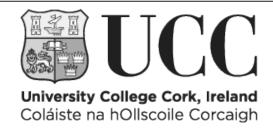


Title	Trial of feasibility and acceptability of routine low-dose aspirin versus early screening test indicated aspirin for pre-eclampsia prevention (TEST study): a multicentre randomised controlled trial		
Authors	Mone, Fionnuala; Mulcahy, Cecilia; McParland, Peter; Breathnach, Fionnuala M.; Downey, Paul; McCormack, Dorothy; Culliton, Marie; Stanton, Alice; Cody, Fiona; Morrison, John J.; Daly, Sean; Higgins, John R.; Cotter, Amanda; Hunter, Alyson; Tully, Elizabeth C.; Dicker, Patrick; Alfirevic, Zarko; Malone, Fergal D.; McAuliffe, Fionnuala M.		
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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. (%)	2 (15.4)	6 (3.5)	5.0 (0.9 – 27.7)
Pre-eclampsia <34-	0(0)	2 (1.2)	-
weeks			
Pre-eclampsia <37-	2 (15.4)	2 (1.2)	15.4 (2.0 – 120)
weeks			
Birthweight <10 <sup>th</sup> centile	4 (30.7)	21 (12.3)	3.2 (0.9 – 11.2)
No. (%)			
Pre-term delivery <34	1 (7.7)	1 (0.6)	14.1 (0.8 – 240)
weeks No. (%)			
NICU admission No.	0 (0)	9 (5.3)	-
(%)			
Outcome No. (%)			
Alive at 6-weeks	13 (100)	169 (98.8)	
Stillbirth	0 (0)	2 (1.2)	
Neonatal death	0 (0)	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.