Table S7
 Summary of findings table for outcome indicators

Patient or population: patients with plaque psoriasis and psoriatic arthritis

Intervention: To facitinib **Comparison:** Placebo

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect	No of Participants	Quality of the evidence
	Assumed risk Placebo	Corresponding risk Tofacitinib	(95% CI)	(studies)	(GRADE)
	Study population		RR 5.21	6401	$\oplus \oplus \oplus \ominus$
PASI 75	88 per 1000	460 per 1000 (361 to 587)	(4.09 to 6.65)	(7 studies)	moderate
	Moderate				
	83 per 1000	432 per 1000			
	Study population	(339 to 552)	RR 12.63	3539	$\oplus \oplus \oplus \oplus$
PASI 90	25 per 1000	321 per 1000	(8.81 to 18.11)	(5 studies)	high
		(224 to 460)			
	Moderate				
	9 per 1000	114 per 1000 (79 to 163)			
PGA 0/1	Study population		RR 4.7	5752	$\oplus \oplus \oplus \oplus$
	111 per 1000	521 per 1000	(4.12 to 5.36)	(5 studies)	high
	Moderate	(457 to 595)	-		
	100 per 1000	470 per 1000	1		
	Study population	(412 to 536)	RR 1.07	4504	$\Delta \Delta \Delta \Delta$
Nasophary ngitis	63 per 1000	68 per 1000 (54 to 85)	(0.86 to 1.34)	(7 studies)	⊕⊕⊕⊝ moderate
	Moderate	(311003)			
	56 per 1000	60 per 1000 (48 to 75)			
URTI	Study population		RR 1.64	4504	$\oplus \oplus \oplus \oplus$
	33 per 1000	54 per 1000 (40 to 75)	(1.2 to 2.25)	(7 studies)	high
	Moderate				
	34 per 1000	56 per 1000 (41 to 77)			
	Study population		RR 2.75	3459	$\oplus \oplus \oplus \oplus$
Hyperchol esterolemi a	16 per 1000	44 per 1000 (26 to 72)	(1.67 to 4.53)	(4 studies)	high
	Moderate	(20 to 12)			
	20 per 1000	55 per 1000 (33 to 91)			
	Study population	·	RR 2.07	3526	$\oplus \oplus \oplus \oplus$
CPK elevation	17 per 1000	35 per 1000 (22 to 57)	(1.28 to 3.34)	(4 studies)	high
	Moderate	(22 to 31)			

	16 per 1000	33 per 1000 (20 to 53)			
Headache	Study population		RR 1.69	3277	$\oplus \oplus \oplus \oplus$
	35 per 1000	59 per 1000 (42 to 83)	(1.2 to 2.4)	(5 studies)	high
	Moderate				
	31 per 1000	52 per 1000 (37 to 74)			

^{*}The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; RR: Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.