

Table 2: Level 1 Checklist for Screening Titles and Abstracts

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

AE= adverse event; CAR = chimeric antigen receptor; CR = complete remission; CRS = cytokine release syndrome; HSCT = hematopoietic stem cell transplant; OS = overall survival; PFS = progression-free survival; PR = partial remission; RCT = randomized controlled trial; WDAE = withdrawal due to adverse event.

^a "Unclear" means that it cannot be ascertained from the title or abstract if the report is potentially relevant to the review.

^b The full-text article of any title or abstract will be retrieved for further review if the response to all screening items above-noted are either "Yes" or "Unclear" by at least one of two independent reviewers.