Supplementary Table S1 Sensitivity analysis randomly assigning participants with missing data on mode of delivery ($n = 19$) to 1 spontaneous or induced vaginal delivery versus cesarean or 2 spontaneous vaginal versus induced vaginal or cesarean, with proportion assigned to each group ranging from 0 to 1.0 in intervals of 0.2

Proportion assigned to cesarean delivery		Cesarean versus vaginal delivery
LDA	Placebo	RR (95% CI)
0	0	1.06 (0.81–1.39)
0	0.2	1.03 (0.79–1.34)
0	0.4	1.00 (0.77–1.30)
0	0.6	0.97 (0.75–1.26)
0	0.8	0.95 (0.73–1.23)
0	1	0.92 (0.71–1.19)
0.2	0	1.08 (0.83–1.41)
0.2	0.2	1.05 (0.80–1.37)
0.2	0.4	1.02 (0.78–1.33)
0.2	0.6	0.99 (0.76–1.29)
0.2	0.8	0.96 (0.74–1.25)
0.2	1	0.94 (0.73–1.21)
0.4	0	1.10 (0.84–1.44)
0.4	0.2	1.07 (0.82–1.39)
0.4	0.4	1.04 (0.80–1.35)
0.4	0.6	1.01 (0.78–1.31)
0.4	0.8	0.98 (0.76–1.27)
0.4	1	0.96 (0.74–1.23)
0.6	0	1.12 (0.86–1.46)
0.6	0.2	1.09 (0.84–1.42)
0.6	0.4	1.06 (0.82–1.37)
0.6	0.6	1.03 (0.79–1.33)
0.6	0.8	1.00 (0.78–1.29)
0.6	1	0.97 (0.76–1.25)
0.8	0	1.14 (0.88–1.49)
0.8	0.2	1.11 (0.85–1.44)
0.8	0.4	1.08 (0.83–1.39)
0.8	0.6	1.05 (0.81–1.35)
0.8	0.8	1.02 (0.79–1.31)
0.8	1	0.99 (0.77–1.27)
1	0	1.16 (0.89–1.51)
1	0.2	1.13 (0.87–1.46)
1	0.4	1.10 (0.85–1.42)
1	0.6	1.07 (0.83–1.37)
1	0.8	1.04 (0.81–1.33)
1	1	1.01 (0.79–1.30)

Abbreviations: CI, confidence interval; LDA, low-dose aspirin; RR, relative risk.