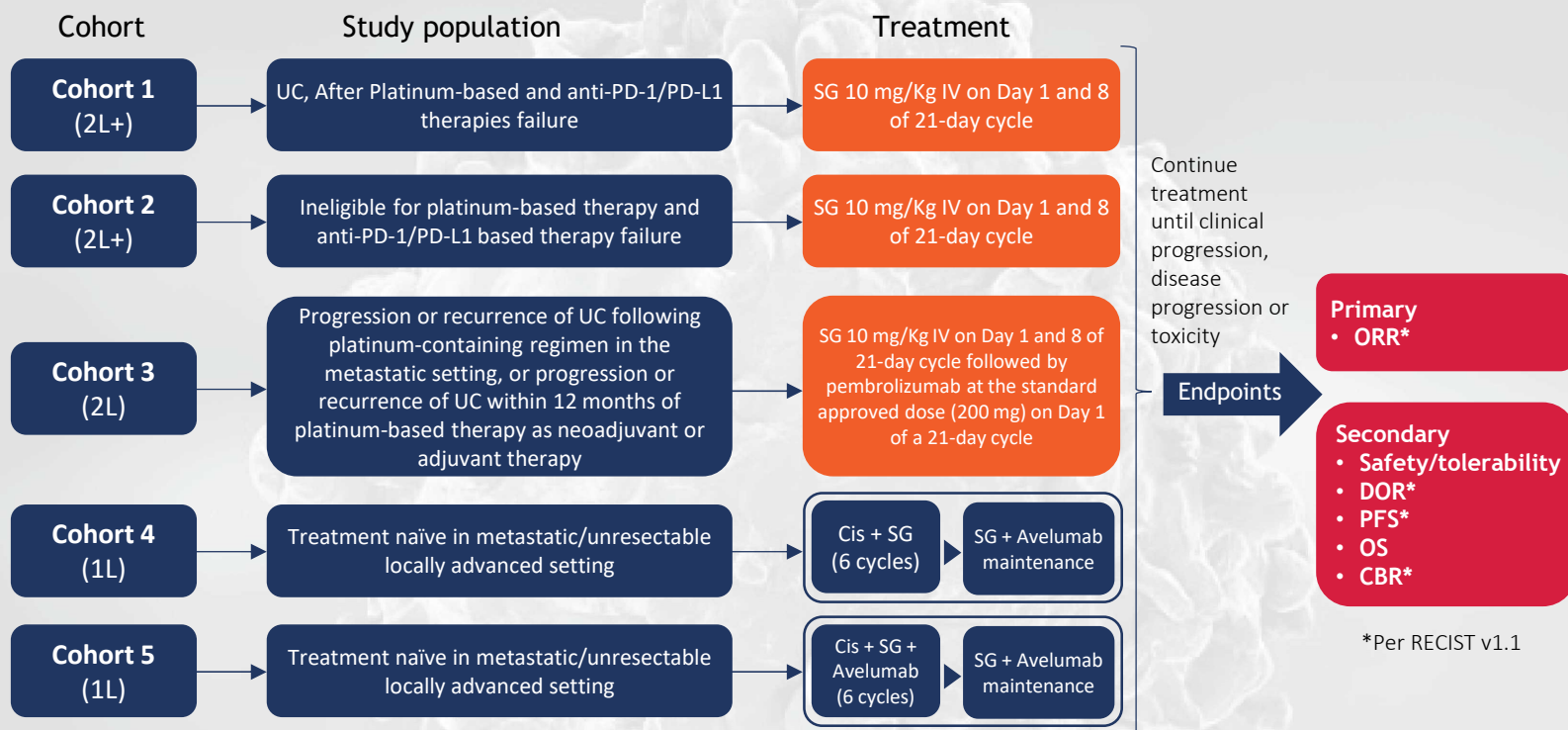


A Phase 2 Open-Label, Study of Sacituzumab Govitecan (SG) in Metastatic Urothelial Cancer (mUC) After Failure of Platinum-Based Regimen or Anti-PD-1/PD-L1 Based Immunotherapy

Study Design

Trial registration number: NCT03547973



Enrollment and key eligibility criteria

Key Inclusion Criteria:

- ECOG of 0 or 1;
- Cohorts 1-3: Creatinine clearance ≥ 30 mL/min; Cohorts 4-5: Creatinine clearance ≥ 50 mL/min
- Adequate organ function and stable brain metastases

Key Exclusion Criteria:

- Immunodeficiency,
- Active Hepatitis B or C,
- Active secondary malignancy

Questions?

- For more information on the TROPHY-01 Study, please contact GileadClinicalTrials@gilead.com

Safety and efficacy have not been demonstrated for the investigational uses described here. There is no guarantee that the investigational therapies or uses will be approved for use.

Visit clinicaltrials.gov for more information. Clinicaltrials.gov: NCT03547973

CBR, clinical benefit rate; Cis, cisplatin; DOR, duration of response; ECOG, Eastern Cooperative Oncology Group; ORR, objective response rate; OS, overall survival; PD-1, programmed cell death-1; PD-L1, programmed death-ligand 1; PFS, progression-free survival; RECIST, Response Evaluation Criteria in Solid Tumors, version 1.1; 1L, first line; 2L, second line

References:

National Institutes of Health. <https://clinicaltrials.gov/ct2/show/NCT03547973>. Accessed December 17, 2021.
Gilead Sciences Data on File.

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Oncology