

Critical appraisal – Randomised controlled trial questions

Tison et al. 2017

Introduction	
Are the aims clearly stated?	Yes
Methods	
Is the study design suitable for the aims?	Yes
Which population was studied?	Convenience sample of 18 commercial dairy farms in Quebec, Canada. 1247 animals started the trial
Were the treatments randomly allocated? If yes, how was the randomisation done?	Yes Coin toss
Were the groups comparable prior to intervention?	Primiparous numbers and DIM not different between groups.
Was the person who administered the interventions blinded?	Not stated but unlikely due to treatments.
Is it clear what measurements were carried out in the study?	Yes
Were the correct measurements chosen? Do they reflect (or are they strongly related to) the outcome of interest?	Yes Yes
Were previously established validated methods used to make the measurements?	No reference was given for the transrectal palpation or vaginoscopy procedures. The scale

(e.g. Glasgow pain score, International Units etc)	used to describe the purulent discharge has been published previously (0-4 scale, a score of 2 or more was considered positive)
What outcomes were measured?	First service conception rate, time to pregnancy up to 300DIM, presence of discharge in the vagina, any bacterial growth cultured from a cytobrush sample from the vagina
Are the outcomes clinically relevant?	Yes
Were the outcomes assessed blind?	Not stated
Are the statistical methods described?	Yes
Was the statistical significance level stated?	Yes
Was the sample size justified?	Yes – sample size calculation was carried out
Was ethical approval obtained?	States that all procedures were approved by the Institutional Animal Care Committee of the University of Montreal
Are the methods described in enough detail that you could repeat them?	Yes
Results	
Were the basic data adequately described?	It would have been good to have more information about the status of the cows within each of the 18 herds
Do the numbers add up? Are all subjects accounted for?	All removals were explained. Almost 1/6 of animals were removed (n=255), mostly because of no insemination at the end of the synchronisation

	protocol (n=156); but 70 cows were excluded from one single herd because of a failure to respect the synchronisation protocol, which is likely to have an impact on the results from that herd.
Was the statistical significance (p value) stated in the results?	Yes
Is this consistent with the methods? (It should be stated in the sample size or power calculation)	Yes
Were any side effects of the intervention reported if applicable?	Not stated
What were the main findings/key results?	Treatment with cephalosporin was associated with a statistically significant improvement in first service conception rate in cows with purulent vaginal discharge (p<0.05), but this was those animals that had purulent discharge at the first examination and not at the second. There was also a reduced time to conception of cows treated with cephalosporin compared with negative controls.
Discussion and conclusion	
What do the main findings/key results mean?	A significant improvement in reproductive performance when treating cows with purulent vaginal discharge with Cephalosporin.
Are the negative findings discussed?	Yes
How are the negative findings interpreted?	Discussed issues of confounding factors, and spontaneous clinical cure.
Does the discussion reflect the results?	Yes
Interpretation	

What are the clinical implications of this study?	A significant improvement in reproductive performance when treating cows with purulent vaginal discharge with Cephapirin.
Are the subjects in the study similar to those in the BET/your own?	Yes
General	
Who funded this study?	Valacta and the Quebec and Canadian Departments of Agriculture – the authors state that these are public sources of money