Supplementary Table 2: Adverse Events Occurring in at Least Two Patients, by Dose Level

	Dose Level 1				Dose Level 2				Dose Level 3				Total on all Dose Levels				
Toxicity grade:	**1**	**2**	**3**	**4**	**1**	**2**	**3**	**4**	**1**	**2**	**3**	**4**	**1**	**2**	**3**	**4**	**Total**
Toxicity:																	
Anemia	1	2	0	0	0	1	1	0	5	4	0	0	6	7	1	0	14
Platelet count decreased	2	0	0	1	1	1	0	1	2	2	2	1	5	3	2	3	13
Creatinine increased	3	0	0	0	1	2	0	0	3	2	0	0	7	4	0	0	11
Hyperglycemia	0	1	0	0	2	1	0	0	5	1	0	0	7	3	0	0	10
Fatigue	1	0	0	0	1	1	0	0	4	1	0	0	6	2	0	0	8
Hypocalcemia	0	0	0	0	1	1	0	0	6	0	0	0	7	1	0	0	8
Dyspnea	2	0	0	0	1	0	0	0	2	2	0	0	5	2	0	0	7
Aspartate aminotransferase increased	1	0	0	0	1	0	0	0	4	0	0	0	6	0	0	0	6
Neutrophil count decreased	0	1	0	0	0	0	1	1	0	1	3	0	0	2	4	0	6
Alkaline phosphatase increased	0	1	0	0	1	1	0	0	1	0	0	0	2	2	0	0	4
Anorexia	3	0	0	0	0	0	0	0	1	0	0	0	4	0	0	0	4
Cough	1	0	0	0	3	0	0	0	0	0	0	0	4	0	0	0	4
Edema limbs	0	0	0	0	1	0	0	0	2	0	0	0	3	0	0	0	3
Lung infection	0	0	0	0	0	1	0	0	1	1	0	0	1	2	0	0	3
Alanine aminotransferase increased	0	1	0	0	1	0	0	0	1	0	0	0	2	1	0	0	3
Blood bilirubin increased	1	0	0	0	0	0	0	0	2	0	0	0	3	0	0	0	3
Hypercalcemia	1	0	0	0	1	0	0	0	1	0	0	0	3	0	0	0	3
Hyperuricemia	0	0	0	0	1	0	0	0	1	0	1	0	2	0	1	0	3
Hypomagnesemia	0	0	0	0	0	0	0	0	3	0	0	0	3	0	0	0	3
Hypophosphatemia	0	0	0	0	0	1	0	0	0	2	0	0	0	3	0	0	3
General disorders & administration site conditions, other	0	0	0	0	1	0	0	0	1	0	0	0	2	0	0	0	2
Erythema multiforme	1	0	0	0	1	0	0	0	0	0	0	0	2	0	0	0	2
Rash acneiform	0	0	0	0	0	0	0	0	1	1	0	0	1	1	0	0	2
Skin and subcutaneous tissue disorders, other	0	0	0	0	1	0	0	0	1	0	0	0	2	0	0	0	2
Diarrhea	0	0	0	0	1	0	0	0	1	0	0	0	2	0	0	0	2
Nausea	1	0	0	0	1	0	0	0	0	0	0	0	2	0	0	0	2

White blood cell decreased	0	0	0	0	0	0	1	0	0	0	1	0	0	0	2	0	2
Alkalosis	0	0	0	0	1	0	0	0	1	0	0	0	2	0	0	0	2
Hyperkalemia	1	0	0	0	0	0	0	0	1	0	0	0	2	0	0	0	2
Hyponatremia	1	0	0	0	1	0	0	0	0	0	0	0	2	0	0	0	2
Peripheral sensory neuropathy	0	0	0	0	1	0	0	0	1	0	0	0	2	0	0	0	2
Acute kidney injury	0	0	0	0	0	0	0	0	1	1	0	0	1	1	0	0	2