

CORRECTION

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Correction to: Sarilumab plus methotrexate in patients with active rheumatoid arthritis and inadequate response to methotrexate: results of a randomized, placebo-controlled phase III trial in Japan

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Following publication of the original article [1], the authors reported an error in Table 2. The data for 'HAQ-DI response (MCID \geq 0.3), n (%) / At week 52' for the 'Sarilumab 150 mg q2w + MTX' and 'Sarilumab 200 mg q2w + MTX' groups should be 41 (50.6) and 37 (46.3), respectively (last row, last two entries of the table).

The corrected table is given below.

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1. Tanaka, et al. Sarilumab plus methotrexate in patients with active rheumatoid arthritis and inadequate response to methotrexate: results of a randomized, placebo-controlled phase III trial in Japan. *Arthritis Res Ther.* 2019;21:79. <https://doi.org/10.1186/s13075-019-1856-4>.

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Table 2 Efficacy results (mITT population)

	Sarilumab			
	Placebo to 150 mg q2w + MTX (n = 41 (n = 14 at week 52)) ^a	Placebo to 200 mg q2w + MTX (n = 40 (n = 15 at week 52)) ^a	150 mg q2w + MTX (n = 81)	200 mg q2w + MTX (n = 80)
Signs and symptoms				
ACR20 response, n (%)				
At week 12	15 (18.5)		54 (66.7) ^{***}	52 (65.0) ^{***}
At week 24	12 (14.8)		55 (67.9) ^{***}	46 (57.5) ^{***}
At week 52	9 (64.3)	10 (66.7)	58 (71.6)	48 (60.0)
ACR50 response, n (%)				
At week 12	5 (6.2)		22 (27.2) ^{***}	25 (31.3) ^{***}
At week 24	8 (9.9)		35 (43.2) ^{***}	31 (38.8) ^{***}
At week 52	8 (57.1)	10 (66.7)	37 (45.7)	38 (47.5)
ACR70 response, n (%)				
At week 12	1 (1.2)		5 (6.2)	15 (18.8) ^{***}
At week 24	3 (3.7)		15 (18.5) ^{**}	12 (15.0) [*]
At week 52	4 (28.6)	3 (20.0)	29 (35.8)	22 (27.5)
ACR components, mean (SD) change from baseline at week 24				
Tender joint count	-9.1 (10.2)		-13.4 (9.9)	-12.4 (11.3)
Swollen joint count	-7.2 (6.7)		-10.6 (8.1)	-9.5 (9.1)
Pain VAS	-22.9 (27.7)		-36.5 (23.4)	-30.2 (23.3)
Physician global VAS	-26.8 (18.4)		-41.8 (21.6)	-43.9 (19.4)
Patient global VAS	-18.3 (22.6)		-32.4 (21.0)	-30.6 (21.9)
HAQ-DI	-0.3 (0.4)		-0.5 (0.5)	-0.6 (0.5)
CRP, mg/l	-1.7 (12.2)		-21.1 (19.5)	-21.3 (18.0)
DAS28-CRP response, mean (SD) change from baseline				
At week 12	-0.8 (1.1)		-2.3 (1.1) ^{***}	-2.3 (1.2) ^{***}
At week 24	-1.5 (1.2)		-2.8 (1.0) ^{***}	-2.8 (1.1) ^{***}
At week 52	-3.1 (1.2)	-2.9 (1.2)	-3.2 (1.2)	-3.2 (1.1)
DAS28-CRP < 2.6, n (%)				
At week 12	3 (3.7)		21 (25.9) ^{***}	27 (33.8) ^{***}
At week 24	6 (7.4)		29 (35.8) ^{***}	32 (40.0) ^{***}
At week 52	7 (50.0)	9 (60.0)	41 (50.6)	43 (53.8)
SDAI, mean (SD) change from baseline				
At week 12	-8.9 (12.0)		-20.7 (11.0) ^{***}	-18.9 (11.6) ^{***}
At week 24	-16.0 (11.6)		-25.2 (11.6) ^{***}	-23.8 (11.3) ^{***}
At week 52	-29.6 (9.9)	-23.4 (12.4)	-29.4 (13.6)	-26.9 (11.5)
SDAI ≤ 3.3, n (%)				
At week 12	0		2 (2.5)	7 (8.8) ^{**}
At week 24	1 (1.2)		5 (6.2)	10 (12.5) ^{**}
At week 52	2 (14.3)	1 (6.7)	19 (23.5)	18 (22.5)
CDAI, mean (SD) change from baseline				
At week 12	-8.7 (11.4)		-18.8 (10.6) ^{***}	-16.8 (10.9) ^{***}
At week 24	-15.7 (11.1)		-23.1 (11.2) ^{***}	-21.7 (10.7) ^{***}
At week 52	-28.4 (9.7)	-21.1 (11.4)	-27.2 (13.1)	-24.8 (10.8)

Table 2 Efficacy results (mITT population) (Continued)

	Sarilumab			
	Placebo to 150 mg q2w + MTX (n = 41 (n = 14 at week 52)) ^a	Placebo to 200 mg q2w + MTX (n = 40 (n = 15 at week 52)) ^a	150 mg q2w + MTX (n = 81)	200 mg q2w + MTX (n = 80)
CDAI ≤ 2.8, n (%)				
At week 12	0		1 (1.2)	5 (6.3)*
At week 24	1 (1.2)		5 (6.2)	8 (10.0)*
At week 52	1 (7.1)	0	17 (21.0)	15 (18.8)
Physical function				
HAQ-DI, mean (SD) change from baseline				
At week 12	-0.1 (0.3)		-0.4 (0.5)***	-0.4 (0.5)***
At week 24	-0.3 (0.4)		-0.5 (0.5)***	-0.6 (0.5)***
At week 52	-0.7 (0.6)	-0.5 (0.3)	-0.6 (0.6)	-0.6 (0.6)
HAQ-DI response (MCID ≥ 0.3), n (%)				
At week 12	19 (23.5)		39 (48.1)**	38 (47.5)**
At week 16	19 (23.5)		37 (45.7)**	37 (46.3)**
At week 24	10 (12.3)		39 (48.1)***	39 (48.8)***
At week 52	9 (64.3)	8 (53.3)	41 (50.6)	37 (46.3)

* $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$ ^aData for combined placebo groups (n=81) shown at weeks 12, 16 and 24. ACR American College of Rheumatology, ACR20/50/70 American College of Rheumatology 20%/50%/70% improvement criteria, CDAI Clinical Disease Activity Index, CRP C-reactive protein, DAS28 Disease Activity Score 28-joint count, HAQ-DI Health Assessment Questionnaire-Disability Index, MCID minimum clinically important difference, mITT modified intent-to-treat, MTX methotrexate, q2w every 2 weeks, SDAI Simplified Disease Activity Index, SD standard deviation, SJC swollen joint count, TJC tender joint count, VAS visual analog scale