## Comparison: Upright position compared with recumbent position in the second stage of labour for women with epidural analgesia

Source: Kibuka M, Thornton JG. Position in the second stage of labour for women with epidural anaesthesia. Cochrane Database Syst Rev. 2017;(2):CD008070.

Quality assessment							No. of participants		Effect			
No. of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Upright position	Recumbent position	Relative (95% CI)	Absolute (95% CI)	Certainty (GRADE)	Importance
Duration	of second	stage labour (	minutes) (from	time of randor	nization to bir	th)						
2	RCTs	serious <sup>a</sup>	serious <sup>b</sup>	not serious	serious <sup>c</sup>	none	199	123	-	MD 22.98 lower (99.09 lower to 53.13 higher)	⊕○○○ VERY LOW	critical
Spontane	ous vagina	al birth										
6	RCTs	serious <sup>d</sup>	serious <sup>b</sup>	not serious	not serious <sup>e</sup>	none	811/2018 (40.2%)	849/1949 (43.6%)	RR 0.97 (0.82-1.14)	13 fewer per 1000 (from 61 more to 78 fewer)	⊕⊕⊖⊖ LOW	critical
Operativ	e birth (cae	sarean or ins	trumental vagin	al)								
6	RCTs	serious <sup>d</sup>	not serious	not serious	not serious <sup>e</sup>	none	1206/2018 (59.8%)	1096/1949 (56.2%)	RR 1.04 (0.89-1.20)	22 more per 1000 (from 62 fewer to 112 more)	⊕⊕⊕⊖ MODERATE	critical
Instrume	ntal vagina	al birth										
6	RCTs	serious <sup>a</sup>	not serious	not serious	not serious <sup>e</sup>	none	993/2018 (49.2%)	923/1949 (47.4%)	RR 1.05 (0.94-1.18)	24 more per 1000 (from 28 fewer to 85 more)	⊕⊕⊕○ MODERATE	critical
Caesarea	n section											
6	RCTs	serious <sup>d</sup>	not serious	not serious	serious <sup>c</sup>	none	213/2018 (10.6%)	173/1949 (8.9%)	RR 1.05 (0.71-1.55)	4 more per 1000 (from 26 fewer to 49 more)	⊕⊕⊖⊖ LOW	critical
Trauma t	o birth can	al requiring su	uturing					'				
3	RCTs	serious <sup>a</sup>	not serious	not serious	not serious <sup>e</sup>	none	1350/1639 (82.4%)	1320/1627 (81.1%)	RR 1.01 (0.89-1.14)	8 more per 1000 (from 89 fewer to 114 more)	⊕⊕⊕⊖ MODERATE	critical
Blood los	s requiring	transfusion						<u>'</u>			<u> </u>	
1	RCT	serious <sup>f</sup>	not serious	not serious	serious <sup>c</sup>	none	63/1556 (4.0%)	52/1537 (3.4%)	RR 1.20 (0.83-1.72)	7 more per 1000 (from 6 fewer to 24 more)	⊕⊕⊖⊖ LOW	critical
Maternal	satisfactio	on with childb	irth experience									
1	RCT	serious <sup>f</sup>	not serious	not serious	not serious <sup>e</sup>	none	963/1556 (61.9%)	973/1537 (63.3%)	RR 0.98 (0.93-1.03)	13 fewer per 1000 (from 19 more to 44 fewer)	⊕⊕⊕○ MODERATE	critical

	Quality assessment						No. of participants		Effect		Containts	
No. of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Upright position	Recumbent position	Relative (95% CI)	Absolute (95% CI)	Certainty (GRADE)	Importance
Apgar score < 7 at 5 minutes												
2	RCTs	serious <sup>a</sup>	not serious	not serious	serious⁵	none	2/1614 (0.1%)	3/1586 (0.2%)	RR 0.66 (0.11-3.94)	1 fewer per 1000 (from 2 fewer to 6 more)	⊕⊕⊖⊝ LOW	critical
Abnormal fetal heart rate patterns requiring intervention												
1	RCT	serious <sup>f</sup>	not serious	not serious	very serious <sup>g</sup>	none	4/58 (6.9%)	2/49 (4.1%)	RR 1.69 (0.32-8.84)	28 more per 1000 (from 28 fewer to 320 more)	⊕○○○ VERY LOW	critical
Low cord	Low cord pH											
2	RCTs	serious <sup>a</sup>	not serious	not serious	not serious	none	9/1581 (0.6%)	25/1578 (1.6%)	RR 0.43 (0.20-0.90)	9 fewer per 1000 (from 2 fewer to 13 fewer)	⊕⊕⊕○ MODERATE	critical
Infant res	Infant resuscitation											
1	RCT	serious <sup>f</sup>	not serious	not serious	serious <sup>c</sup>	none	92/1556 (5.9%)	91/1537 (5.9%)	RR 1.00 (0.75-1.32)	0 fewer per 1000 (from 15 fewer to 19 more)	⊕⊕○○ LOW	critical
Perinatal death												
1	RCT	serious <sup>f</sup>	not serious	not serious	serious⁵	none	1/1556 (0.1%)	0/1537 (0.0%)	RR 2.96 (0.12-72.69)	0 fewer per 1000 (from 0 fewer to 0 fewer)	⊕⊕○○ LOW	critical

CI: confidence interval; MD: mean difference; RCT: randomized controlled trial; RR: risk ratio.

- a Most of the pooled effect derived from studies with a moderate or high risk of bias but without a substantial proportion (i.e. with < 50%) from studies with a high risk of bias.
- <sup>b</sup> CI is imprecise.
- 6 Most of the pooled effect derived from studies with a moderate or high risk of bias but with a substantial proportion (i.e. > 50%) from studies with a high risk of bias.
- d Small sample size and/or few events.
- <sup>e</sup> Confidence interval crossing the line of no effect but precise (not downgraded).
- f Single study with design limitations.
- <sup>g</sup> Wide confidence interval crossing the line of no effect, and a small sample size.