

# Insurance Status, Neurotoxicity, and Clinical Outcomes After CAR-T Therapy in a Highly Socially Vulnerable Cohort

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## INTRODUCTION

- Chimeric antigen receptor T-cell (CAR-T) therapy has transformed outcomes in relapsed or refractory hematologic malignancies.
- Despite these advances, disparities in outcomes remain a concern among socioeconomically disadvantaged populations.
- Data evaluating the impact of socioeconomic factors on **post-infusion outcomes** in socially vulnerable CAR-T populations remain limited.
- We evaluated a diverse cohort of CAR-T recipients treated at a large urban academic cancer center.

## AIM

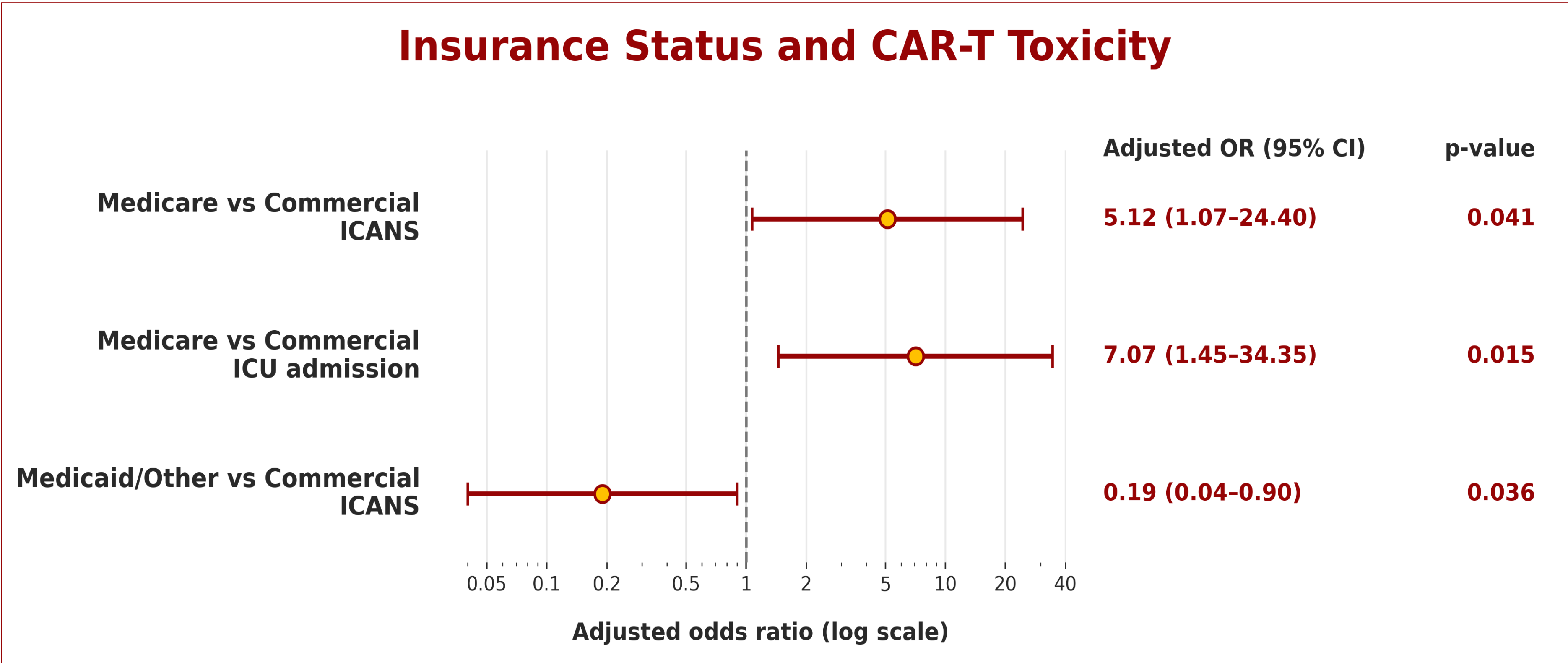
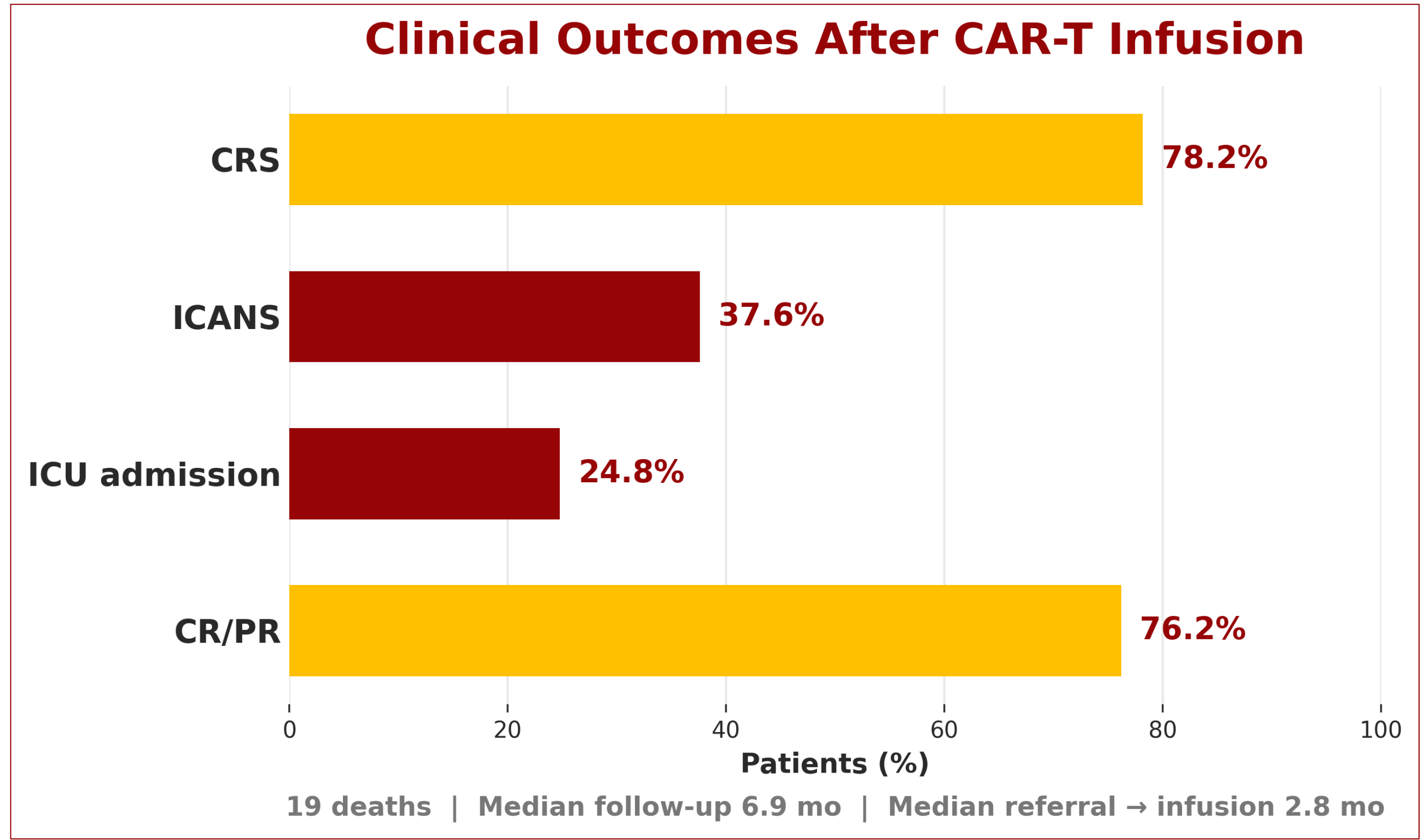
- To evaluate associations between **socioeconomic characteristics** and key clinical outcomes among patients receiving commercial CAR-T therapy.
- Outcomes of interest included **CRS, ICANS, ICU admission, treatment response, overall survival, and time from referral to infusion**.

## METHODS

- Retrospective cohort study of **101 patients** who received commercial CAR-T therapy at **USC Norris Comprehensive Cancer Center from 2022–2025**.
- CAR-T products included **axi-cel, brexu-cel, and cilta-cel**.
- Socioeconomic variables included **race/ethnicity, insurance status, Social Vulnerability Index (SVI), primary language, and distance from the transplant center**.
- Clinical outcomes included **CRS, ICANS, ICU admission, treatment response, overall survival, and time to infusion**.
- Logistic regression was used for binary outcomes, ordinal logistic regression for toxicity grades, and Cox proportional hazards models for survival and time-to-infusion outcomes.
- Each socioeconomic variable was evaluated in a **separate multivariable model** to minimize multicollinearity.
- Models were adjusted for **age, sex, and diagnosis subtype**; statistical significance was defined as **p < 0.05**.

## RESULTS

| Selected Cohort Characteristics  |                          |
|----------------------------------|--------------------------|
| CHARACTERISTIC                   | OVERALL COHORT (N = 101) |
| Age, median (IQR)                | 62 (49–70) yr            |
| Male                             | 64 (63.4%)               |
| Hispanic                         | 55 (54.5%)               |
| Non-English primary language     | 43 (42.6%)               |
| Commercial insurance             | 50 (49.5%)               |
| Overall SVI, median (IQR)        | 93 (74–96)               |
| Highest SVI quartile             | 73 (72.3%)               |
| Distance to center, median (IQR) | 22.4 (10.3–113) mi       |
| DLBCL                            | 44 (43.6%)               |
| ALL                              | 21 (20.8%)               |
| Multiple myeloma                 | 22 (21.8%)               |



| Adjusted Socioeconomic Associations Across Outcomes |     |             |             |          |          |                  |
|-----------------------------------------------------|-----|-------------|-------------|----------|----------|------------------|
|                                                     | CRS | ICANS       | ICU         | Response | Survival | Time to infusion |
| Race                                                | NS  | NS          | NS          | NS       | NS       | NS               |
| Ethnicity                                           | NS  | NS          | NS          | NS       | NS       | NS               |
| Insurance                                           | NS  | SIGNIFICANT | SIGNIFICANT | NS       | NS       | NS               |
| Overall SVI                                         | NS  | NS          | NS          | NS       | NS       | NS               |
| SVI Q4                                              | NS  | NS          | NS          | NS       | NS       | NS               |
| Distance                                            | NS  | NS          | NS          | NS       | NS       | NS               |
| Language                                            | NS  | NS          | NS          | NS       | NS       | NS               |

Models adjusted for age, sex, and diagnosis subtype

## CONCLUSION

- In this highly socially vulnerable, predominantly Hispanic cohort of CAR-T recipients, socioeconomic factors were **not significantly associated with treatment response, overall survival, or time to infusion** after adjustment.
- Insurance status was associated with select toxicity outcomes:** Medicare was associated with higher odds of **ICANS and ICU admission**, while Medicaid/Other insurance was associated with lower odds of ICANS compared with commercial insurance.
- Findings should be considered **hypothesis-generating** given the limited sample size, short follow-up, wide confidence intervals, and potential residual confounding.

### Future Considerations

- Future studies should evaluate disparities across the **entire CAR-T care pathway—including referral, insurance authorization, leukapheresis, and infusion—to identify potential barriers that may occur before patients receive therapy**.

## REFERENCES

Ahmed N, Shahzad M, Shippey E, et al. Socioeconomic and racial disparity in chimeric antigen receptor T cell therapy access. *Transplant Cell Ther*. 2022;28(7):358–364. doi:10.1016/j.jtct.2022.04.008.

Boloori A, Nategh E, Su CT. Association of social vulnerability index and chimeric antigen receptor T-cell therapy administration, 2018–2023. *Oncologist*. 2025;30(9):oyaf236. doi:10.1093/oncolo/oyaf236.

Dragoi D, Cusworth S, Oldham L, et al. CAR T access and outcomes in large B-cell lymphoma according to ethnicity and socioeconomic deprivation in the UK. *Br J Haematol*. 2025;206(4):1178–1185. doi:10.1111/bjh.19997.

Kansagra A, Farnia S, Majhail N. Expanding access to chimeric antigen receptor T-cell therapies: challenges and opportunities. *Am Soc Clin Oncol Educ Book*. 2020;40:1–8.

Snyder S, Chung KC, Jun MP, et al. Access to chimeric antigen receptor T cell therapy for diffuse large B cell lymphoma. *Adv Ther*. 2021;38:4659–4674.

Alqazaqi R, Schinke C, Thanendrarajan S, et al. Geographic and racial disparities in access to chimeric antigen receptor-T cells and bispecific antibodies trials for multiple myeloma. *JAMA Netw Open*. 2022;5:e2228877.