

Global Disparities in Access to CAR-T Cell Therapy for Hematologic Malignancies

Emerging Challenges in Equitable Cellular Therapy Delivery

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INTRODUCTION

Chimeric antigen receptor T-cell (CAR-T) therapy has transformed outcomes in relapsed and refractory hematologic malignancies, producing durable responses in aggressive lymphomas, leukemias, and multiple myeloma. However, access to CAR-T therapy remains highly unequal worldwide. We reviewed emerging evidence regarding **geographic**, **economic**, **infrastructural**, and **regulatory** disparities affecting global CAR-T access.

METHODS

A structured literature review was performed evaluating access to CAR-T therapy in hematologic malignancies.

Sources:

- 1

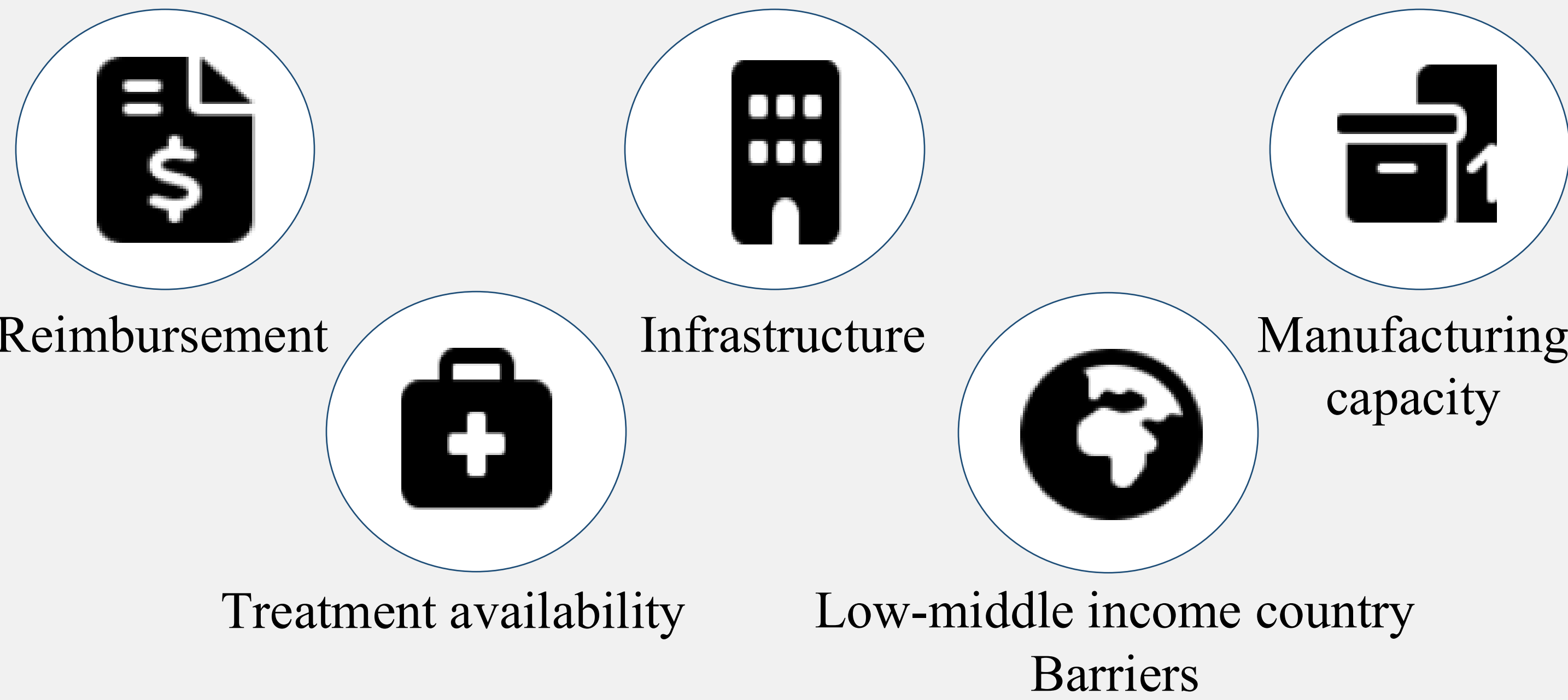
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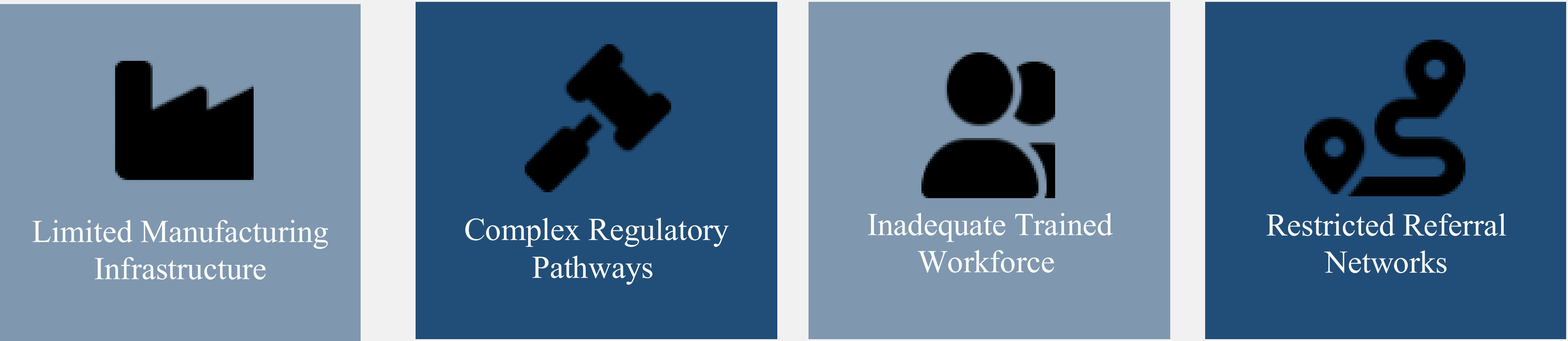
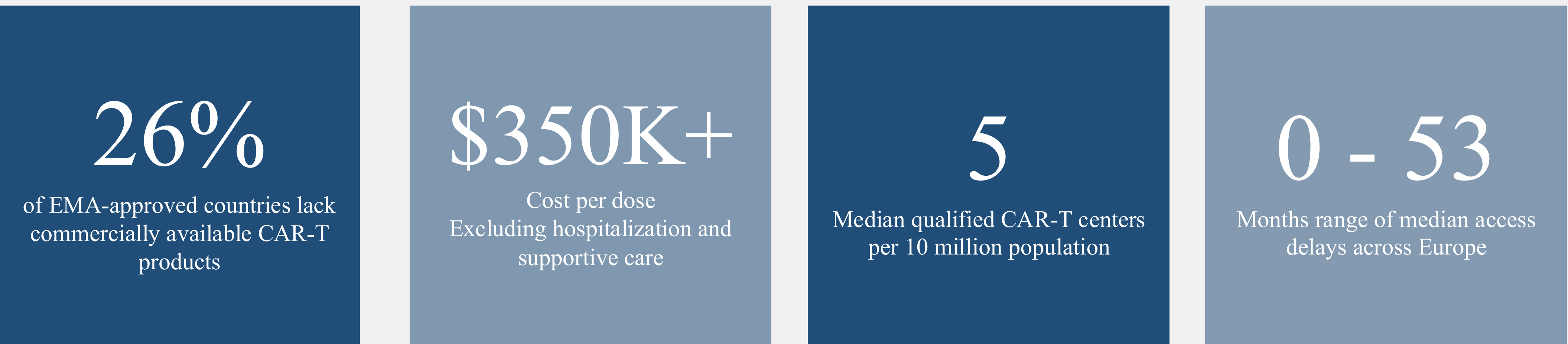
Conference Proceedings
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Health Policy analysis

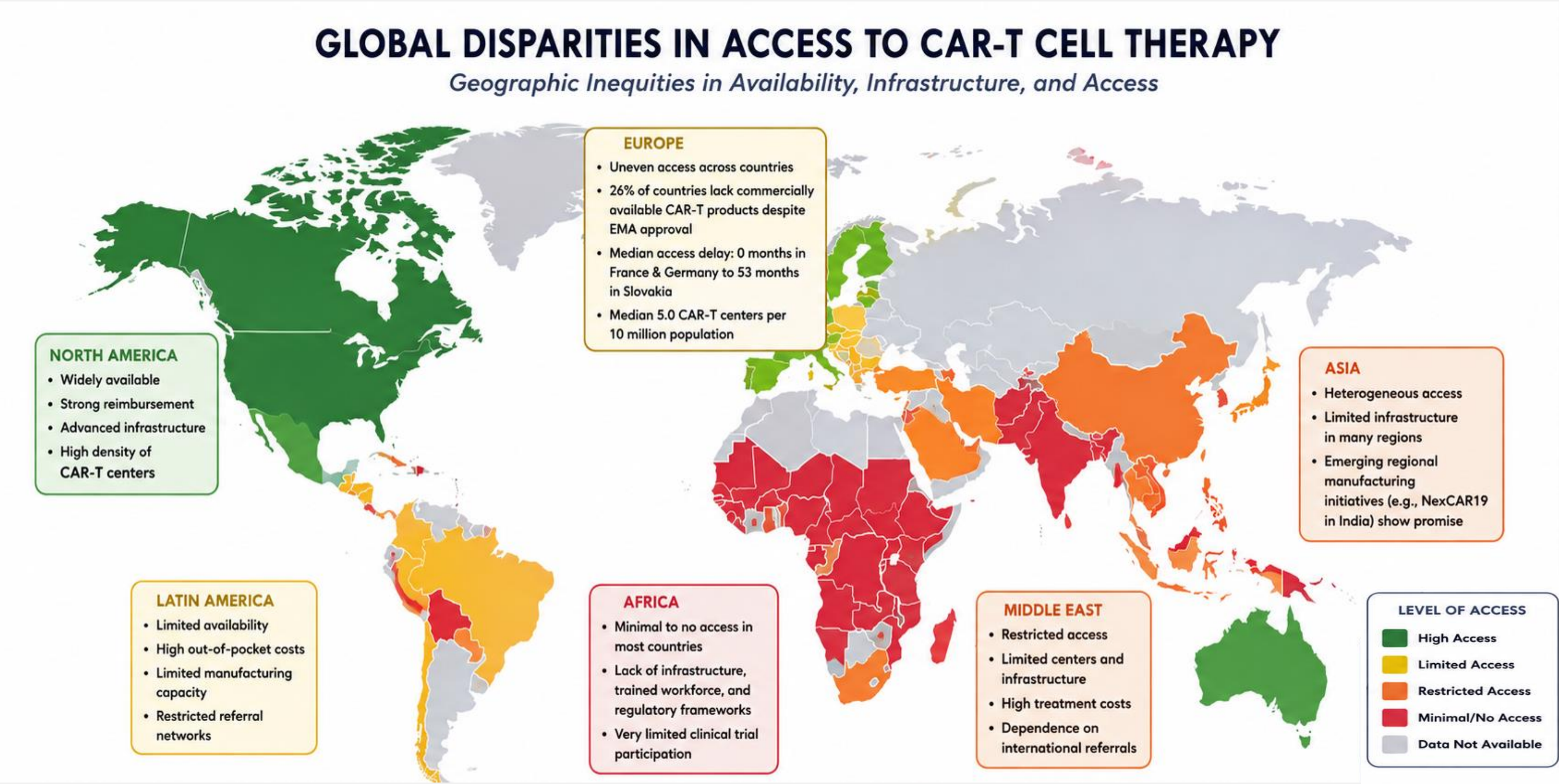
DOMAINS ASSESSED



RESULTS



More than 1500+ CAR-T clinical trials have been initiated globally, but remain concentrated in high-income countries



CONCLUSION

- CAR-T therapy remains inaccessible to large populations because of intersecting economic, infrastructural, regulatory, and workforce barriers.
- Access disparities are present even among high-income countries and are magnified in LMICs.
- Equity-oriented delivery will require health-system planning beyond product approval alone

ACTIONABLE SOLUTIONS

- Regional manufacturing**
Build local GMP capacity and reduce reliance on centralized export models
- Streamlined regulation**
Coordinate approvals, quality standards, and cross-border policy frameworks
- Innovative reimbursement**
Use sustainable coverage models, risk-sharing and outcome-linked payments
- Decentralized delivery**
Expand qualified centres, workforce training, referrals and follow-up care

REFERENCES

- Hartmann J, et al. *EMBO Mol Med*. 2017;9:1183-1197.
- Gagelmann N, et al. *Lancet Haematol*. 2022;9:e786-e795.
- Odstrcil MS, et al. *Blood Rev*. 2024;63:101136.
- Pennings ERA, et al. *HemaSphere*. 2026.

