Quality assessment								Number of patients		Effect		
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Acetyl-L- carnitine	placebo	Relative (95% CI)	Absolute (95% Cl)	Quality	Importance
ADHD (follow up: m	ean 52 weeks; asse	essed with: Conr	ers' Parents)									·
1	randomised trials	very serious	not serious	not serious	serious <sup>2</sup>	none	24	27	-	MD <b>2.8 fewer</b> (7.58 fewer to 1.98 more)		CRITICAL
ADHD (follow up: m	ean 52 weeks; asse	essed with: Conr	ers' Teachers)									
1	randomised trials	very serious	not serious	not serious	serious <sup>2</sup>	none	24	27	-	MD <b>0.5 more</b> (5.08 fewer to 6.08 more)		CRITICAL
Quality of life – not i	reported			•								
-	-	-	-								_	CRITICAL
Community particip	ation and meaningfu	Il occupation – n	ot reported									
-	-	-	-		-						-	CRITICAL
Adaptive functioning	g (post-treatment) (fo	ollow up: mean 5	52 weeks; assesse	ed with: VABS –	full scale)							
1	randomised trials	very serious	not serious	not serious	serious <sup>2</sup>	none	24	27	-	MD <b>8.2 more</b> (0.04 fewer to 16.44 more)		IMPORTAN
Adaptive functioning	g (follow up: mean 5	2 weeks; assess	sed with: VABS -	socialization sca	lle)							•

Quality assessment							Number of patients			Effect		
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Acetyl-L- carnitine	placebo	Relative (95% CI)	Absolute (95% Cl)	Quality	Importance
1	randomised trials	very serious	not serious	not serious	serious <sup>2</sup>	none	24	27		MD <b>11.3 more</b> (2.18 more to 20.42 more)		IMPORTANT

1. Risk of selection and detection bias

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

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