

Table SI. Adverse events.

Adverse events	Cohort 1 (N=31)		Cohort 2 (N=40)	
	Any, n (%)	Grade ≥ 3 , n (%)	Any, n (%)	Grade ≥ 3 , n (%)
Hypertension	5 (16.1)	0 (0.0)	5 (12.5)	1 (2.5)
Nausea	6 (19.3)	2 (6.5)	4 (10.0)	0 (0.0)
Diarrhea	3 (9.7)	1 (3.2)	9 (22.5)	1 (2.5)
Fatigue	3 (9.7)	0 (0.0)	7 (17.5)	0 (0.0)
Decreased appetite	3 (9.7)	0 (0.0)	5 (12.5)	0 (0.0)
Peripheral neuropathy	11 (35.4)	3 (9.7)	3 (7.5)	1 (2.5)
Neutropenia	8 (25.8)	2 (6.5)	18 (45.0)	3 (7.5)
Thrombocytopenia	3 (9.7)	1 (3.2)	2 (5.0)	0 (0.0)
Leukopenia	9 (29.0)	2 (6.5)	18 (45.0)	3 (7.5)
Proteinuria	0 (0.0)	0 (0.0)	3 (7.5)	0 (0.0)