Erratum and discussion of propensity score reweighting

© Austin Nichols austinnichols@gmail.com

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1 Erratum

Nichols (2007) described estimating the probability an observation receives a binary "treatment" as a function of observable variables X (using e.g. logit or probit), and using the estimated probabilities of treatment or "propensity scores" $\hat{\lambda}$ to reweight the data (as an alternative to matching). Section 3.4 neglects to mention that the weights $\hat{\lambda}/\left(1-\hat{\lambda}\right)$ should only be applied to the control group in order to make the mean of each variable in the matrix X (i.e. those variables included in the propensity score model) approximately equal across the treatment and control groups. The examples in Section 3.5 also neglect this restriction on propensity-based weights.

That is, the line

$$g w=ps/(1-ps)$$

which generates a weight equal to $\hat{\lambda}/\left(1-\hat{\lambda}\right)$ should be followed by the command

(where _tr is a treatment indicator and _ps the propensity score). This makes the weight equal one for observations receiving treatment (e.g. those belonging to a union, or having completed college), i.e. those having _tr==1. It is also advisable to scale weights within the treatment and control groups so that the reweighted proportions are similar to those observed in the original sample. In fact, the reweighting of the control group to resemble the treatment group is only one of several plausible reweighting schemes, and a regression of outcomes on a treatment indicator using this weight can be considered an estimate of the average treatment effect on the treated (ATT).

2 Alternative weighting schemes

The pair of commands generating weights can be replaced by the single command

with the same result. A rescaled weight to approximately preserve proportions in treatment and control would be

noting in passing that multiplying treatment weights by p/(1-p) where p is the proportion of the sample receiving treatment, or multiplying control weights by (1-p)/p, or multiplying treatment weights by p and control weights by (1-p), all produce identical results if weights are themselves rescaled to sum to N (note that Stata internally rescales aweights to sum to N). The weight $\widehat{\lambda}/\left(1-\widehat{\lambda}\right)$ for untreated

"control" observations reweights the distribution of observable characteristics included in the logit or probit model to be like that of the treated group. A weighted regression of outcome on treatment is thus a comparison of means across treatment and control groups, but the control group is reweighted to represent the average outcome that treatment group would have exhibited in the absence of treatment. That is, every control group observation is contributing to an estimate of the mean counterfactual outcome for all treated observations (rather than specific observations being matched).

An alternative weighting scheme of the form

```
su _tr
g w2=cond(_tr,(1-_ps)/_ps*r(mean)/(1-r(mean)),1)
```

reweights the distribution of observables in the treatment group to be like that of the control group. A comparison of means across (reweighted) treatment and control groups, for example using a weighted regression of an outcome variable on the treatment indicator, is then an estimate of the average treatment effect on the controls (ATC). The treatment group is reweighted to represent the average outcome that control group would have exhibited in the presence of treatment.

One method of computing an estimate of the average treatment effect for the population (ATE) is to take the weighted mean of these two estimates, with the weight attached to the ATT equal to the proportion receiving treatment and the weight attached to the ATC equal to one minus the proportion receiving treatment.

An alternative estimate of the ATE is available. First, note that the outcome under treatment for the whole population, i.e. the mean outcome if every unit received treatment, can be estimated by a weighted mean of outcomes in the treatment group with weights $1/\hat{\lambda}$ (Brunell and Dinardo 2004). Similarly, the outcome under control for the whole population, i.e. the mean outcome if every unit received no treatment, can be estimated by a weighted mean of outcomes in the control group with weights $1/\left(1-\hat{\lambda}\right)$. The weights for both groups are given by

```
su _tr
g w3=cond(_tr,1/_ps*r(mean)/(1-r(mean)),1/(1-_ps))
```

and an ATE estimate is then simply a weighted comparison of means (e.g. via a regression). One problem that is exacerbated in this scheme is measurement error in the estimated propensity score: as Dinardo (2002) writes: "small errors in estimating $\rho(x)$ can produce potentially large errors in the weights. Since the weight is a nuisance parameter from the viewpoint of estimating a density or a specific moment of the distribution, this is not a straightforward problem."

A fourth reweighting scheme

```
g w4=cond(_tr,(1-_ps),_ps)
```

minimizes the observable distance between treatment and control groups in the sense that a test statistic for the difference in means (the Hotelling test) is zero (and the weighted groups are of equal size, so the mean of the treatment indicator is one half), but a difference in means using this weight is not so readily interpreted as an average treatment effect. Nevertheless, simulation evidence not presented here indicates it may be very effective (in the sense of having small bias and MSE) in estimating the average treatment effect, especially when estimated probabilities are near zero or one. ¹ It also exhibits good robustness to omitted variables in the selection equation (the first stage logit or probit).

See Lunceford and Davidian (2004) and Busso, DiNardo, and McCrary (2008) for additional discussion of construction of weights and rescaling, including a asymptotically variance-minimizing choice.

¹Note that estimated propensities near zero or one represent a possible violation of the condition required for matching or reweighting that the probability of treatment is bounded away from zero and one. In this case, it is advisable to restrict to a subpopulation in which estimated propensities are never near zero or one and reestimate. The densities of propensities near the zero and one boundaries should be estimated using kdens, with boundary correction options, available from SSC.

3 Result of reweighting

The results of reweighting are clear in a Hotelling test or an equivalent linear discriminant model. The example below² using hotelling and regress give identical F statistics, but the regress approach allows relaxing of the assumption of equal variance across groups via the vce(robust) option.

```
webuse nlswork, clear
keep if year==77
local x "collgrad age tenure not_smsa c_city south nev_mar"
hotelling 'x', by(union)
regress union 'x'
regress union 'x', vce(robust)
logit union 'x'
predict _ps if e(sample)
summarize union if e(sample)
local p=r(mean)
generate w3=cond(union, 'p'/_ps/(1-'p'),1/(1-_ps))
hotelling 'x' [aw=w], by(union)
regress union 'x' [aw=w]
regress union 'x' [aw=w], vce(robust)
regress ln_wage union 'x' [aw=w3], vce(robust)
```

Note that the F statistic drops from 20 to 0.1 after reweighting (18 to 0.1 using heteroskedasticity-robust statistics), and the weighted means of each individual variable look much closer. The last regression of log(wage) on union using the inverse probability weights based on propensity scores gives an estimate of the effect of union membership on wages, over both union and nonunion workers, suggesting that an individual would earn fourteen percent more in 1977 as a union member than as a nonunion worker, on average. Using instead weights generated by

```
g w1=cond(union, 'p'/(1-'p'), _ps/(1-_ps))
regress ln_wage union 'x' [aw=w1], vce(robust)
```

gives an estimate of the effect of union membership on wages for union members, suggesting that union members earned fourteen and one half percent more in 1977 than they would have as nonunion workers.

What is not clear from hotelling or regress is that even if the means of X variables are equal in the reweighted sample, that does not imply that their distributions are similar. As long as treatment status can be inferred from higher moments of the X variables, we have not fully controlled for the observable differences across treatment and control groups. In practice, however, reweighting to make means match seems to make the distributions of observables very similar, for the same reason that matching on the propensity score does.

The difference in distributions is most cleanly observable in the distribution of estimated propensity scores, but can be seen in the individual variables (e.g. tenure in the example below; note that kdens is available from SSC).

```
webuse nlswork, clear
keep if year==77
local x "collgrad age tenure not_smsa c_city south nev_mar"
logit union 'x'
predict _ps if e(sample)
kdens _ps if union, bw(.03) ll(0) ul(1) gen(f1 x) nogr
kdens _ps if !union, bw(.03) ll(0) ul(1) gen(f0) at(x) nogr
label var f0 "pdf of propensities for unweighted non-union (control) obs"
label var f1 "pdf of propensities for unweighted union (treatment) obs"
```

²Note that this extract of the National Longitudinal Survey of Young Women 14-26 years of age in 1968 does not include sample weights, but in general we would prefer to convolve the weights by multiplying our reweighting factor by the sample weights.

```
line f1 f0 x, leg(col(1)) name(unwtd, replace)
summarize union if e(sample)
local p=r(mean)
generate w3=cond(union, 'p'/(1-'p'), _ps/(1-_ps))
kdens _ps if union [aw=w3], bw(.03) ll(0) ul(1) gen(g1 x1) nogr
kdens _ps if !union [aw=w3], bw(.03) ll(0) ul(1) gen(g0) at(x1) nogr
label var g0 "pdf of propensities for reweighted non-union (control) obs"
label var g1 "pdf of propensities for reweighted union (treatment) obs"
line g1 g0 x1, leg(col(1)) name(rewtd, replace)
kdens tenure if union [aw=w3], bw(1.5) ll(0) gen(td) at(tenure) nogr
label var td "Density for union members"
kdens tenure if !union [aw=w3], bw(1.5) ll(0) gen(cd) at(tenure) nogr
label var cd "Density for nonunion reweighted to resemble union members"
line td cd tenure, sort leg(col(1))
```

Matching on the propensity score ensures the distributions of estimated propensity scores are virtually identical in (matched) treatment and control groups, especially if matching models are iterated until balance is achieved, but reweighting does not. For example, if the distribution of some variable (including propensity scores) is bimodal in the control group and single-peaked in the treatment group, those properties will typically still be observable in the reweighted data. Nevertheless, reweighting achieves much of the balancing achievable via matching on the propensity score.

The last approach to reweighting always achieves the smallest difference in means, with an F statistic as close to zero as is feasible given machine precision, but the distributions of observable characteristics and estimated propensity scores are very similar under all these approaches to reweighting. See Iacus, King, and Porro (2008) for an alternative matching method that controls, up to user-chosen specified levels, "for all imbalances in central absolute moments, comoments, coskewness, interactions, nonlinearities, and other multidimensional distributional differences between treated and control groups."

4 Uses of reweighting

The propensity-based reweighting approach is at the heart of the method proposed by Dinardo, Fortin, and Lemieux (1996). The paradigmatic example of that approach uses two years of data, estimates the probability that an observation is in the first year or the second, then reweights the second year's observations by $\hat{\lambda}/\left(1-\hat{\lambda}\right)$ so that the distributions are nearly equal across the two years. Changes in means or distributions (of some outcome variables) in the reweighted data are then interpreted as estimates of change had the means of the X variables not changed over time.³

A similar method could be applied to estimate the proportion of a wage gap observed across men and women or white and nonwhite workers that are attributable to characteristics, along the lines of the oaxaca method described by Jann (2008) and related methods referenced there. The connections between these methods is discussed by e.g. Dinardo (2002) and Lemieux (2002).

The reweighting approach extends easily to a polychotomous categorical treatment variable, by considering the analogy to the Dinardo, Fortin, and Lemieux (1996) approach applied to multiple years. For example, each subsequent year's data can be reweighted to have observable characteristics similar to the first year, or each year can be reweighted to match some other base year's distribution. In the same way, observations receiving various levels of treatment can be reweighted to match some base category (the choice of base category may affect the interpretation of results).

Extensions of the reweighting approach to the case of a continuous treatment are also possible using the generalized propensity score approach of Hirano and Imbens (2004), described by Bia and Mattei (2008). The generalized propensity score r(t|x) is the density of treatment conditional on X = x, estimated as $\hat{r}(t,x)$.

³See also Altonji, Bharadwaj, and Lange (2008) for a recent paper dealing not only with differences in distributions in samples, but sample attrition and missing values.

5 Missing data, and a conjecture

Note that this kind of reweighting could also be employed to correct for bias due to missing data. For example, the distributions of variables observed for both survey respondents and nonrespondents (i.e. potential stratification variables) can be adjusted via reweighting to look similar. In that case, the hope is that the unobservable survey responses of nonrespondents will be suitably captured by reweighted respondents. An alternative approach, imputing responses to nonrespondents, is a form of matching. Depending on the type of imputation, this may be propensity score matching, or nearest neighbor matching, or exact matches on observables (also known as hotdeck imputation).

The standard approach to missing data is to multiply impute responses (see e.g. Carlin, Galati, and Royston 2008). It is natural to wonder whether multiple imputation could not also be fruitfully applied to the imputation of hypothetical counterfactual outcomes (the unobserved outcomes of treatment cases when in the control group, or outcomes of control cases when in the treatment group).

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7 References

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