Quality assessment							Number of patients		Effect			
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Clonidine	placebo	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
ADHD symptoms:	conduct (follow up: 6	6 weeks; assess	ed with: Parent C	onnor's score –	conduct scale)							
1	randomised trials	very serious	not serious	not serious	serious <sup>2</sup>	none	9	10	-	MD <b>7.4 fewer</b> (10.34 fewer to 4.46 fewer)		
ADHD symptoms:	mpulsive hyperactiv	vity (follow up: 6	weeks; assessed	with: Parent Co	nnor's score – l	mpulsive hyperactive sca	le)					
1	randomised trials	very serious	not serious	not serious	serious <sup>3</sup>	none	9	10	-	MD <b>2.6 fewer</b> (6.54 fewer to 1.34 more)		

Quality assessment							Number of patients		Effect			
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Clonidine	placebo	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	randomised trials	very serious	not serious	not serious	serious <sup>3</sup>	none	9	10		MD <b>24.7 fewer</b> (49.35 fewer to 0.05 fewer)		
ADHD symptoms (clinician rated) (follow up: 6 weeks; assessed with: CGI)												
1	randomised trials	very serious	not serious	not serious	serious <sup>3</sup>	none	9	10	-	MD <b>1.8 fewer</b> (3.11 fewer to 0.49 fewer)		
Much or very much improved (follow up: 6 weeks; assessed with: CGI)												
1	randomised trials	very serious	not serious	not serious	serious <sup>2</sup>	none	7/9 (77.8%)	0/10 (0.0%)	RR 16.50 (1.07 to 253.40)	0 fewer per 1000 <sup>4</sup> (from 0 fewer to 0 fewer)		
Quality of life – not reported												
-	-	-	-			-					-	
Community participa	Community participation and meaningful occupation – not reported											
-	-	-				-					-	

1.

2.

Risk of selection and selective outcome reporting bias 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). 3.

Absolute risk value is 0 as no events of interest occurred for this outcome 4.

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