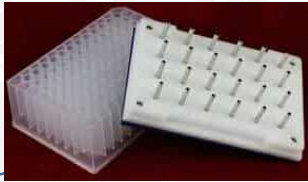
Continuously Enhanced CRO Core Competitiveness

- Molecular Cloud
- Rabbit Monoclonal Antibodies
- Combinatorial Library
- CRO Automation

Solving Biologics Drug Industry Challenge

Protein Purification Solution

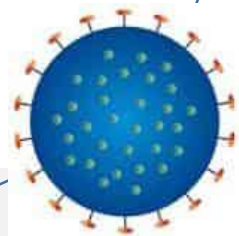
96 well Magnetic plate



Magnetic Box



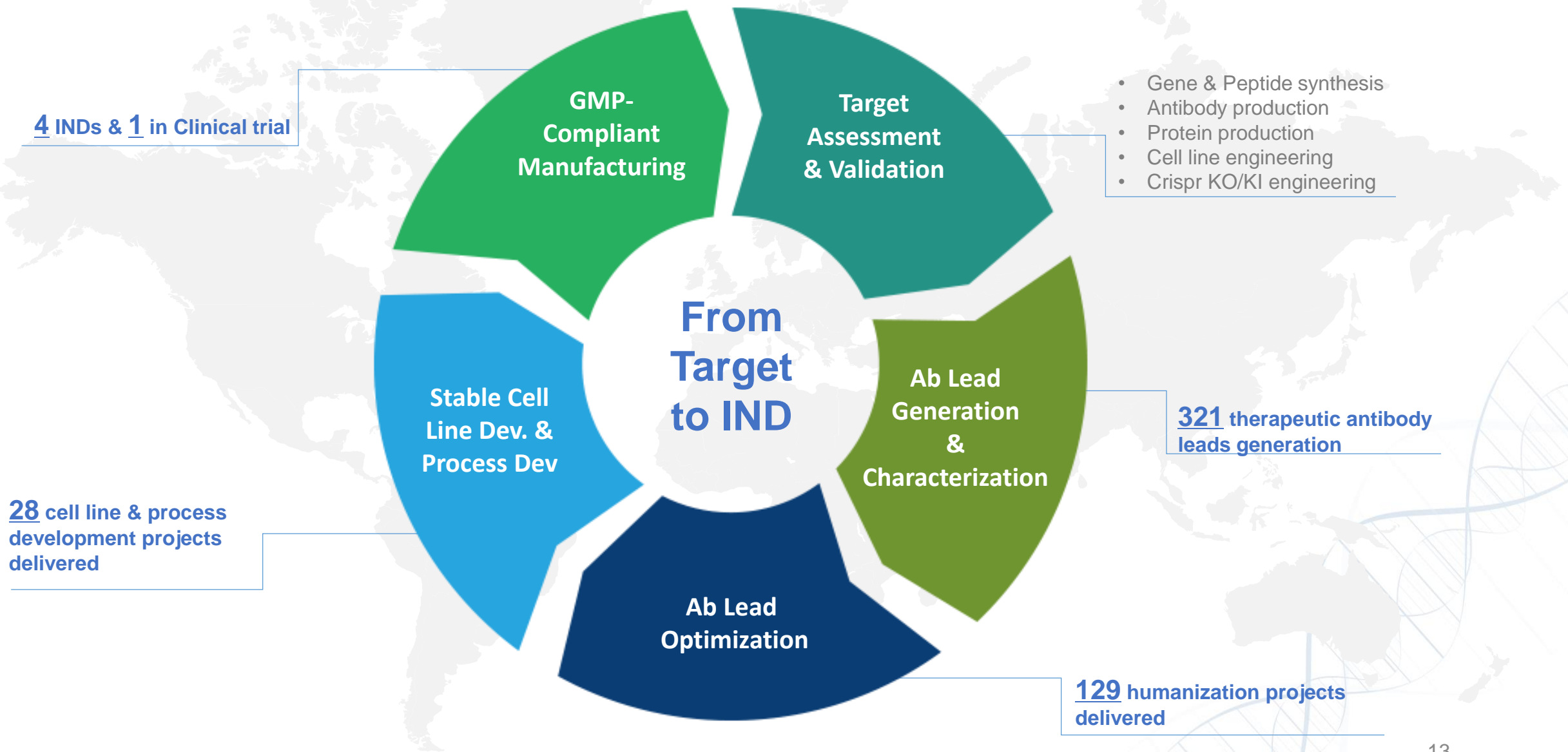
AmMag SA



Magnetic Beads

- Collaboration with Amgen, We developed the **innovative Mag beads capture instrument** for antibody purification
- Able to **provide high through put capability** without prior centrifugation and filtration of samples.
- Becoming the **first player in the commercial magnetic bead technology market** to address this need
- **Priced at USD70K** per unit.

Biologics CDMO Builds up Advanced Tech Platform



SMAB Platform to Enable Client Success

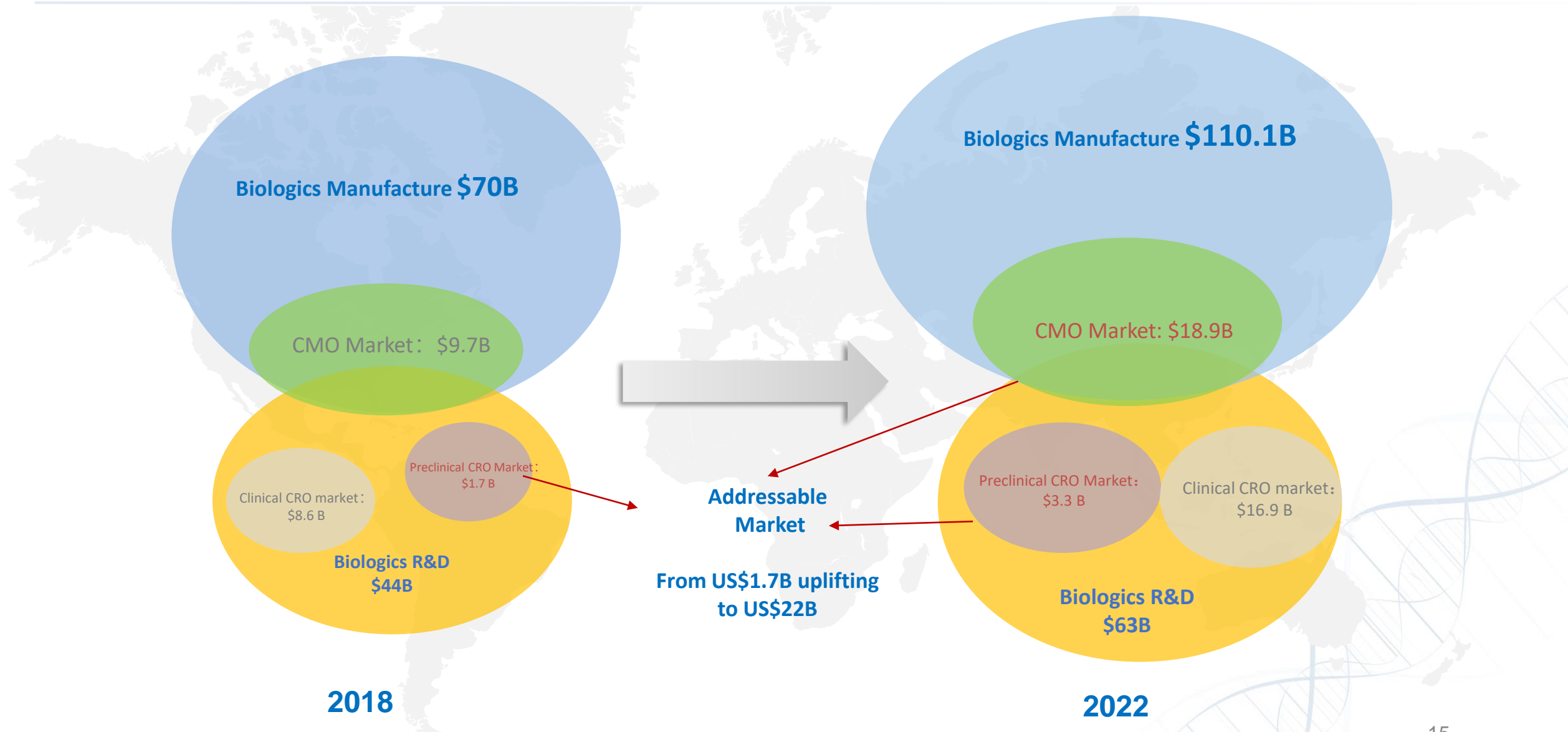
Bispecific Antibody Platform-SMAB

- **Naïve:** sequence mutation not needed, minimum gene editing;
- **Outstanding developability**, as good as Monoclonal Ab;
- **Low immunogenicity, less side effects**;
- Flexible format: **≥ 2 targets/epitopes**;
- Unique **molecular flexibility** for precise adjustment of dual target effects

Commercial Collaboration

- **3 Partners** on SMAB Platform;
- License and further development of **2 SMAB-based antibody drug molecules**;
- License and development of **1 biosimilar**;
- Strategic partnership on **PD, PK & toxicity assessment**;
- Strategic partnership on full-humanized mouse **antibody discovery**

Promising Market Prospectus for Biologics CDMO



Investment in CDMO GMP Facilities to Fuel up Growth



CDMO R&D Labs

1. From an idea to preclinical

- Full cycle of Services
- Antibody Discovery,
- Cell line development, Formation and engineering, assay development
- Preclinical development



GenScript Science Park



CDMO GMP Manufactory Facility

2. Enabling Clinical Trials

- Compliant to GMP regulation in US, CN and EU
- For Clinical I/II

3. Extending to Commercial manufacturing (incl: plasmid & virus)

- Compliant to GMP regulation in US, CN and EU
- For Clinical III and Commercial manufacturing

Committed to “Make the Best Enzyme”

Comprehensive R&D Competences

- **R&D capacity** to enable the launch of new products efficiently
- Latest Gene-editing technology (Crispr/Cas9) enhanced **Comprehensive expression platform**

Production Facility Building up & Optimization

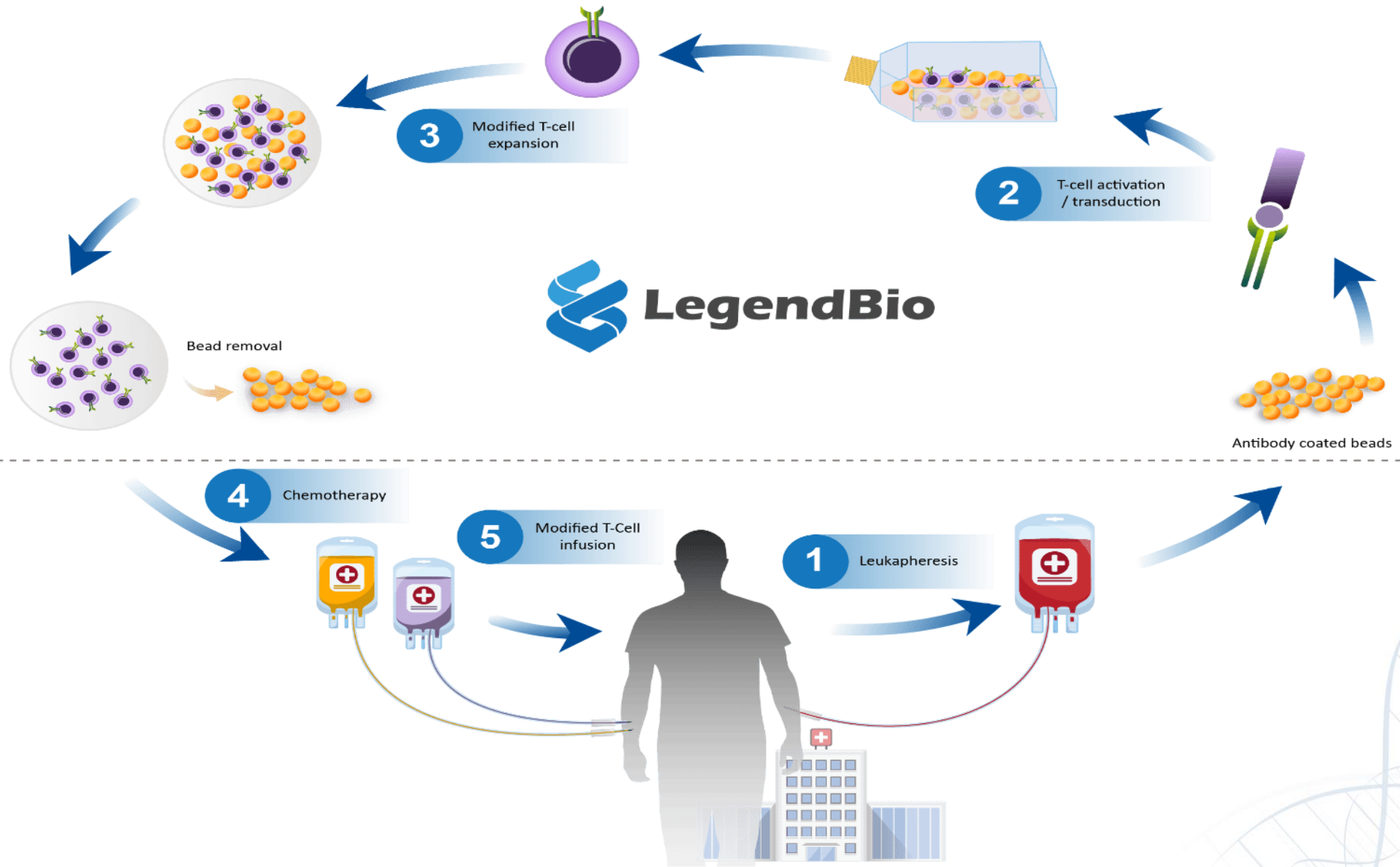
- **Production capacity reached 150,000 standard tons.**
- **Optimization of production processes** will improve the competitiveness of our products



Enhance Commercial Competences

- **Recruited industrial experts to lead the commercial team**
- **Establish long-term partnership** with influential customers

Introduction of CAR-T Cell Therapy



Cell Therapy – Helping to improve the lives of patients worldwide

US & China Regulatory Approval

- **Received FDA & CFDA clearance** to initiate clinical studies with JNJ-68284528 / LCAR-B38M for treatment of Multiple Myeloma;
- FDA has granted the **Orphan Drug Designation** to JNJ-68284528 / LCAR-B38M
- EMA has granted the **PRIME designation** to JNJ-68284528 / LCAR-B38M

US & China Clinical Progress

- US Phase 1b clinical trial has been running as planned, **first batch of milestone payment US\$25M received**
- China **Phase 2 Confirmatory Trial** has been initiated since Feb 2019

US & China GMP Facility

- **China GMP facility** has been up and running and LCAR-B38M for Phase 2 clinical trial has been initiated as planned;
- **US GMP facility** for pivotal study and commercial supply will be ready for operation in Q2, 2019 as planned.

Build Up World Class Management Team for Legend

| Function |  US | China  |
|-----------------------------|--|---|
| CEO | Yuan Xu - Merck, Gilead, Novartis, Amgen, GSK, Genentech | |
| R&D | Qiong Wang AstraZeneca, NCI | Frank Fan, Simon Wu GenScript |
| Clinical | Syed Rizvi Celgene, Novartis, Merck | Tracy Luo Amgen, AstraZeneca |
| Commercial | Steve Gavel Celgene, Millennium, IMS Health, Amgen | Chong Yang Roche, Bayer, Novartis |
| Manufacturing | Jason Hamilton, John Tomtishen Novartis, Fox Chase Cancer Center | David He Boehringer Ingelheim |
| Global Quality | Alan Kick - Celgene, Dendreon, Pfizer, JNJ, Roche | |
| Global Regulatory | Yuhong Qiu - Novartis, JNJ | |
| Global Business Development | Meeta Chatterjee – Merck, Schering-Plough | |

Global Clinical Trial Updates¹

US & Europe Clinical Trial Updates¹

Overview

- Phase: 1b/2
- Purpose: Phase 1b to test safety and to determine the dosage of phase 2; Phase 2 to test efficacy

Planned Enrollment

- 110 participants

US Trial Location-Recruiting

- Sarah Cannon Research Institute – Nashville, TN
- Mount Sinai Medical Center – New York, NY
- Levine Cancer Institute – Charlotte, NC
- University of Chicago – Chicago, IL

Worldwide Trial Location-Not yet recruiting

- United States (13 additional sites)
- Belgium (3 sites)
- France (3 sites)
- Israel (2 sites)
- Netherlands (3 sites)
- Spain (3 sites)

Current Status

- Patient Treatments on going as planned

China Clinical Trial Updates¹

Overview

- Phase: 2
- Purpose: To test safety and efficacy

Planned Enrollment

- 60 participants

China Trial Location-Recruiting

- 上海交通大学医学院附属瑞金医院
- 浙江大学医学院附属第一医院

China Trial Location-Not yet recruiting

- 江苏省人民医院
- 西安交通大学第二附属医院
- 四川大学华西医院
- 福建医科大学附属协和医院
- 北京大学第三医院
- 上海长征医院

Current Status

- Patient Treatments triggered off as planned

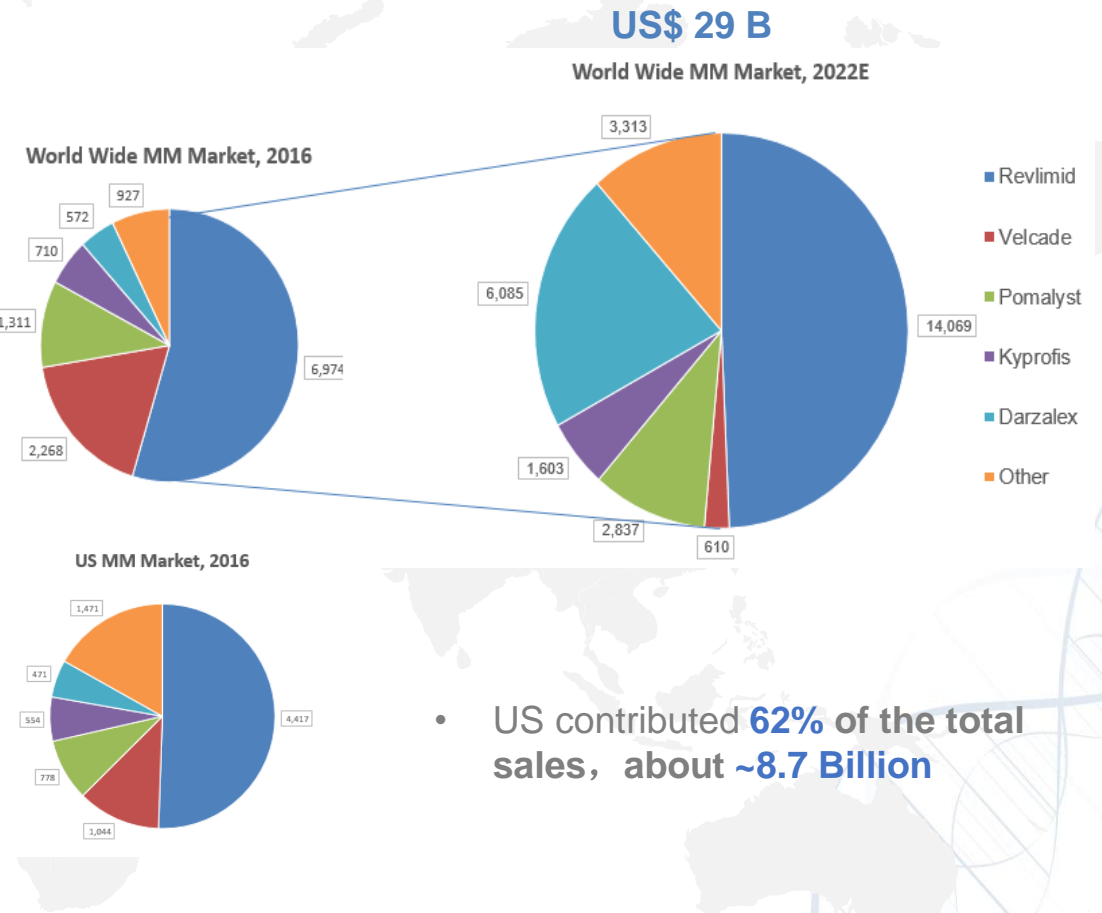
1: as of 20 Mar2019

2. For more information please visit:<https://www.clinicaltrials.gov/ct2/show/record/NCT03548207?recrs=a&cond=Multiple+Myeloma&spons=Janssen&draw=2&rank=3&view=record>
<http://www.chinadrugtrials.org.cn/eap/clinicaltrials.searchlistdetail>

Multiple Myeloma Prevalence and Market Potential

230,000 5-year worldwide prevalence³
1% of worldwide new cancer cases³
2.1% of all US cancer deaths⁴

50.7%
 5-year survival rate⁴

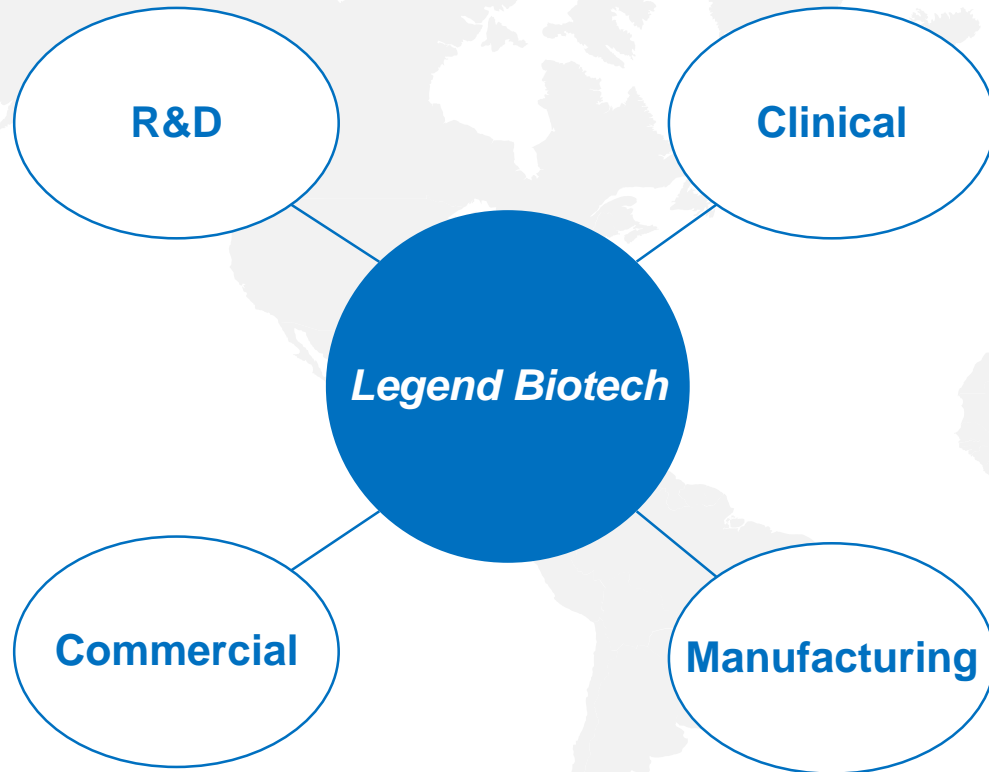


- US contributed **62%** of the total sales, about **~8.7 Billion**

1. 来源: EVP MM treatment regimen approvals; GlobalData Pharmaceutical Intelligence <https://pharma.globaldata.com/HomePage/Index>
 2. Lub S et al. *Oncotarget*.2016;7:6521-37.
 3. Multiple Myeloma Research Foundation. Available: <https://themmrf.org/multiple-myeloma/what-is-multiple-myeloma>. Accessed January 4, 2019.
 4. Ruzafa J et al. *Pharmacoeconom Drug Saf*.2016;25(8):871-9.
 5. Cancer Stat Facts: Myeloma. National Cancer Institute: Surveillance, Epidemiology, and End Results Program. Available: <https://seer.cancer.gov/statfacts/html/mulmy.html>. Accessed January 4, 2019..

Fully Integrated Capability and Global Footprint

Fully Integrated Capability



Legend Global R&D Strategy

Core Technology

CAR

TCR

Product Platform

Autologous
Allogeneic

Disease Areas

- Hematologic Malignancies
- Solid Tumors
- Infectious Diseases
- Autoimmune Diseases

2019-20 Cell Therapy Pipeline Advancement

Current Pipeline

**LCAR-B38M Ph 2
Trials in US/EU/China**

Worldwide Collaboration with



Future Pipeline

AML

Lymphoma

**Gastric
Cancer**

**Ovarian
Cancer**

**Infectious
Disease**

First-in-Human Studies: Next Generation CAR-T/TCR-T/Allogeneic Platforms

Clinical Plan

Multiple IIT programs in China

Multiple IND programs in US

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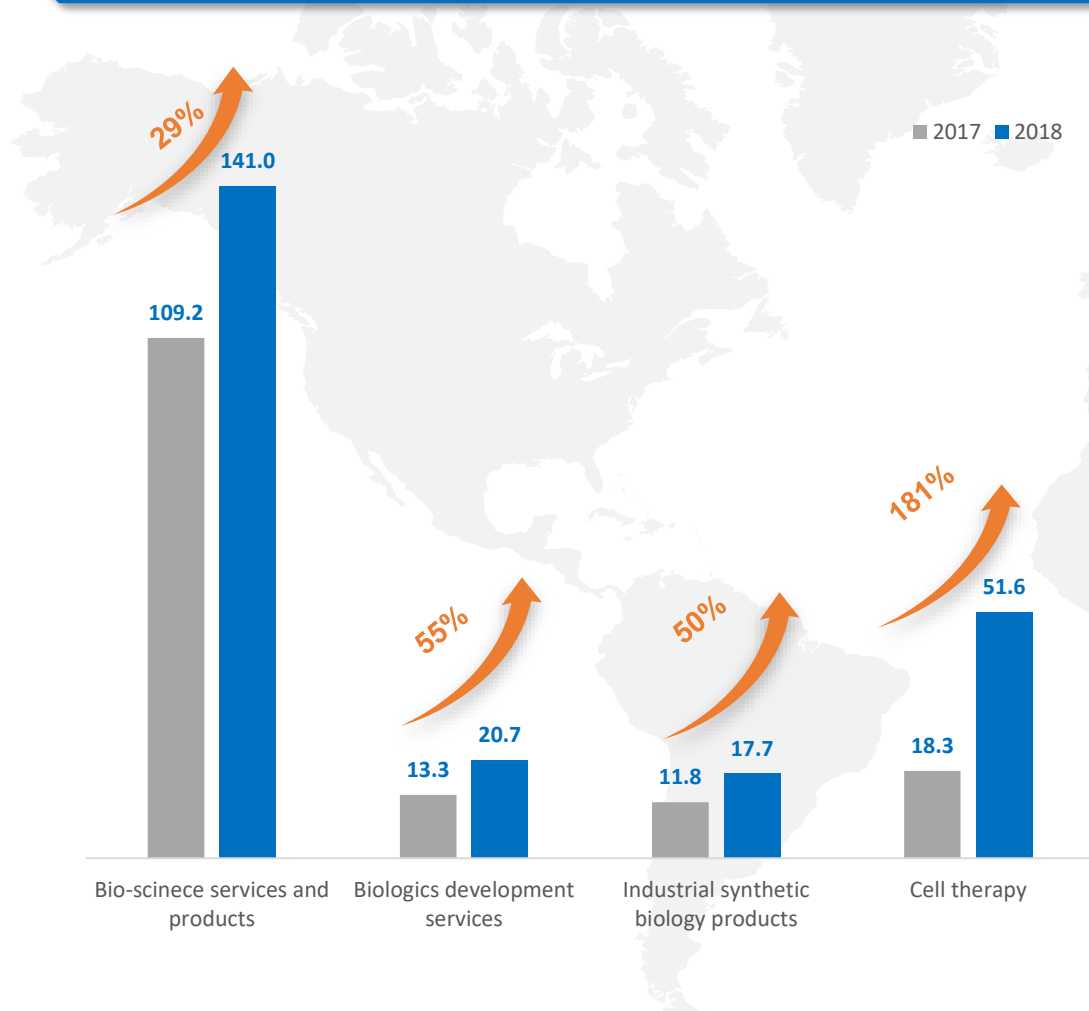
| | 2017 (US\$M) | 2018 (US\$M) | % Change |
|---------------------------|-----------------|-----------------|----------|
| Revenue | 152.6 | 231.0 | 51.3% |
| Gross Profit | 104.6 | 158.5 | 51.6% |
| Gross Margin | 68.5% | 68.6% | |
| Net Profit | 27.0 | 20.8 | (23.1 %) |
| R&D | 18.1 | 74.1 | 309% |
| CAPEX | 32 | 79.6 | 148.8% |
| Cash Positon ¹ | 127.3 | 577.3 | 353.7% |

- **Fast growth in top line even excluding legend portion, recorded 30+% growth in 2018;**
- **Solid and stable gross margin;**
- **Significant increase in research and development expense;**
- **Intensive capital investing in legend and other innovation business;**
- **Strong cash position.**

1. Cash Position=Financial assets at fair value through profit or loss+Cash and cash equivalents+Pledged short-term deposits + Available for sale investments

51%+ Growth in Revenue and Gross Profit

Revenue by Business Segments(US\$ M)

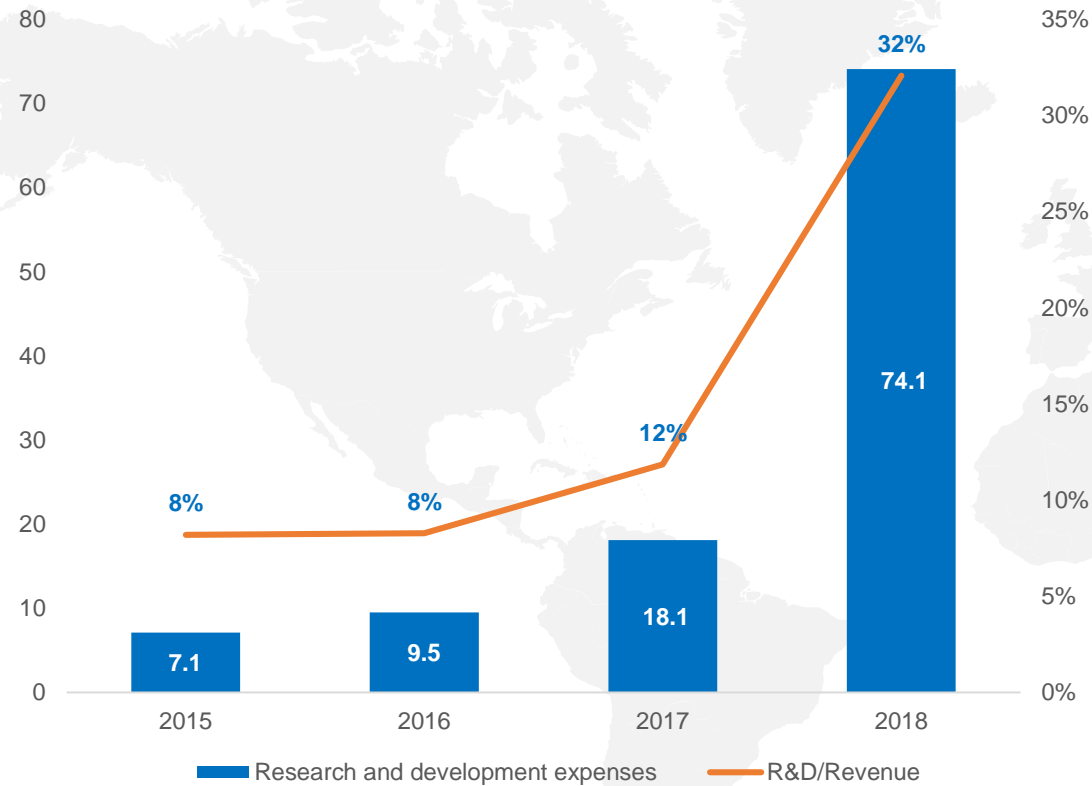


A Strong, Consistent, Sustainable Business

- **4 consecutive years** of revenue growth since listing, showing strong momentum;
- **Continued growth & success in 4 focused** business segments
- **Grow revenue in Bio-Science** services and products segment **above market average**
- Create value through **strategic collaboration in biologics development** services segment
- Sustainable revenue contribution from upfront payment and milestone payment of **JNJ collaboration**

Significant Increase in R&D

R&D Investment (US\$ M)

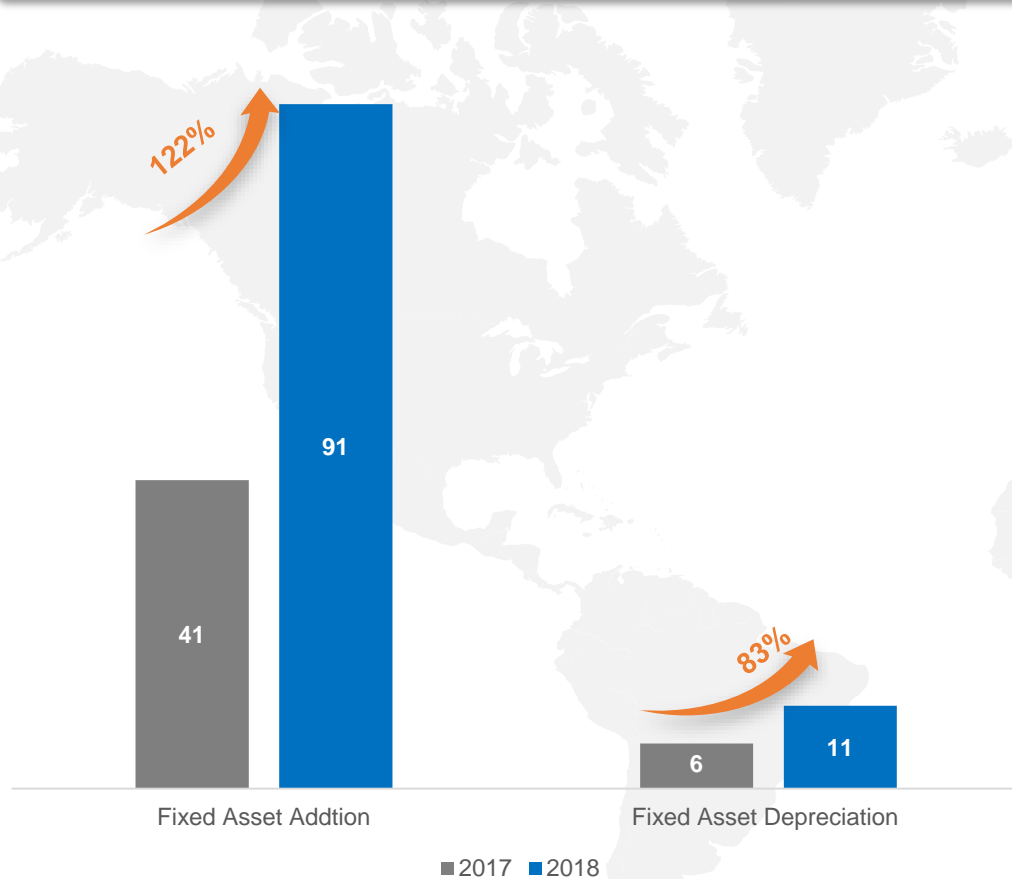


Prioritized R&D Investment, Foundational to our strategy

- Invested US\$74M+ in R&D, increased by 3 times ;
- 4 year CARG is at 118.53%;
- Invest in Cell therapy US\$55M+
 - Groundbreaking cell therapy platform for the treatment of cancer;
 - Lead program, LCAR-B38M (BCMA), IND program in both US and China;
 - Multiple additional programs under development;
- Invest in Bio-Science and other innovation;
 - Development of gene synthesis chip and magnetic beads
 - Portfolio optimization

Intensively Capital Expenditure in 2018

2018 Fixed Assets¹ Increase and Depreciation (US\$ M)



Capital Expenditure increased by 149% to US\$79.6M

➤ Invest in Bio-Science

- High throughput and automation renovation in gene synthesis;
- New production system implantation of industry oligo;
- Biological GMP facility build up;

➤ Invest in Cell therapy

- GMP facilities readiness for Clinical trial both in US and China;
- Research and development equipment;

➤ Infrastructures

- Zhenjiang land and construction;
- Nanjing Legend land and construction;
- Warehouse and staff facilities;

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GenScript

Make Research Easy

Thanks

GenScript Biotech Corporation

Make People and Nature Healthier through Biotechnology

For More Information: <https://www.genscript.com/>

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