

# INPLASY

## Effect of FGFR Alteration on Prognosis of Urothelial Carcinoma Patients with Immune Checkpoint Inhibitors: A Meta-Analysis

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### ADMINISTRATIVE INFORMATION

**Support** - None.

**Review Stage at time of this submission** - Completed but not published.

**Conflicts of interest** - None declared.

**INPLASY registration number:** INPLASY202430081

**Amendments** - This protocol was registered with the International Platform of Registered Systematic Review and Meta-Analysis Protocols (INPLASY) on 019 March 2024 and was last updated on 19 March 2024.

### INTRODUCTION

**Review question / Objective** The aim of this study was to compare prognosis and response to immune checkpoint inhibitors (ICIs) between fibroblast growth factor receptor (FGFR)-altered urothelial carcinoma (UC) patients and FGFR-wildtype UC patients.

**Condition being studied** UC patients treated with ICIs.

### METHODS

**Participant or population** UC patients treated with ICIs.

**Intervention** UC patients with FGFR alteration.

**Comparator** UC patients without FGFR alteration.

**Study designs to be included** Cohort studies.

**Eligibility criteria** None.

**Information sources** Pubmed, Embase, Medline, Cochrane Library and ClinicalTrial.gov were systematically searched prior to 1st February, 2024 in order to identify published articles in English.

**Main outcome(s)** Overall survival (OS) and tumor response assessment.

**Quality assessment / Risk of bias analysis** Newcastle-Ottawa Scale.

**Strategy of data synthesis** In assessing the relationship between OS and FGFR alteration in UC with ICIs, hazard ratios (HR) with corresponding 95% confidence intervals (CI) were employed. For the association between tumor response assessment and FGFR alteration in UC with ICIs, odds ratios (OR) with 95% CI were utilized. Heterogeneity among enrolled studies was gauged via Cochrane's Q test and Higgins I<sup>2</sup> statistic. Absence of obvious heterogeneity (P>0.05 for Cochrane's Q test and I<sup>2</sup><50%) led to

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the application of fixed-effect models; otherwise, random-effect models were employed. The statistical significance of pooled results was evaluated using the Z-test. Statistical analysis was conducted via Review Manager (version 5.3; The Cochrane Collaboration) and Stata (version 12.0; Stata Corporation).

**Subgroup analysis** Subgroup analyses were performed based on lines of ICI treatment and FGFR3 status.

**Sensitivity analysis** Sensitivity analysis measured the stability and robustness of pooled results using the leave-one-out method.

**Country(ies) involved** China.

**Keywords** immune checkpoint inhibitors; fibroblast growth factor receptor; urothelial carcinoma.

#### **Contributions of each author**

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