



**BeyondSpring**  
P H A R M A C E U T I C A L S

**Turning the Tide in Cancer Therapy**  
*Plinabulin - Redefining Oncology Efficacy and Tolerability*

# Disclaimer



This presentation has been prepared for informational purposes only. No money or other consideration is being solicited, and if sent in response, will not be accepted. This presentation shall not constitute an offer to sell, or the solicitation of an offer to buy, any securities, nor shall there be any sale of these securities in any state or jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state or jurisdiction. The Company is not under any obligation to make an offering. It may choose to make an offering to some, but not all, of the people who indicate an interest in investing. The information included in any registration statement will be more complete than the information the Company is providing now, and could differ in important ways.

This presentation and any accompanying oral commentary contain forward-looking statements about BeyondSpring Inc. (“BeyondSpring” or the “Company”). Forward-looking statements are based on our management’s beliefs and assumptions and on information currently available to our management, including those described in the forward-looking statements and risk factors sections of the Company’s 10-K filed on March 25, 2026 and other filings with the United States Securities and Exchange Commission (SEC).

Such statements are subject to known and unknown risks, uncertainties, and other factors that may cause our or our industry’s actual results, levels of activity, performance, or achievements to be materially different from those anticipated by such statements. In some cases, you can identify forward-looking statements by terminology such as “may,” “will,” “should,” “expects,” “plans,” “anticipates,” “believes,” “estimates,” “predicts,” “potential,” “intends,” or “continue,” or the negative of these terms or other comparable terminology. Forward-looking statements contained in this presentation include, but are not limited to, (i) statements regarding the timing of anticipated clinical trials for our product candidates and our research and development programs; (ii) the timing of receipt of clinical data for our product candidates; (iii) our expectations regarding the potential safety, efficacy, or clinical utility of our product candidates; (iv) the size of patient populations targeted by our product candidates and market adoption of our product candidates by physicians and patients; and (v) the timing or likelihood of regulatory filings and approvals.

Except as required by law, we assume no obligation to update these forward-looking statements publicly, or to update the reasons why actual results could differ materially from those anticipated in the forward-looking statements, even if new information becomes available in the future.

The market data and certain other statistical information used throughout this presentation are based on independent industry publications, governmental publications, reports by market research firms or other independent sources. Some data are also based on our good faith estimates. Although we believe these third-party sources are reliable, we have not independently verified the information attributed to these third-party sources and cannot guarantee its accuracy and completeness. Similarly, our estimates have not been verified by any independent source.

By attending this presentation, you acknowledge that you will be solely responsible for your own assessment of the market and our market position and that you will conduct your own analysis and be solely responsible for forming your own view of the potential future performance of our business.

# Company Highlights



## BYSI: Oncology-focused Global Biotech

- **Company Overview**

Founded 2010 | New Jersey-based, Nasdaq-listed since 2017

- **Late-Stage Clinical Asset**

First-in-Class anti-cancer agent  
Plinabulin

- **Equity owner of SEED Therapeutics**

Clinical-Stage Molecular Glue Co. with investments from Eli Lilly & Eisai



## Plinabulin: Durable Anti-cancer Efficacy and Bone Marrow Protection

- **Unique MoA**

GEF-H1–mediated dendritic-cell (DC) maturation and chemotherapy-induced neutropenia (CIN) mitigation

- **Clinical Efficacy & Safety**

700+ cancer patients treated; Phase 3 OS benefit in 2L/3L EGFR WT NSCLC

- **Long-Term IP Protection**

Composition-of-matter protection through 2036 (potentially extendable through 2041)



## DUBLIN-4: Confirmatory Global Phase 3 in Post-ICI NSCLC

- **Mechanism-based Population**

Post-ICI, non-squamous NSCLC with no actionable driver alternations

- **Clear Regulatory Pathway**

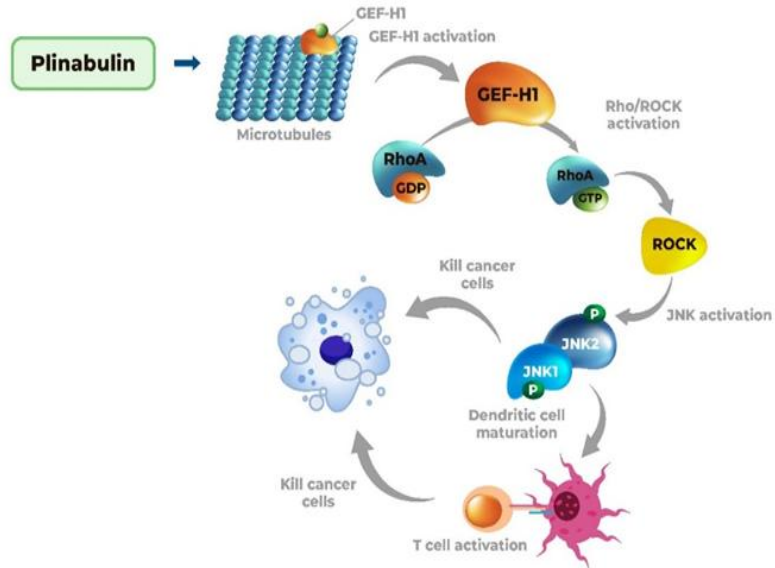
442-patient confirmatory Phase 3; OS primary endpoint

- **Experienced Team & Partners**

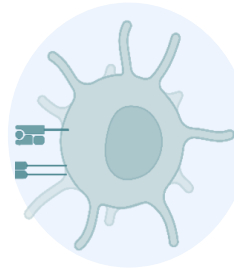
Proven global execution and successful NDA experience

# Plinabulin: Dual Immunomodulatory and Neutropenia-Mitigating Activity

Plinabulin releases GEF-H1 from microtubules, promoting dendritic-cell maturation and reducing CIN<sup>1-6</sup>

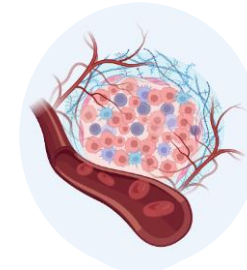


DC Maturation & APC Activation



Targets Acquired Resistance from ICI

Tumor Vasculature Modulation



Targets Non-squamous NSCLC

Increase Chemotherapy Treatment Cycles & Maintain Dose Intensity



Reduces Chemotherapy-induced Neutropenia

1. La Sala et al., *Chem* 5(11): 2969-2986 (2019); 2. Kashyap et al., *Cell Reports* 28(13): 3367-3380 (2019); 3. Choi et al. *Cell* 189 (2): 461-477 (2026); 4. Tonra JR (2020) *Cancer Chemother Pharmacol* 85:461; 5. Blayney DW (2020) *JAMA Oncol* 6(11):e204429; 6. Chan DCH (2021) *Blood Advances* 5(16):3120

# DUBLIN-4 Targets a Large, Underserved Post-ICI NSCLC Population

Despite strong first-line success, treatment options after progression remain limited

ICI annual sales at ~\$60 billion;  
Post-ICI Unmet Need



Plinabulin overcomes ICI resistance  
via DC maturation & APC activation

Limited Treatment Options for Post-ICI NSCLC without Driver Alteration



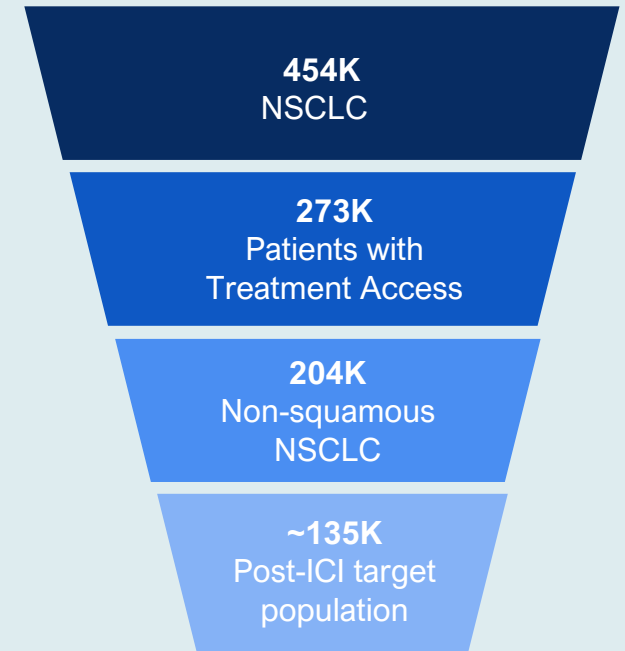
Docetaxel

- Median OS: ~9-11 months
- 40% experience severe neutropenia

12 Phase 3 Studies with no  
OS Benefit vs. Docetaxel

- 8 global trials including anti-PD-(L)1 combos **did not improve OS vs. docetaxel**<sup>1, 2</sup>
- 4 global trials including ADC drugs **did not improve OS vs. docetaxel**

Projected NSCLC Patient Pool  
(US + EU<sup>3</sup>)



Potential patient pool ~135K  
(2037 estimated<sup>4</sup>)

1. Malinou J et al., ASCO 2024; 2. Dragnev KH et al. ASCO 2025;

3. EU5 represents Germany, France, United Kingdom, Italy and Spain; 4.2037E represents the estimated patient pool at projected peak sales, assumed seven years post-launch.

# DUBLIN-3: Phase 3 Foundation for DUBLIN-4

## DUBLIN-3 Study: 2L/3L EGFR WT NSCLC after progression on platinum therapy

N=559

R

Single-Blind • Randomized  
1:1 ratio; 21-days per cycle

**DP:**

Docetaxel  
(75 mg/m<sup>2</sup>, day 1)  
+ **Plinabulin**  
(30 mg/m<sup>2</sup>, day 1, 8)

**D:**

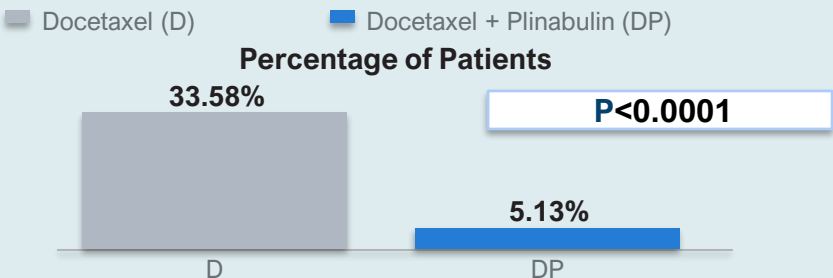
Docetaxel  
(75 mg/m<sup>2</sup>, day 1)  
+ Placebo (day 1, 8)

- **Geography**  
~87% China / ~13% U.S. & Australia
- **Limited Prior PD-(L)1 Exposure**
- **Primary Endpoint**  
Overall Survival — Met

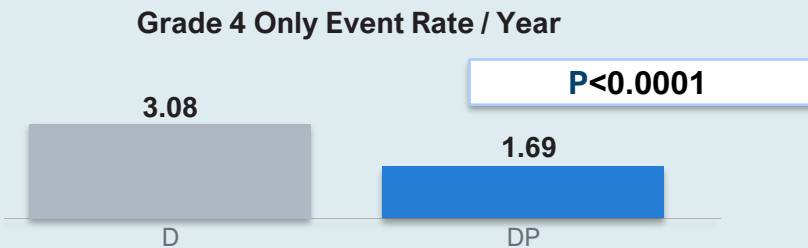
## Established Safety Benefit

### DP with significantly lower Grade 4 Neutropenia

- All Cycles Day 8 - Day 8 is ANC nadir for docetaxel



### DP has more cycles of treatment; DP with significantly lower exposure-adjusted Grade 4 AEs



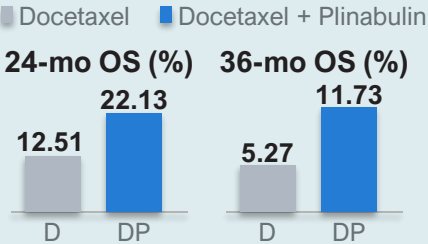
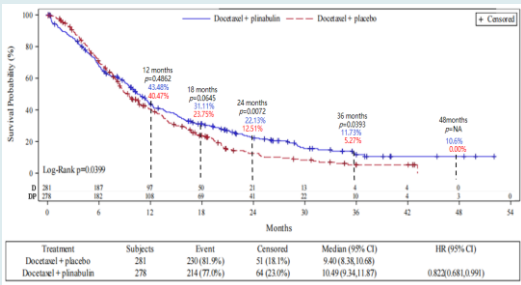
# Clinical Efficacy Supports DUBLIN-4 Plinabulin-Mechanism Patient Selection

DUBLIN-3 efficacy in post-ICI subgroup signal and Study 303 prospective study data provide the clinical rationale for DUBLIN-4 in a Plinabulin-mechanism targeted population:

- Non-squamous NSCLC + Progressed on ICI

## DUBLIN-3 Phase 3 (ITT)

- Global randomized study
- EGFR WT 2L/3L NSCLC (N=559)
- OS HR=0.82, log rank p=0.0399

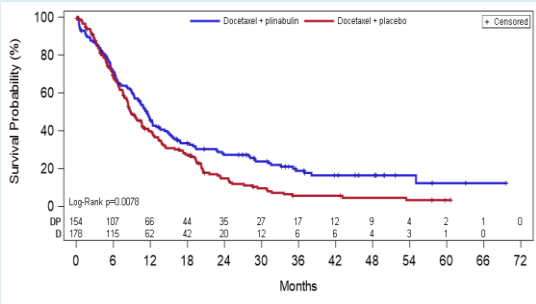


Phase 3 long-term OS benefit

1. DUBLIN-3: *Lancet Respir Med* 2024

## DUBLIN-3 Non-squamous

- Non-squamous (N=332)

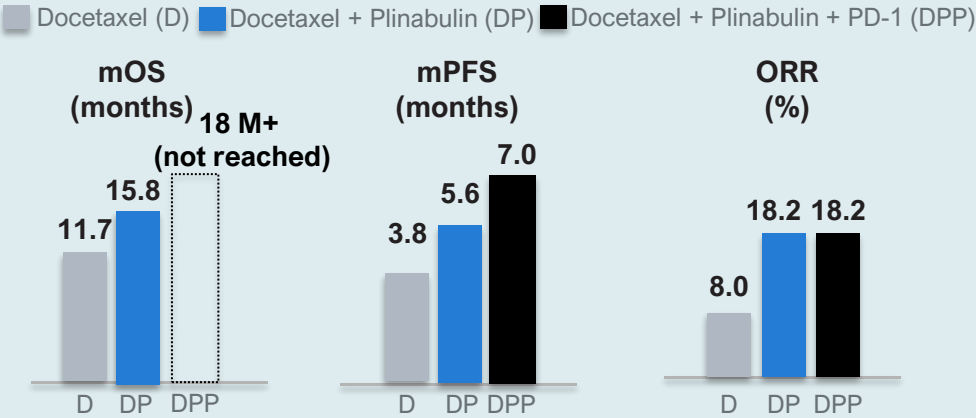


- mOS: 11.37 vs. 8.81 mo
- OS HR 0.72
- Log rank p=0.0078

mOS extension of 2.5 months

## DUBLIN-3 Post Hoc<sup>1</sup> + Study 303 Prospective<sup>2</sup> in NSCLC Post-ICI

1. **DUBLIN-3 Post-hoc:** Non-squamous, post-ICI EGFR WT NSCLC (D: n=25; DP n=22)
2. **Study 303 Prospective:** post-ICI NSCLC (n=47)



### DUBLIN-3 (DP vs. D)

OS HR 0.55; PFS HR 0.67; ORR 18.2% vs 8.0%.

### Study 303

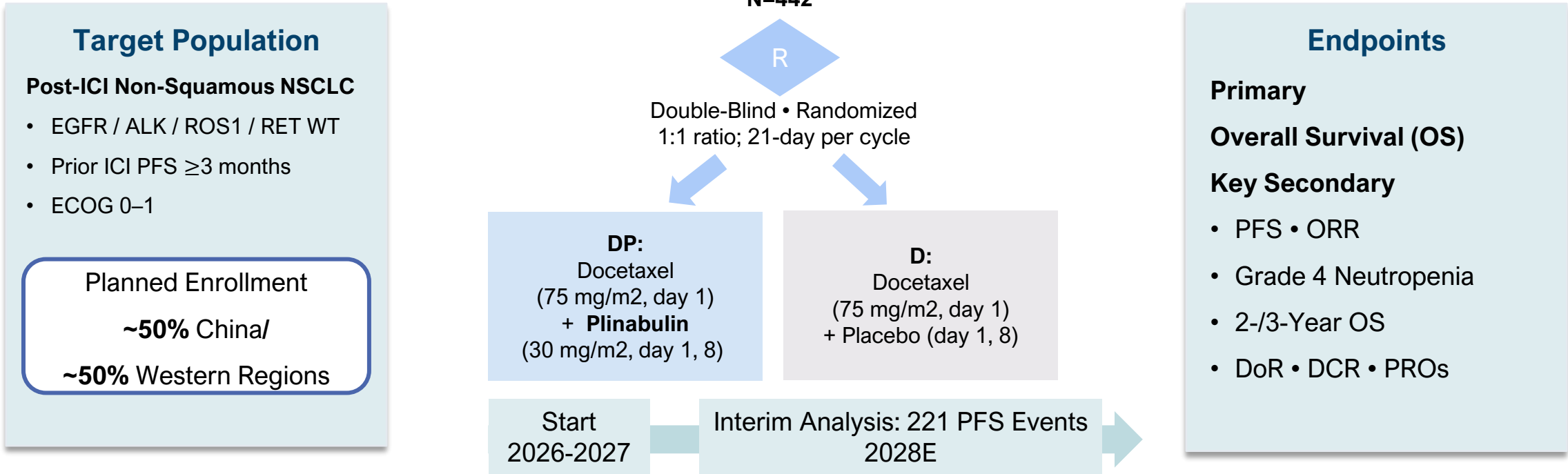
2-year OS 58%; DCR 80%; mPFS 7.0 M, ORR 18.2%.

Data from both studies showed superior OS, PFS, and ORR vs. historical data of SOC docetaxel in a similar population (Docetaxel: mOS 11.8 M, mPFS 3.8 M, ORR 12.8%<sup>3</sup>)

2. 303 Study: *ASCO* 2026; 3. *Thoracic Oncology* 43(3): 260-272 (2024)

# DUBLIN-4 at a Glance: Confirmatory Global Phase 3 Study, FDA Fast Track Designation

Docetaxel + Plinabulin vs. Docetaxel in Post-ICI, Driver-Negative Non-Squamous NSCLC



## Why We Can Deliver DUBLIN-4

**Experienced Team**  
Global oncology & regulatory experience

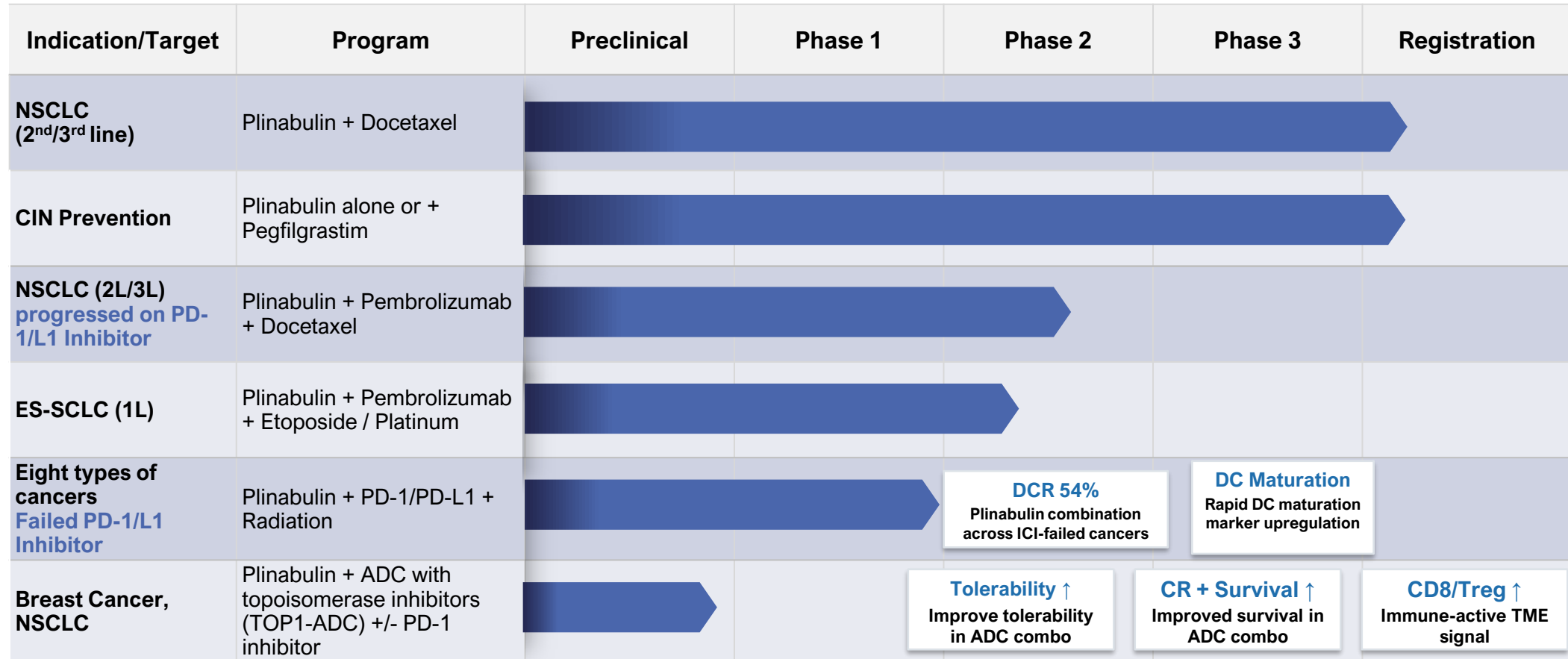
**Phase 3 Track Record**  
2 completed Phase 3 programs

**Established Clinical Network**  
Sites, investigators & CRO partners

**DUBLIN-3 Know-How**  
Directly informs DUBLIN-4 execution

# Plinabulin: One Drug, Multiple Clinical Pipelines

700+ patients treated with Plinabulin



# Plinabulin: Clear Path to Value Creation in NSCLC & Beyond

From Clinical Evidence to DUBLIN-4 and Commercial Potential



## DIFFERENTIATED ASSET

Clinical & Safety Evidence

- Novel GEF-H1 mechanism
- >700 pts treated; Positive Phase 3 OS
- Increased chemotherapy treatment cycles and significant reduction in CIN

## PHASE 3 OPPORTUNITY

DUBLIN-4 – MoA targeted  
NSCLC post-ICI

- 442-patient global confirmatory study with OS as primary endpoint
- Interim analysis expected 2028

## VALUE CREATION

Market & Commercial Path

- Large post-ICI patient population
- Long-term commercial potential

# Experienced Leadership and SAB Support to Drive Successful Clinical Execution



**Min Qiu**  
Chief Executive Officer

Biopharma leader with 10+ years in strategy, financing, business development at public traded companies and China Merchants Bank.



**June Lu, PhD**  
Chief Scientific Officer

R&D leader with 26 years of experience at Endocyte/Novartis. Areas include cancer, autoimmunity, drug resistance, CAR-T and RLT.



**Na Li**  
Chief Financial Officer

Executive with 20+ years in FP&A, treasury, financing at BeyondSpring and public companies.



**John Mao, PhD**  
SVP, Development

Biopharma leader with 30+ years in global development and regulatory strategy at Foresee and Cytokinetics.



**Helein Li, MD**  
VP, Clinical Science

Clinical development leader with 20+ years in trials, regulatory strategy, and safety at BeyondSpring, Sanofi, Roche.



**Hao Zhang**  
Senior Director, CMC

CMC leader with nearly 10 years in manufacturing, supply chain, and regulatory execution.



**Joy Jia**  
VP, Finance

Finance leader with 10+ years of expertise in SEC reporting, SOX, FP&A, audits, capital markets, and controls.



**John Ruckdeschel, MD**  
FOX Chase Cancer Center

Oncology leader with 30+ years of experience. Prior CEO at Moffitt, Karmanos, Mississippi cancer centers.



**Adam Grippin, MD**  
MD Anderson Cancer Center

Physician-scientist advancing therapeutics, immunotherapy, and dendritic cell activation strategies.



**Stephen Lin, MD**  
MD Anderson Cancer Center

Thoracic oncology physician-scientist advancing radiotherapy, immunotherapy, and biomarker-driven research in lung cancer.



**Trevor Feinstein, MD**  
Piedmont Cancer Institute

Thoracic oncologist and research leader with expertise in lung cancer. Head of Lung at OneOncology in the US.



**Alfred Zippelius, MD**  
University Hospital Basel

Translational oncology leader in cancer immunotherapy, checkpoint resistance, ADC, and cellular therapies.



**Michel Steinmetz, PhD**  
Paul Scherrer Institute

Structural biology leader with 20+ years in microtubule research, cryo-EM, biophysics, and drug discovery.



**William Decker, PhD**  
Baylor College of Medicine

Immunotherapy expert with 20+ years in dendritic cell biology and cellular therapies.

# Complementary Board with Finance, Strategic and Business Excellence



**Brendon Delaney**

Director

Seasoned commercial leader with 25+ years' experience, leading global oncology product launches at Constellation, Immunomedics, Celgene, Novartis, and Genentech.



**Jen Majeti, PhD, MBA**

Vice Chairman

Biotech investor and pharma/biotech executive with 20+ years' research and business experience, leading global collaborations at Erasca, Roche, BioDuro, and Amgen.



**Lan Huang, PhD**

Co-Founder & Chairman

Biotech entrepreneur with 20+ years across U.S. and China; founder of multiple companies, MSKCC-trained scientist with *Science/Nature* publications, inventor with multiple patents.



**Mathew Kirkby, MA**

Director

Seasoned finance leader with over 20 years of global banking experience, with senior roles at HSBC, CIMB, RBS, and ABN AMRO across Asia and Europe.



**Patrick Fabbio**

Director

Seasoned CFO with 25+ years of financial leadership across public and private life science companies, including Protara, Progenics, Catalent, WindMIL, and Sanofi.



**Sihai Xu, MBA**

Director

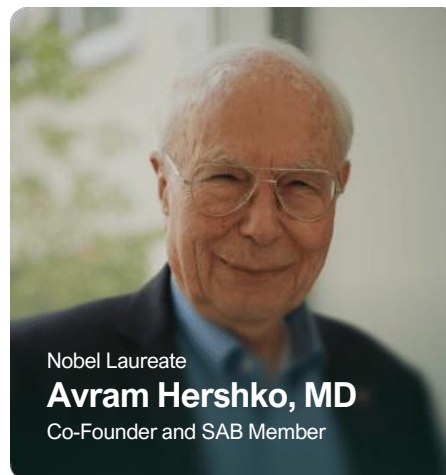
Finance leader with nearly 30 years' experience in China, CFO and board member of public traded companies. Advisor for pre-IPO, IPO, and restructuring transactions across China and Hong Kong.

# **Create The Glue Capture The Protein Eliminate The Disease**

De Novo Molecular Glues for Protein Targets  
Beyond the Reach of Traditional Therapies

# SEED Was Co-founded By Pioneers in Targeted Protein Degradation and Eli Lilly

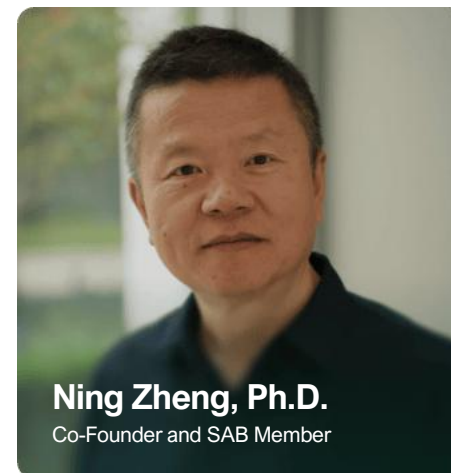
- SEED's differentiation with Target-centric approach to use novel E3s to degrade undruggable targets
  - Scientific capability to use >600 E3 ligases, which consists only 2 structural classes of HECT and RING, solved first by co-founders.
  - Novel E3s or RITE3 use can increase tissue selectivity and scale to reach full therapeutic potential of molecular glues.
- Combined leadership record of 40 INDs and 12 NDAs
- Balanced by an experienced board across finance, risk, legal, and drug development



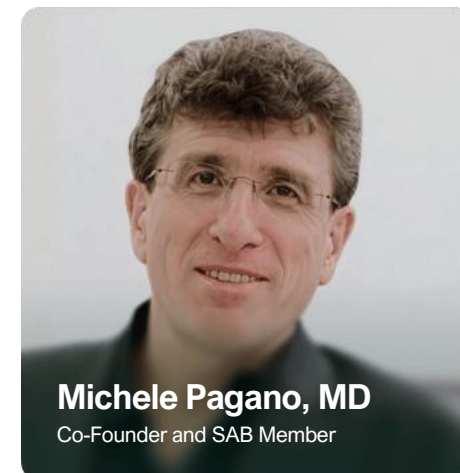
Nobel Laureate who discovered the ubiquitin–proteasome system



Solved the first HECT E3 ligase structure; 20+ years in therapeutic dev.



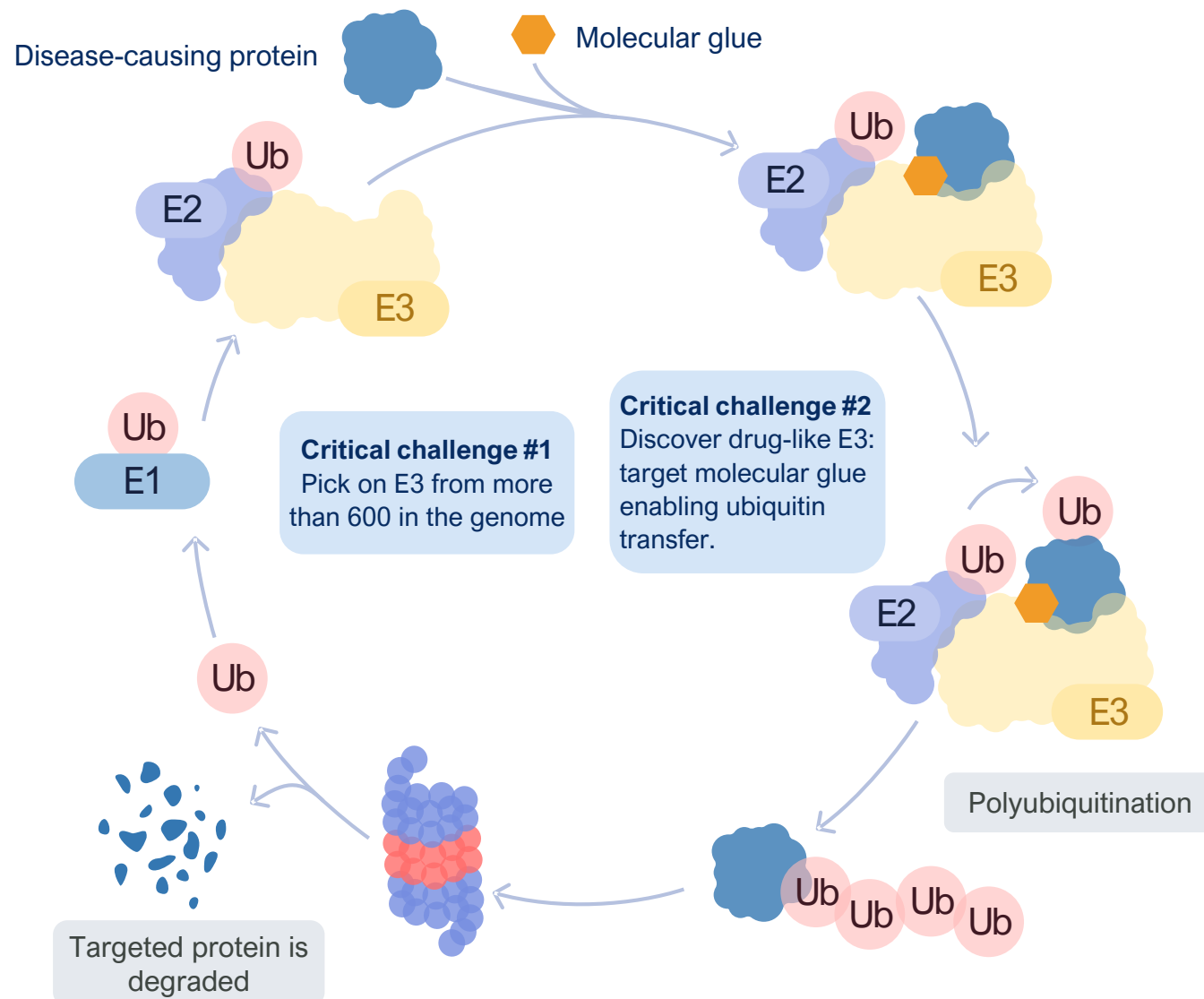
Solved the first RING E3 ligase and coined the term “molecular glue”



Defined SCF ubiquitin ligase and core cell-cycle ubiquitin biology

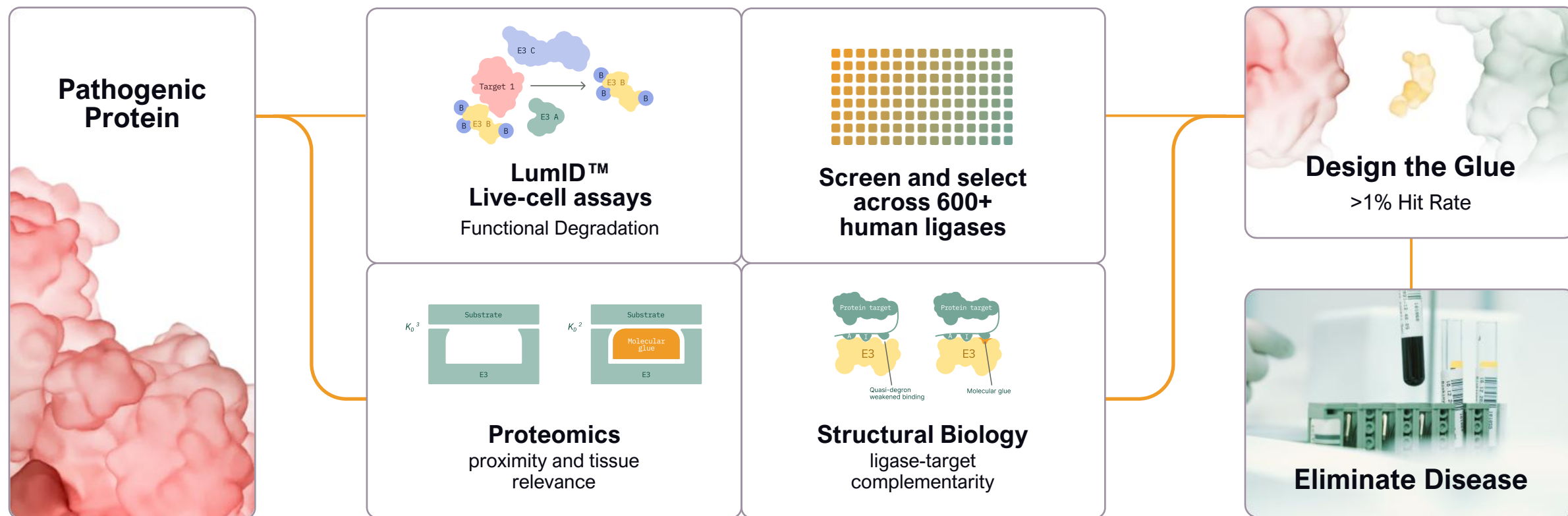
# Elimination, Not Inhibition, Is The Next Therapeutic Paradigm

The ubiquitin-proteasome system (UPS) evolved to tag and destroy aberrant proteins with exquisite selectivity, compounded by molecular glues for precision



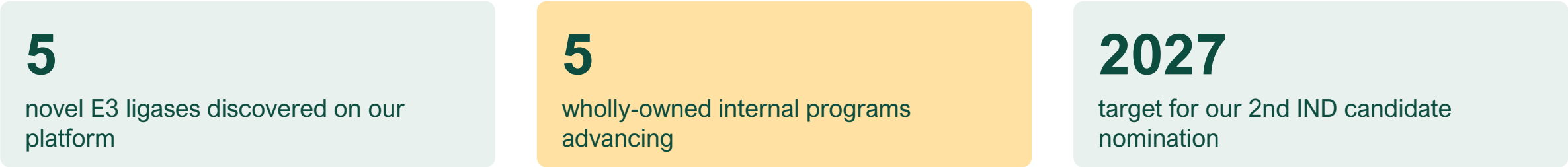
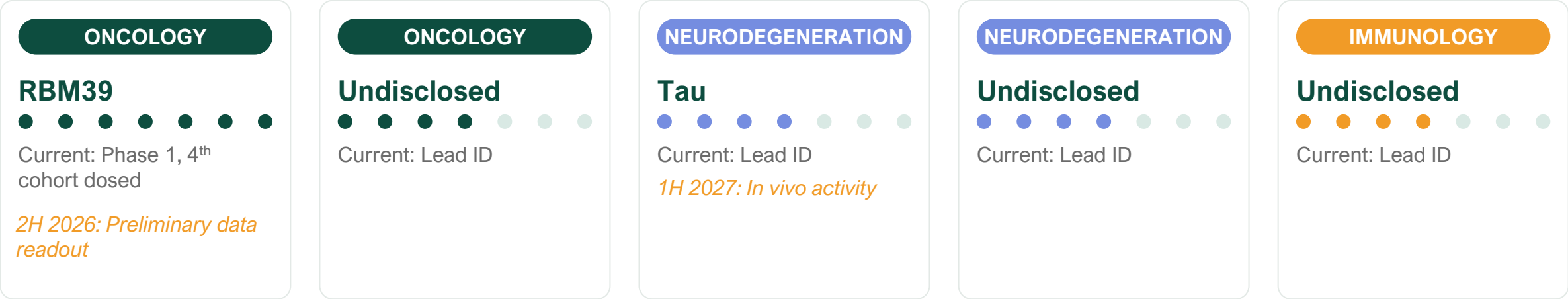
# SEED Built The TPD 2.0 Engine To Find The Optimal Ligase and Degrade Any Disease-causing Protein

SEED's RITE3™ platform unlocks access to more than 600 ligases encoded in the human genome. Then designs custom molecular glues to bind them. More ligases. Better matches. Higher hit rates.



# SEED's Internal Pipeline Spans Three Therapeutic Areas

SEED Therapeutics is advancing 5 wholly-owned programs in oncology, neurodegeneration, and immunology, powered by 4 novel E3 ligases discovered on our proprietary platform.



# RBM39 is a Master Regulator of Oncogenic Splicing

## BIOLOGY

### Why it matters

- Master regulator of oncogenic RNA splicing programs essential for tumor survival
- Splicing machinery is non-enzymatic and historically undruggable

## MECHANISM

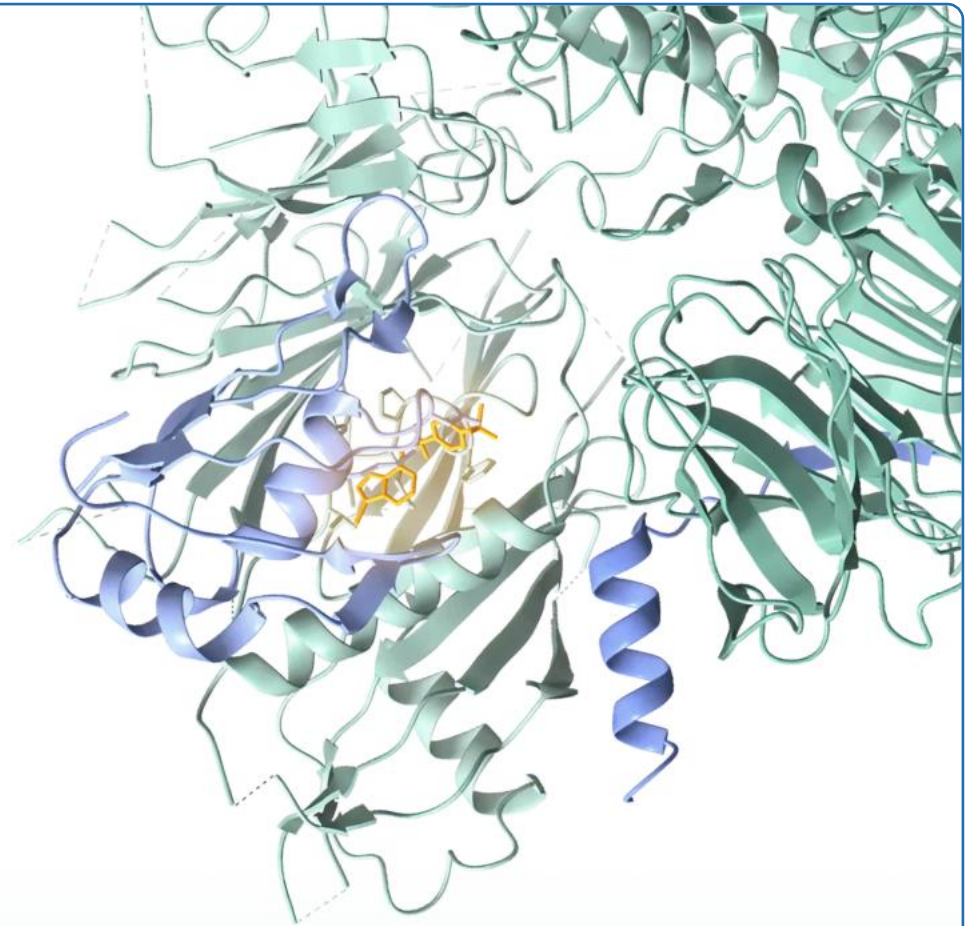
### Why it works

- Molecular glue-mediated degradation via DCAF15 selectively eliminates RBM39
- Induces lethal mis-splicing in tumors while sparing normal cells via redundancy

## VALIDATION

### Why it's de-risked

- Genetic + pharmacologic degradation drives strong anti-tumor effects
- Broad dependency across Ewing sarcoma and multiple solid tumors



RBM39-DCAF15 complex structurally solved → enables next-gen precision degraders

\*Structure of DCAF15-RBM39 complex solved by Prof. Zheng

# Clinical Success Path for ST-01156: Validated Target, Differentiated Molecule, Proven Team

Three independent layers of evidence (target, molecule, team) support a high probability of clinical success.

## 1 VALIDATED TARGET

### RBM39

*Biology, structure, and genetics all confirm RBM39 as a tractable oncology target.*

- **Mechanism:** RNA splicing regulator
- **Structural proof:** DCAF15 E3 ligase resolved
- **Genetics:** depletion slows tumor progression and improves survival in vivo
- **Disease relevance:** overexpressed across multiple cancers, linked to poor survival
- **Clinical derisking:** Recursion's REC-1245 shows linear PK and on-target engagement

## 2 DIFFERENTIATED MOLECULE

### ST-01156

*Best-in-class profile vs. E7820 and REC-1245, with brain penetrance unique to the class.*

- **Optimized chemistry:** improved potency and metabolic stability, no hERG inhibition
- **Tumor-selective:** degrades RBM39 in tumor, not in normal tissue
- **Wide therapeutic window:** favorable PK profile and target engagement.
- **Tolerability-oriented dosing schedule:** 5 days on / 2 days off schedule supports normal tissue recovery and sustained activity
- **CNS-penetrant:** engineered for brain penetrance

## 3 PROVEN TEAM

### Execution

*Approval-experienced leaders running the program at the top U.S. and China cancer centers.*

Dana-Farber

MD Anderson

Memorial Sloan Kettering

Mass General

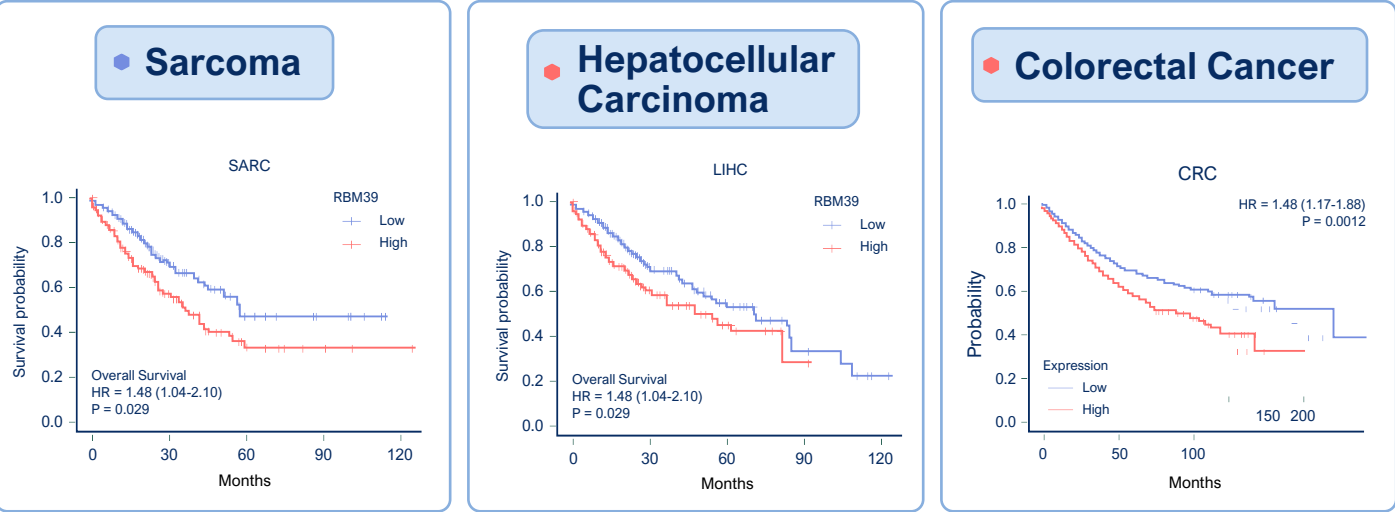
City of Hope

Hoag

Wuhan Union  
Hospital, China

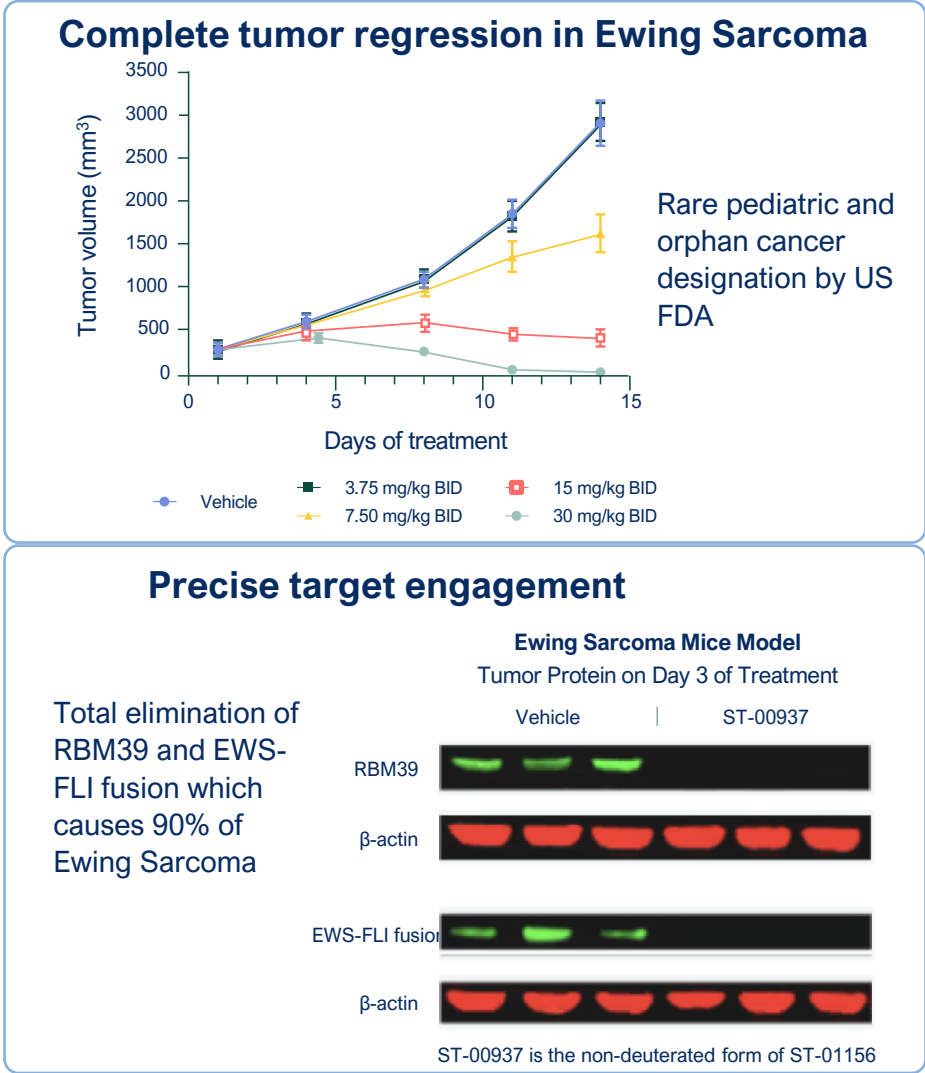
**10+** targeted-oncology approvals  
across the senior development team

# ST-01156: RBM39 Opportunity Across >1 Million Addressable Patients And Supported by Complete Tumor Regression and FDA Designations



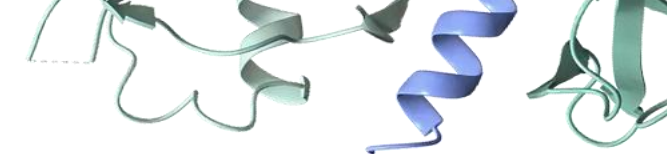
RBM39 overexpression leads to poor patient survival

● Pediatric tumors      ● High prevalence tumors



J Cell Mol Med. 2022;26:4859–4871.  
J Cancer. 2025; 16(7): 2233-2249.

# Highly Experienced Board of Directors



**Lan Huang, PhD**

Co-Founder, CEO and Chairman

Pioneer in E3 structure; Serial biotech entrepreneur with 20+ years of drug development experience, including NDA-ready assets



**James Tonra, PhD**

President, CSO and Director

20+ years of drug discovery experience that led to 5 NDAs; Ex leadership role in Regeneron, Millennium, ImClone, Kadmon, and BYSI



**Linus Lin, PhD**

Director

AVP of Molecular Discovery Capabilities at Lilly Global; Ex leadership role in Lilly Chorus, Lilly China R&D Center, WuXi AppTec, and Merck



**Yoshiharu Mizui, PhD**

Director

Founder and President of Eisai Innovations, Inc.; former Global Business Development and Strategy Head in Eisai's Oncology Business Group



**Jackson Tai**

Director

Retired board member for Lilly, HSBC Holdings, Mastercard; Former DBS Bank CEO, former J.P. Morgan & Co. investment banker; Expert in finance and risk

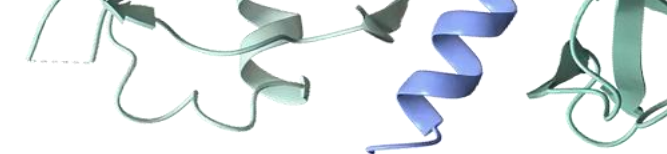


**Ko-Yung Tung, JD**

Director

Former Eisai director, World Bank General Counsel, and lecturer at Harvard and Yale Law School; Expert in law and international business

# World-Class Scientific Advisory Board



## Founding Science: TPD & Ubiquitin Biology



### Avram Hershko, MD

*Co-Founder*

**Godfather' of TPD; 2004 Nobel Laureate;** Distinguished Professor at the Technion; co-discoverer of ubiquitin-mediated protein degradation.



### Ning Zheng, PhD

*Co-Founder*

**HHMI Investigator;** Professor of Pharmacology, **University of Washington.** Pioneered RING Finger E3 structure; coined 'molecular glue' (2007).



### Michele Pagano, MD

*Co-Founder*

**HHMI Investigator;** Chair of Biochemistry, **NYU Medical School.** Global thought leader in TPD biology, ubiquitin-proteolysis, and cancer therapeutics.

## Clinical Development & Oncology



### George Demetri, MD

*SAB Chair, Clinician with Track Record for NDA*

Harvard Medical School Professor; Director, Sarcoma & Bone Oncology at **Dana-Farber** and Brigham and Women's Hospital. Led Gleevec's development; co-founded IDRX (\$1B GSK acquisition).

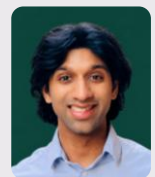


### Baohui Han, MD

*Lung Cancer Expert*

Professor & Honorary Director of Respiratory Medicine at **Shanghai Chest Hospital;** former VP & Executive Dean at Chinese Academy of Lung Cancer. 300+ publications in targeted therapy and immunotherapy.

## Emerging Technology: AI & Neuroscience



### Pranam Chatterjee, PhD

*AI Expert*

Assistant Professor at **University of Pennsylvania;** **AI-driven molecular design** pioneer developing generative models for functional biologics. Co-founded ventures spanning fertility, RNA medicine, and oncology.



### Mansuo Shannon, PhD

*Neuroscience Expert*

Pharmaceutical executive with 18+ years at Bayer, Eli Lilly, Roche/Chugai, and Merck; **AskBio Chief Scientific Officer.** **Developed CNS drugs,** including Lilly's approved Alzheimer's drug.

## Business & Corporate Leadership



### Alan Roemer, MBA, MPH

*Business*

Life sciences executive and board leader: launched multibillion-dollar biotech, **enabled 9 drug approvals, raised ~\$2B,** and led 5 IPOs.

# Breakthrough Investments Highlight the Value of Molecular Glues and Degraders



## Discovery stage TPD assets

Upfront payments of \$35M - \$60M and \$500M - \$5B in potential milestones.



Novartis Sticks With Orionis in Second Deal: \$40M upfront with total value up to \$1.4B



## Pre-IND / IND stage TPD assets

\$100- \$300M in upfront payments and up to \$2B potential milestones.



Gilead eyes Kymera's 'adhesive' cancer drug in up to \$750M deal and \$85M upfront



## Clinical stage TPD assets (Phases I & II)

\$150M - \$650M in upfront payments, \$350M investment and \$2.1B in potential milestones.



Nurix and Roche deal: \$700 M upfront for Nurix's BTK degrader with total value up to \$2.3B

# SEED Therapeutics – Category-defining Molecular Glue Company

## 1 Molecular glue field with clinical validation + value recognition

Molecular glue and TPD validation keeps building: Revolution Medicines (~\$38B market cap), the Orionis–Novartis collaboration at discovery stage (\$40M upfront, up to \$1.4B), and the Nurix–Roche transaction for clinical stage BTK degrader (\$700M upfront, up to \$2.3B).

## 2 Best-in-class RITE3™ platform for multiple molecular glues

Validated by global pharma collaborations and investments following multi-year diligence. Beyond ST-01156, the RITE3™ platform has already produced additional clinic-bound programs, the wholly-owned Tau degrader and a second oncology program, creating multiple future value inflections.

## 3 Experienced team with NDA and M&A track record

Founders are pioneers in targeted protein degradation, and the development team has a track record of 40 INDs and 12 NDAs. A deep board and advisory bench add finance, legal, drug-development, M&A, and commercialization expertise. Global clinical synergies across the U.S. and China to accelerate development with cost efficiency.

## 4 Clinical proof of mechanism in RBM39 target engagement

ST-01156, novel RBM39 degrader: Rapid enrollment with leading U.S. institutions, such as Dana-Farber, MSK and MD Anderson. Enrolling the third ST-01156 dose cohort, with target engagement already confirmed below the dose range where anti-tumor activity is expected. Tumor responses are anticipated in cohort 4 and are possible in the current cohort.

## 5 Efficient path to speedy approval + commercial upside

ST-01156's Ewing sarcoma or neuroblastoma offers an efficient route to human proof of concept and a potential Priority Review Voucher, while colorectal, hepatocellular, and endometrial cancers represent large long-term markets.

The model echoes Parabilis' record \$670M biotech IPO: a rare-disease beachhead with parallel large-indication development.

## 6 Upcoming value-inflection catalysts

ST-01156 Phase 1b completion potentially supporting accelerated NDA in orphan indications in 2029, and Phase 2 pivotal study of larger indications, and a second IND candidate in 2027 and its preclinical development.



**BeyondSpring**  
P H A R M A C E U T I C A L S

# Thank You

General Inquiries: [GENERAL@beyondspringpharma.com](mailto:GENERAL@beyondspringpharma.com)

Investor: [IR@beyondspringpharma.com](mailto:IR@beyondspringpharma.com)

Media: [PR@beyondspringpharma.com](mailto:PR@beyondspringpharma.com)

[www.beyondspringpharma.com](http://www.beyondspringpharma.com)