



Commentary

Commentary: Assessing long-run deworming impacts on education and economic outcomes: a comment on Jullien, Sinclair and Garner (2016)

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Introduction

Jullien, Sinclair and Garner (2016)¹ (henceforth JSG) state that they seek to ‘appraise the methods’ of three recent papers that estimate long-run impacts of mass deworming on educational or economic outcomes. This commentary focuses on their discussion of Baird, Hicks, Kremer and Miguel (2016)² (henceforth Baird). We welcome scrutiny of our work, and appreciate the opportunity to discuss JSG.¹

Baird² finds evidence of gains in some educational and labour outcomes 10 years after a deworming programme in 75 Kenyan primary schools. Some gains are found in the full sample, and others among either males or females, in ways that are sensible given the context, e.g. there are gains in manufacturing employment among males but not females, fewer of whom work in this sector in Kenya.

Below we discuss JSG’s claim that the evidence in Baird² is unreliable. It is not surprising that any two scholars might interpret a body of results differently, but JSG¹ make a series of claims that appear overstated or are somewhat misleading. Due to word limits, we discuss some points here and others in the [Supplementary Appendix](#) (available as [Supplementary data](#) at *IJE* online).

Discussion of JSG

JSG¹ do not make a substantive critique that the results in Baird are inaccurate or not robust to alternative specifications, presumably because they did not identify such issues. Rather, they make a methodological critique, namely that the results in Baird² are unreliable due to potentially selective reporting of positive results (JSG,¹ Table 3). We have several responses.

First, JSG¹ do not present any statistical evidence of selective reporting. They acknowledge that their claims are instead based on ‘a narrative analysis’. Like witchcraft, it is easy to make claims about selective reporting, but difficult to prove—or disprove—whether it has occurred. In fact, most patterns presented in their tables lean against systematic reporting bias.

To start, across working paper versions of Baird² the same basic set of outcomes are presented with only minor adjustments (typically in response to suggestions by colleagues or journal referees). Furthermore, many of the robustness checks in Baird²—examining alternative outcomes or statistical specifications, and multiple testing adjustments—are included precisely to address reporting concerns. If the main results reported in Baird² were simply

false-positives, then perhaps only roughly 5% of all tests in Baird² and its lengthy appendix would be significant at 95% confidence, but the proportion is an order of magnitude higher. If we had attempted to ‘cherry-pick’ results, the proportion of significant results should have risen across versions of Baird, but it remains entirely stable (see JSG,¹ Appendix 3).

JSG¹ focus on the fact that Baird² present results for the full sample as well as by gender. Since the gender breakdown was not present in the first, incomplete, 2011 working paper version, JSG¹ imply that any discussion of gender per se constitutes evidence of selective reporting. (Note that, beyond gender, there is little subgroup reporting in Baird.²)

Yet there are ample conceptual rationales for considering impacts for women and men separately. It is standard in economics to disaggregate labour market analysis by gender (Bertrand 2011),³ especially for young adults, given the effects of childbearing. An influential contribution, Pitt, Rosenzweig and Hassan (2012),⁴ cited in Baird,² makes a theoretical case and presents evidence that educational impacts of health investments are likely to be larger among females, and labour impacts larger for males, in a low-income setting. Any a priori analytical plan would have specified this subgroup analysis. JSG’s dismissal of the results for females trivializes the importance of gender in low-income countries like Kenya, where women and men face starkly different economic opportunities.

Another aspect of the selective reporting discussion relates to the multiple testing adjustments in Baird.² In contrast to Baird,² JSG¹ claim (JSG,¹ Table 6) that the main results are not robust to multiple testing correction. We found this discussion to be among the least informative parts of JSG.¹ The data in the ‘Effect robust to adjustment for multiple inference?’ column ignores the fact that the adjusted *P*-values corresponding to the ‘No’ values range from 0.07 to 0.13, in other words near traditional significance levels even after adjustment (see our Table 1). Baird² report these values but JSG¹ opt not to mention them. By overemphasizing small changes in *P*-values around the arbitrary 0.05 threshold, JSG¹ create the impression that results are fragile when that is not the case. Moreover, deworming effects on the most comprehensive measure of living standards in Baird,² the meals eaten outcome, remain significant at the 0.05 level even after adjustment.

The final column in JSG’s Table 6 (‘Effect consistent across related outcomes?’) is also largely uninformative. The goal of the multiple testing adjustment is to account for a set of results; cherry-picking one outcome in a broader family that is not significant and highlighting it as evidence of a lack of robustness, as JSG¹ do, is less scientific. For example, for the Baird² finding that males who

received more deworming work more in manufacturing jobs, JSG¹ argue that there are no related outcomes with statistically significant effects; however, in fact these men also have significantly higher labour earnings.

JSG¹ also critique approaches to the presentation of results in Baird² that are standard in economics and other social sciences, but do not conform to norms in their own field. For instance, JSG¹ mention—at least a dozen times!—whether or not results are reported in Baird’s abstract, and emphasize that the height results are not reported there. Yet it is not surprising that this result did not make it into Baird’s abstract: the structure of economics abstracts is not standardized, unlike public health articles, and they are short, typically 100–250 words (the Baird² abstract has 146 words). Instead, economics articles usually summarize results in the introduction. The height results in Baird² are reported in the introduction, as well as in the main text and tables.

JSG¹ also emphasize that Baird² lack a pre-analysis plan. Whereas it is true that Baird² did not register a pre-analysis plan, such plans were until recently largely unknown in economics, and the American Economic Association RCT registry was only established in 2013.

At times, JSG¹ appeal to Cochrane review results (Taylor-Robinson *et al.* 2015)⁵ to bolster their case. However, the Cochrane review results are problematic (Montresor *et al.* 2016),⁶ with an incomplete sample of studies, improper selective exclusion of a study that shows weight gains (e.g. Stephenson *et al.* 1993),⁷ and an underpowered statistical test. Croke *et al.* (2016)⁸ show that mass deworming leads to child weight gains at the community level.

Conclusion

The issue of selective reporting raised by JSG¹ is potentially important, but the evidence presented in JSG¹ does not change our interpretation of Baird². It is JSG’s right to interpret the evidence in their own way, of course, but we cannot help but feel that a more even-handed discussion would have been more productive for scientific progress. A more scientific assessment would discuss Baird’s strengths as well as weaknesses, for example: the value of its long-term longitudinal data, which allow estimation of the benefit-cost ratio for mass deworming and suggest that long-run income gains might be 100 times the (small) initial cost. A more even-handed appraisal would not cherry-pick null results to highlight, present multiple testing adjustments in a tendentious fashion (in JSG¹ Table 6) or summarily dismiss analysis by gender in this context. The discussion could have mentioned a methodological strength of Baird,² namely, the fact that two orthogonal

Table 1 Unadjusted *P*-values and adjusted *q*-values, added to Jullien *et al.* Table 6 (2016)¹

Outcomes reported in the abstract		Effect robust to adjustment for multiple inference?		
		Unadjusted <i>P</i> -value	JSG claim	Adjusted <i>q</i> -value
Men	'stay enrolled for more years of primary school'	0.022	No	0.071
	'work 17% more hours each week'	0.017	No	0.083
	'spend more time in non-agricultural self-employment'	0.066	Remains borderline	0.133
	'spend more time in manufacturing'	0.015	No	0.083
	'miss one fewer meal per week'	0.003	Yes	0.031
Women	'one quarter more likely to have attended secondary school'	0.022	No	0.084
	'reallocating time from traditional agriculture into cash crops'	0.031	No	0.103
	'reallocate time from traditional agriculture into non-agricultural self-employment'	0.025	No	0.103

All but the last column reproduced from Jullien *et al.* Table 6 (2016).¹ The last column includes the adjusted false discovery rate (FDR) *q*-values from the Supplementary Appendix of Baird *et al.* (2016).²

sources of variation—a cost-sharing experiment carried out in a random subset of schools (which lowered deworming drug take-up), and variation in exposure to cross-school treatment spillovers—both reinforce the main results.

It may be worth stepping back and thinking about the broader public policy debate regarding deworming. The decision to fund mass deworming should be based on comparing its expected costs and benefits, so even a small probability that the effects in Baird² are present would make the cost effectiveness analysis favourable. To be very concrete, was the Indian Government's recent decision to carry out mass school-based deworming—at pennies per dose (using safe and approved drugs) in areas with widespread infections—misguided and not 'informed by robust evidence', as JSG¹ suggest? It appears that even JSG¹ agree that deworming might be sensible and cost-effective in such a setting, when they write: 'If a community in a given setting has a high prevalence of untreated worm infections, then mass-deworming programmes may well be an effective way to reach and treat a large number of children'.

The long-run benefits found in Baird,² Croke (2014)⁹ and Ozier (2016),¹⁰ as well as Bleakley (2007),¹¹ medium-run schooling impacts reported in Miguel and Kremer (2004)¹² and the short-run child weight gains in Croke *et al.* (2016),⁸ taken together may lead mass deworming to go from being merely 'very cost-effective' to 'extremely cost-effective'. Either way, the logic of mass drug deworming administration in endemic regions appears as clear today as it was when the World Health Organization began supporting this policy decades ago.

Supplementary Data

Supplementary data are available on the *IJE* website.

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