

## BACKGROUND

Recent results from several clinical trials evaluating chemoimmunotherapy combination as neoadjuvant therapy have shown improved pathologic complete response (pCR) in some cancer types other than PGC. Thus, this phase II study (Neo-PLANET) was conducted to evaluate the efficacy and safety of camrelizumab plus chemoradiation as neoadjuvant therapy for locally advanced PGC.

#### Neo-PLANET (NCT03631615) is a single center, single arm, open-label phase II study.

- Primary endpoint: pCR
- survival), safety profile

<ul> <li>Patients with</li> <li>resectable gastric cancer; cT3-4aN+M0; proximal stomach; adenocarcinoma;</li> <li>age 18-75;</li> <li>KPS&gt;=80</li> </ul>	nec	Induction neochemotherapy XELOX 1 cycle	<u>Concurrent</u> <u>neochemoradiation</u> Radiotherapy 45Gy / 25 days Capecitabine 850mg/m2 , bid , po ,5W	Consolidation neochemotherapy XELOX 1 cycle	<u>Surgery</u> Total	Total
				gastrectomy, D2 lymph node dissection		

#### > Patient Characteristics

Between September 2018 and December 2020, 36 patients v enrolled. Patients' baseline characteristics are indicated in

### Completion of Neoadjuvant Treatment and Surger

Thirty-two (88.9%) patients completed neoadjuvant camrelizumab as planned. One patient with progressive disc and three with adverse events failed to complete the full cyc of camrelizumab.

Beyond one patient with liver metastasis, one with peritone metastasis and one patient who refused surgery, 33 (91.7%) underwent radical surgery. R0 resection was achieved in all patients undergoing surgery.

## **STUDY DESIGN**

Secondary endpoints: MPR (major pathologic response) rate , PFS (progression-free survival), OS (overall

# RESULTS

	Table 1. Patient Charac	atient Characteristics		
	Characteristics	Patients, n (%) (n=36)		
	Median age, yrs (range)	65.5 (35-72)		
were n table 1.	Male	28 (77.8)		
	ECOG PS 0	36 (100.0)		
ry	cT ■cT3 ■cT4a	6 (16.7) 30 (83.3)		
sease ycles eal (6)	cN ■cN0 ■cN+	0 36 (100.0)		
	Location of primary tumor Gastroesophageal junction Stomach body	19 (52.8) 17 (47.2)		
	Lauren classification Intestinal Diffuse Mixed NA	19 (52.8) 5 (13.9) 10 (27.8) 2 (5.6)		

#### > Efficacy

In the full analysis set, 12 (33.3%) patients achieved pCR in both primary tumor and lymph nodes; 28 (77.8%) achieved lymph node negative; 16 (44.4%) achieved MPR (table 2). Patients are still being followed up, and survival data such as PFS and OS will be further disclosed.

 Table 2. Clinical Outcomes

Characteristics	Patients, n (%) (n=36)	Adverse E	
Becker-TRG <ul> <li>la</li> <li>lb</li> <li>II</li> </ul>	12 (33.3) 4 (11.1) 8 (22.2)	Lymphocy White bloc	
<ul><li>III</li><li>Missing</li></ul>	9 (25.0) 3 (8.3)	Anemia Reactive of Platelet co	
pCR MPR	12 (33.3) 16 (44.4)		
урТ •урТ0 •урТ1 •урТ2 •урТ3 •урТ4	12 (33.3) 4 (11.1) 5 (13.9) 8 (22.2) 4 (11.1)	Neutropen Vomiting Nausea Hyperglyc	
ypN •ypN0 •ypN1 •ypN2 •ypN3	28 (77.8) 3 (8.3) 1 (2.8) 1 (2.8)	Hypocalc Weight lo Pruritus Hypothyr	

# CONCLUSIONS

Camrelizumab combined with chemoradiation in the neoadjuvant setting showed promising results in patients with locally advanced PGC, and further investigation is warranted in a phase III clinical trial.

## REFERENCES

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## > Safety

Among 36 patients who received at least one dose of the protocol treatment, treatment emergent adverse events (TEAEs) of any grade occurred in 36 (100%) patients and grade 3/4 were observed in 29 (80.56%) patients (table 3).

#### Patients, n (%) (n=36) events Any Grade Grade 3 35 (97.2) 24 (66.7) 3 (8.3) yte count decreased 26 (72.2) od cell decreased 2 (5.6) 25 (59.4) capillary hyperplasia 25 (69.4) 1 (2.8) 19 (52.8) ount decreased 0 0 17 (47.2) nia 0 0 16 (44.4) 0 0 1 (2.8) 15 (41.7) 1 (2.8) 6 (16.7) cemia 0 6 (16.7) aemia 0 5 (13.9) 0 SS 5 (13.9) 0 oidism 4 (11.1) 0 0

#### Table 3. TEAEs that occurred in ≥10% of patients