This paper is concerned with the evolution of endosymbiont virulence in the *Wolbachia-Drosophila melanogaster* system. The authors used experimental evolution (and artificial selection) to determine how a high temperature affects endosymbiont virulence and consequently host survival. A high temperature was hypothesized to select for decreased endosymbiont virulence (measured as density, as well as octomom copy number). In contrast to the authors’ expectations, they find that density, as well as octomom copy number, increased during the course of the experiment in control and treatment lines. It is argued that this outcome could be due to the artificial manipulation of egg number (which is set a 100 for each generation) and/or directional selection for high density/octomom number variants.

This is an interesting study, because it can shed light on the reasons why high virulent endosymbiotic strains can persist in nature and it can give some more insight into the way populations may be affected when exposed to a high virulent endosymbiotic strain, which is of particular interest from an applied perspective, where virulent strains are used to control the spread of disease through insect vectors.

It is unfortunate that this virulent strain of *Wolbachia* does not actually occur in nature. This reduces the ecological and evolutionary relevance of the study. Overall, the results reported are negative, which is disappointing for the authors considering the amount of work that has been done. It would have been better if the authors could have included treatments where eggs were not artificially selected, and if “outbred” lines would have been included as a control, because I think this would have helped in understanding why you obtained the results that you got. Alternatively, additional experiments could have been performed (backcrossing to the parental generation) to tease apart the different explanations provided for the results obtained.

You mention that ‘Results contribute to the ongoing debate on the relationship between wMelPop octomom copy number, density and virulence’. I think that more details need to be provided in the introduction as to what is actually debated in this regard. Who has found what and when, how do previous findings contradict each other, and more importantly, how can new light be shed to resolve this debate?

What is the likelihood that reduced virulence would/would not evolve in this particular host strain/Wolbachia strain combination? As you rightfully point out in your MS, you see an overall increase in virulence, so there is a response, but you have only used 1 Drosophila strain (mixing different isofemale lines) and 1 Wolbachia strain. Even if your results would have been positive, it would have been very difficult to generalize your findings. Moreover, if more *Drosophila* strains had been tested, perhaps the results would have been different. I think this should be discussed in the manuscript.
Line 23: Explain what octomom copies are (i.e., copies of a *Wolbachia* DNA region containing 8 genes).

Line 28: I think this statement is rather misleading, because you are also proposing other explanations for your results, and you do not provide any evidence (even suggestive) that intra-host selection could play a role in your experiment. I suggest you just state that you discuss your results ‘with respect to the evolutionary causes of wMelPop virulence’.

Line 40-42: This statement requires several references (assuming the theoretical and experimental work were described in separate papers).

Line 60-61: Why was the repetition of the octomom proposed as the genomic basis of wMelPop high density and virulence? Because selected lines with different octomom copy numbers determined *Wolbachia* titers and strength of lethality of the phenotype (Chrostek, Texeira 2015 PLoS Biol). Please explain this in the MS.

Line 62-63: And why were these links called into question? Please explain.

Lines 64-65: If the strain is not known to occur in nature, what is the incentive for doing the experiment from an evolutionary point of view? To better understand the link between virulence and vertical transmission? To see whether extreme environmental conditions can mitigate the negative fitness effects from an applied point of view (as this *Wolbachia* strain is used to reduce virus transmission in vector insects)? This needs to be explained more clearly in the introduction.

Lines 67-68: The authors have altered the environmental conditions by increasing the temperature (experimental evolution), but have also enforced late reproduction (artificial selection). In experimental evolution experiments, only the environment is altered, and there is no further selection by the experimenter of specific individuals for producing the next generation. I would suggest that the authors describe their experiments as such: experimental evolution in conjunction with artificial selection.

Line 109: Why were these generations chosen to perform measurements?

Lines 108-111: Why were eggs used either for assessing survival or *Wolbachia*-related traits? Why not do both measurements at each generation? As you mention 100 eggs are a subset (line 103), this should be feasible by collecting more eggs. Please explain this in the manuscript.

Figure 1: This figure clearly shows when measurements were taken, but aren’t the paraquat lines your ‘manipulated’ control (where you do not expect selection to occur?). The control lines indicated here are actually the lines that should experience selection. This is confusing. Were
there also control lines added that did not experience any treatment, because that would be your actual, unmanipulated, control?

Lines 116-120: The number of lines that went extinct is quite high (n = 8 and 5 for control and paraquat, respectively) considering the number of lines that were started at the beginning of the experiment (n = 20, 10 for control and 10 for paraquat). Would it have been possible to collect more eggs (e.g., after 8 days of age) to at least ensure survival of the lines? Or to have had more replicates within each line (by collecting/distributing multiple egg batches in different vials)?

Line 168: The rationale for doing this experiment needs to be explained in the introduction, because it come out of the blue here.

Line 202: The fact that survival is affected would indeed suggest that there would be some selection pressure, but the question is how strong this selection pressure really is. As you mention later on (line 301), while survival was affected, it still remained high at 8 days. I wonder if selecting 100 eggs at the age of 8 days could have counteracted the effects you would have expected under your experimental evolution regime (high temperature), because it reduced the strength of selection.

Line 208: There is a trend though in the interaction between generation and treatment, that is interesting. Might be worth to highlight this.

Lines 233-237: How can you explain this increase in density and octomom number in control and paraquat lines? You start explaining this in the next section, but it makes more sense to me to discuss this here (or results and discussion should be reported in separate sections, rather than together). The observed differences over time could be due to the fact that genetic similarity increases again during the course of the experiment (isofemale lines were first crossed, but then new isofemale lines were set up for the experiment from the mixed parental generation). This could have been tested if unmanipulated control lines were included (with or without artificial selection to tease apart the effect of temperature on the one hand and late reproduction on the other). Another way to test this would have been to backcross the selected lines to the parental strain. This would also allow you to tease apart whether the observed changes are due to the host nuclear genomic background or the wMelPop genome (as discussed in Rohrscheib et al 2016 PLoS Pathogens).

Line 284: If genetic drift was at play, you would expect a random outcome (increase/stable/decrease virulence) in different replicated experimental evolution lines (both treatment and control), right? You would expect that the interaction between generation and treatment was significant (and there is a trend). If I am right, this would mean that you can
exclude random drift as an explanation for your findings. I would rather think that the specific genetic background of the line affects the course of virulence evolution.

Line 298-301: If you had included unselected lines that went through the experimental evolution experiment, without the selection of eggs at age 8, you would have been able to tease this apart.

**General comments**

David Monnin and colleagues investigated virulence evolution of the symbiotic *Wolbachia* strain wMelPop in *Drosophila* under high temperature (at which the strain is assumed to be virulent) and in the presence or absence of paraquat. Their hypotheses were well-supported by theoretical expectations regarding the evolution of reduced virulence of vertically transmitted symbionts. The introduction is well-written with clear predictions regarding the experiment. These predictions were, however, not confirmed by their experiment and their main hypothesis - the evolution of reduced virulence - was not confirmed. The discussion provides the alternative explanation that intra-host selection might be important here, and I agree that it might possibly play a role here.

I do have some major concerns about how the evolution of virulence was measured. First, from the manuscript it is not clear which survival parameter you compare between G3 and G9. Time until half of the flies died? mean survival time? I think survival analysis would be the best option here. I also wonder if using only fly survival of wMelPop infected flies is a good proxy for virulence? I think that virulence is best measured as the reduction in survival compared to an uninfected control. This way you also correct in some way for measuring survival on different days (is it possible that e.g. differences in food quality between days can be a confounding factor?). Ideally you would even isolate different *Wolbachia* lineages and re-infect flies to test everything at once (but maybe this is technically not possible?). My feeling is that possibly some patterns were not found because of the way virulence was measured.

I also have a general suggestion on providing more information on the population dynamics of wMelPop. I think some information is crucial here to understand how selection might act on these bacteria. Is there a population bottleneck at reproduction/how many bacterial cells are transmitted to the egg?, What is the number of
generations within a host? Especially in the light of intra-host selection this might be very important.

Specific comments

Line 60: do you have more functional information on the genes contained in ‘octomom’?

Line 64: Here it is indicated that this Wolbachia strain is not known to occur in nature. Could you then give a little more explanation where it comes from?

Line 91: I think it is better to mention the experimental temperature in the next section.

Line 103: Why do you use unsupplemented medium for the paraquat lines in the egg-laying phase of the experiment?

Line 104: Is it known if wMelPop affects reproduction of the flies? If so, this might actually be a stronger selective force for reduced virulence than longevity.

Line 168: Maybe best to quickly describe here why you do this (it is only clear when reading the last section of the discussion).

Line 202: In the absence on mortality from non-infected flies it is very difficult to say that this mortality is induced by the Wolbachia.

Line 213: “virulence possibly decreased subsequently” is a bit weird here, as this is shown not to be the case in line 217.

Line 215: Can you statistically test differences in extinction rates for different conditions?
Remarks Monnin et al., 2020

Experimental evolution of virulence and associated traits in a Drosophila melanogaster – Wolbachia symbiosis

Abstract
- Line 17: why is it important to mention “high rearing temperatures”. Does the infectivity of *Wolbachia* depend on temperature?
- Line 23: what are octomom copies?
- Line 23: to which conditions are referred? You mean high temperature and enforced late reproduction?
- Line 25: are correlated with what? With each other?

Introduction
- Line 34: trade-off. You mention that parasites face a trade-off to maximize their transmission. In the next sentences first a positive correlation is mentioned and then this trade-off is described. For me it was not directly clear what the trade-off was. I think it would be more clear if you directly mention the players in this trade-off before you explain it: “… face a trade-off between the transmission rate and its virulence”.
- Line 45: inter-host and intra-host. Can you explain what you mean with this two types of selection?
- Line 56: I think it would be a good idea to mention here a bit more about what is known about the virulence of *Wolbachia* under different temperatures.

Material and Methods
- Line 99: What is paraquat exactly?
- Line 126: How was the extinction rate calculated? What do you mean with the extinction rate was measured both before and after the last survival measurement (G9)?
- Line 139: “300 µL of AB solution was added” (instead of were added)?
- Line 172: “supplemented with water (control) or paraquat (as described above)” (instead of water, paraquat)?
- Why was generation 9 chosen to compare survival and *Wolbachia* density?
- Is there a reason why specific generations were picked to measure survival (G3 and G9) or density and octomom copy number (G2, G9, G12 and G17)?

Results and discussion
- Line 202 + fig 2a: Is it possible to plot both control and paraquat separately in this figure. I think this figure would be more interesting to show.
- Line 294-295: Inter-host and intra-host selection. For me it is not clear what the different selection pressures are in these two types of selection.

General remarks
- More information about inter-host and intra-host selection is needed.
+ Experiment is well explained.
- The surprising results are elaborately explained. Multiple possible explanations are formulated and discussed.