- 1 Long-Term Health-Related Quality of Life in Patients with Plaque Psoriasis
- 2 Treated with Certolizumab Pegol: Three-Year Results from Two
- 3 Randomised Phase 3 Studies (CIMPASI-1 and CIMPASI-2)

4 SUPPORTING INFORMATION

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1 SUPPLEMENTARY TABLES AND FIGURES

	CZP 200 mg Q2W (N=183)			C	ZP 400 mg (2W (N=173)	All CZP (N=356)				Placebo (N=97)		
	Baseline	Week 16	Week 48	Week 144	Baseline	Week 16	Week 48	Week 144	Baseline	Week 16	Week 48	Week 144	Baseline	Week 16
DLQI scores, me	ean ± SD											•		
Total DLQI	14.3 ± 7.4	4.3 ± 5.7	3.9 ± 5.4	4.3 ± 5.8	13.7 ± 6.9	3.9 ± 5.8	2.8 ± 5.2	4.1 ± 6.3	14.0 ± 7.1	4.1 ± 5.7	3.4 ± 5.3	4.2 ± 6.0	13.4 ± 7.8	10.3 ± 6.8
Symptoms and Feelings	4.2 ± 1.5	1.4 ± 1.4	1.4 ± 1.4	1.5 ± 1.5	4.1 ± 1.4	1.3 ± 1.4	1.1 ± 1.4	1.4 ± 1.6	4.2 ± 1.4	1.4 ± 1.4	1.2 ± 1.4	1.5 ± 1.6	4.0 ± 1.5	3.2 ± 1.6
Leisure	2.7 ± 2.0	0.7 ± 1.3	0.6 ± 1.2	0.6 ± 1.3	2.4 ± 1.9	0.6 ± 1.2	0.5 ± 1.3	0.7 ± 1.5	2.5 ± 2.0	0.6 ± 1.2	0.6 ± 1.3	0.7 ± 1.4	2.5 ± 2.2	1.7 ± 1.7
Daily Activities	3.2 ± 1.9	0.9 ± 1.3	0.8 ± 1.2	0.8 ± 1.3	3.2 ± 1.7	0.8 ± 1.4	0.6 ± 1.3	0.9 ± 1.6	3.2 ± 1.8	0.8 ± 1.3	0.7 ± 1.3	0.8 ± 1.5	3.1 ± 1.8	2.3 ± 1.6
Work and School	0.9 ± 1.0	0.2 ± 0.6	0.2 ± 0.4	0.2 ± 0.6	0.9 ± 1.0	0.2 ± 0.7	0.2 ± 0.6	0.2 ± 0.7	0.9 ± 1.0	0.2 ± 0.6	0.2 ± 0.5	0.2 ± 0.6	0.9 ± 1.1	0.6 ± 0.9
Personal Relationships	2.0 ± 1.9	0.7 ± 1.3	0.5 ± 1.1	0.6 ± 1.2	1.9 ± 1.9	0.6 ± 1.4	0.4 ± 1.2	0.6 ± 1.4	1.9 ± 1.9	0.7 ± 1.3	0.5 ± 1.2	0.6 ± 1.3	1.8 ± 2.1	1.6 ± 1.8
Treatment	1.3 ± 1.1	0.4 ± 0.7	0.3 ± 0.6	0.4 ± 0.7	1.2 ± 1.1	0.3 ± 0.7	0.2 ± 0.6	0.4 ± 0.8	1.3 ± 1.1	0.4 ± 0.7	0.3 ± 0.6	0.4 ± 0.7	1.2 ± 1.0	0.9 ± 0.9
EQ-5D-3L														
EQ-5D-3L score														
Mobility	123 (67.2)	152 (83.1)	153 (83.6)	144 (78.7)	125 (72.3)	143 (82.7)	142 (82.1)	135 (78.0)	248 (69.7)	295 (82.9)	295 (82.9)	279 (78.4)	73 (75.3)	70 (72.2)
Self-Care	165 (90.2)	171 (93.4)	170 (92.9)	169 (92.3)	153 (88.4)	163 (94.2)	163 (94.2)	161 (93.1)	318 (89.3)	334 (93.8)	333 (93.5)	330 (92.7)	91 (93.8)	89 (91.8)
Usual Activities	125 (68.3)	150 (82.0)	158 (86.3)	151 (82.5)	115 (66.5)	143 (82.7)	147 (85.0)	140 (80.9)	240 (67.4)	293 (82.3)	305 (85.7)	291 (81.7)	65 (67.0)	64 (66.0)
Pain/Discomfort	33 (18.0)	100 (54.6)	106 (57.9)	101 (55.2)	42 (24.3)	108 (62.4)	110 (63.6)	96 (55.5)	75 (21.1)	208 (58.4)	216 (60.7)	197 (55.3)	16 (16.5)	28 (28.9)
Anxiety/ Depression	93 (50.8)	131 (71.6)	125 (68.3)	119 (65.0)	87 (50.3)	132 (76.3)	125 (72.3)	123 (71.1)	180 (50.6)	263 (73.9)	250 (70.2)	242 (68.0)	50 (51.5)	52 (53.6)
EQ VAS score,	65.9 ±	77.5 ±	78.6 ±	78.3 ±	66.1 ±	78.5 ±	81.5 ±	78.3 ±	66.0 ±	78.0 ±	80.0 ±	78.3 ±	66.2 ±	68.2 ±
mean ± SD	22.7	18.0	17.8	17.6	22.2	17.2	15.3	17.0	22.4	17.6	16.7	17.3	21.2	20.2
SF-36 scores, m	ean ± SD	P									P			
Physical Functioning	47.7 ± 9.6	51.2 ± 8.5	51.6 ± 8.3	50.2 ± 9.3	49.1 ± 9.4	51.3 ± 9.7	51.6 ± 9.1	50.4 ± 9.8	48.4 ± 9.5	51.3 ± 9.1	51.6 ± 8.7	50.3 ± 9.5	48.5 ± 8.8	49.6 ± 9.1
Role Physical	46.7 ± 9.5	50.3 ± 7.9	50.4 ± 8.5	49.0 ± 9.5	47.3 ± 10.0	50.5 ± 8.9	51.0 ± 8.9	50.3 ± 9.0	47.0 ± 9.7	50.4 ± 8.4	50.7 ± 8.7	49.6 ± 9.3	47.1 ± 8.8	47.8 ± 8.5
Bodily Pain	45.3 ± 10.4	52.2 ± 10.0	52.2 ± 9.5	50.7 ± 10.2	46.6 ± 11.1	53.4 ± 10.2	53.5 ± 9.9	51.7 ± 10.8	45.9 ± 10.7	52.8 ± 10.1	52.8 ± 9.7	51.2 ± 10.5	45.5 ± 10.7	47.0 ± 10.3
General Health	47.1 ± 10.1	50.7 ± 9.6	49.7 ± 9.8	49.6 ± 10.3	47.5 ± 9.3	50.3 ± 9.9	51.0 ± 10.2	49.0 ± 10.2	47.3 ± 9.7	50.5 ± 9.7	50.3 ± 10.0	49.3 ± 10.3	46.4 ± 10.4	47.2 ± 10.4
Mental Health	46.8 ± 10.7	51.7 ± 9.6	51.3 ± 10.1	50.0 ± 10.8	46.3 ± 10.1	50.5 ± 9.9	51.1 ± 9.9	50.2 ± 10.5	46.5 ± 10.4	51.1 ± 9.8	51.2 ± 10.0	50.1 ± 10.6	44.9 ± 12.0	46.0 ± 11.8

Table S1. Patient-reported outcomes to Week 144 (LOCF)

Role Emotional	46.8 ± 10.4	50.3 ± 8.9	50.1 ± 9.1	49.5 ± 10.1	46.1 ± 10.8	49.8 ± 9.6	50.7 ± 9.0	49.7 ± 9.6	46.4 ± 10.6	50.1 ± 9.2	50.4 ± 9.0	49.6 ± 9.8	46.7 ± 10.3	47.3 ± 9.7
Social Functioning	45.4 ± 11.1	51.0 ± 8.9	50.5 ± 9.6	49.9 ± 9.8	45.1 ± 11.0	50.7 ± 9.2	51.6 ± 8.9	50.1 ± 10.0	45.2 ± 11.0	50.9 ± 9.1	51.1 ± 9.3	50.0 ± 9.9	45.0 ± 11.7	45.6 ± 10.8
Vitality	48.5 ± 9.8	53.0 ± 9.7	53.7 ± 9.8	52.8 ± 10.6	47.6 ± 10.0	51.9 ± 10.2	53.7 ± 10.3	52.3 ± 10.6	48.1 ± 9.9	52.5 ± 9.9	53.7 ± 10.0	52.5 ± 10.6	47.2 ± 9.7	47.8 ± 10.1
PCS	46.9 ± 9.3	51.2 ± 8.5	51.2 ± 8.5	50.1 ± 9.6	48.5 ± 9.2	51.9 ± 9.6	52.1 ± 9.0	50.6 ± 9.4	47.7 ± 9.3	51.5 ± 9.1	51.7 ± 8.7	50.3 ± 9.5	47.7 ± 8.9	48.8 ± 9.3
MCS	46.9 ± 11.0	51.4 ± 9.8	51.1 ± 10.3	50.4 ± 11.1	45.5 ± 10.6	50.1 ± 10.2	51.3 ± 9.8	50.2 ± 10.9	46.2 ± 10.8	50.8 ± 10.0	51.2 ± 10.1	50.3 ± 11.0	45.3 ± 12.6	45.9 ± 12.3
WPAI scores, m	ean ± SD													
Absenteeism ^a	4.5 ± 16.4	5.2 ± 14.5	5.3 ± 14.7	5.2 ± 15.8	3.4 ± 13.9	8.1 ± 21.0	4.3 ± 14.9	7.6 ± 21.6	4.0 ± 15.2	6.7 ± 18.0	4.8 ± 14.8	6.3 ± 18.9	3.9 ± 10.1	6.5 ± 20.2
Presenteeism ^a	19.6 ± 25.6	10.0 ± 17.6	9.4 ± 19.0	9.2 ± 17.5	19.6 ± 22.7	6.9 ± 14.4	5.6 ± 15.0	10.0 ± 19.6	19.6 ± 24.2	8.4 ± 16.2	7.5 ± 17.2	9.6 ± 18.5	18.8 ± 22.2	19.1 ± 23.7
Overall Work	23.1 ±	14.1 ±	13.0 ±	13.3 ±	22.1 ±	13.3 ±	9.2 ± 20.6	16.5 ±	22.6 ±	13.7 ±	11.1 ±	14.9 ±	20.8 ±	25.0 ±
Impairment ^a	29.2	22.3	22.7	22.3	25.5	24.7	9.2 ± 20.0	27.4	27.4	23.5	21.8	25.0	24.4	28.2
Overall Activity Impairment	33.0 ± 30.6	11.9 ± 19.3	10.7 ± 19.1	12.0 ± 20.0	33.6 ± 28.8	9.8 ± 19.9	8.3 ± 18.3	13.4 ± 24.0	33.3 ± 29.7	10.9 ± 19.6	9.5 ± 18.7	12.7 ± 22.0	29.9 ± 25.8	28.0 ± 27.3

All CZP arm includes all patients randomised to certolizumab pegol (CZP) at baseline. Data are presented as last observation carried forward (LOCF) for all patients randomised to CZP at baseline 12345678 according to PsA status; patients who did not achieve a ≥50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aAssessed in employed patients only; patient numbers: CZP 200 mg Q2W N=130; CZP 400 mg Q2W N=127; all CZP N=257; Placebo N=68. Patients with no baseline value were not included in these analyses. CZP: certolizumab pegol; DLQI: Dermatology Life Quality Index; EQ-5D-3L: EuroQol 5-Dimensions 3-Level; EQ VAS: EuroQol Visual Analogue Scale; LOCF: last observation carried forward; PASI: Psoriasis Area and Severity Index; Q2W: every two weeks; SD: standard deviation; SF-36: 36-Item Short Form Survey; WPAI: Work Productivity and Activity Impairment.

		Previous Bio	logic Therapy		No Previous Biologic Therapy					
	Baseline	Week 16	Week 48	Week 144	Baseline	Week 16	Week 48	Week 144		
	N=121	N=114	N=102	N=70	N=235	N=230	N=197	N=153		
DLQI										
DLQI 0/1, n (%)	2 (1.7)	63 (55.3)	65 (63.7)	43 (61.4)	5 (2.1)	108 (47.0)	117 (59.4)	87 (56.9)		
DLQI domain score of 0, n (%)										
Symptoms and Feelings	1 (0.8)	42 (36.8)	52 (51.0)	34 (48.6)	1 (0.4) ^a	65 (28.3)	87 (44.2)	65 (42.5)		
Leisure	23 (19.0)	90 (78.9)	80 (78.4)	58 (82.9)	45 (19.0)ª	158 (68.7)	157 (79.7)	118 (77.1)		
Daily Activities	10 (8.3)	78 (68.4)	76 (74.5)	51 (72.9)	21 (8.9) ^a	139 (60.4)	137 (69.5)	109 (71.2)		
Work and School	53 (43.8)	104 (91.2)	94 (92.2)	67 (95.7)	97 (40.9) ^a	185 (80.4)	176 (89.3)	129 (84.3)		
Personal Relationships	35 (28.9)	90 (78.9)	86 (84.3)	59 (84.3)	76 (32.1) ^a	163 (70.9)	167 (84.8)	123 (80.4)		
Treatment	32 (26.4)	81 (71.1)	80 (78.4)	54 (77.1)	86 (36.3) ^a	171 (74.3)	168 (85.3)	125 (81.7)		
EQ-5D-3L,										
EQ-5D-3L score of 1, n (%)										
Mobility	77 (63.6)	96 (84.2)	84 (82.4)	49 (70.0)	171 (72.8)	194 (84.3)	173 (87.8)	128 (83.7)		
Self-Care	105 (86.8)	109 (95.6)	97 (95.1)	66 (94.3)	213 (90.6)	218 (94.8)	188 (95.4)	147 (96.1)		
Usual Activities	77 (63.6)	100 (87.7)	90 (88.2)	56 (80.0)	163 (69.4)	188 (81.7)	175 (88.8)	133 (86.9)		
Pain/Discomfort	16 (13.2)	67 (58.8)	67 (65.7)	38 (54.3)	59 (25.1)	143 (62.2)	131 (66.5)	101 (66.0)		
Anxiety/Depression	68 (56.2)	96 (84.2)	81 (79.4)	53 (75.7)	112 (47.7)	164 (71.3)	140 (71.1)	113 (73.9)		
EQ VAS score, mean ± SD	66.1 ± 23.0	80.5 ± 16.9	83.5 ± 14.6	79.8 ± 16.5	65.9 ± 22.2	78.4 ± 16.4	82.2 ± 13.8	82.6 ± 13.4		
WPAI										
WPAI scores, mean ± SD										
Absenteeism ^b	4.4 ± 16.4	6.8 ± 17.4	3.0 ± 11.7	5.1 ± 18.3	3.8 ± 14.6	6.2 ± 17.4	4.4 ± 13.8	4.1 ± 14.8		
Presenteeism ^b	19.2 ± 24.5	8.1 ± 17.6	5.5 ± 14.0	6.0 ± 11.4	19.8 ± 24.1	7.4 ± 14.8	5.6 ± 14.7	8.8 ± 17.8		
Overall Work Impairment ^b	22.5 ± 28.1	13.2 ± 23.6	7.7 ± 18.0	10.8 ± 20.8	22.7 ± 27.1	12.5 ± 22.3	9.2 ± 19.1	12.2 ± 22.2		
Overall Activity Impairment	33.3 ± 30.9	9.8 ± 19.4	5.9 ± 13.7	8.6 ± 16.4	33.2 ± 29.1	10.3 ± 19.1	7.1 ± 14.8	9.0 ± 18.6		

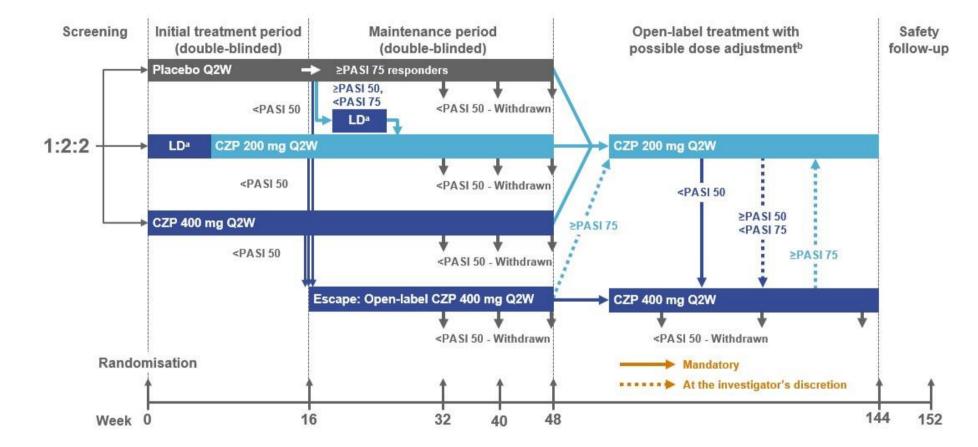
Table S2. Patient-reported outcomes by previous biologic therapy use to Week 144 (OC) 1

Data are presented as observed for all patients randomised to certolizumab pegol (CZP) at baseline according to previous biologic use; patients who did not achieve a \geq 50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aN=237; ^bAssessed in employed patients only; patient

numbers, previous biologic therapy: Baseline N=84; Week 16 N=79; Week 48 N=76; Week 144 N=45; no previous biologic therapy: Baseline N=173; Week 16 N=172; Week 48 N=147; Week 144 N=120. DLQI: Dermatology Life Quality Index; EQ-5D-3L: EuroQol 5-Dimensions 3-Level; EQ VAS: EuroQol Visual Analogue Scale; OC: observed case; Q2W: every two weeks;

3456 WPAI: Work Productivity and Activity Impairment.

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1 **Figure S1.** The initial, maintenance and open-label periods of the CIMPASI-1 and CIMPASI-2 phase 3 trials

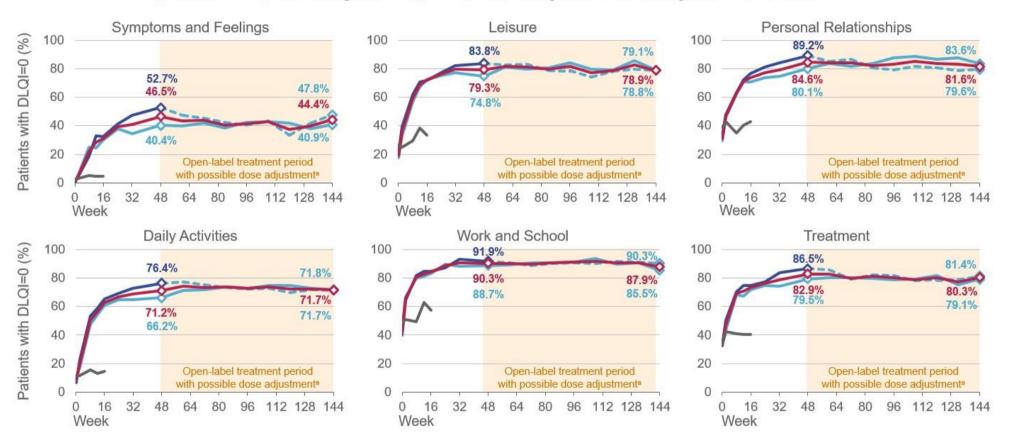
^aDose adjustments were permitted through Weeks 60–132; dose escalation was mandatory in patients not achieving ≥50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50), and at the investigator's discretion in patients achieving PASI 50 but not PASI 75; patients who had received certolizumab pegol (CZP) 400 mg every two weeks (Q2W) for at least 12 weeks could have had their dose reduced, at the investigator's discretion, if they achieved PASI 75, and were mandatorily withdrawn in they did not achieve PASI 50; ^bPatients entering the open-label period from the CZP 400 mg Q2W escape arm continued to receive CZP 400 mg Q2W but may have had their dose reduced to CZP 200 mg Q2W at Week 48, at the discretion of the investigator, if they achieved PASI 75, CZP; certolizumab pegol: LD; loading dose; PASI: Psoriasis Area and Severity Index; O2W; every two weeks.



→ All CZP → CZP 200 mg Q2W

• .◇• CZP 400 mg Q2W → CZP 200 mg Q2W -

- Placebo



Data are presented as observed for all patients according to initial randomisation group; CZP-randomised patients who did not achieve a \geq 50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aDose adjustments were mandatory or at the investigator's discretion, based on PASI response. Number of patients with assessments at each visit are identical to those presented in **Figure 1**, except for Week 0. Week 0: CZP 200 mg Q2W, n=184; CZP 400 mg Q2W, n=174; all CZP, n=358; placebo, n=98. CZP: certolizumab pegol; DLQI: Dermatology Life Quality Index; PASI: Psoriasis Area and Severity Index; Q2W: every two weeks.

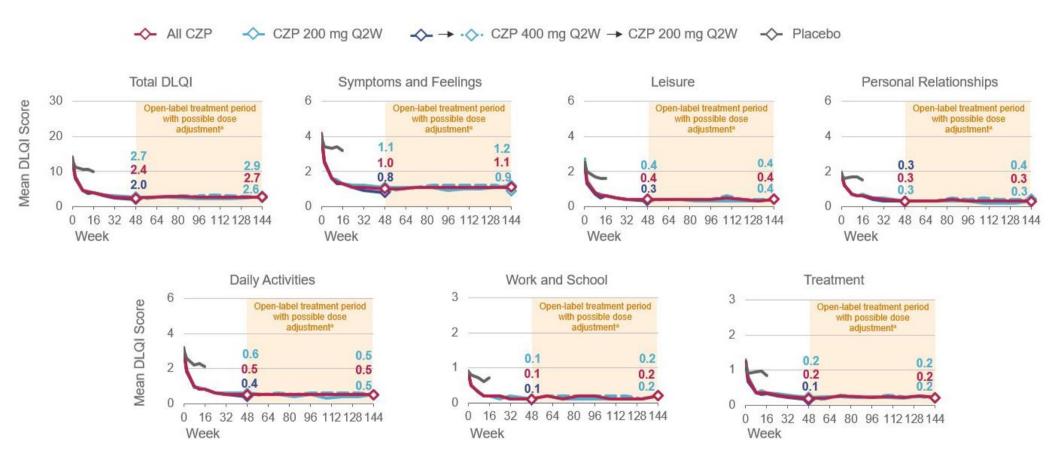
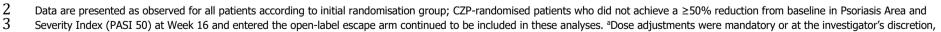


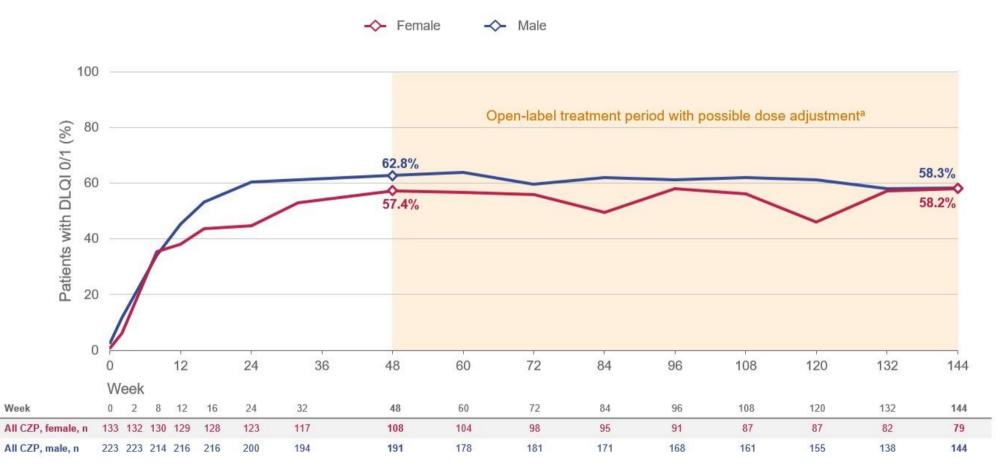
Figure S2b. Absolute total and domain DLQI scores to Week 144 (OC) 1



Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aDose adjustments were mandatory or at the investigator's discretion, based on PASI response. Number of patients with assessments at each visit are identical to those presented in Figure 2 at all timepoints. CZP: certolizumab pegol; DLQI: Dermatology Life Quality

4 5 Index; PASI: Psoriasis Area and Severity Index; Q2W: every two weeks.

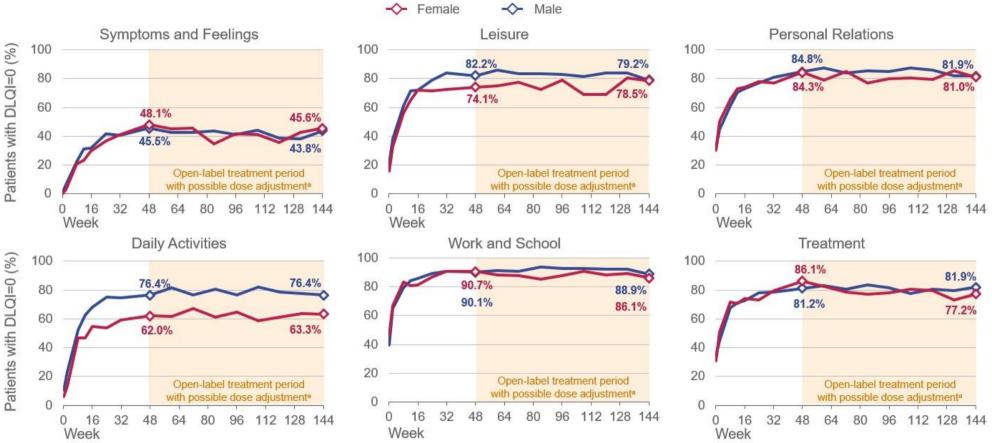




2 3 Data are presented as observed for all patients randomised to CZP at baseline according to sex; patients who did not achieve a ≥50% reduction from baseline in Psoriasis Area and Severity

Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aDose adjustments were mandatory or at the investigator's discretion, based on

4 PASI response. CZP: certolizumab pegol; PASI: Psoriasis Area and Severity Index; Q2W: every two weeks.



1 **Figure S3b.** DLQI domain score=0, CZP-randomised patients by sex to Week 144 (OC)



Data are presented as observed for all patients randomised to CZP at baseline according to sex; patients who did not achieve a \geq 50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aDose adjustments were mandatory or at the investigator's discretion, based on

PASI response. Number of patients with assessments at each visit are identical to those presented in **Figure S3A**, except for Males, Week 0 (n=225). CZP: certolizumab pegol; DLQI: Dermatology

Life Quality Index; PASI: Psoriasis Area and Severity Index; Q2W: every two weeks.



1 Figure S3c. EQ-5D-3L score=1 and mean EQ VAS, CZP-randomised patients by sex to Week 144 (OC)

Data are presented as observed for all patients randomised to CZP at baseline according to sex; patients who did not achieve a \geq 50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aDose adjustments were mandatory or at the investigator's discretion, based on PASI response. CZP: certolizumab pegol; EQ-5D-3L: EuroQol 5-Dimensions 3-Level; EQ VAS: EuroQol Visual Analogue Scale; PASI: Psoriasis Area and Severity Index; PsA: psoriatic arthritis; Q2W: every two weeks.

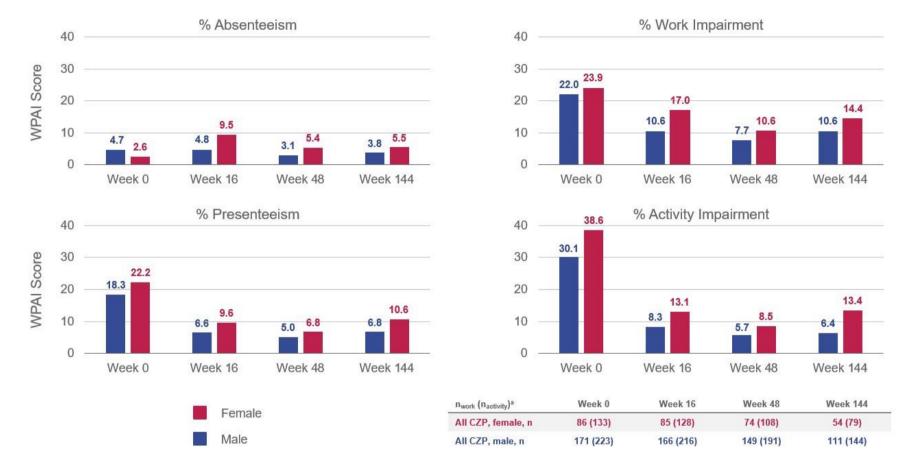
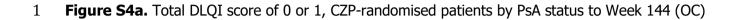


Figure S3d. Mean WPAI, CZP-randomised patients by sex (OC) 1

Data are presented as observed for all patients randomised to CZP at baseline according to sex; patients who did not achieve a \geq 50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. From Week 48, dose adjustments were mandatory or at the investigator's discretion,

3 4 based on PASI response. a The 'nwork' category includes absenteeism, presenteeism and overall work impairment; 'nativity' includes overall activity impairment. CZP: certolizumab pegol; PASI: Psoriasis

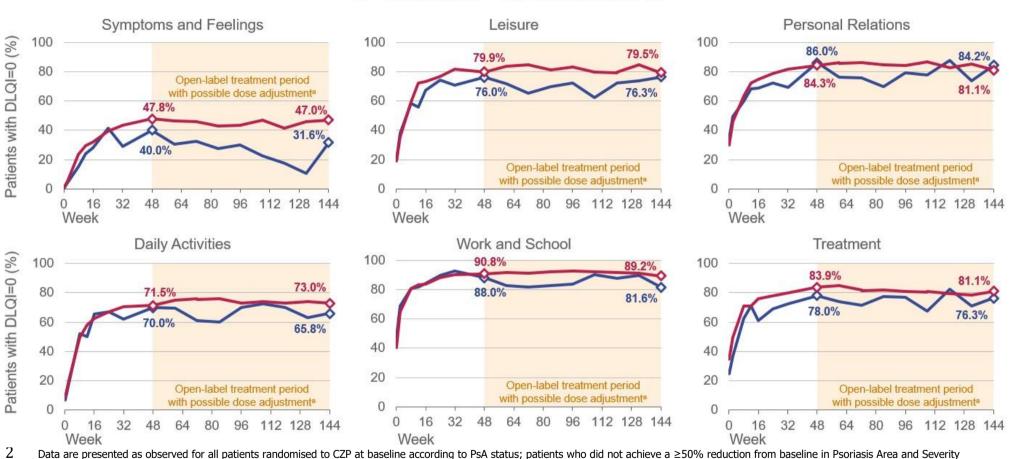
5 Area and Severity Index; Q2W: every two weeks.





Data are presented as observed for all patients randomised to CZP at baseline according to PsA status; patients who did not achieve a ≥50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. aDose adjustments were mandatory or at the investigator's discretion, based on

2 3 4 PASI response. CZP: certolizumab pegol; PASI: Psoriasis Area and Severity Index; PsA: psoriatic arthritis; Q2W: every two weeks.



No Concomitant PsA

Figure S4b. DLQI domain score=0, CZP-randomised patients by PsA status to Week 144 (OC)

Concomitant PsA

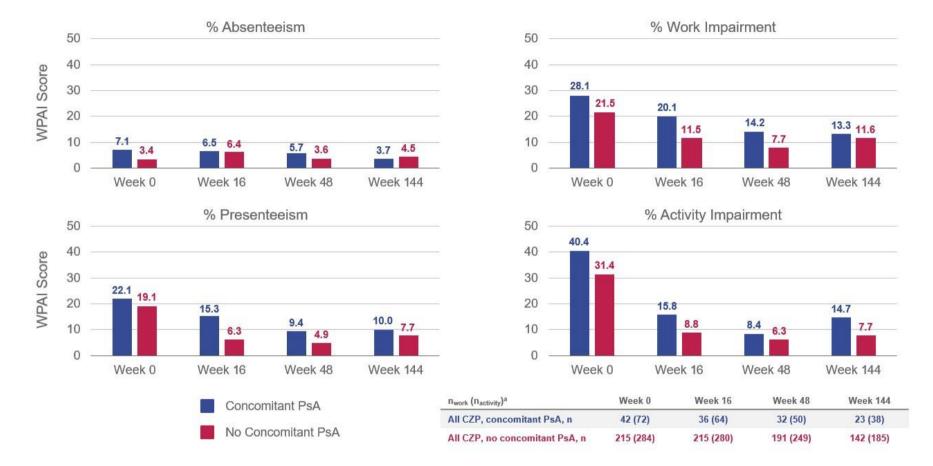
Data are presented as observed for all patients randomised to CZP at baseline according to PsA status; patients who did not achieve a \geq 50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aDose adjustments were mandatory or at the investigator's discretion, based on PASI response. CZP: certolizumab pegol; DLQI: Dermatology Life Quality Index; PASI: Psoriasis Area and Severity Index; PsA: psoriatic arthritis; Q2W: every two weeks. Number of patients with assessments at each visit are identical to those presented in **Figure S4A**, except for those with no concomitant PsA, Week 0 (n=286).



1 Figure S4c. EQ-5D-3L score=1 and mean EQ VAS, CZP-randomised patients by PsA status to Week 144 (OC)

Data are presented as observed for all patients randomised to CZP at baseline according to PsA status; patients who did not achieve a \geq 50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. ^aDose adjustments were mandatory or at the investigator's discretion, based on PASI response. CZP: certolizumab pegol; EQ-5D-3L: EuroQol 5-Dimensions 3-Level; EQ VAS: EuroQol Visual Analogue Scale; PASI: Psoriasis Area and Severity Index; PsA: psoriatic arthritis;

O2W: every two weeks.



1 Figure S4d. Mean WPAI, CZP-randomised patients by PsA status (OC)

Data are presented as observed for all patients randomised to CZP at baseline according to PsA status; patients who did not achieve a \geq 50% reduction from baseline in Psoriasis Area and Severity Index (PASI 50) at Week 16 and entered the open-label escape arm continued to be included in these analyses. From Week 48, dose adjustments were mandatory or at the investigator's discretion, based on PASI response. ^aThe 'n_{work}' category includes absenteeism, presenteeism and overall work impairment; 'n_{activity}' includes overall activity impairment. CZP: certolizumab pegol; PASI: Psoriasis Area and Severity Index; PsA: psoriatic arthritis; Q2W: every two weeks.