

# Advancing Real-World Evidence in Diffuse Large B-Cell Lymphoma: Insights from EHR-Derived Data in the UK

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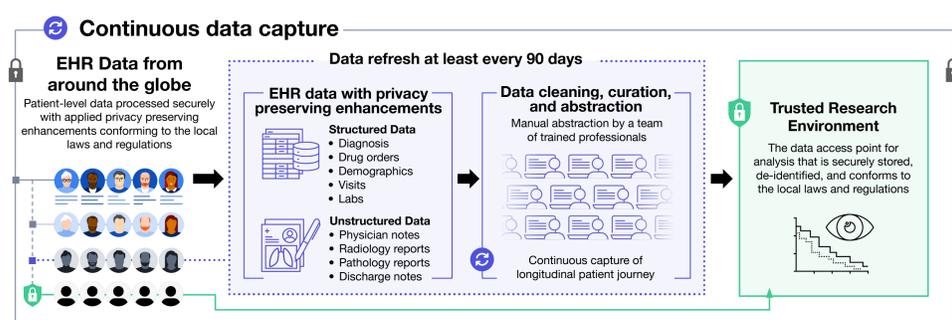
## Background

- Diffuse large B-cell lymphoma (DLBCL) is the most common aggressive Non-Hodgkin Lymphoma in the UK, and is associated with substantial morbidity and mortality. There is a growing need for region-specific real-world data (RWD) to understand variations in disease characteristics, healthcare delivery, and outcomes
- We aimed to describe the clinical characteristics and treatment patterns of patients with DLBCL in the UK using comprehensive, deidentified electronic health record (EHR) data

## Methods

- Data source:** The UK-based, EHR-derived deidentified Flatiron Health Research Database, comprising longitudinal patient-level data originated from a range of National Health Service (NHS) clinics and curated via technology-enabled abstraction<sup>1,2</sup>
- Setting:** The study included 1265 patients at least 18 years of age, diagnosed with DLBCL between January 2011 and June 2025
- Main outcome measures:** Descriptive disease characteristics, treatment patterns, and overall survival (OS) were estimated using Kaplan-Meier methods

**Figure 1. Overview of the Process for EHR-Derived RWD Curation**



## Results

- Data completeness was high: 76% patients had defined cell of origin (COO), 56% of patients had a recorded cytogenetic test with *MYC* being the most frequently tested
- Median OS from initial diagnosis in the full cohort was 6.0 years (95% CI: 5.0-7.1)

**Table 1. Demographics of Patients Included in the Study**

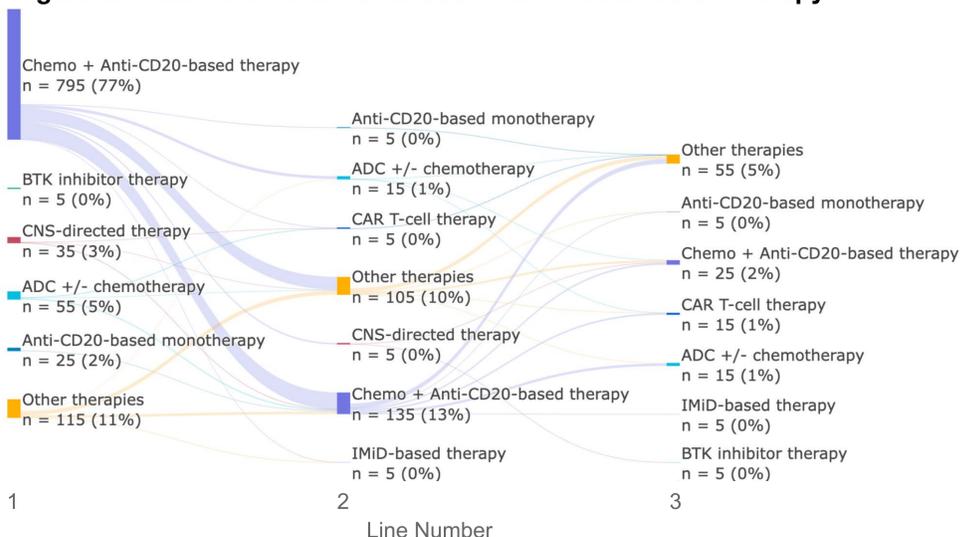
Characteristic	Full cohort <sup>a</sup> (n = 1265)	Treated cohort <sup>a</sup> (n = 1035)
Age, median (IQR), y	69 (58, 77)	68 (57, 76)
Disease stage at initial diagnosis, No. (%)		
I	125 (16)	105 (15)
II	105 (14)	95 (14)
III	115 (15)	105 (15)
IV	415 (55)	375 (55)
Unknown	505	335
ECOG at index, No. (%) <sup>b</sup>		
0-1	35 (58)	535 (71)
2+	25 (42)	215 (29)
Unknown	1205	285
Cell of origin, No. (%)		
Germinal center B-cell	495 (52)	425 (54)
Activated B-cell	465 (48)	365 (46)
Unknown	305	245
Extranodal disease, No. (%)		
Yes	905 (71)	715 (69)
No	365 (29)	315 (31)
<i>MYC</i> cytogenetic test at index, No. (%) <sup>c</sup>		
<i>MYC</i> +	105 (15)	55 (15)
<i>MYC</i> -	615 (85)	305 (85)
Not tested	555	675

ECOG, Eastern Cooperative Oncology Group; IQR, interquartile range <sup>a</sup>To ensure privacy protection, all patient counts are rounded up to 10 with 5 subtracted <sup>b</sup>ECOG from 30 days prior to 7 days after index <sup>c</sup>Cytogenetic test results from 30 days prior to 90 days after index

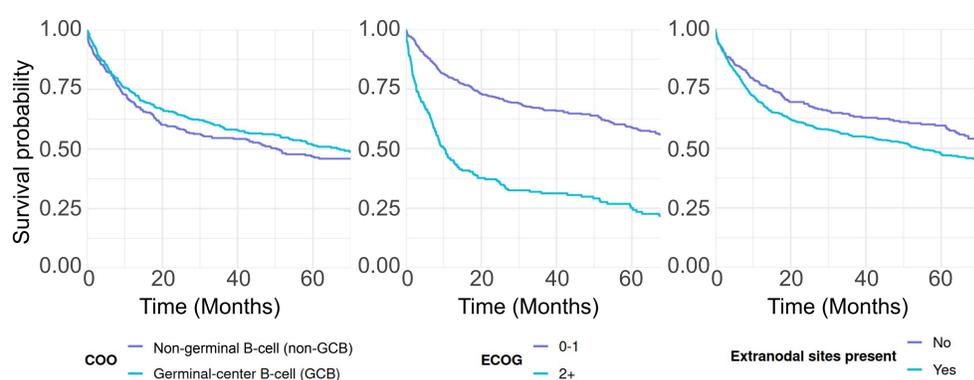
## Results (continued)

- From the 1035 treated patients:
  - 65 (6%) received a transplant, and 45 (4%) received CAR T cell therapy.
  - Median OS was 7.3 years (95% CI: 6.1-9.0) from start of treatment
  - Survival was lower in patients with more advanced stage, non-germinal B-cell as COO, and extranodal sites present at initial diagnosis (Figure 3)

**Figure 2. Treatment Patterns Across First Three Lines of Therapy**



**Figure 3. OS Stratified by COO, ECOG, and Presence of Extranodal Sites from Time of Treatment Initiation**



The UK DLBCL dataset reflects **expected clinical patterns** and offers insights to inform and optimise DLBCL management

## Future Directions

- Harmonised oncology EHR-derived datasets have been developed at the regional level to enable cross-country comparisons of patients
- The quality and completeness of technology-enabled abstraction data in the UK are high and enable detailed and insightful analyses, going beyond the standard EHR-derived datasets

## References

- Flatiron Health. Database Characterization Guide. Flatiron.com. Published March 18, 2025. Accessed October 1, 2025. <https://flatiron.com/database-characterization>
- Adamson B et al. *ESMO Real World Data Digit Oncol*. doi.org/10.1016/j.esmorw.2025.100113

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