

Glofitamab in Relapsed/Refractory Large B Cell Lymphoma – Real-World Experience in Hong Kong

Dr. Ryan CW Ho

MBBS (Hons), MRes(Med), MRCP (UK), FHKCP, FHAKM (Medicine)

Division of Haematology, Medical Oncology and HSCT

Department of Medicine, Queen Mary Hospital

Disclosures

- None

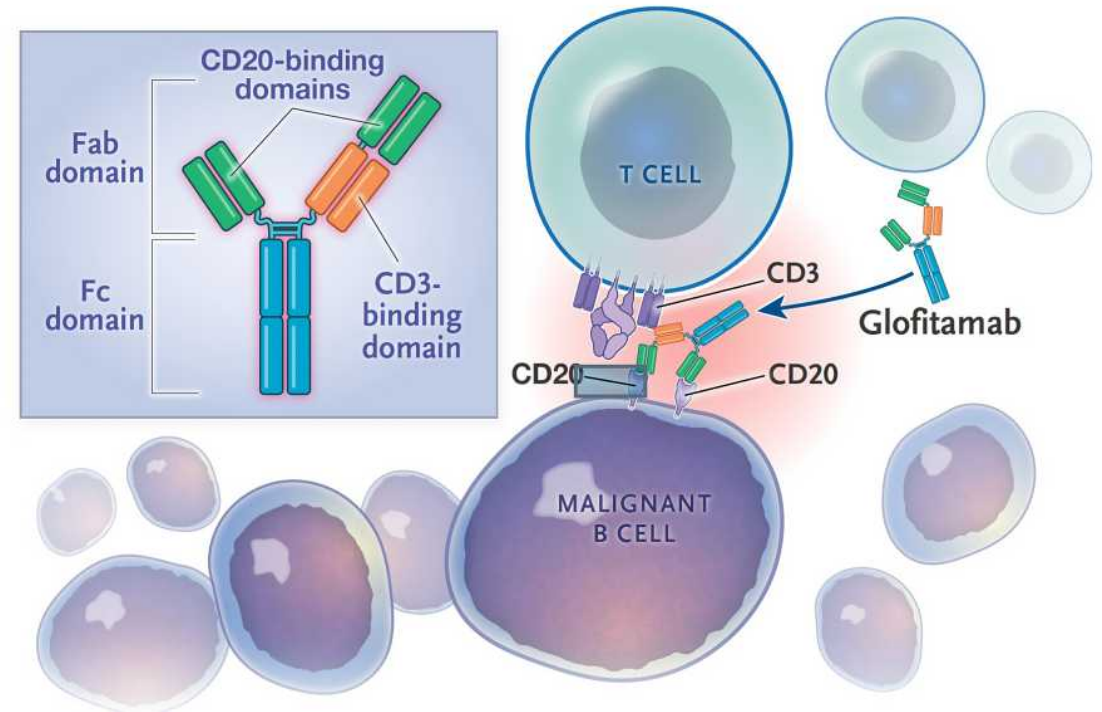
Acknowledgement

- Dr. Thomas Chan
- Prof. Eric Tse
- Prof. YL Kwong

- Roche HK – Compassionate Use Programme for Glofitamab

Introduction

- Bi-specific antibody against CD20 x CD3 for R/R DLBCL
- R/R DLBCL
- FDA-approved since June 2023





Total Patients
Analyzed

18



Study Centres



Queen Mary Hospital



Pamela Youde
Nethersole Eastern
Hospital



Data Collection
Period

2021-07-30
to
2025-08-12



Follow-up Duration

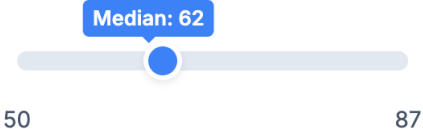
13.2 months
(Median)

Range: 1 - 47.3 months

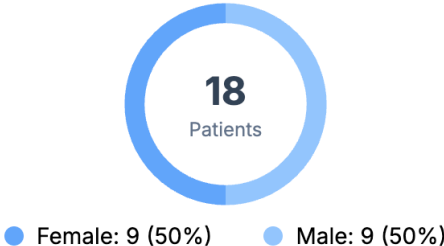
The Present Study

Demographics

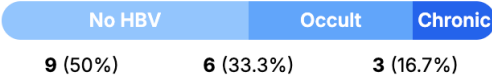
Age at Treatment Start



Sex Distribution



Hepatitis B Status



*Note: 1 patient had detectable HBV DNA <10 IU/mL.

Disease Characteristics

83.3%

Stage III-IV

100%

Extranodal Sites

38.9%

Bulky Disease

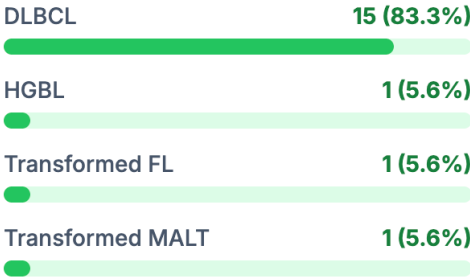
11.1%

Bone Marrow Inv.

0%

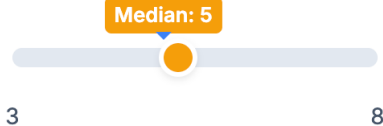
CNS Involvement

Histology

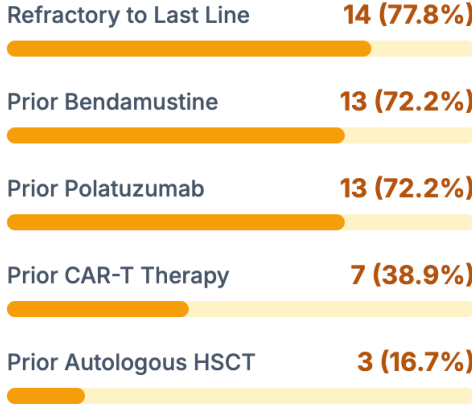


Prior Therapy

Prior Lines of Therapy



Prior Treatment Exposures



Key Characteristics of the Cohort

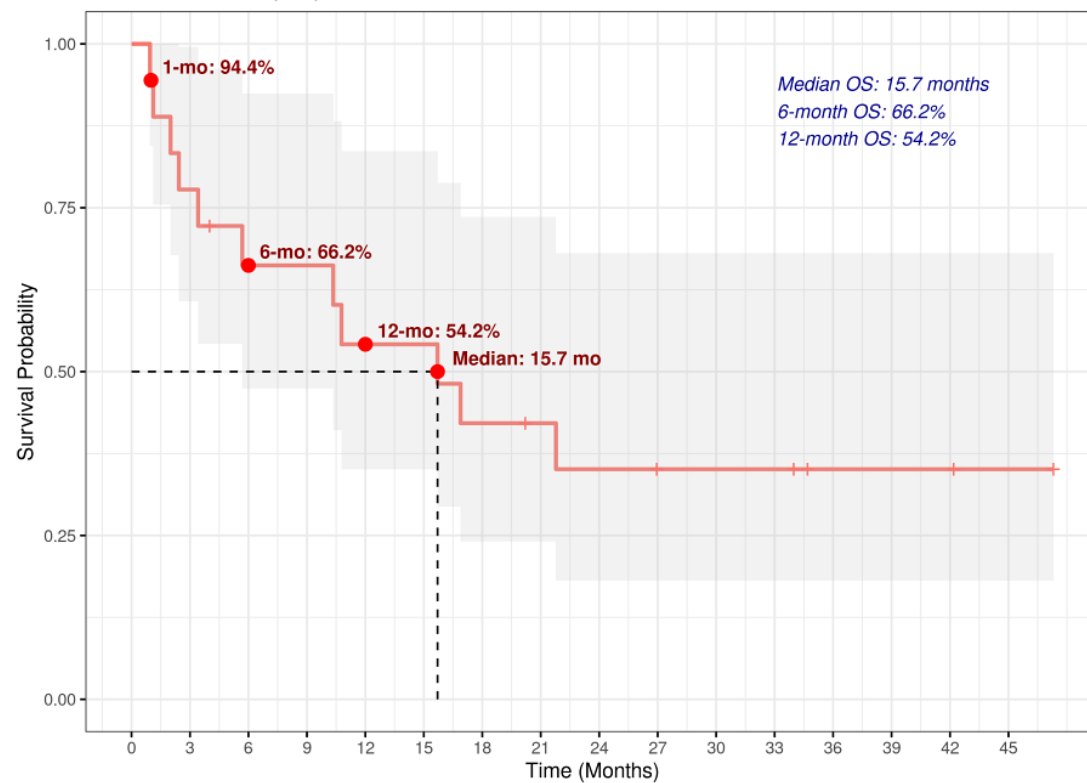
Glofitamab in Hong Kong - Our Early Experience

Each bar represents one patient's time on study

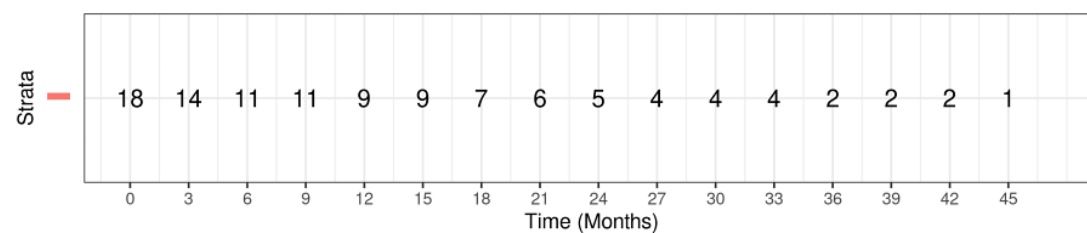


The HK Glofitamab Cohort in Swimmer's Plot

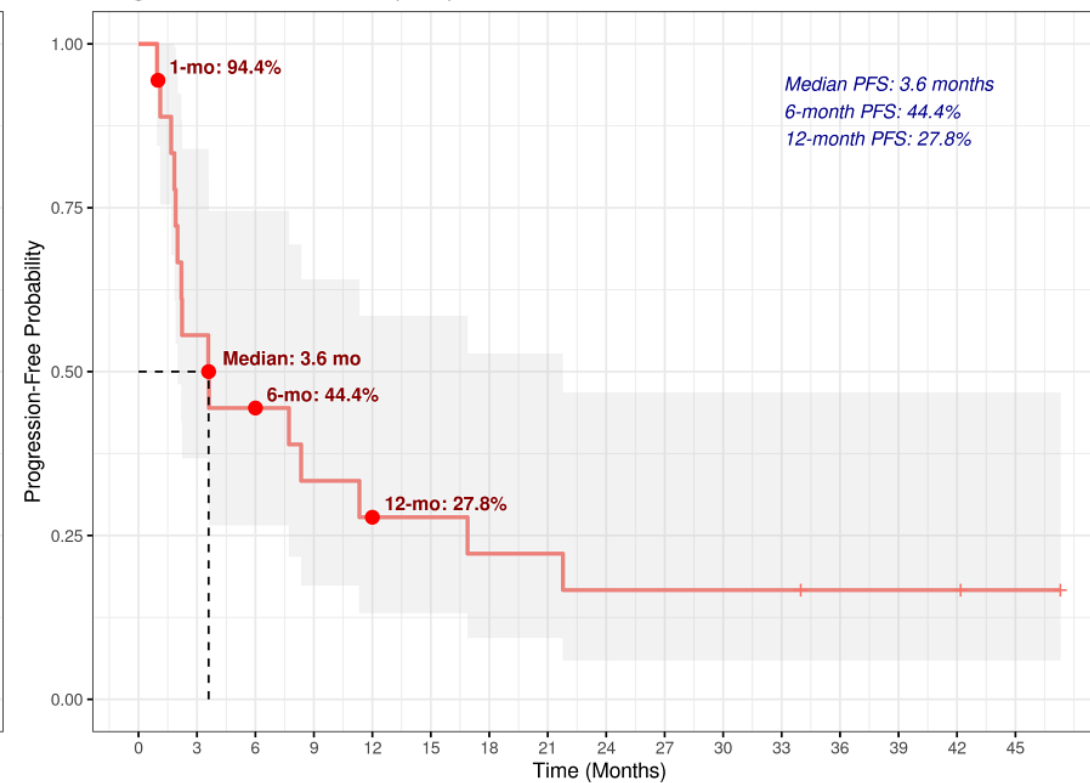
Overall Survival (OS) of the Glofitamab Cohort



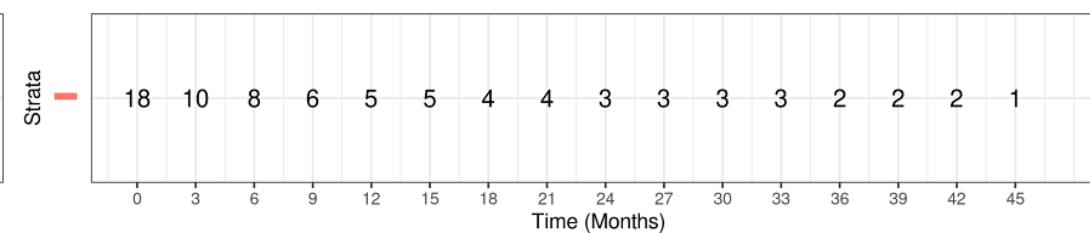
Number at risk



Progression-Free Survival (PFS) of the Glofitamab Cohort



Number at risk



Response & Survival

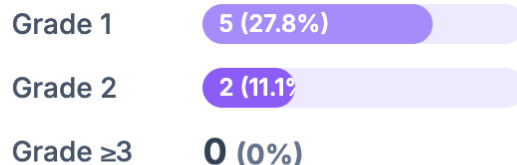
Safety

Cytokine Release Syndrome (CRS)

38.9%

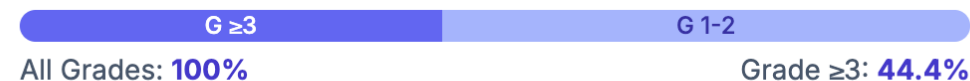
Any Grade CRS
(n=7)

Grade Breakdown

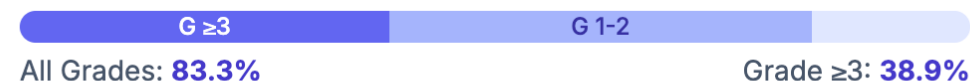


Hematologic Toxicities

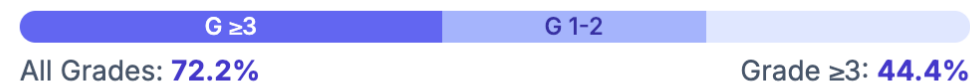
Anemia



Thrombocytopenia



Neutropenia



Immune Effector Cell-Associated Neurotoxicity (ICANS)

5.6%

Any Grade ICANS
(n=1)

5.6%

Grade ≥3 ICANS
(n=1)

Hong Kong Data in a Global Context

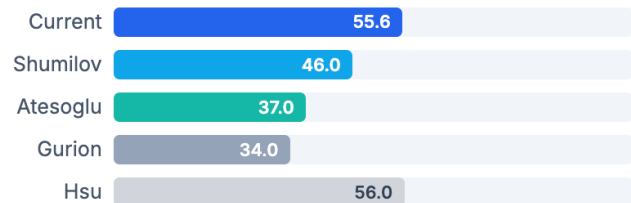
Metric	Current Study (N=18)	Shumilov '25 (N=70)	Atesoglu '23 (N=43)	Gurion '25 (N=35)	Hsu '24 (N=51)
ORR (%)	55.6	46	37	34	56
CR rate (%)	44.4	27	21	14	23
Median PFS (mo)	3.6	3.6	3.3	2.0	3.2
Median OS (mo)	15.7	5.7	8.8	4.0	8.4
CRS (All Grades, %)	38.9	39	28	37	N/A*
ICANS (All Grades, %)	5.6	10	7	11	6
Prior CAR-T (%)	38.9	13	14	23	8

*Hsu 2024 reported CRS events, not percentage of patients, and is not directly comparable.

■ Current Study (N=18) ■ Shumilov '25 (N=70) ■ Atesoglu '23 (N=43) ■ Gurion '25 (N=35) ■ Hsu '24 (N=51)

Efficacy Outcomes

Overall Response Rate (ORR %)



Complete Response (CR) Rate %

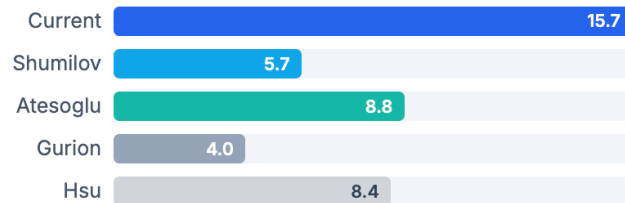


Survival Outcomes (Median Months)

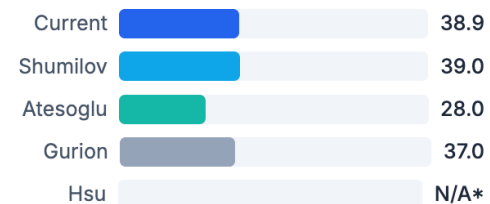
Progression-Free Survival (PFS)



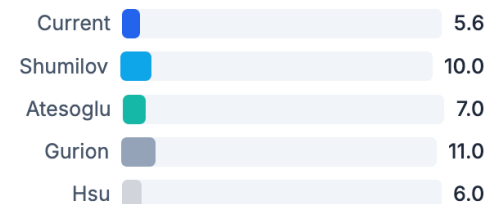
Overall Survival (OS)



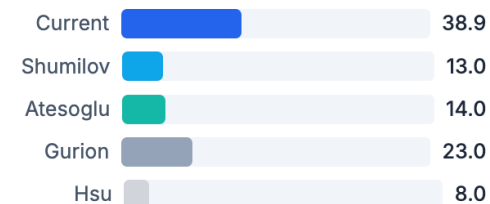
CRS (All Grades %)



ICANS (All Grades %)



Prior CAR-T Exposure %



Comparing HK Data with International Reports

Conclusion

- Real world efficacy on par with international experience
- Extra-nodal disease not a barrier to response
- HBV-DNA negativity provides sufficient clearance
- One instance of severe ICANS with NCSE

Thank you!