



Revolutionizing CAR T-Cell Therapy

JANUARY 2024

OTCQB: KRBP

Kiromic.com



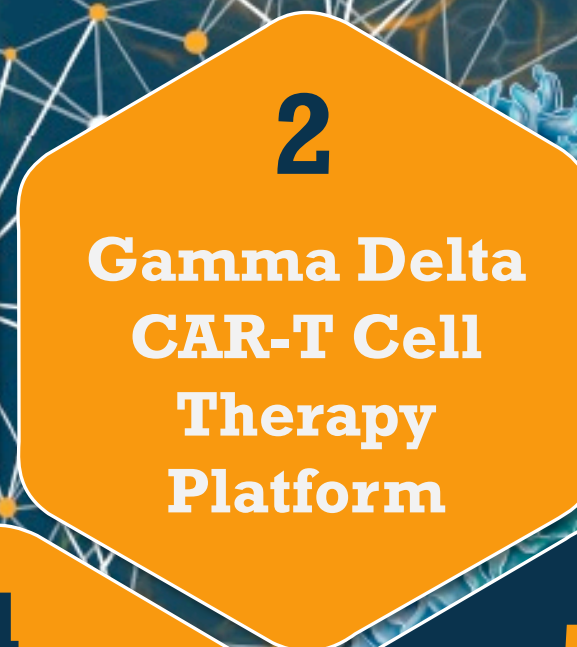
Forward Looking Statements

This presentation contains forward-looking statements that involve substantial risks and uncertainties. Kiromic makes such forward-looking statements pursuant to the safe harbor provisions of the United States Private Securities Litigation Reform Act, Section 21E of the Securities Exchange Act of 1934, as amended, and other federal securities laws. All statements other than statements of historical facts are forward-looking statements. In some cases, you can identify forward-looking statements by terms such as: “will,” “potential,” “could,” “can,” “believe,” “intends,” “continue,” “plans,” “expects,” “anticipates,” “estimates,” “may,” or the negative of these terms or other comparable terminology. These forward-looking statements include, but are not limited to, statements regarding: Kiromic’s current and anticipated IND applications including statements regarding the scope of and timing for submission of an IND application; the Deltacel™ product platform; the sponsored research agreement and the data that will be generated as a result of such collaboration; the timing for submitting and activating Kiromic’s IND applications; the benefits of utilizing non-genetically engineered Gamma Delta T cells as our first in-human study; Kiromic’s ability to achieve its objectives; and the timing for the initiation and successful completion of Kiromic’s clinical trials of its product candidates. These forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause actual results, levels of activity, performance, or achievements to be materially different from the information expressed or implied expressed or implied by these forward-looking statements. These risks and uncertainties include, but are not limited to, the risks and uncertainties discussed in our Annual Report on Form 10-K for the year ended December 31, 2022, and as detailed from time to time in our SEC filings. You should not rely upon forward-looking statements as predictions of future events. Although we believe that the expectations reflected in the forward-looking statements are reasonable, we cannot guarantee that the future results, levels of activity, performance, or events and circumstances reflected in the forward-looking statements will be achieved or occur. Moreover, neither we nor any other person assumes responsibility for the accuracy and completeness of the forward-looking statements. Such forward-looking statements relate only to events as of the date of this press release. We undertake no obligation to update any forward-looking statements except to the extent required by law.

Contents

- **The Kiromic Difference and Market Opportunity**
 - Diamond AI™ (Artificial Intelligence)
 - Gamma Delta T-cell (GDT) Therapy:
Mechanism of Action (MOA), Product Pipeline, cGMP Manufacturing
 - Current Status and Path Forward

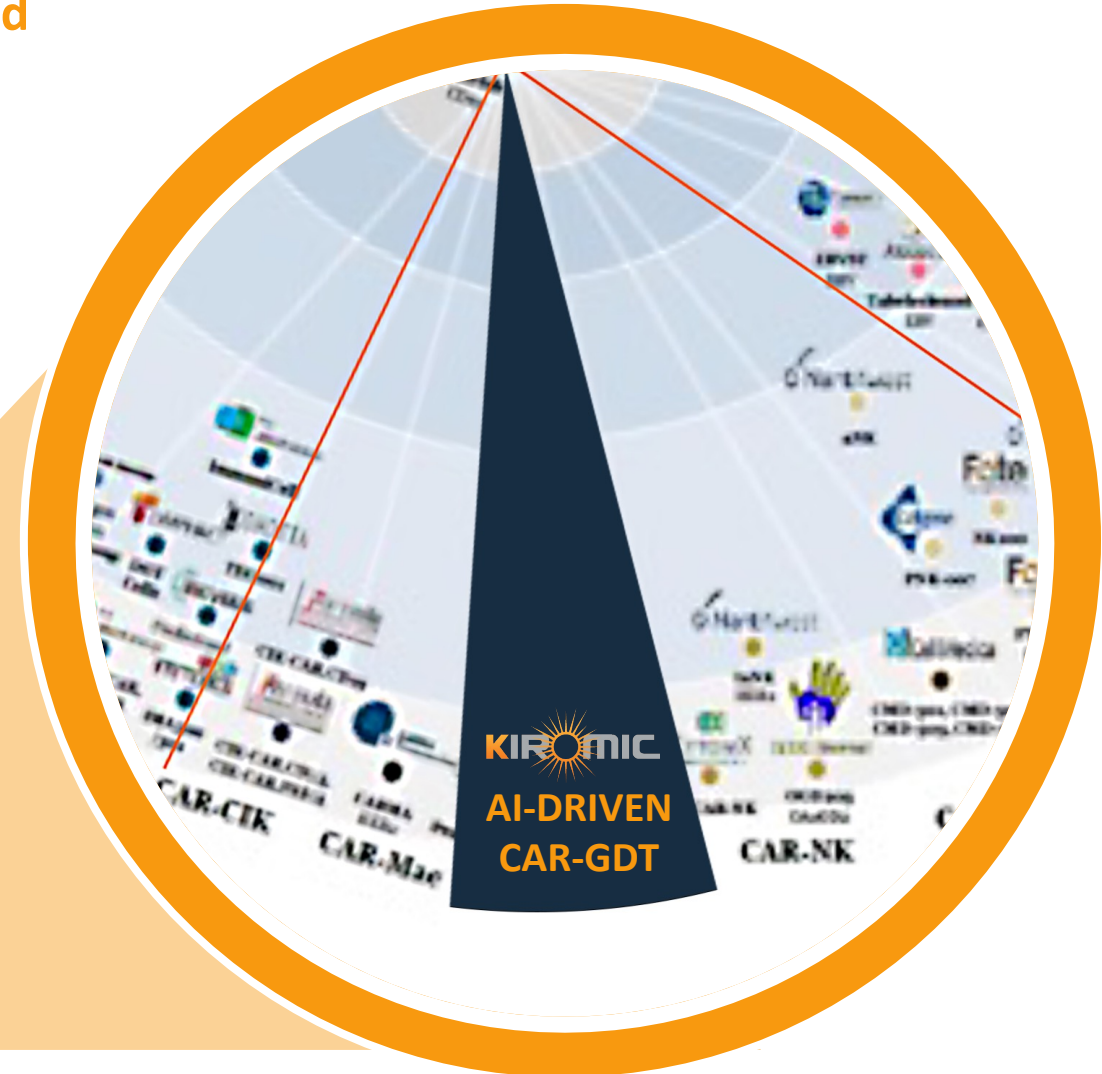
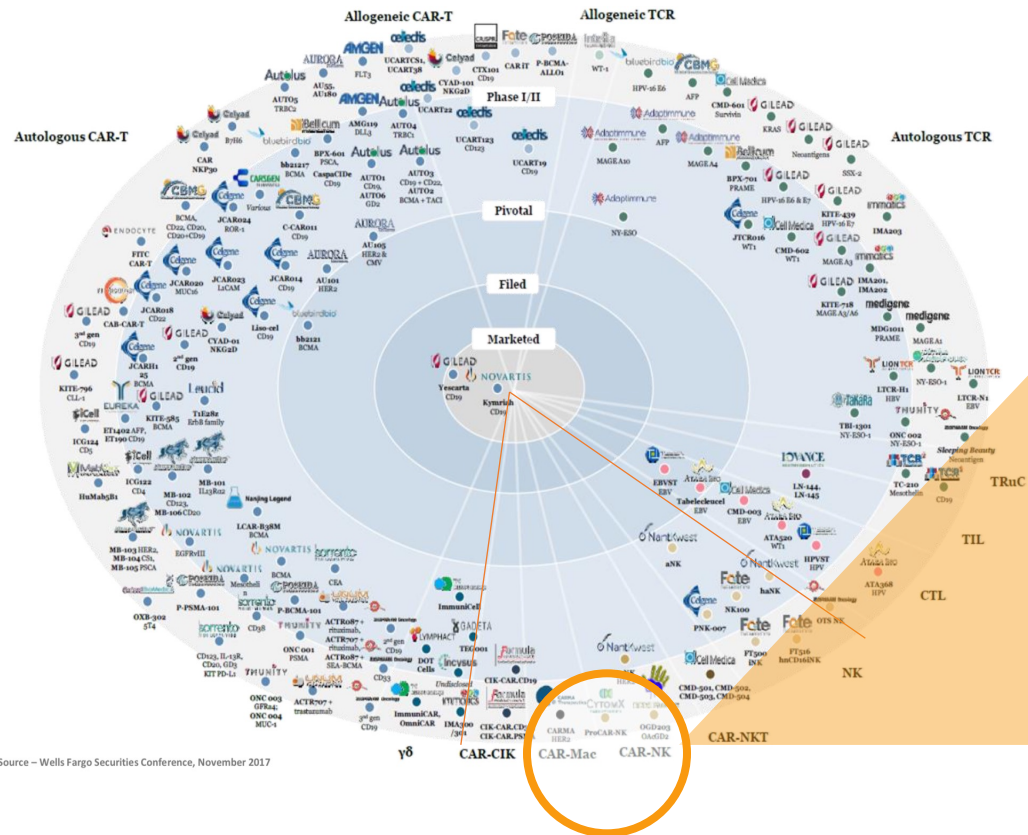
Kiromic BioPharma
is an allogeneic Gamma
Delta T-cell therapy
company
featuring unique,
proprietary, end-to-end
bioinformatic, AI
targeting, and
manufacturing
technologies to address
solid tumors



¹American Cancer Society, Cancer Facts & Figures,
2022. <https://www.cancer.org/research/cancer-facts-statistics.html>

Strategic Competitive Landscape

8 Known Companies in the Gamma Delta T Cell Therapy space.
No Known Competitors with AI-driven Technology Combined
with a Gamma Delta CAR-T Delivery Platform.



**Global CAR T-Cell
Therapy Market
by 2027¹ (USD)**

**\$33+
Billion**

90%

**of Cancers Are
Solid Malignancies²**

¹ Global CAR T-Cell Therapy Market, By Product Type, By Tumor Type, By Indication, By Treatment Type, By Targeted Antigen, By End User, By Region, Competition, Forecast and Opportunities, 2017-2027 (ReportLinker)

² American Cancer Society, Cancer Facts & Figures, 2022. <https://www.cancer.org/research/cancer-facts-statistics.html>

Competitive Difference

Allogeneic Gamma Delta Based T-Cell Therapies

Superior specificity for multiple solid tumors

- **Potential broad treatment** for solid malignancies that express Kiromic-developed biomarkers such as Iso-mesothelin.
- Solid tumors represent approx. 90% of new cancer diagnoses but finding specific targets to treat them has been challenging.
- Kiromic tackles the issue by identifying new cancer specific targets.

Vertical Integration

- In-house **cGMP manufacturing**
- In-house **QC/EM lab**
- In-house **product and process development** (R&D and MSAT)

Superior Efficacy from $\gamma\delta$ T cells

- **Strong efficacy** in pre-Clinical animal models.
- In solid tumors, the benefit of infiltrating conventional T cells may vary.
- In contrast, GDT cells are the infiltrating immune cells most likely to be associated with positive outcomes, as shown in an analysis of 18,000 tumors from 39 indications¹

Lower Costs/ Greater Access²

- 1. Outpatient treatment** means reduced hospitalization and other treatment related costs.
- 2. Lower projected cost** increases patient and health care professional access to these therapies, and potentially provides important quality-of-life benefits for patients as well.

1. Gentles AJ, Newman AM, Liu CL, *et al.* The prognostic landscape of genes and infiltrating immune cells across human cancers. Nat Med. 2015 Aug;21(8):938-945.

2. Maziarz RT. CAR T-cell therapy total cost can exceed \$1.5M per treatment. Cell Therapy Next; May 29, 2019.

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Artificial Intelligence and Bioinformatic Analytic Discovery & Development Platform

Algorithms and Large-Scale Genomics Analysis for Target Prediction



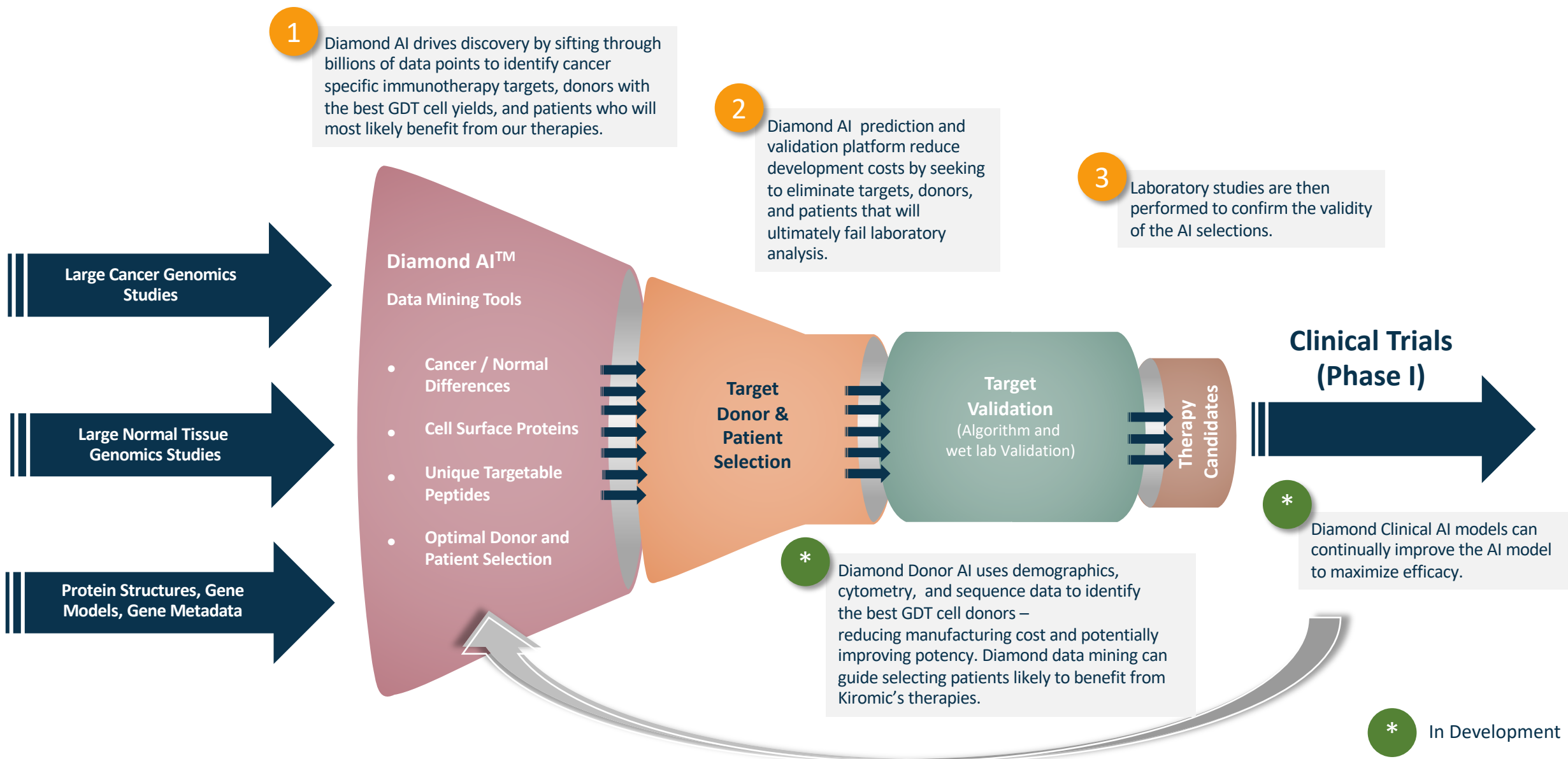
A.I. integrated with each stage of the Kiromic therapy production lifecycle

Discovering New Multi-tumor Targets

Identifying Optimal Donors and Patients to Maximize the Therapy Success

The Kiromic **Difference** - **Diamond AI™** Target Discovery Platform

Diamond AI™ target discovery platform powers innovation and significantly reduces development time and cost.



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Kiromic GDT Cell Therapy Pipeline

Multiple Indications

Deltacel™

Unmodified, off-the-shelf product candidate targeting stress ligands on cancer cells
Initial indication: NSCLC in combination with targeted, low-dose radiation

Isocel™

Engineered off-the-shelf product candidate targeting a tumor-specific variant of Mesothelin in Ovarian Cancer, Mesothelioma, and Pancreatic Cancer

Procel™

Engineered off-the-shelf product candidate targeting PDL-1+ tumors

Allogeneic
Healthy Donors

Off-the-Shelf
Cryopreserved

Innate
Rapid Acting

**Potent
and Safe**

**Commercially
Viable**

GDT: 1-5% of
circulating T-cells



Deltacel: Non-Viral Gamma Delta T-Cell Development

**Kiromic
Proprietary
In-house GDT
Cell Isolation
and Expansion**

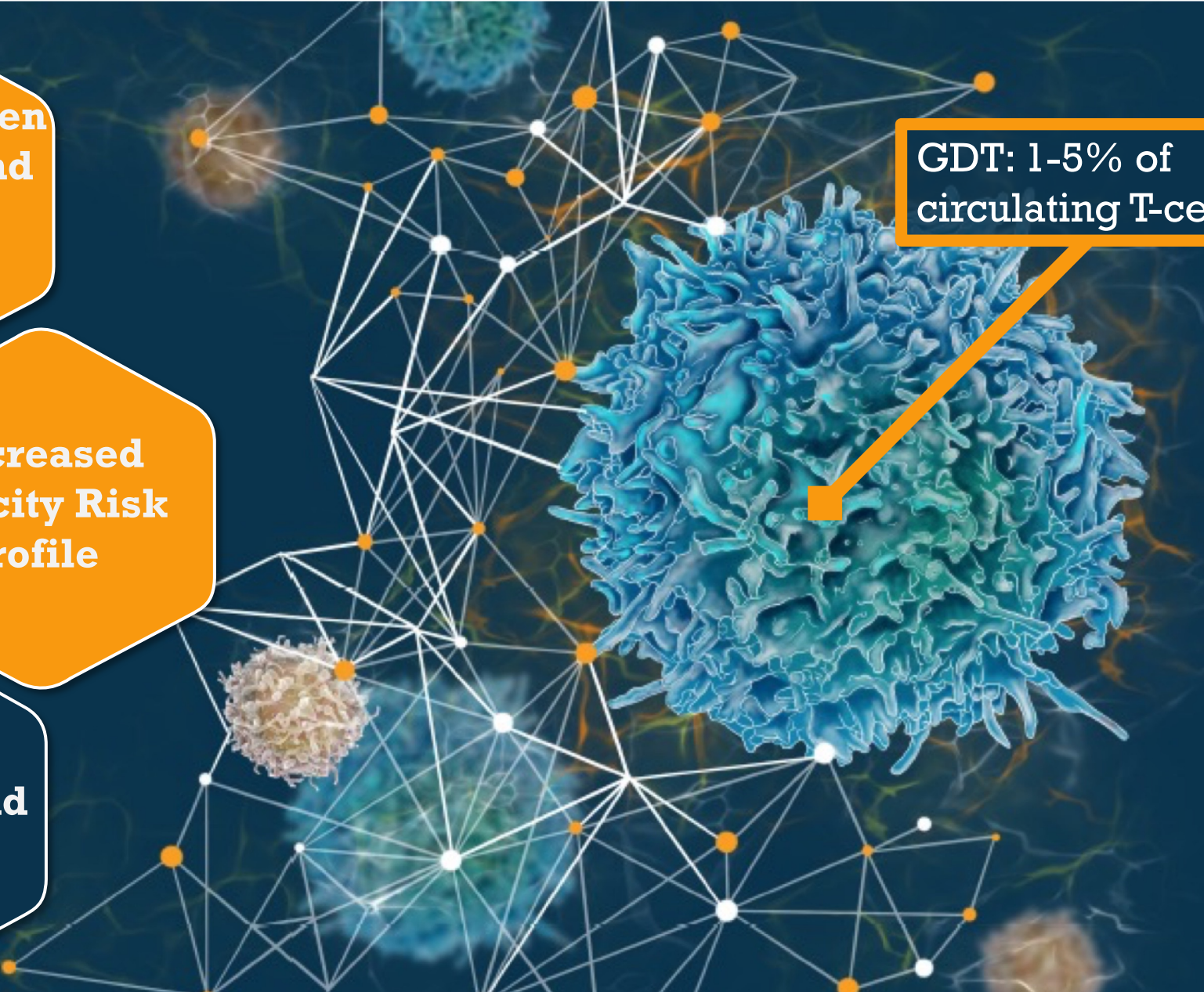
**Bridge between
the Innate and
Adaptive
Immune
Response**

**Rapid Response
to Attack
Cancer Cells**

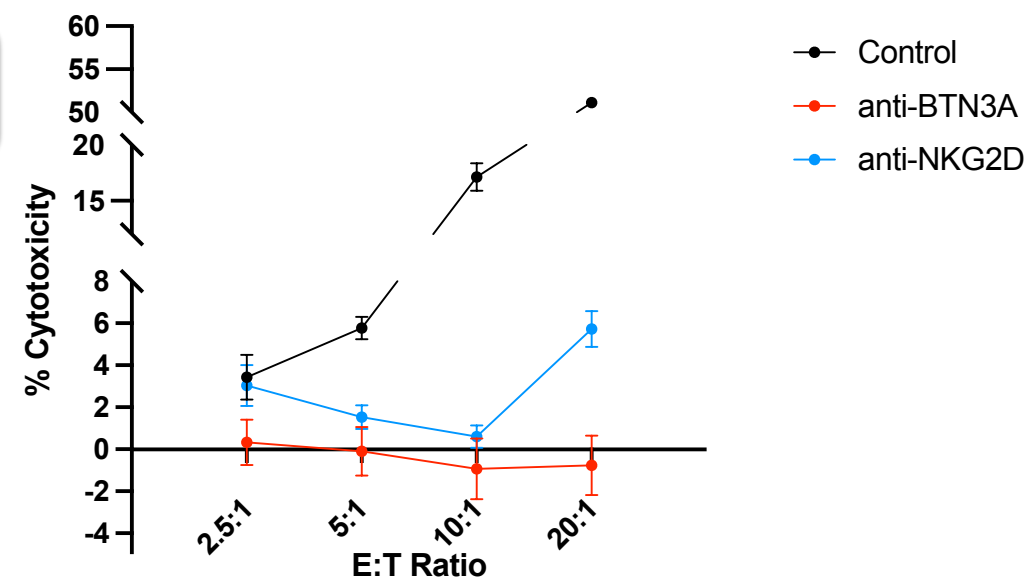
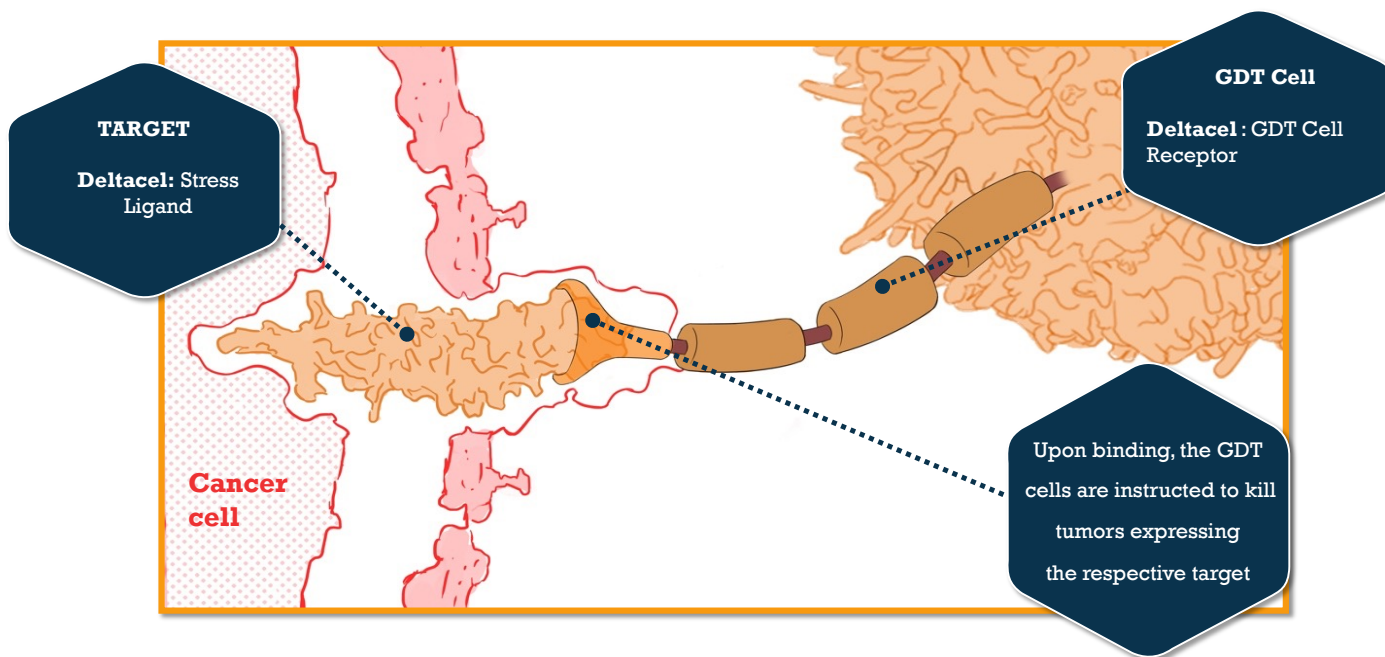
**Decreased
Toxicity Risk
Profile**

**Virus Free
Expansion and
Production**

**GDT: 1-5% of
circulating T-cells**

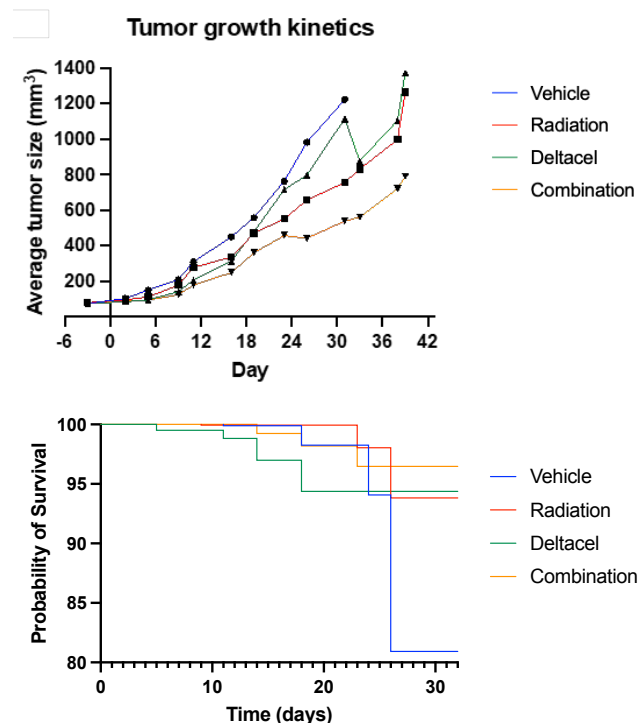


GDT Cell Therapy Mechanism of Action: Targeting Unique **Identifiers** on Tumor Tissues



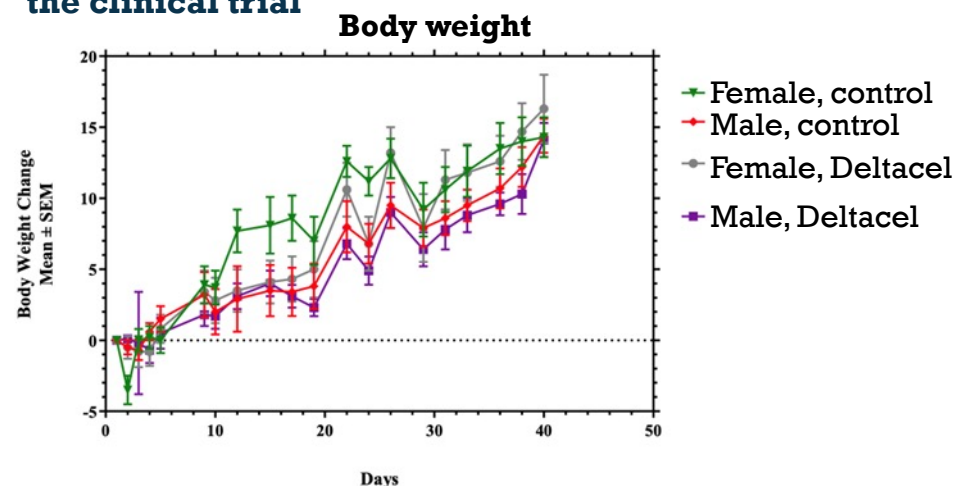
KB-GDT-01 T-Cell Therapy (Deltacel) Strong Efficacy

Deltacel™ effectively controls established A549 NSCLC tumors in immunocompromised mice when combined with a low-dose radiation



KB-GDT-01 T-Cell Therapy (Deltacel) Strong Safety

Deltacel™ does not cause any macroscopic or microscopic toxicity, even when given at over 8x the maximum dose that will be tested in the clinical trial

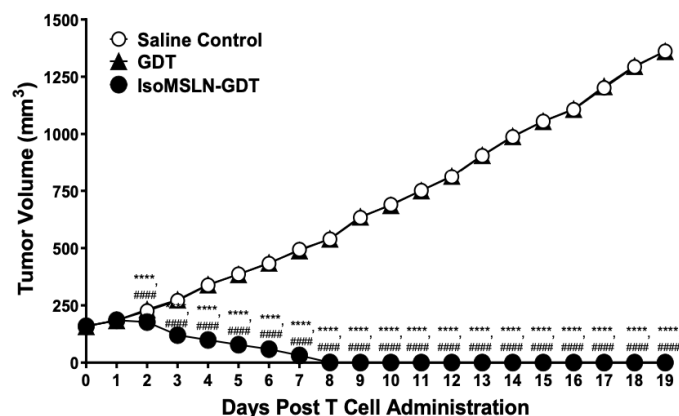


- ✓ There were no treatment-associated impacts on body weights, food consumption, or cage-side/clinical observations. Macroscopic evaluations at necropsy did not identify any evidence of test article-related toxicity. Microscopic histopathological evaluations showed no evidence of Deltacel-related toxicity
- ✓ Clinical pathology evaluations determined that all fluctuations among individual and mean values of tested analytes were considered sporadic, and not related to the administration of Deltacel.
- ✓ Plasma cytokine analysis showed that Deltacel administration did not result in the overproduction of inflammatory cytokines which may pose a safety concern.

GDT CAR T-Cell Therapy (Isocel™)* Strong Efficacy

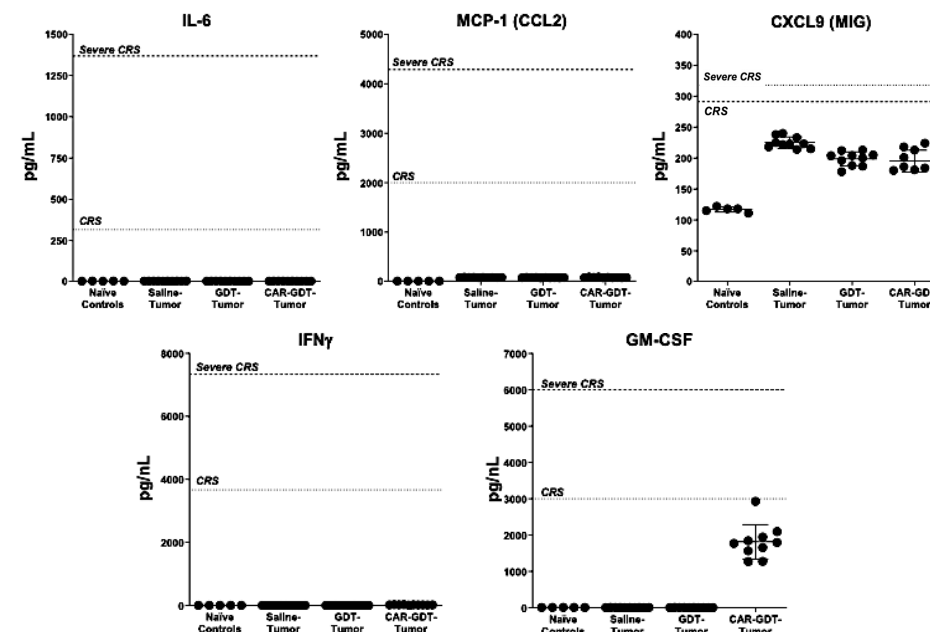
Isocel™ eradicates established NCI-H226 pleural epithelioid mesothelioma and prevents tumor growth in a model of recurrence.

Tumor eradication



GDT CAR T-Cell Therapy (Isocel™)* Strong Safety

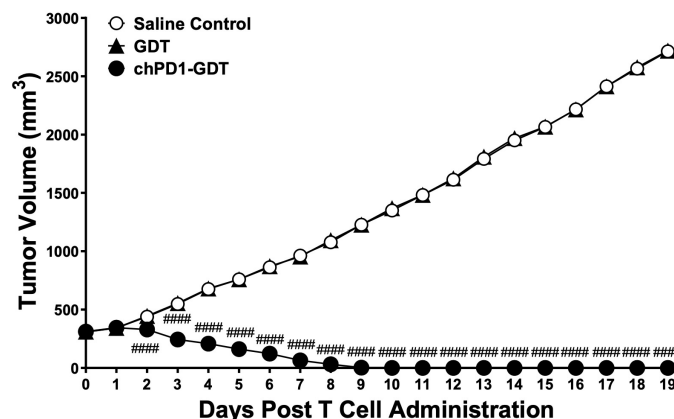
Isocel™ does not lead to cytokine level increases modeled to cause severe CRS or CRS, with circulating cell numbers regulated by objective response.



GDT chPD1 T-Cell Therapy (Procel™)* Strong Efficacy

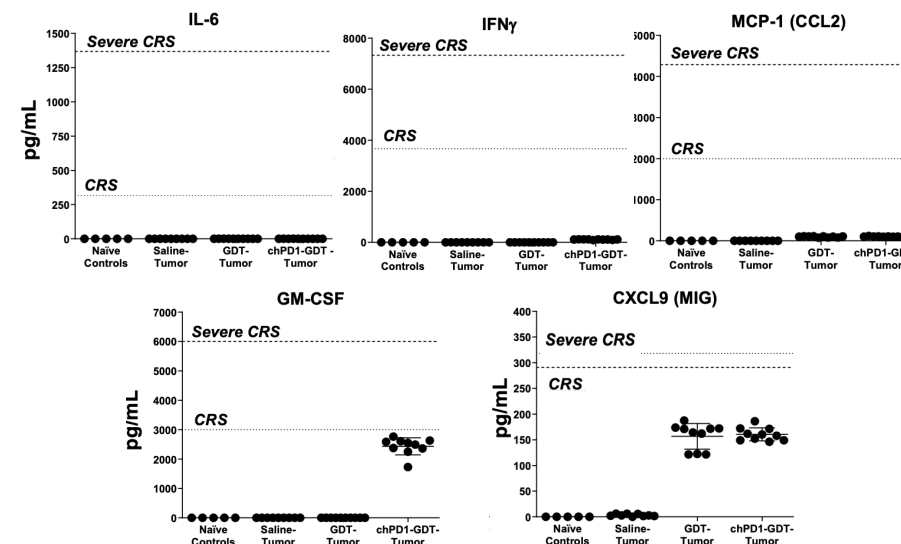
Procel™ eradicates established NCI-H226 pleural epithelioid mesothelioma and extends survival.

Tumor eradication








GDT chPD1 T-Cell Therapy (Procel™)* Strong Safety

Procel™ does not lead to cytokine level increases modeled to cause severe CRS or CRS, with circulating cell numbers regulated by objective response.



*Preclinical models: nude mice with subcutaneous NCI-H226 cells injections

Pipeline

Clinical Trial Candidate	Target	Pre-Clinical	Phase 1
Deltacel-01 Deltacel™ in combination with low-dose radiation Allogeneic, Non-Viral, Non-engineered off-the-shelf GDT therapy	 <small>THE UNIVERSITY OF TEXAS</small> MDAnderson Cancer Center	Multiple Tumor Cell Markers	First Patient Dosed 12/13/2023
New IND Isocel™ in combination with low-dose radiation Allogeneic, off-the-shelf, GDT CAR-T therapy	 <small>THE UNIVERSITY OF TEXAS</small> MDAnderson Cancer Center	Mesothelin Isoform <i>KRBP Proprietary Target</i>	H1 2025* Expected Beginning of Activation Process for Clinical Trial
New IND ALEXIS - ISO-1 Isocel™ Allogeneic, off-the-shelf, GDT CAR-T therapy	 <small>THE UNIVERSITY OF TEXAS</small> MDAnderson Cancer Center	Mesothelin Isoform <i>KRBP Proprietary Target</i>	H1 2025* Expected Beginning of Activation Process for Clinical Trial
New IND Procel™ in combination with low-dose radiation Allogeneic, off-the-shelf, GDT CAR-T therapy	 <small>THE UNIVERSITY OF TEXAS</small> MDAnderson Cancer Center	PD-L1	H1 2025* Expected Beginning of Activation Process for Clinical Trial
New IND ALEXIS - PRO-1 Procel™ Allogeneic, off-the-shelf, GDT CAR-T therapy	 <small>THE UNIVERSITY OF TEXAS</small> MDAnderson Cancer Center	PD-L1	H1 2025* Expected Beginning of Activation Process for Clinical Trial

* Subject to obtaining sufficient financing to support the progression of the development of those additional clinical trial candidates.

In-House cGMP Manufacturing Creates De-Risked Value

**Clinical-Grade,
cGMP-Compliant
Cell Therapy
Manufacturing
Space**

**Dedicated
cGLP
Microbiology
and QC Lab**

**Dedicated
Product
Development
Suite**

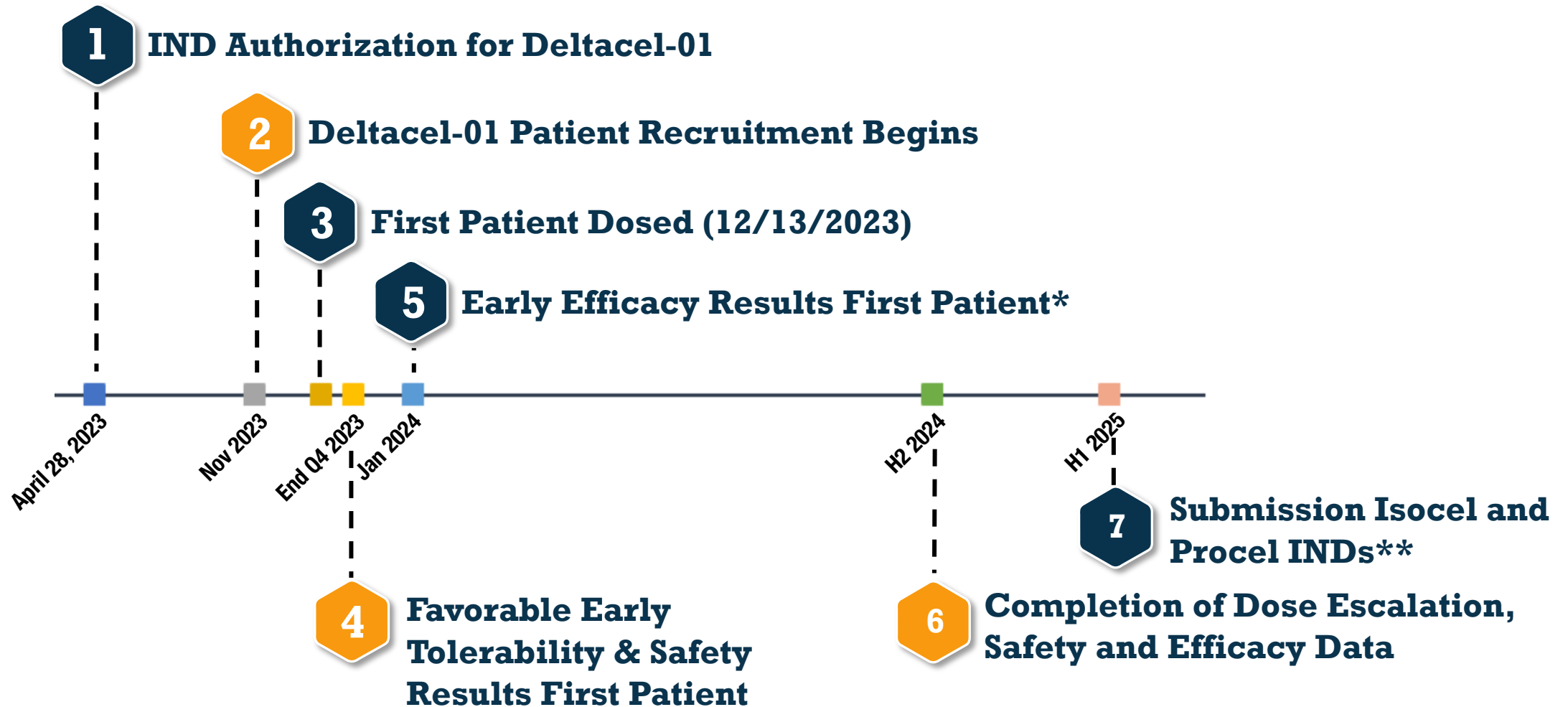
**34,000 sq ft
Facility
Operations**

**12,000 sq ft
R&D Laboratory
&
Manufacturing
Space**

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Recent and Upcoming Milestones*



*The milestones and timing of completion are based upon the company's current expectations in consultation with its partners and vendors.

** Subject to obtaining sufficient financing to support the progression of the development of those additional clinical trial candidates.

Leadership Team

**Pietro
Bersani**
CPA, CGMA

CEO



Deloitte.

ARTHUR
ANDERSEN

**Leonardo
Mirandola**
Ph.D.

**CSO/INTERIM
COO**



**Scott
Dahlbeck**
MD, Pharm.D.

COSO



Texas Tech Univ
Health Science
Center



University of TX
Health Science
Center Houston



University of
Nebraska College
Medical Center of Pharmacy



**Brian
Hungerford**
CPA, CGMA

CFO

Deloitte.



accenture



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**Michael
Nagel**

Chairperson

**Pietro
Bersani**
CPA, CGMA

Director

**Pam
Misajon**

**Independent
Director**

**Michael
Catlin**

**Independent
Director**



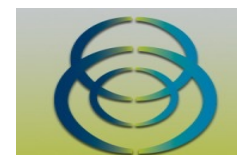
Fuel Systems Solutions



Deloitte.



ARTHUR
ANDERSEN



Summary Balance Sheet & Cap Table

Balance Sheet Data (As of September 30, 2023)	As Reported
Cash and Cash Equivalents	\$4,379,500
Working Capital	(\$11,710,200)
Total Assets	\$14,233,500
Total Stockholders' Deficit	(\$4,487,400)

Cap Table (As of September 30, 2023)	Common Stock Equivalents
Common Stock	1,176,290
Restricted Stock Units (\$4.58 Weighted average grant date fair value)	13,729
Options (\$101.04 Weighted average exercise price)	18,093
Warrants	15,416
Convertible Preferred Share Shares (\$14MM principal & \$6.50 share conversion)	2,153,846
Convertible Notes (\$7.2MM principal & \$6.50 share conversion) (\$2.4MM principal & \$5.00 share conversion)	1,587,692
Fully Diluted Common Shares	4,965,036





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