

## **GlycoNex Inc. 2024 Investor Conference**





synergistic effects and escape from drug resistance in cancer therapy.

#### Using Anti-Glycan Antibodies to Overcome Protein Therapy Resistance





### **Pipelines**



Antibodies targeting tumor-associated carbohydrate antigens

#### Innovative programs of ADC-centric anti-glycan therapy







### 2023-2024 Accomplishments

### • GNX102 (First-in-Class Antibody)

• Phase 1 Completion in the US and TW: Excellent safety profile and good patient tolerance.

### • GNX1021 (ADC)

- Demonstrated preclinical efficacy in gastric cancer models
- Good tolerability in rats and comparable to benchmark ADC
- Manufacturing-ready

### • Denosumab Biosimilar

- Phase 1 completion in Japan
- Phase 3 trial ready to launch



# GNX102

### Finished Phase-1 clinical trial



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## GNX102 phase1 clinical trial result

46 patients#小細胞肺癌10 kinds of tumors用癌10 kinds of tumors大腸直腸癌Two dosing designs子宮頸癌US and Taiwan子宮上皮癌Excellent safety result膀胱癌



# GNX1021 (ADC)

Scheduled to enter phase 1 clinical trial in 2026 Q2 (Taiwan, Japan)



### Antibody-Drug Conjugate, ADC

#### 抗體藥物複合體結構

在單株抗體(標靶)上,以連接子攜帶化療藥物, 可將藥物精準投放在腫瘤細胞上,攜帶的化療藥 物可以多達4個甚至8個。可達到比過去化療更 好的毒殺效果,又能避免傷害正常組織。







# One Dose of GNX1021 (ADC) can suppress the growth of gastric cancer

GNX102 單株抗體之腫瘤抑制效果(多劑量)

GNX1021 ADC 之腫瘤抑制效果(單劑量)





### Strategic Focus and Timeline for 2025 and Beyond

#### GNX1021 development

- ➢ Preclinical Phase (Q4 2024 Q3 2025)
  - Complete Toxicology Studies (by Q3 2025)
  - Manufacturing and CMC (by Q4 2025)
  - Data Package Preparation for IND Submission (Q4 2025)
- ► IND Submission (Q1 2026)
- Phase 1 Clinical Study Launch (Q2 2026)





# Denosumab biosimilar (SPD8)

### Finished phase-1 Scheduled to enter phase-3 in 2024 Q4

### Denosumab biosimilar (SPD8) Phase-1 results

- 110 patients
- Similar PK data
- No serious side effects
- Equivalent safety





### Strategic Focus and Timeline for 2025 and Beyond

#### Denosumab Biosimilar

- Indication: Osteoporosis and preventing bone complications associated advanced cancer
- Phase 3 clinical trials are scheduled for initiation in Q4 2024, aimed at way for regulatory approval and commercialization by 2027.
- > Plans for multi-country expansion into other Asian markets are in

#### Market Potential:

Positioned to capture significant market share in the biosimilar space where demand for affordable biologic alternatives is rising.



# 200L pipeline

### Phase-1 drug production

- For cost and time efficiency

### Antibody Manufacture Capability





抗體開發

量產細胞株開發



分析方法建立



產程開發



臨床試驗抗體製造 200 L pipeline



- Phase-1 drug production
- Complied with Japan GMP regulations

治験薬の製造管理、品質管理等に関する基準

#### GlycoSH: Anti-glycan Antibody Bank





## **Pipeline Strategy & Growth Potential**

#### • Diversified Pipeline:

GlycoNex's pipeline spans multiple therapeutic areas to balance risk and maximizing

- innovative drugs vs. biosimilars
- oncology and bone related diseases

#### • Innovative Drug Development:

Our first-in-class and next-generation antibody programs (GNX102, GNX1021, GNX201, positioned to tap into high-value markets with unmet medical needs.

#### • Biosimilar Development:

The development of Denosumab biosimilar in Japan provides a blueprint for future where biosimilars are expected to see rapid adoption.



### **Financial information**

Publicly Traded since 2012/12 company code 4168	Unit: TWD, thousands 2024/06/30
Paid-in capital	1,086,401
Total Assets	1,416,154
Net Assets	1,156,291
Property, plant and equipment	834,339



# GlycoNex Inc.

# THANK YOU!