

Supplementary Table S1. Baseline and follow-up characteristics of MTX dose escalation patients stratified by achievement of 3V-remission.

	3V-remission achieved n=63	3V-remission not achieved n=130	<i>p</i> -value
Characteristics at baseline (T-1)			
Age at diagnosis (years)	57 (51-68)	52 (43-63)	0.016
Sex (F)	46/63 (73.0)	106/130 (81.5)	0.192
Positive RF	38/63 (60.3)	80/130 (61.5)	0.876
Positive ACPA	33/63 (52.4)	75/130 (57.7)	0.537
ANA ≥1:160	9/63 (14.3)	36/130 (27.7)	0.046
Joint erosions	8/63 (12.7)	33/130 (25.4)	0.06
MTX weekly dose (mg)	10 (7.5-10)	10 (7.5-10)	0.137
CRP (mg/dl)	0.9 (0.34-2.7)	1.5 (0.4-3.9)	0.096
ESR (mm/h)	26 (16-49)	35 (17-51)	0.197
TJC	4 (2-12)	5 (2-11)	0.443
SJC	3 (1-10)	2 (1-10)	0.693
Characteristics at the moment of treatment change (T0)			
MTX dose	15 (15-15)	15 (15-15)	0.094
CRP (mg/dl)	0.7 (0.3-1.3)	1.4 (0.6-3.7)	<0.001
ESR (mm/h)	26 (14-37)	26 (16-41)	0.581
TJC	2 (1-10)	3 (1-10)	0.499
SJC	2 (0-5)	2 (0-5)	0.595
Characteristics 6 months after MTX escalation (T1)			
CRP (mg/dl)	0.3 (0.1-0.5)	1.2 (0.4-3.0)	<0.001
ESR (mm/h)	20 (11-29)	27 (14-43)	0.003
TJC	0 (0-0)	2 (0-4)	<0.001
SJC	0 (0-0)	3 (1-4)	<0.001

Data are shown as median (interquartile range) or number (percentage).

MTX: methotrexate; CRP: C-reactive protein; ESR: erythrocyte sedimentation rate; TJC: tender joint count; SJC: swollen joint count; b/tsDMARD: biologic/targeted synthetic disease-modifying anti-rheumatic drug.