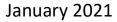




Antibody Therapeutics for Immunological Diseases



Industry leaders spearheading the way















Sean Bohen, MD. PhD. Board of directors

CEO of Olema Oncology; Former CMO of AstraZeneca

Previously:Genentech
Stanford University



Former CEO of Castlight Health.

Previously:
Achaogen
Genentech
Renovis

Judy Chou, PhD. President & CEO Board of directors

Former Global head/SVP of Biotech at Bayer.

Previously:
Pfizer/Medivation
Tanvex Biopharma
Genentech
Wyeth
Abbott

Harvard Medical School

Patrick Yang, PhD. Chairman

Former EVP of technical operations at Juno Therapeutics.

Previously:
Roche
Genentech
Merck & Co.

Stephen Juelsgaard, DVM, JD. Board of directors

Former EVP, General Counsel & chief compliance officer at Genentech.

Previously:Rio Grande Neurosciences
Wilson Sonsini

Corsee Sanders, PhD. Board of directors

Member of Board of Trustee, Fred Hutchinson Former EVP of Development at Juno Therapeutics.

> Previously: Roche Genentech

Joe McCracken, DVM. Head of business development

Former VP & global head of development & licensing at Roche.

Previously:Genentech
Aventis Pharma



PSGL-1, the master regulator of late stage T-cell life cycle

DAY 1 |||||||| DAY 5 ||||| DAY 7 |||||| DAY 7 ||||| DAY 7 ||||| DAY 7 |||||| DAY 10

DEFENSE

The body's defense mediating immune cells against infectious diseases and cancer

PSGL-1 REGULATION

HOMEOSTASIS

The body follows its natural path of T-cell death, remaining healthy

Naïve Attack of Activation Expansion of Conventional Infectious Effector T-cells activated T-cells

T-Cell Pathogen

No apoptosis induced

Mechanism is I

through PSGL-1

Apoptosis Selective

Selective apoptosis of late-stage activated T-cells

Mechanism is Independent to cell migration and has no impact to platelet function

blood 2004 104: 3933-3942 Prepublished online June 15, 2004; doi:10.118/blood-2003-05-1679

Cross-linking of P-selectin glycoprotein ligand-1 induces death of activated T cells

Shu-Ching Chen, Chiu-Chen Huang, Chung-Liang Chien, Chung-Iluan Jeng, Ho-Ting Su, Evelyn Chiang, Meng-Li

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Validated across multiple indications

Status of Pipeline projects

	Product Candidate	Indication	Pre-Clinical	Phase 1	Phase 2	Phase 3
PSGL-1 Programs	Neihulizumab (AbGn-168H)	Steroid Refractory aGVHD	Filing fast-track/breadesignation expecte strong single dose re	d in 2021; Initial		
		Anti-TNF and/or anti- Integrin refractory UC	Phase 2a data demo drug-free stage	nstrated durable resp	onse even at	
		Plaque psoriasis and Psoriatic arthritis	· ·	PSGL-1 Mab MOA; Povelopment of AbGn-10		
	Leiolizumab (AbGn-268)	Multiple autoimmune diseases	Demonstrated increase in potency			

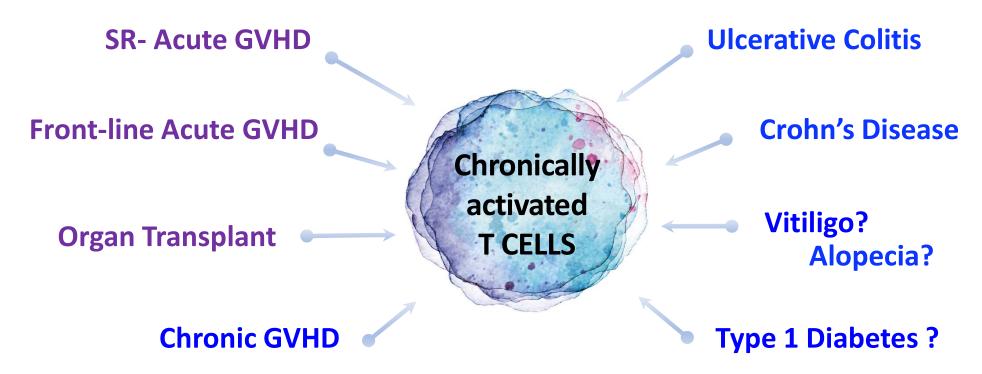


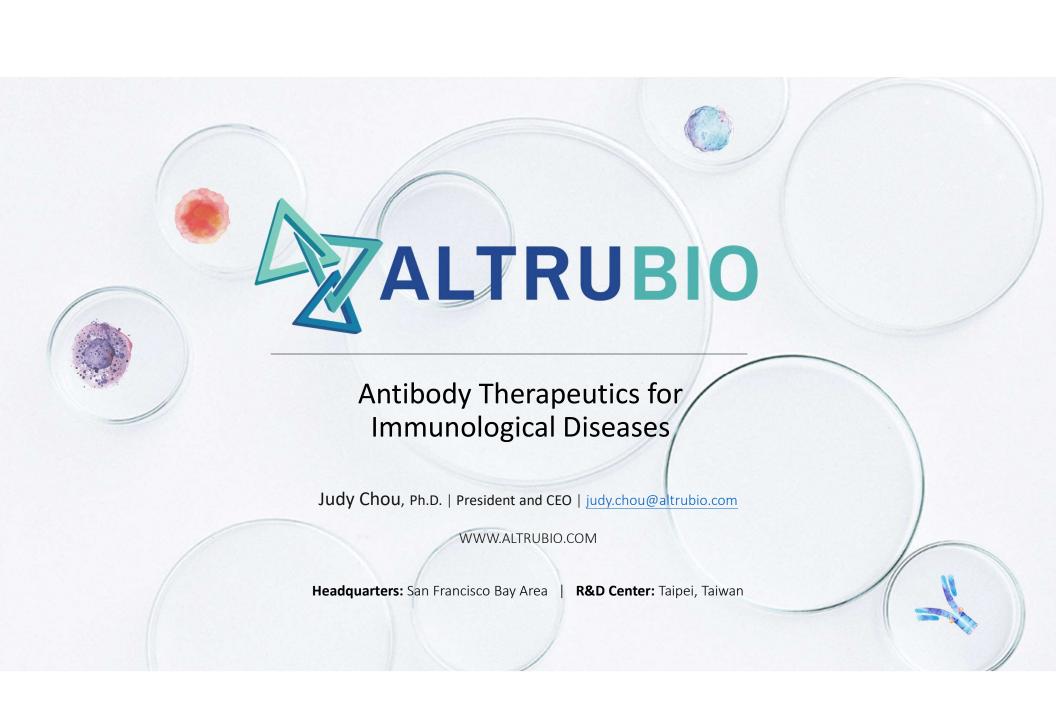




Multiple Opportunities for Indication Expansion

PSGL-1 agonists: Neihulizumab (AbGn-168H) IV Leiolizumab (AbGn-268) SubQ





FORWARD-LOOKING STATEMENTS

This presentation contains forward-looking statements, including: statements about our expectations regarding the potential benefits, activity, effectiveness and safety of our product candidates; our expectations with regard to the results of our clinical studies, preclinical studies and research and development programs, including the timing and availability of data from such studies; our preclinical, clinical and regulatory development plans for our product candidates, including the timing or likelihood of regulatory filings and approvals for our product candidates; and our expectations with regard to our ability to acquire, discover and develop additional product candidates and advance such product candidates into, and successfully complete, clinical studies. These statements involve substantial known and unknown risks, uncertainties and other factors that may cause our actual results, levels of activity, performance or achievements to be materially different from the information expressed or implied by these forward-looking statements. We may not actually achieve the plans, intentions or expectations disclosed in our forward-looking statements, and you should not place undue reliance on our forward-looking statements. Actual results or events could differ materially from the plans, intentions and expectations disclosed in the forward-looking statements we make. The forward-looking statements in this presentation represent our views as of the date of this presentation. We anticipate that subsequent events and developments will cause our views to change. However, while we may elect to update these forward-looking statements at some point in the future, we have no current intention of doing so except to the extent required by applicable law. You should, therefore, not rely on these forward-looking statements as representing our views as of any date subsequent to the date of this presentation. This presentation concerns product candidates that are under clinical investigation and which has not yet been approved for marketing by the U.S. Food and Drug Administration. It is currently limited by Federal law to investigational use, and no representation is made as to its safety or effectiveness for the purposes for which it is being investigated.



