	Part A*		Part B							
PK Parameters	First Dose 6th Dose		First Dose				Last dose			
	100 mg every 2	100 mg every 2	100 mg every 2	100 mg every 4	50 mg every 4 weeks	25 mg every 4	100 mg every 2	100 mg every 4	every 4	25 mg every 4 weeks
	weeks	weeks	weeks	weeks		weeks	weeks	weeks	weeks	
n	14	11	29	29	28	31	26	22	21	28
C _{max} (µg/mL)	7.50 (2.91)	20.01 (4.40)	6.60 (3.03)	6.73 (2.73)	2.99 (1.51)	1.86 (1.11)	15.68 (7.07)	9.34 (4.31)	4.26 (1.73)	2.93 (1.29)
t _{max} (day)	3.9	4.0	4.9	5.0	5.4	5.0	5.0	4.0	5.0	4.0
	[2.2, 13.8]	[3.0, 7.0]	[2.8, 10.7]	[3.0, 13.9]	[2.9, 12.6]	[2.9, 27.8]	[0, 9.7]	[0, 7.1]	[0, 7.0]	[2.8, 11.7]
$AUC_{(t1-t2)}$	80.63	232.43	66.03	107.88	49.38	29.08	188.67	172.04	83.52	52.02 (22.03)
(µg.day/mL)†	(25.66)	(48.90)	(31.62)	(43.05)	(23.15)	(10.65)	(90.43)	(86.17)	(35.17)	
t _{1/2} (day)	NA	18.3 (4.5)	NA	NA	NA	NA	19.0 (4.3)	16.0 (4.1)	16.1 (4.8)	14.9 (4.5)
R _{Cmax}	NA	2.7 (1.1)	NA	NA	NA	NA	2.6 (1.0)	1.5 (0.9)	1.7 (0.6)	1.6 (0.6)
R _{AUC(0-14d)}	NA	2.7 (0.5)	NA	NA	NA	NA	3.0 (1.2)	1.6 (0.8)	1.8 (0.6)	1.7 (0.4)

Supplemental Table 2. Derived sirukumab pharmacokinetic parameters

Data presented as mean (SD) or median [range] unless noted otherwise. * Data from one of the study sites that participated in Part A were excluded from all pharmacokinetics analyses due to questionable data integrity. $\dagger AUC_{(t1-t2)}$, area under the serum concentration versus time curve between 2 defined sampling timepoints (AUC_{0-14d} was reported for the every-2-week treatment group; AUC_{0-28d} was reported for q4w treatment groups); PK, pharmacokinetic; C_{max}, observed maximum serum concentration; NA, not available; R_{AUC(t1-t2)}, accumulation ratio calculated from AUC_(t1-t2) after the last dose of multiple doses and AUC_(t1-t2) after the first dose; R_{Cmax}, accumulation ratio calculated from C_{max} after the last dose of multiple doses and C_{max} after the first dose; t_{1/2}, terminal half-life; t_{max}, observed time to reach maximum serum concentration.