Analysis Plans in Economics

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- Completely standard practice in medical trials and required in fact by the FDA

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"It is possible, for example, to estimate whether the average treatment effect is positive or negative for various subgroups of interest. One immediate charge against such a procedure is data mining. A sufficiently determined examination of any trial will eventually reveal some subgroup for whom the treatment yielded a significant effect of some sort, and there is no general way of adjusting standard errors to protect against the possibility. In drug trials, the FDA rules require that analytical plans be submitted prior to trial, and at least one economic experiment—moving to opportunity—has imposed similar rules on itself, see the protocol by Feins and McInnis (2001). I am not arguing against post-trial subgroup analysis, only that any special epistemic status (as in "gold standard," "hard," or "rigorous" evidence) possessed by RCTs does not extend to subgroup analysis if only because there is no general guarantee that a new RCT on post-experimentally defined subgroups will yield the same result."

On the other hand...

Deaton: "Randomization in the tropics":

"It is clearly absurd to discard data because we do not know how to analyze it with sufficient purity. Indeed, many important findings have come from post-trial analysis of experimental data, both in medicine and in economics, for example of the negative income tax experiments of the 1960s. None of which resolves the concerns about data-mining. In large-scale, expensive, trials, a zero or very small result is unlikely to be welcome, and there is likely to be overwhelming pressure to search for some subpopulation that gives a more palatable result."

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- How to solve data mining? Ex-ante designated one table as "key results" table and one number, the average standardized effects for 12 main indicators, as key metric of success.

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- Some thoughts:
 - *Endline vs. midline.* We did a new endline analysis plan again after having looked at midline data, but before looking at endline data. This seems totally kosher to me: this is fresh data.
 - One-sided tests. My view is we should be allowed to specify one-sided p-values in our analysis plans. At least we would get something in return for pre-specifying, since in many cases – like my education example – that's how we react anyway!

Some examples from my current experience

Targeting the Poor (with Alatas Banerjee, Hanna, and Tobias)

Setting:

- Study in Indonesia on how to identify poor households for anti-poverty programs
- Three treatments: PMT, Community Targeting, and Hybrid (to prevent elite capture of community process)

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 - Paper ended up being about something very different than we thought. (No elite capture; about different local welfare metrics)
 - Luckily in this case our analysis plan was sufficiently general that it was OK. But usually that won't be true. How to retain this flexibility? What if we had been required to report tons of null results on elite capture for parts of the tree we didn't explore?

Some examples from current experience

Taxes in Pakistan (with Khwaja and Khan)

- Setting:
 - Multi-year RCT in Pakistan on tax collectors
 - Treatment is pay-for-performance incentives for tax collectors (3 versions thereof), vs. control

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- Doing analysis plan rapidly, but hard to think out every detail.
- Is it better to do a not-perfect analysis plan that one at all? Will we be hamstrung in our analysis in the future?

Some thoughts

- When pre-analysis plans seem most helpful.
 - When you have a party with strong vested interest. Get them to sign off up front!
 - When you suspect subgroup analysis is going to be a big deal. See Deaton.
 - Specifying your specifications. Usually a good idea.
 - Specifying control variables for balance checks.
 - When you have a simple outcome variable or can define one; or, alternatively, when you have a zillion outcome variables and are worried about data mining.

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- When it can be harder... and why we should be careful not to necessarily penalize people for not having them.
 - When things happen in real time (aside: how does medicine deal with this? optimal stopping rules? I don't know)
 - When you are testing theories
 - When your understanding of what you're testing evolves as you move through the project