

TABLE 1

Question: Should antibiotics (amoxicillin/cefdinir) be used in children with uncomplicated severe acute malnutrition?

Settings: Community

Number of studies	Quality assessment						Number (%) of patients		Effect		Quality	Importance
	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Antibiotics (amoxicillin/cefdinir)	Control	Relative (95% CI)	Absolute		
Mortality up to 6 weeks (follow-up median 6 weeks)												
1	Randomized trials	Serious ^a	No serious inconsistency	No serious indirectness	No serious imprecision	None	82/1847 (4.4)	68/920 (7.4%)	RR 1.46 (1.19 to 1.80)	34 more per 1000 (from 14 more to 59 more)	+++ MODERATE	CRITICAL
Time to recovery												
1	Randomized trials	Serious ^a	No serious inconsistency ²	No serious indirectness	No serious imprecision	None	-	-	MD -0.50 (-2.11 to 1.11)	-	+++ MODERATE	CRITICAL
Weight gain (g/kg/day)												
1	Randomized trials	Serious ^a	No serious inconsistency ^b	No serious indirectness	No serious imprecision	None	-	-	MD 0.55 (0.18 to 0.92)	-	+++ MODERATE	CRITICAL
Mortality up to 12 weeks (follow-up median 12 weeks)												
1	Observational studies	Very serious ^c	No serious inconsistency ^b	No serious indirectness	Serious ^d	None	13/498 (2.6)	34/1955 (1.7)	RR 1.50 (0.80 to 2.82)	-	+ VERY LOW	CRITICAL
Recovery at 12 weeks (follow-up median 12 weeks)												
1	Observational studies	Very serious ^{c,e}	No serious inconsistency ^b	No serious indirectness	No serious imprecision	None	417/498 (83.7)	1673/1955 (85.6)	RR 0.98 (0.94 to 1.02)	-	+ VERY LOW	CRITICAL

CI: confidence interval; RR: relative risk; MD: mean difference.

^a Double blind study conducted in a setting of high HIV prevalence and most children had kwashiorkor. The response to antibiotics could be modified by these two factors.

^b Only one study.

^c Comparison of two different cohorts from different parts of Malawi. There were also significant differences in baseline characteristics between the cohorts.

^d Few events and wide confidence intervals.

^e Participants and researchers not blinded to the interventions.