Quality assessment							Number of patients		Effect			
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Methylphenidate	placebo	Relative (95% CI)	Absolute (95% CI)	Quality	Importance
ADHD (follow up:	mean 16 weeks; as	ssessed with: (	Connors' ADHD ir	ndex [parent rate	ed])							•
1	randomised trials	not serious	not serious	not serious	serious 1	none	61	61		MD <b>3.3 fewer</b> (6.79 fewer to 0.19 more)	⊕⊕⊕○ MODERATE	CRITICAL
ADHD (follow up:	mean 16 weeks; as	ssessed with: (	Connors' ADHD ir	idex [teacher rat	ted])							•
1	randomised trials	not serious	not serious	not serious	serious <sup>1</sup>	none	61	61	-	MD <b>4.1 fewer</b> (7.57 fewer to 0.63 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
Hyperactivity (follo	w up: mean 16 we	eks; assessed	with: Conners' hy	peractivity scale	e [parent rated])							-1
1	randomised trials	not serious	not serious	not serious	serious <sup>2</sup>	none	61	61	-	MD <b>1.5 fewer</b> (3.44 fewer to 0.44 more)	⊕⊕⊕○ MODERATE	CRITICAL
Hyperactivity (follo	w up: mean 16 we	eks; assessed	with: Conners' hy	peractivity scale	e [teacher rated	1)						-
1	randomised trials	not serious	not serious	not serious	serious 1	none	61	61	-	MD <b>2.6 fewer</b> (4.68 fewer to 0.52 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
'Improved' or 'bette	ı er' (follow up: mear	n 16 weeks; as	ssessed with: Clini	cal Global Impre	essions-Improve	ement)	l					
1	randomised trials	not serious	not serious	not serious	serious <sup>1</sup>	none	24/61 (39.3%)	4/61 (6.6%)	RR 6.00 (2.21 to 16.26)	328 more per 1000 (from 79 more to 1000 more)	⊕⊕⊕○ MODERATE	CRITICAL
Quality of life – no	t reported	ı			1		1					,

Quality assessment								Number of patients		Effect		
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Methylphenidate	placebo	Relative (95% CI)	Absolute (95% CI)	Quality	Importance
-	-	-	-	-	-	-					-	CRITICAL
Community participation and meaningful occupation – not reported												
-	1	1	-	1	1	-					-	CRITICAL
Weight (follow up: mean 16 weeks; assessed with: kg)												
1	randomised trials	not serious	not serious	not serious	serious <sup>2</sup>	none	61	61	-	MD <b>4.2 kg fewer</b> (10.25 fewer to 1.85 more)	⊕⊕⊕⊖ MODERATE	IMPORTANT

Sample size less than optimal information size (<400 for continuous outcomes or <300 for dichotomous outcomes).

Confidence intervals cross one minimally important difference. Sample size less than optimal information size (<400 for continuous outcomes or <300 for dichotomous outcomes).