Quality assessment								Number of patients		Effect		
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Antioxidant	placebo	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Mental health (all levels of learning disabilities) (follow up: mean 104 weeks; assessed with: DMR [sum of cognitive scores])												
1	randomised trials	serious 1	not serious	not serious	serious <sup>2</sup>	none	No significant differences in DMR cognitive scores scores between antioxidant and placebo groups					CRITICAL
Mental health (all levels of learning disabilities) (follow up: mean 104 weeks; assessed with: Severe impairment battery)												
1	randomised trials	serious 1	not serious	not serious	serious <sup>2</sup>	none	No significant differences in Severe Impairment Battery scores between antioxidant and placebo groups					CRITICAL

Quality assessment							Number of patients		Effect			
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Antioxidant	placebo	Relative (95% CI)	Absolute (95% Cl)	Quality	Importance
Quality of life	- not reported											
-	-	-	-	-	-	-					-	CRITICAL
Community participation and meaningful occupation – not reported												
-	-	-	-	-	-	-					-	CRITICAL
Adaptive functioning (all levels of learning disabilities) (follow up: mean 104 weeks; assessed with: Brief Praxis Test)												
1	randomised trials	serious 1	not serious	not serious	serious <sup>2</sup>	none	No significant dif	ferences in Brie	f Praxis Test se	cores between antioxidant and placebo		IMPORTANT
Adaptive func	tioning (all levels	s of learning	disabilities) (follo	w up: mean 104	weeks; assess	ed with: DMR [sum o	f social skills])					
1	randomised trials	serious 1	not serious	not serious	serious <sup>2</sup>	none	No significant differences in DMR sum of social scores scores between antioxidant and placebo groups					IMPORTANT
Adaptive functioning (all levels of learning disabilities) (follow up: mean 104 weeks; assessed with: Bristol Activities of Daily Living Scale )												
1	randomised trials	serious <sup>1</sup>	not serious	not serious	serious <sup>2</sup>	none	No significant difi antioxidant and p			IMPORTANT		
Any serious a	Any serious adverse event (incapacitation and/or inability to sustain daily activities) (all levels of learning disabilities) (follow up: mean 104 weeks; assessed with: [ITT/analysed as randomised])											
1	randomised trials	serious <sup>1</sup>	not serious	not serious	serious <sup>2</sup>	none	14/29 (48.3%)	11/29 (37.9%)	RR 1.27 (0.70 to 2.32)	<b>102 more per 1000</b> (from 114 fewer to 501 more)		IMPORTANT

- 1. Risk of selective outcomes bias.
- Sample size less than optimal information size (<400 for continuous outcomes or <300 for dichotomous outcomes).</li>

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