

Table 3: Level 2 Checklist for Screening Full-Text Articles and Study Reports

Reviewer: _____		Date: _____	
Ref ID:			
Author:			
Publication Year:			
Did the study include:	Yes (Include)	No (Exclude)	
A. The population of interest:			
• Adults (mean age of ≥ 18 years) with r/r-DLBCL	<input type="checkbox"/>	<input type="checkbox"/>	
• Mixed, with ≥ 80% being adults with r/r-DLBCL	<input type="checkbox"/>	<input type="checkbox"/>	
• Pediatric or young adult (≤ 25 years) with r/r-ALL	<input type="checkbox"/>	<input type="checkbox"/>	
• Mixed, with ≥ 80% being pediatric or young adult (≤ 25 years) with r/r-ALL	<input type="checkbox"/>	<input type="checkbox"/>	
B. The intervention of interest:			
• Tisagenlecleucel alone	<input type="checkbox"/>	<input type="checkbox"/>	
• Tisagenlecleucel together with drug interventions	<input type="checkbox"/>	<input type="checkbox"/>	
• Tisagenlecleucel together with HSCT	<input type="checkbox"/>	<input type="checkbox"/>	
C. A comparator of interest:			
• Other Car T-cell therapies	<input type="checkbox"/>	<input type="checkbox"/>	
• Conventional salvage therapy	<input type="checkbox"/>	<input type="checkbox"/>	
• No Comparator	<input type="checkbox"/>	<input type="checkbox"/>	
D. Outcome(s) of interest:			
• Objective efficacy outcomes (e.g., CR, PR, OS, PFS, etc.)	<input type="checkbox"/>	<input type="checkbox"/>	
• Quality of life	<input type="checkbox"/>	<input type="checkbox"/>	
• Safety outcomes such as Grade ≥ 3 AEs (e.g., CRS, prolonged cytopenias, infections and infestations, febrile neutropenia, neurological effects including hallucination and dysphasia, etc.)	<input type="checkbox"/>	<input type="checkbox"/>	
E. A study design of interest:			
• RCTs	<input type="checkbox"/>	<input type="checkbox"/>	
• Non-randomized controlled trials	<input type="checkbox"/>	<input type="checkbox"/>	
• Single-arm studies	<input type="checkbox"/>	<input type="checkbox"/>	
• Cohort studies	<input type="checkbox"/>	<input type="checkbox"/>	
• Case-control studies	<input type="checkbox"/>	<input type="checkbox"/>	
F. Notes:			
G. Selected for inclusion in the review ^a		Yes <input type="checkbox"/>	No <input type="checkbox"/>
H. Reason(s) for exclusion		<input type="checkbox"/> Irrelevant population <input type="checkbox"/> Irrelevant intervention <input type="checkbox"/> Irrelevant comparator <input type="checkbox"/> Irrelevant outcomes <input type="checkbox"/> Irrelevant study design <input type="checkbox"/> Other (specify):	

AE= adverse event; CAR = chimeric antigen receptor; CR = complete remission; HSCT = hematopoietic stem cell transplant; OS = overall survival; PFS = progression-free survival; PR = partial remission; RCT = randomized controlled trial; r/r-DLBCL = relapsed/refractory diffuse large B-cell lymphoma; r/r-ALL = relapsed/refractory B-cell acute lymphoblastic lymphoma.

^a Both reviewers must answer “Yes” to all questions for inclusion at the full-text level. If there is a discrepancy between the reviewers, disagreements will be resolved by discussion or with the involvement of a third reviewer, if necessary.