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Title	Trial of feasibility and acceptability of routine low-dose aspirin versus early screening test indicated aspirin for pre-eclampsia prevention (TEST study): a multicentre randomised controlled trial
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Outcome	Screen positive; Aspirin Group 3A N=13	Screen negative; No Aspirin Group 3B N=171	OR (95% CI)
Preeclampsia No. (%) Pre-eclampsia <34-weeks Pre-eclampsia <37-weeks	2 (15.4) 0 (0) 2 (15.4)	6 (3.5) 2 (1.2) 2 (1.2)	5.0 (0.9 – 27.7) - 15.4 (2.0 – 120)
Birthweight <10th centile No. (%)	4 (30.7)	21 (12.3)	3.2 (0.9 – 11.2)
Pre-term delivery <34 weeks No. (%)	1 (7.7)	1 (0.6)	14.1 (0.8 – 240)
NICU admission No. (%)	0 (0)	9 (5.3)	-
Outcome No. (%) Alive at 6-weeks Stillbirth Neonatal death	 13 (100) 0 (0) 0 (0)	 169 (98.8) 2 (1.2) 0 (0)	

Table S2 - Secondary outcome measures in Group 3 (screen and treat)

(Expressed as average and standard deviation unless otherwise stated)

Note: ORs are not presented when number of events is 0 in the Screen –positive group.