

Exploratory biomarker analysis of the phase 3 RATIONALE 305 trial: First-line (1L) tislelizumab (TIS) + chemotherapy (CT) vs placebo (PBO) + CT for advanced gastric cancer/gastro-oesophageal junction adenocarcinoma (GC/GEJC)

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ABSTRACT

Background

In the randomised phase 3 RATIONALE 305 trial (NCT03777657), patients (pts) with advanced GC/GEJC treated with 1L TIS + CT had significantly improved overall survival (OS) vs PBO + CT. We present exploratory biomarker and molecular subtyping results from RATIONALE 305 derived using RNA sequencing (RNAseq) and whole exome sequencing (WES) data.

Methods

RNAseq (507/997, 50.9%) of baseline (BL) tumor tissue was used to assess gene expression signatures (GES). GES subgroups were defined by median cutoff. WES (235/997, 23.6%) of BL tumor tissue and matching blood samples was performed to assess tumor mutational burden (TMB), human leukocyte antigen (HLA) genotyping, significantly mutated genes, and cytoband amplifications (amp). Associations between biomarker status and OS were evaluated. All *P* values are descriptive.

Results

BL characteristics in the GES- or WES-evaluable pts were similar to the ITT population. High inflammation (e.g. cytotoxic T cells [CTL]) and low immunosuppression (e.g. neutrophils) GES were associated with improved OS for TIS + CT vs PBO + CT (**Table**). High clonal

TMB was linked to improved OS with TIS + CT vs PBO + CT (**Table**) while total TMB showed no clear association. Non-HLA B27 supertype or *TP53* wildtype or 20q13.13 non-amp were associated with improved OS benefit with TIS + CT vs PBO + CT (**Table**). Additional biomarker analyses including immunohistochemistry and molecular subtyping results will be presented.

Conclusion

This exploratory biomarker analysis found that high CTL GES and low neutrophil GES, as well as clonal TMB, certain HLA genotypes, *TP53* wildtype, and cytoband non-amp were all associated with OS benefit in 1L TIS + CT vs PBO + CT treated pts with advanced GC/GEJC.

Table

Population	Subpopulation	TIS + CT vs PBO + CT			
		N	OS HR (95% CI)	P value	Interaction P value
GES evaluable		507			
	CTL high	254	0.69 (0.52, 0.91)	0.009	0.063
	CTL low	253	1.00 (0.76, 1.32)	0.999	
	Neutrophil high	254	1.01 (0.76, 1.33)	0.970	0.044
	Neutrophil low	253	0.67 (0.50, 0.89)	0.005	
WES evaluable		235			
	High cTMB (90 th percentile)	24	0.37 (0.14, 0.95)	0.038	0.125
	Low cTMB (90 th percentile)	211	0.80 (0.59, 1.08)	0.144	
	HLA B27	52	1.30 (0.70, 2.41)	0.401	0.045
	HLA non-B27	183	0.64 (0.46, 0.89)	0.007	
	TP53 mutation ^a	148	0.96 (0.67, 1.38)	0.825	0.025
	TP53 wildtype	84	0.48 (0.29, 0.78)	0.003	
	20q13.13 non-amp	205	0.64 (0.47, 0.88)	0.005	0.002
	20q13.13 amp	30	2.58 (1.15, 5.79)	0.021	

^aPts with TP53 amplification not included