

Supplementary Table S2. Phase I patient disposition

Parameter, <i>n</i> (%)	Group I				Group II				
	Nintedanib, 100 mg bid	Nintedanib, 150 mg bid	Nintedanib, 200 mg bid	Total	Nintedanib, 50 mg bid	Nintedanib, 100 mg bid	Nintedanib, 150 mg bid	Nintedanib, 200 mg bid	Total
Discontinued from trial medication	6 (100)	3 (100)	4 (100)	13 (100)	3 (100)	4 (100)	4 (100)	8 (100)	19 (100)
Progressive disease	2 (33.3)	1 (33.3)	1 (25.0)	4 (30.8)	0	2 (50.0)	1 (25.0)	1 (12.5)	4 (21.1)
AEs	4 (66.7)	2 (66.7)	3 (75.0)	9 (69.2)	3 (100)	2 (50)	2 (50)	7 (87.5)	14 (73.7)
DLT	2 (33.3)	1 (33.3)	0	3 (23.1)	0	1 (25.0)	0	2 (25.0)	3 (15.8)
Other AE	2 (33.3)	1 (33.3)	3 (75.0)	6 (46.2)	3 (100)	1 (25.0)	2 (50.0)	5 (62.5)	11 (57.9)
Non-compliant with protocol	0	0	0	0	0	0	0	0	0
Lost to follow-up	0	0	0	0	0	0	0	0	0
Refused to continue taking trial medication	0	0	0	0	0	0	0	0	0
Other	0	0	0	0	0	0	1 (25.0)	0	1 (5.3)

Abbreviations: AE, adverse events; DLT, dose-limiting toxicity

