# **Contact Us**



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# **Descartes-08**

Descartes-08 is an investigation product not yet approved by the US Food and Drug Administration (FDA).

Descartes-08 is an autologous T-cell product expressing a chimeric antigen receptor (CAR) directed against B-cell maturation antigen (BCMA) on plasma cells, plasma blasts, and plasmacytoid dendritic cells.

Descartes-08 is a cell therapy that uses mRNA to transiently express the BCMA-directed CAR proteins.



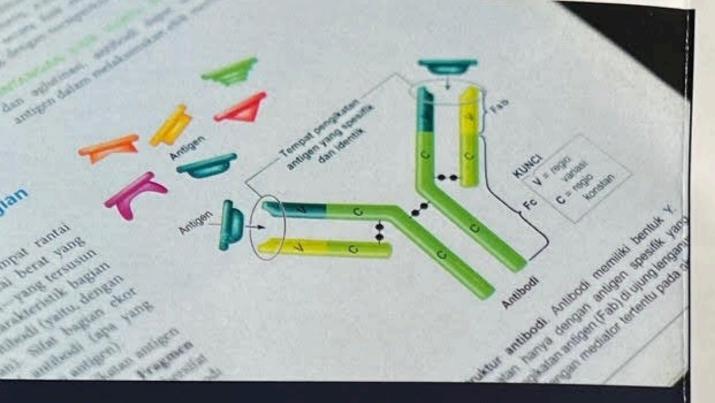


Descartes-08 for Children, Adolescents, and Young Adults with Autoimmune Diseases

www.cartesiantherapeutics.com

Pioneering Cell Therapy for Autoimmunity

www.cartesiantherapeutics.com



# mRNA Cell Therapy

Cartesian's mRNA approach to CAR-T therapies is designed to expand the reach of potent cell therapy products to address autoimmunity.



### No Lymphodepletion

No associated cytopenia or chemotherapy toxicities



### **Administered Outpatient**

Convenient dosing schedule



**Delivered at Therapeutic Levels** 

Designed to deliver dose levels without uncontrolled proliferation



## **Transient Cell Modification**

Does not carry risk of genomic integration



Phase 1/2 dose-escalation basket study evaluating the safety, tolerability, and early efficacy of Descartes-08 in four pediatric indications:

- Juvenile Dermatomyositis (JDM)\*
- Childhood-onset Lupus (cSLE)
- ANCA-associated Vasculitis (AAV)
- Juvenile Myasthenia Gravis (JMG)

\*JDM has received Rare Pediatric Disease Designation for DC-08

## Study Design

#### Part 1

Intrapatient Dose Level Escalation
DL 1-> DL 2-> DL 3
1x/week inpatient infusions over 6 weeks

#### Part 2



Maximum Tolerated Dose from Part 1 administered 1x/week over 6 weeks in an outpatient setting

### Part 1 - Initial Safety Cohort

- Up to 18 participants
- If a DLT occurs in the first 3 participants, study converts to a 3+3 dose-escalation design

#### Part 2 - Basket Expansion Phase

- Up to 10 participants per cohort
- Weekly dosing at the MTD established in Part 1

# **Eligbility Criteria**

- Patients must be at least 12 years of age
- Definitive diagnosis for cSLE, AAV, JDM, or JMG
- Diagnosis ≥12 months prior to screening required
- Must not have weakness sufficient to require wheelchair
- No history or presence of active CNS lupus or other CNS disease
- Concomitant immunosuppressive drugs must be deemed necessary by the investigator

To learn more about the full eligibility criteria for the study, please contact us by email <a href="helios001@cartesiantx.com">helios001@cartesiantx.com</a>

