



Viva Biotech 2020 Annual Results

Shanghai · China Mar. 31, 2021

DISCLAIMER



Forward-Looking Statements

Certain information set forth in this presentation contains "forward-looking statements" which are not historical facts, but instead are predictions about future events based on our beliefs as well as assumptions made by and information currently available to our management. Although forward-looking statements contained in this presentation are based upon what management of the Company believes are reasonable assumptions, future events could turn out differently from those anticipated in such statements. There can be no assurance that forward-looking statements will prove to be accurate as such forward-looking statements necessarily involve known and unknown risks and uncertainties, which may cause results and future events in future periods to differ materially from any projections of future performance or result expressed or implied by such forward-looking statements. Accordingly, you are strongly cautioned not to place undue reliance on forward-looking statements. All information provided in this presentation is as of the date of this presentation and are based on assumptions that we believe to be reasonable as of this date the Company undertakes no obligation to update forward-looking statements if circumstances or management's estimates or opinions should change except as required by applicable securities laws or listing rules.

Use of Non-IFRS and Adjusted Non-IFRS Financial Measures

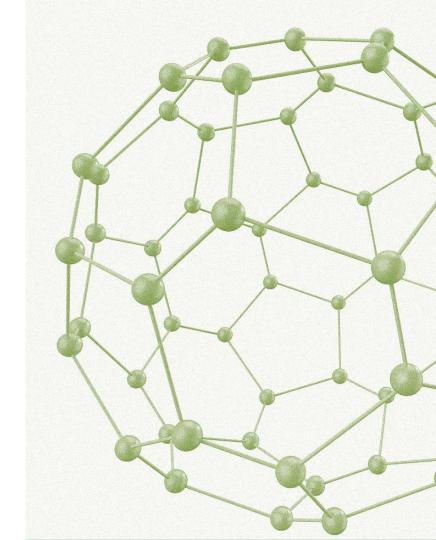
Except for Group's consolidated financial statements which are presented in accordance with the International Financial Reporting Standards ("IFRS"), the Company has provided adjusted Non-IFRS statements to supplement, which are not required by, or presented in accordance with, the IFRS. Adjusted Non-IFRS statements includes, but is not limited to, (i) the adjusted Non-IFRS net profit, (ii) adjusted Non-IFRS net profit margin, and (iii) adjusted Non-IFRS earnings per share (excluding Listing expenses and fair value loss on financial liabilities at FVTPL). These adjusted Non-IFRS statements are provided to allow Company's management and investors the understand and assess the underlying business performance, operating trends and financial performance by eliminating the impact of certain unusual and non-recurring items that the Group does not consider indicative of the performance of the Group's business. However, the presentation of these non-IFRS financial measures is not intended to be considered in isolation or as a substitute for the financial information prepared and presented in accordance with the IFRS. You should not view the adjusted results on a stand-alone basis or as a substitute for results under the IFRS.

Our Mission

Our mission is to become a cradle for promising biotechnology companies from all over the world.

We operate a leading structure-based, integrated drug discovery platform with a number of world-leading technologies and a scalable business model to share the upside of our customers' IP value.

We are dedicated to establishing a one-stop service platform for global innovative drug R&D and manufacturing clients.





Industry Overview

Evolving Trend of CXO



03

Vertical integration biotech sector

CXO shares IP value, breaking through the ceiling of traditional service model

01

Global outsourcing drives the increasing demand for China-based CXO Capital barriers and

technical barriers higher for late comers

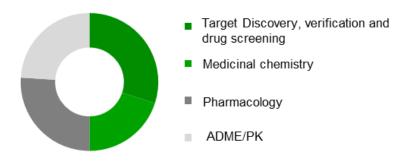
Specialization vs. generalization

SBDD is the Mainstream of Novel Drug Discovery

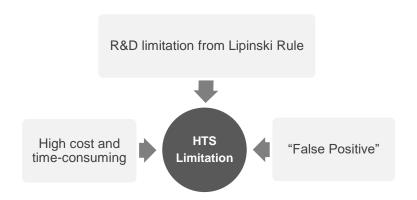


R&D Outsourcing=Discovery/Preclinical CRO->CDMO->Clinical CRO

Breakdown of Global Drug Discovery and IND-enabling Studies Outsourcing Market



Rapid Affinity Screening Commonly Used in SBDD Is Gradually Replacing High-throughput Screening



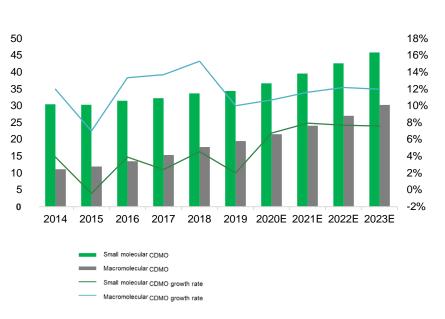
Sources: CRL、安信证券、Cancer Drug Design and Discovery (2016)

Biotech Drives Growth of CRO/CDMO Market

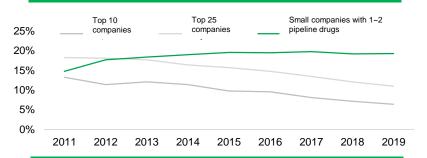


R&D Outsourcing=Discovery/Preclinical CRO->CDMO->Clinical CRO

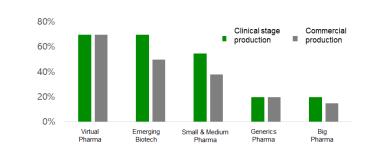
Global CDMO Market Size and Growth Rate 2014~2023E (Billion USD)



Distribution of Global Drug R&D Pipelines Proportion of Small Company Pipelines



Demands of Different Pharma & Bioteches for Outsourcing Production



Financing and M&A of Global Innovative Drug Industry



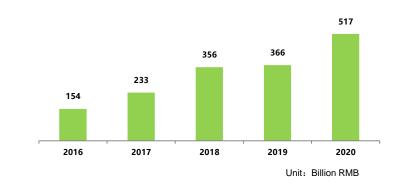
Major M&As in the Global Biomedical Industry



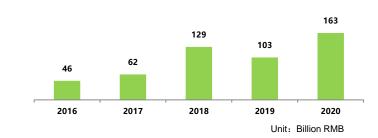
Multiple Successful Listings of Global Biomedical Enterprises



Investment and Financing in Global Health Care Industry (2016-2020)



Investment and Financing in China Health Care Industry (2016-2020)



^{*}Figures above are the total amount of fundraising, Sources: Public news



Business Highlights

Business Highlights





CFS Services

CFS-Drug Discovery Service has delivered over **21,000** protein complex structures and **1,584** independent drug targets

Drug Discovery Service backlog contract amount surged by **73.9%**, reaching RMB**607** million

R&D investment totaled RMB66 million, 46.7% YOY



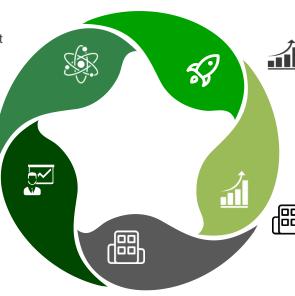
Customers & Backlogs

The total number of customers is 1252 CFS-Drug Discovery Service: More than 543 customers worldwide, repeating customers remains over 85% CFS-CDMO Business(Langhua): 709 customers, repeating customers remains 100%



Strategic Acquisitions

Langhua Pharmaceutical: November 2020 SYNthesis: March 2021



EFS Incubation Portfolio

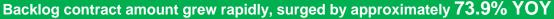
Reviewed **834** projects **Made 20** new investments **10** reinvestment on existing portfolio companies **125** pipelines in total

Employees & Facilitiess

No. of employees increased to **1619** Drug Discovery Service: area of lab. & offices increased to **24,000** sq.m. CFS-CDMO Business(Langhua): The production capacity in 2020 is **671**m³

Internal Growth: Drug Discovery Service





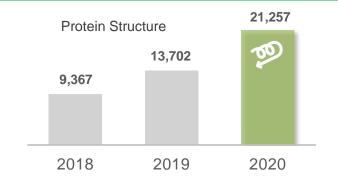


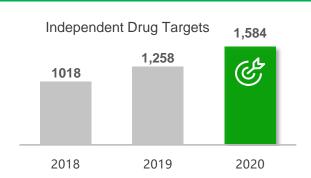


CFS-Drug Discovery Service backlog contract amount **+55%**EFS backlog contract amount **+139%**

The number of protein structures delivered +7,555 and the number of independent drug targets delivered +326

11





CFS-Drug Discovery Service: Optimization of Customer Structure

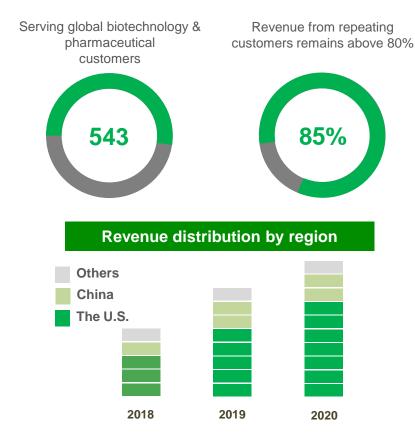


Served Top 10 global pharmaceutical companies Pfizer U NOVARTIS 🖨 MERCK Johnson Johnson BAYER E R gsk GlavoSmithKline SANOFI J Obbvie Takedo Served 35 "Fierce Biotech's Fierce 15" promising biotech companies FORMA FILEDADELITYCES MANUTOR SET ALL SYMMOKARdia

Padlock SYR S () REVOLUTION Full Therapeutics IDEA/A RELAY

TANGO therapeutics:

Beam mana (Included Interapeutics dewpoint) Prelude THERAPEUTICS DEWPOINT AND THE THERAPEUTICS

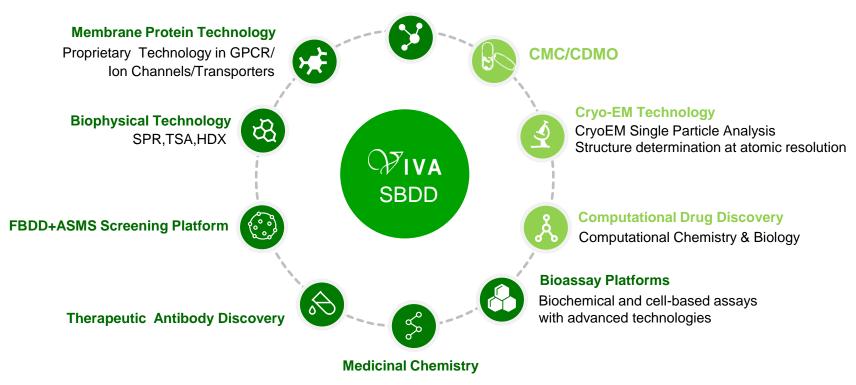


Continued Investment in New Technology Platforms



During the reporting period, the Company's R&D investment totaled RMB66 million, representing a YOY increase of approximately 46.7%

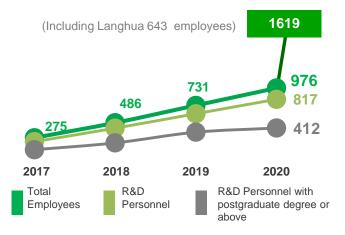
Protein Production & Structural Biology



Rapid Increase in Employees & Scale of Facilities







- In Sept 2020, signed investment agreement with Qiantang, Hangzhou
- In Jul 2020, successfully entered into a bid for a new property in Zhoupu, Shanghai
- In Apr 2020, Chengdu New Drug Incubation and Production R&D Center commenced construction
- In Feb 2020, acquired the property located in Shanghai.

Strategic M&A: Construction of an Integrated Platform





2020.11

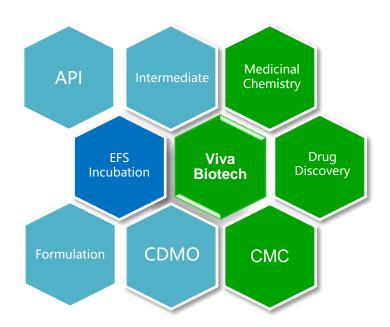
Completed the very substantial acquisition of 80% of the equity interest in Zhejiang Langhua Pharmaceutical Co., Ltd.



2021.3

Completed the acquisition of 100% equity of SYNthesis med chem (Hong Kong).

- ◆ Diversifying regional coverage, enhancing business development in North America/Asia/Europe/Australia
- Strengthening the existing CMC/CDMO business and expanding into formulation production
- ◆ One-stop services from R&D to production



CFS-CDMO: Langhua Business Highlights







Financial Highlights

- Revenue RMB1,518 million, an increase of 22.67% YoY
- Net profit was RMB171 million, an increase of 134.38% YoY
- CDMO business achieved sales of RMB875 million, with an increase of RMB310 million, and a growth rate of 54% YoY
- The commercialization of reserve varieties and the increasing demand for commercial varieties have become the main reason for the growth



Customer Base

- A total of **709** customers have been served, and the retention rate of the top 10 customers is **100%**
- Independently produced 95 varieties of API and CDMO projects



- The production capacity in 2020 is **671 m³**, and it is expected to increase to **1,500 m³** in 2023
- 3 major laboratories (Shanghai, Ningbo, Taizhou),
- Develop continuous reactions, enzyme catalysis, photo-redox reactions, and establish a technology platform for regular formulation and special formation

2020 Langhua Pharma Projects







Completed 48 CDMO projects, 10 out of which had gross profits exceeding 5 million

1

The sewage pretreatment workshop, MVR, desalination dryer, membrane treatment equipment, etc. were put into use, and the upgrading of the sewage station was completed

A new process was put into use in the spironolactone workshop, and the registration of levofloxacin in the United States was completed

3

The civil engineering construction of Workshop T02 (4 floors above the ground) has been completed and is **in process of installing equipment**

4

19 new varieties (12 APIs and 7 advanced intermediates) obtained the environmental permits

5

Accelerate the Construction of CDMO Platform



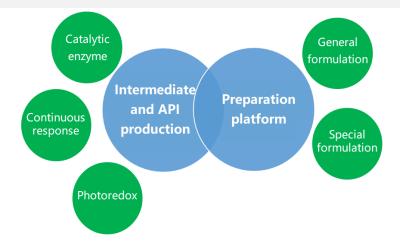


Develop innovative technologies and establish a formulation platform

- Develop green chemical technologies such as continuous reactions, enzyme catalysis and photo-redox reactions to produce pharmaceutical intermediates and APIs
- Establish a technology platform for regular formulation and special formation (such delayed release, nanometer, liposome, etc.)

Integrated one-stop service covering all stages of production

- Continue to increase investment in R&D capabilities and expand existing R&D centers in Taizhou and Ningbo
- Establish a Shanghai R&D center staffed with 300~400 people, including drug intermediates, API and formulation R&D center, covering the process development, scale-up production and application of intermediates and drugs at all stages of preclinical, clinical and commercialization of APIs and formulations



Optimize EHS, Quality, Automatic Production System, Expand Production Capacity



Consolidate the EHS system and improve the quality system

- Continue to increase investment in EHS and to benchmark international advanced standard
- Continue to improve the quality control system to meet the audit requirements of various countries on GMP





Enhance the automation in our workshops to Improve efficiency and utilization

- The production capacity in 2020 is 671m³, and it is expected to increase to 1,500 m³ in 2023 so as to significantly increase production Estimated capex of RMB 960 million
- Introduce the use of the latest equipment to improve the level of automation and efficiency

EFS: Further Expand Incubation Portfolio



2020 New Added Companies











































Reviewed 834 Projects in 2020

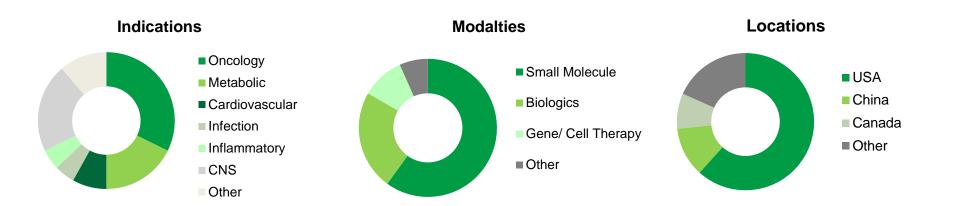
Invested **21** New Companies

Additional Invested 10 Companies

Companies at SPA

Diversified Modalities & Locations & Indications





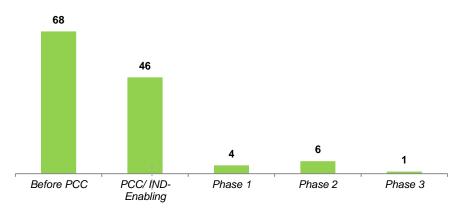
As of Dec 31, 2020, Viva Biotech had a total of 67 portfolio start-ups, the average agreed shareholding ratio was 19.31%.

^{*} As of Dec 31, 2020, Based on Accumulated Incubation Companies

Fast Growth in Pipelines/ R&D Progress



125 Portfolio Pipelines in Total



*As of 12, 31, 2020, Based on Accumulated Incubation Companies

About 55% entered PCC/IND-enabling stage; 11 pipelines at clinical stage







Released Phase 2 Clinical

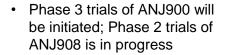
Clinical Trial

Data of AR882

First patient dosed in Phase 1













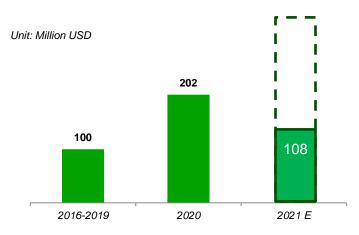
 FDA granted IND approval for Q-1802: Clinical trial application accepted by NMPA

Note: The R&D progress of the incubated project comes from its publicly released press release or announcement.

Rapid Increase in Follow-on Investment



Follow-on Investment Increased Rapidly



Portfolio Follow-on Investment

In 2020, **35** new investment institutions participated in the follow-on financing of Viva portfolio companies



A+ Round 20 MN USD



C Round



C Round



A+ Round



A Round 42 MN USD

2020



B Round 100 MN USD



A Round 10 MN CAD

2021

Note: The financing progress of the incubated project comes from its publicly released press release or announcement.

Dogma Therapeutics Reached the Acquisition Agreement with AZ



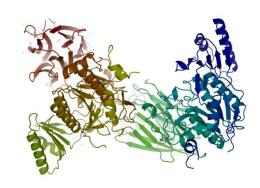
Investment Time: January 2017

Equity ownership: 14.40%

Reached the Acquisition Agreement on Oral PCSK9 Inhibitor with AstraZeneca in September 2020



- PCSK9 is a highly-sought-after target for the industry in past ten years;
- Structure-enabled discovery makes it possible for success;
- Speedy lead optimization completed in 1.5 years;
- First small molecule PCSK9 inhibitors with excellent properties were discovered;



(PCSK9 protein structure)

Constantly Optimize Innovative Biotech Ecosystem





2021 Partnership Summit

- **√60+** portfolio companies
- **√230+** global investors attended
- **√300+** 1-on-1 meetings

VBI DEMO DAY 2020

- **√6+8** Portfolio Companies
- **√200+** Investors Each Demo Day

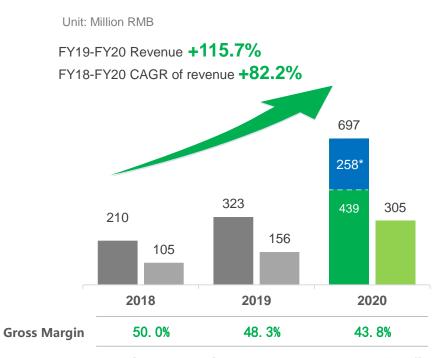




Financial Performance

Revenue and Gross Margin





Rapid Growth in Revenue and Gross Profit

*During the reporting period, the revenue contribution from the consolidated statement of the acquisition of Langhua Pharma was RMB25.8 million

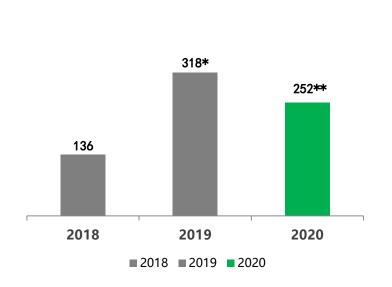




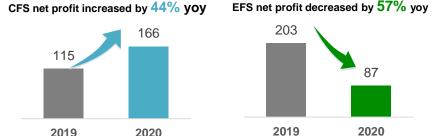
Net Profit



Net Profit of CFS & EFS



RMB '000	2019	2020
Net profit excluding extraordinary profit of loss	318,019*	252318**
Net profit of EFS business***	203,268	86,729
FVTPL Gain	212,700	106,941
Disposal Gain	11,684	6,303
Share of loss	(1,908)	(487)
VBI Expense	(15,266)	(26,028)
EFS Income Tax Expenses	(3,942)	0
Net profit of CFS business	114,751	165,589



*2019 net profit of 265.87 million has been adjusted to deduct non-recurring costs and add back non-recurring income, which include: the one-time-costs preference share financial liabilities of 34.33 million and the listing fee of 17.91 million, to reach the adjusted net profit of 318.02 million

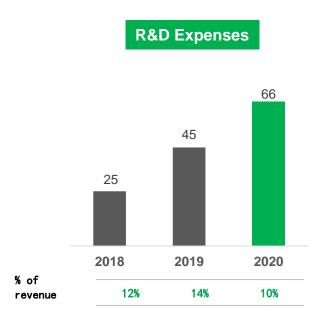
**2020 net loss of 378.87 million has been adjusted to deduct non-recurring costs and add back non-recurring income, which include: the fair value loss on financial liabilities at FVTPL of 547.05 million, interest expenses of the debt components of the Convertible Bonds of 52.12 million, transaction costs relating to the derivative component of the Convertible Bonds of 5.07 million, gain on repurchase of the Convertible Bonds of 4.45 million, transaction costs relating to the acquisition of 15.58 million and amortization of acquired assets of 15.82 million

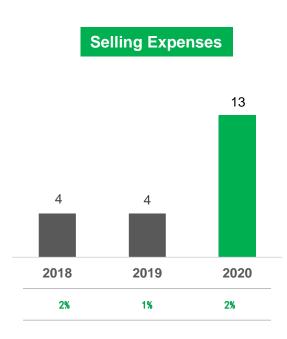
***Revenue from EFS business is calculated based on fair value gain on financial assets fair value through profit or loss and other gains and losses relating to incubation business.

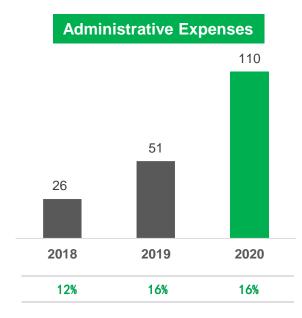
R&D, Selling & Administrative Expenses



(As of the end of 2020, Unit: million)







Fund raising activities to accelerate the Construction of an Integrated Platform



2020.2



Issued **US\$180 million** convertible bonds due 2025 guaranteed by the Company.

2020.12



Issued **US\$280 million** 1.00% convertible bonds due 2025 guaranteed by the Company

Such financing activities provided sufficient financial support for the Company in terms of business expansion and integration in the downstream of the industry chain, its business development and daily operations.

(The Group's financial cost during the reporting period was approximately RMB 62.1 million, mainly due to the increase in interest on the convertible bond, etc.)

2020.7



The Company placed an aggregate of 130,000,000 shares at a placing price of HK\$8.15 per placing share, raising net proceeds of approximately HK\$1,050 million.



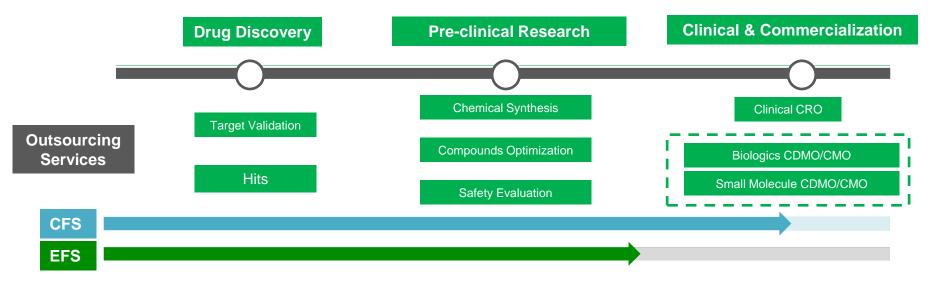
Recommended a final dividend of HKD1.0 cents per share by cash with a total amount of approximately **HKD 20 million**.



Strategic Development

Future Development Strategies





To explore downstream of novel drugs R&D industrial chain:

- · Cooperation with top enterprises in various industry segments
- To avoid negative factors including high investment and timeconsuming in industrial landscape
- To build up capability of providing quality services to customers in all-stage rapidly

To expand the scale and depth of industrial capital investment continuously:

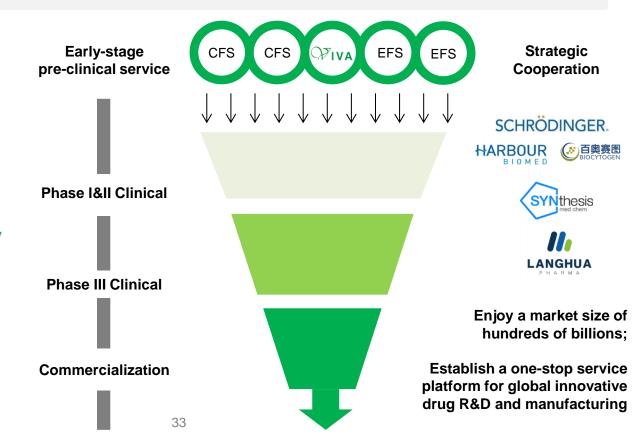
- · Cooperation with several top early stage VC institutions
- · Co-investment, contribution and investment research
- Industrial investment fund led by Viva to invest in middle or even late stage novel drugs R&D enterprises/projects

Build up Service Capabilities by "Following the Startups"



R&D Outsourcing=Discovery/Preclinical CRO->CDMO->Clinical CRO

- Working along with CFS/EFS clients from the beginning; Establish services going downstream
- Build through strategic collaboration with cost efficiency
- High quality client base of startups with integrated services





Q&A Session







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