

Эконометрика, 2017-2018, 1 модуль

Семинары 1 - 2

2.04.18 и 9.04.18 для

Группы Э_Б2015_Э_3

Семинарист О.А.Демидова

Критика М.Фридменом стандартной функции потребления, раздел 8.5.

1) (Доугерти, 8.7) В некоторой экономике дисперсия переменного дохода составляет 0.5 от дисперсии постоянного дохода, склонность к потреблению товаров кратковременного пользования за счет постоянного дохода составляет 0.6, а расходы на товары длительного пользования отсутствуют. Каким будет значение мультипликатора, полученного на основе построения «наивной» регрессионной зависимости потребления от дохода, и каково его истинное значение?

2) (Доугерти, раздел 8)

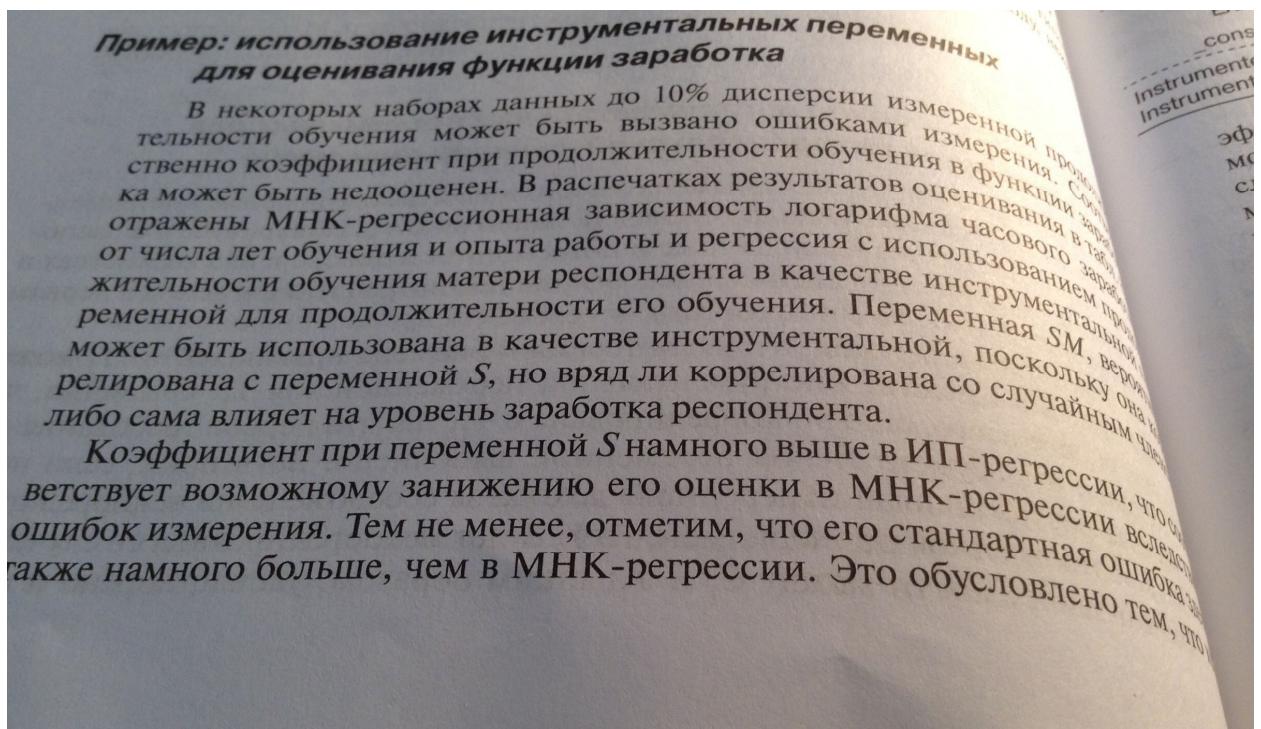


Таблица 8.2

reg LGEARN S EXP

Source	SS	df	MS				
Model	50.9842581	2	25.492129				
Residual	135.723385	537	.252743734				
Total	186.707643	539	.34639637				

LGEARN	Coef.	Std. Err.	t	P > t	[95% Conf. Interval]
S	.1235911	.0090989	13.58	0.000	.1057173 .141465
EXP	.0350826	.0050046	7.01	0.000	.0252515 .0449137
_cons	.5093196	.1663823	3.06	0.002	.1824796 .8361596

Number of obs = 540
F(2,537) = 100.86
Prob > F = 0.0000
R-squared = 0.2731
Adj R-squared = 0.2704
Root MSE = .50274

ivreg LGEARN EXP (S=SM)
instrumental variables (2SLS) regression

Source	SS	df	MS				
Model	46.9446075	2	23.4723038				
Residual	139.763036	537	.260266361				
Total	186.707643	539	.34639637				

LGEARN	Coef.	Std. Err.	t	P > t	[95% Conf. Interval]
S	.1599676	.0252801	6.33	0.000	.1103076 .2096277
EXP	.0394422	.0058092	6.79	0.000	.0280306 .0508537
_cons	-.0617062	.4061769	-0.15	0.879	-.8595966 .7361841

Number of obs = 540
F(2,537) = 28.38
Prob > F = 0.0000
R-squared = 0.2514
Adj R-squared = 0.2486
Root MSE = .51016

Instrumented: S
Instruments: EXP SM

Таблица 8.4

. ivreg LGEARN EXP ASVABC MALE ETHBLACK ETHHISP
(S=SM SF SIBLINGS LIBRARY)
Instrumental variables (2SLS) regression

Source	SS	df	MS				
Model	64.4915831	6	10.7485972				
Residual	122.21606	533	.229298424				
Total	186.707643	539	.34639637				
LGEARN	Coef.	Std. Err.	t	P > t	[95% Conf.		
S	.111379	.0476886	2.34	0.020	.0176984	Inter	
EXP	.0258798	.0081187	3.19	0.002	.0099313		
ASVABC	.0092263	.007991	1.15	0.249	-.0064714		
MALE	.2619787	.0429283	6.10	0.000	.1776492		
ETHBLACK	-.0121846	.0822942	-0.15	0.882	-.1738454		
ETHHISP	.0457639	.0955115	0.48	0.632	-.1418612		
_cons	.2258512	.3887468	0.58	0.562	-.5378125		

Instrumented: S

Instruments: EXP ASVABC MALE ETHBLACK ETHHISP SM SF SIBLINGS LIBRARY

. estimates store EARNIV

. reg LGEARN S EXP ASVABC MALE ETHBLACK ETHHISP

Source	SS	df	MS				
Model	65.490707	6	10.9151178				
Residual	121.216936	533	.227423895				
Total	186.707643	539	.34639637				
LGEARN	Coef.	Std. Err.	t	P > t	[95% Conf.		
S	.0883257	.0109987	8.03	0.000	.0667196	Inter	
EXP	.0227131	.0050095	4.53	0.000	.0128724		
ASVABC	.0129274	.0028834	4.48	0.000	.0072633		
MALE	.2652878	.042235	6.28	0.000	.1823203		
ETHBLACK	.0077265	.0715863	0.11	0.914	-.1328994		
ETHHISP	.0536544	.0937966	0.57	0.568	-.1306019		
_cons	.4002952	.1663149	2.41	0.016	.0735821		

люч «constants»
ен имеет раз-
сл одинако-

Окончание табл. 8.4

estimates store EARNOLS
hausman EARNIV EARNOLS, constant
— Coefficients —

	(b) EARNIV	(B) EARNOLS	(b - B) Difference	sqrt(diag (V_b - V_B)) S.E.
S	.111379	.0883257	.0230533	.0464029
EXP	.0258798	.0227131	.0031667	.0063889
ASVABC	.0092263	.0129274	-.0037011	.0074527
MALE	.2619787	.2652878	-.0033091	.0076842
ETHBLACK	-.0121846	.0077265	-.019911	.0405924
ETHHISP	.0457639	.0536544	-.0078904	.018018
_cons	.2258512	.4002952	-.174444	.3513736

b = consistent under Ho and Ha; obtained from ivreg
B = inconsistent under Ha, efficient under Ho;
obtained from regress

Test: Ho: difference in coefficients not systematic

$$\chi^2(7) = (b - B)'[(V_b - V_B)^{-1}](b - B) = 0.25$$

$$Prob > \chi^2 = 0.9999$$

5.3.2 Medical expenditures with one endogenous regressor

We consider a model with one endogenous regressor, several exogenous regressors, and one or more excluded exogenous variables that serve as the identifying instruments.

The dataset is an extract from the Medical Expenditure Panel Survey (MEPS) of individuals over the age of 65 years, similar to the dataset described in section 3.2.1. The equation to be estimated has the dependent variable `ldrugexp`, the log of total out-of-pocket expenditures on prescribed medications. The regressors are an indicator for whether the individual holds either employer or union-sponsored health insurance (`hi_empunion`), number of chronic conditions (`totchr`), and four sociodemographic variables: age in years (`age`), indicators for whether female (`female`) and whether black or Hispanic (`blhisp`), and the natural logarithm of annual household income in thousands of dollars (`linc`).

We treat the health insurance variable `hi_empunion` as endogenous. The intuitive justification is that having such supplementary insurance on top of the near universal Medicare insurance for the elderly may be a choice variable. Even though most individuals in the sample are no longer working, those who expected high future medical expenses might have been more likely to choose a job when they were working that would provide supplementary health insurance upon retirement. Note that Medicare did not cover drug expenses for the time period we study.

We use the global macro `x2list` to store the names of the variables that are treated as exogenous regressors. We have

```
. * Read data, define global x2list, and summarize data
. use mus06data.dta
. global x2list totchr age female blhisp linc
. summarize ldrugexp hi_empunion $x2list
```

Variable	Obs	Mean	Std. Dev.	Min	Max
ldrugexp	10391	6.479668	1.363395	0	10.18017
hi_empunion	10391	.3796555	.4853245	0	1
totchr	10391	1.860745	1.290131	0	9
age	10391	75.04639	6.69368	65	91
female	10391	.5797325	.4936256	0	1
blhisp	10391	.1703397	.3759491	0	1
linc	10089	2.743275	.9131433	-6.907755	5.744476

6.3.3 Available instruments

We consider four potential instruments for `hi_empunio`. Two reflect the income status of the individual and two are based on employer characteristics.

The `ssratio` instrument is the ratio of an individual's social security income to the individual's income from all sources, with high values indicating a significant income constraint. The `lowincome` instrument is a qualitative indicator of low-income status. Both these instruments are likely to be relevant, because they are expected to be negatively correlated with having supplementary insurance. To be valid instruments, we need to assume they can be omitted from the equation for `ldrugexp`, arguing that the direct role of income is adequately captured by the regressor `linc`.

The `firmsz` instrument measures the size of the firm's employed labor force, and the `multlc` instrument indicates whether the firm is a large operator with multiple locations. These variables are intended to capture whether the individual has access to supplementary insurance through the employer. These two variables are irrelevant for those who are retired, self-employed, or purchase insurance privately. In that sense, these two instruments could potentially be weak.

```

* Summarize available instruments
* summarize sratio lowincome multlc firmsz if linc!=.

```

Variable	Obs	Mean	Std. Dev.	Min	Max
<code>ssratio</code>	10089	.5365438	.3678175	0	9.25062
<code>lowincome</code>	10089	.1874319	.3902771	0	1
<code>multlc</code>	10089	.0620478	.2412543	0	1
<code>firmsz</code>	10089	.1405293	2.170389	0	50

We have four available instruments for one endogenous regressor. The obvious approach is to use all available instruments, because in theory this leads to the most efficient estimator. In practice, it may lead to larger small-sample bias because the small-sample biases of IV estimators increase with the number of instruments (Hahn and Hausman 2002).

At a minimum, it is informative to use `correlate` to view the gross correlation between endogenous variables and instruments and between instruments. When multiple instruments are available, as in the case of overidentified models, then it is actually the partial correlation after controlling for other available instruments that matters. This important step is deferred to sections 6.4.2 and 6.4.3.

6.3.4 IV estimation of an exactly identified model

We begin with IV regression of `ldrugexp` on the endogenous regressor `hi_empunion`, instrumented by the single instrument `ssiratio`, and several exogenous regressors.

We use `ivregress` with the `2sls` estimator and the options `vce(robust)` to control for heteroskedastic errors and `first` to provide output that additionally reports results from the first-stage regression. The output is in two parts:

```
. * IV estimation of a just-identified model with single endog regressor
. ivregress 2sls ldrugexp (hi_empunion = ssiratio) $x2list, vce(robust) first
```

First-stage regressions

	Coef.	Robust Std. Err.	t	P> t	[95% Conf. Interval]
hi_empunion					
totchr	.0127865	.0036655	3.49	0.000	.0056015 .0199716
age	-.0086323	.0007087	-12.18	0.000	-.0100216 -.0072431
female	-.07345	.0096392	-7.62	0.000	-.0923448 -.0545552
blhisp	-.06268	.0122742	-5.11	0.000	-.08674 -.0386201
linc	.0483937	.0066075	7.32	0.000	.0354417 .0613456
ssiratio	-.1916432	.0236326	-8.11	0.000	-.2379678 -.1453186
_cons	1.028981	.0581387	17.70	0.000	.9150172 1.142944

Number of obs = 10089
F(6, 10082) = 119.18
Prob > F = 0.000
R-squared = 0.0761
Adj R-squared = 0.0755
Root MSE = 0.4672

Instrumental variables (2SLS) regression

	Coef.	Robust Std. Err.	z	P> z	[95% Conf. Interval]
ldrugexp					
hi_empunion	-.8975913	.2211268	-4.06	0.000	-1.330992 -.4641908
totchr	.4502655	.0101969	44.16	0.000	.43028 .470251
age	-.0132176	.0029977	-4.41	0.000	-.0190931 -.0073421
female	-.020406	.0326114	-0.63	0.531	-.0843232 .0435113
blhisp	-.2174244	.0394944	-5.51	0.000	-.294832 -.1400167
linc	.0870018	.0226356	3.84	0.000	.0426368 .1313668
_cons	6.78717	.2688453	25.25	0.000	6.260243 7.314097

Number of obs = 10089
Wald chi2(6) = 2000.86
Prob > chi2 = 0.000
R-squared = 0.0640
Root MSE = 1.3177

Instrumented: hi_empunion
Instruments: totchr age female blhisp linc ssiratio

6.3.6 Testing for regressor endogeneity

The preceding analysis treats the insurance variable, `hi_empunion`, as endogenous. If instead the variable is exogenous, then the IV estimators (IV, 2SLS, or GMM) are still consistent, but they can be much less efficient than the OLS estimator.

The Hausman test principle provides a way to test whether a regressor is endogenous. If there is little difference between OLS and IV estimators, then there is no need to instrument, and we conclude that the regressor was exogenous. If instead there is considerable difference, then we needed to instrument and the regressor is endogenous. The test usually compares just the coefficients of the endogenous variables. In the case of just one potentially endogenous regressor with a coefficient denoted by β , the Hausman test statistic

$$T_H = \frac{(\hat{\beta}_{IV} - \hat{\beta}_{OLS})^2}{\hat{V}(\hat{\beta}_{IV} - \hat{\beta}_{OLS})}$$

is $\chi^2(1)$ distributed under the null hypothesis that the regressor is exogenous.

Before considering implementation of the test, we first obtain the OLS estimates to compare them with the earlier IV estimates. We have

```
. * Obtain OLS estimates to compare with preceding IV estimates
. regress ldrugexp hi_empunion $x2list, vce(robust) .

Linear regression                               Number of obs =   10089
                                                F(   6, 10082)    876.85
                                                Prob > F          0.0000
                                                R-squared         0.1770
                                                Root MSE         1.236
```

ldrugexp	Coef.	Robust Std. Err.	t	P> t	[95% Conf. Interval]	
hi_empunion	.0738788	.0259948	2.84	0.004	.0229435	.1248141
totchr	.4403807	.0093633	47.03	0.000	.4220268	.4587346
age	-.0035295	.001937	-1.82	0.068	-.0073264	.0002675
female	.0578055	.0253651	2.28	0.023	.0080848	.1075262
blhisp	-.1513068	.0341264	-4.43	0.000	-.2182013	-.0844122
linc	.0104815	.0137126	0.76	0.445	-.0163979	.037361
_cons	5.861131	.1571037	37.31	0.000	5.553176	6.169085

The OLS estimates differ substantially from the just-identified IV estimates given in section 6.3.4. The coefficient of `hi_empunion` has an OLS estimate of 0.074, greatly different from the IV estimate of -0.898 . This is strong evidence that `hi_empunion` is endogenous. Some coefficients of exogenous variables also change, notably, those for `age` and `female`. Note also the loss in precision in using IV. Most notably, the standard error of the instrumented regressor increases from 0.026 for OLS to 0.221 for IV, an eightfold increase, indicating the potential loss in efficiency due to IV estimation.

The `hausman` command can be used to compute T_H under the assumption that $\hat{V}(\hat{\beta}_{IV} - \hat{\beta}_{OLS}) = \hat{V}(\hat{\beta}_{IV}) - \hat{V}(\hat{\beta}_{OLS})$; see section 12.7.5. This greatly simplifies analysis because then all that is needed are coefficient estimates and standard errors from separate IV estimation (IV, 2SLS, or GMM) and OLS estimation. But this assumption is too strong. It is correct only if $\hat{\beta}_{OLS}$ is the fully efficient estimator under the null hypothesis of exogeneity, an assumption that is valid only under the very strong assumption that model errors are independent and homoskedastic. One possible variation is to perform an appropriate bootstrap; see section 13.4.6.

The postestimation `estat endogenous` command implements the related Durbin-Wu-Hausman (DWH) test. Because the DWH test uses the device of augmented regressors, it produces a robust test statistic (Davidson 2000). The essential idea is the following. Consider the model as specified in section 6.2.1. Rewrite the structural equation (6.2) with an additional variable, v_1 , that is the error from the first-stage equation (6.3) for y_2 . Then

$$y_{1i} = \beta_1 y_{2i} + x'_{1i} \beta_2 + \rho v_{1i} + u_i$$

Under the null hypothesis that y_2 is exogenous, $E(v_{1i} u_i | y_{2i}, x_{1i}) = 0$. If v_1 could be observed, then the test of exogeneity would be the test of $H_0: \rho = 0$ in the OLS regression of y_1 on y_2 , x_1 , and v_1 . Because v_1 is not directly observed, the fitted residual vector

\hat{v}_i from the first-stage OLS regression (6.3) is instead substituted. For independent homoskedastic errors, this test is asymptotically equivalent to the earlier Hausman test. In the more realistic case of heteroskedastic errors, the test of $H_0: \rho = 0$ can still be implemented provided that we use robust variance estimates. This test can be extended to the multiple endogenous regressors case by including multiple residual vectors and testing separately for correlation of each with the error on the structural equation.

We apply the test to our example with one potentially endogenous regressor, `hi_empunio`, instrumented by `ssiratio`. Then

```
. * Robust Durbin-Wu-Hausman test of endogeneity implemented by estat endogenous
. ivregress 2sls ldrugexp (hi_empunio = ssiratio) $x2list, vce(robust)

Instrumental variables (2SLS) regression                                Number of obs =   10089
                                                                    Wald chi2(6)      2000.86
                                                                    Prob > chi2       0.0000
                                                                    R-squared         0.0640
                                                                    Root MSE         1.48177
```

ldrugexp	Coef.	Robust Std. Err.	z	P> z	[95% Conf. Interval]	
hi_empunio	-.8975913	.2211268	-4.06	0.000	-1.330992	-.4641908
totchr	.4502655	.0101969	44.16	0.000	.43028	.470251
age	-.0132176	.0029977	-4.41	0.000	-.0190931	-.0073421
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```
Instrumented: hi_empunio
Instruments: totchr age female blhisp linc ssiratio

. estat endogenous
Tests of endogeneity
Ho: variables are exogenous

Robust score chi2(1)      =   24.935   (p = 0.0000)
Robust regression F(1,10081) =   26.4333   (p = 0.0000)
```

The last line of output is the robustified DWH test and leads to strong rejection of the null hypothesis that `hi_empunio` is exogenous. We conclude that it is endogenous.

We obtain exactly the same test statistic when we manually perform the robustified DWH test. We have

```
. * Robust Durbin-Wu-Hausman test of endogeneity implemented manually
. quietly regress hi_empunio ssiratio $x2list
. quietly predict v1hat, resid
. quietly regress ldrugexp hi_empunio v1hat $x2list, vce(robust)
. test v1hat

( 1) v1hat = 0
      F( 1, 10081) =   26.43
      Prob > F =   0.0000
```

4) (Демешев, Борzych, 18.1)

Величины X_i равномерны на отрезке $[-a; 3a]$ и независимы. Есть несколько наблюдений, $X_1 = 0.5$, $X_2 = 0.7$, $X_3 = -0.1$.

1. Найдите $\mathbb{E}(X_i)$ и $\mathbb{E}(|X_i|)$.
2. Постройте оценку метода моментов, используя $\mathbb{E}(X_i)$.
3. Постройте оценку метода моментов, используя $\mathbb{E}(|X_i|)$.
4. Постройте оценку обобщённого метода моментов используя моменты $\mathbb{E}(X_i)$, $\mathbb{E}(|X_i|)$ и взвешивающую матрицу.

$$W = \begin{pmatrix} 2 & 0 \\ 0 & 1 \end{pmatrix}$$

5. Найдите оптимальную теоретическую взвешивающую матрицу для обобщённого метода моментов
6. Постройте двухшаговую оценку обобщённого метода моментов, начав со взвешивающей матрицы W

Эконометрика, 2017-2018, 1 модуль

Семинары 1 - 2

2.04.18 и 9.04.18 для

Группы Э_Б2015_Э_3

Семинарист О.А.Демидова

Критика М.Фридменом стандартной функции потребления, раздел 8.5.

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2) (Доугерти, раздел 8)

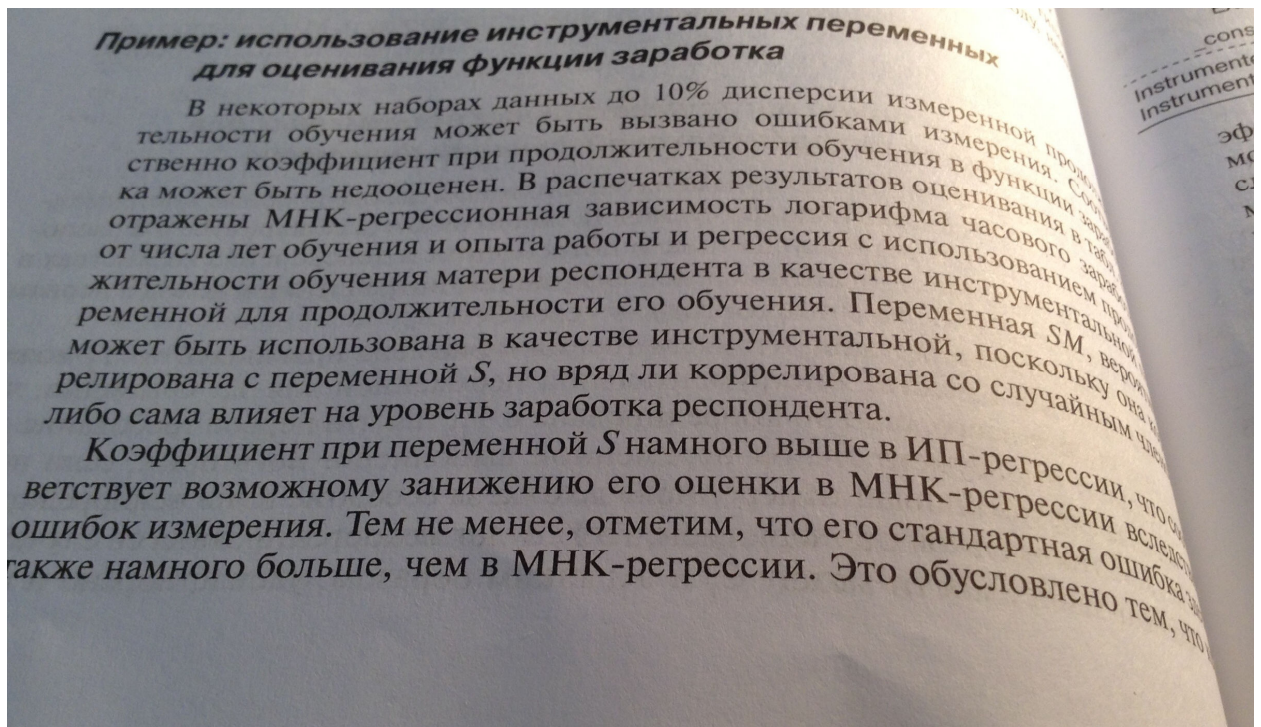


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LGEARN

Coef. Std. Err. t P>|t| [95% Conf. Interval]

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Coef. Std. Err. t P>|t| [95% Conf. Interval]

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EXP .0394422 .0058092 6.79 0.000 .0280306 .0508537

_cons -.0617062 .4061769 -0.15 0.879 -.8595966 .7361841

Instrumented: S

Instruments: EXP SM

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_cons	.2258512	.3887468	0.58	0.562	-.5378125		

Number of obs = 540
F(6,533) = 37.7
Prob > F = 0.000
R-squared = 0.346
Adj R-squared = 0.338
Root MSE = .478

Instrumented: S
Instruments: EXP ASVABC MALE ETHBLACK ETHHISP SM SF SIBLINGS LIBRARY

. estimates store EARNIV
. reg LGEARN S EXP ASVABC MALE ETHBLACK ETHHISP

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ETHHISP	.0536544	.0937966	0.57	0.568	-.1306019		
_cons	.4002952	.1663149	2.41	0.016	.0735821		

Number of obs = 540
F(6,533) = 47.99
Prob > F = 0.000
R-squared = 0.353
Adj R-squared = 0.345
Root MSE = .478

люч «constants»
ен имеет раз-
сл одинако-

Окончание табл. 8.4

estimates store EARNOLS
hausman EARNIV EARNOLS, constant
— Coefficients —

	(b) EARNIV	(B) EARNOLS	(b - B) Difference	sqrt(diag (V_b - V_B)) S.E.
S	.111379	.0883257	.0230533	.0464029
EXP	.0258798	.0227131	.0031667	.0063889
ASVABC	.0092263	.0129274	-.0037011	.0074527
MALE	.2619787	.2652878	-.0033091	.0076842
ETHBLACK	-.0121846	.0077265	-.019911	.0405924
ETHHISP	.0457639	.0536544	-.0078904	.018018
_cons	.2258512	.4002952	-.174444	.3513736

b = consistent under Ho and Ha; obtained from ivreg
B = inconsistent under Ha, efficient under Ho;
obtained from regress

Test: Ho: difference in coefficients not systematic

$$\chi^2(7) = (b - B)'[(V_b - V_B)^{-1}](b - B) = 0.25$$

$$Prob > \chi^2 = 0.9999$$

5.3.2 Medical expenditures with one endogenous regressor

We consider a model with one endogenous regressor, several exogenous regressors, and one or more excluded exogenous variables that serve as the identifying instruments.

The dataset is an extract from the Medical Expenditure Panel Survey (MEPS) of individuals over the age of 65 years, similar to the dataset described in section 3.2.1. The equation to be estimated has the dependent variable `ldrugexp`, the log of total out-of-pocket expenditures on prescribed medications. The regressors are an indicator for whether the individual holds either employer or union-sponsored health insurance (`hi_empunion`), number of chronic conditions (`totchr`), and four sociodemographic variables: age in years (`age`), indicators for whether female (`female`) and whether black or Hispanic (`blhisp`), and the natural logarithm of annual household income in thousands of dollars (`linc`).

We treat the health insurance variable `hi_empunion` as endogenous. The intuitive justification is that having such supplementary insurance on top of the near universal Medicare insurance for the elderly may be a choice variable. Even though most individuals in the sample are no longer working, those who expected high future medical expenses might have been more likely to choose a job when they were working that would provide supplementary health insurance upon retirement. Note that Medicare did not cover drug expenses for the time period we study.

We use the global macro `x2list` to store the names of the variables that are treated as exogenous regressors. We have

```
. * Read data, define global x2list, and summarize data
. use mus06data.dta
. global x2list totchr age female blhisp linc
. summarize ldrugexp hi_empunion $x2list
```

Variable	Obs	Mean	Std. Dev.	Min	Max
ldrugexp	10391	6.479668	1.363395	0	10.18017
hi_empunion	10391	.3796555	.4853245	0	1
totchr	10391	1.860745	1.290131	0	9
age	10391	75.04639	6.69368	65	91
female	10391	.5797325	.4936256	0	1
blhisp	10391	.1703397	.3759491	0	1
linc	10089	2.743275	.9131433	-6.907755	5.744476

6.3.3 Available instruments

We consider four potential instruments for `hi_employment`. Two reflect the income status of the individual and two are based on employer characteristics.

The `ssratio` instrument is the ratio of an individual's social security income to the individual's income from all sources, with high values indicating a significant income constraint. The `lowincome` instrument is a qualitative indicator of low-income status. Both these instruments are likely to be relevant, because they are expected to be negatively correlated with having supplementary insurance. To be valid instruments, we need to assume they can be omitted from the equation for `ldrugexp`, arguing that the direct role of income is adequately captured by the regressor `linc`.

The `firmsz` instrument measures the size of the firm's employed labor force, and the `multlc` instrument indicates whether the firm is a large operator with multiple locations. These variables are intended to capture whether the individual has access to supplementary insurance through the employer. These two variables are irrelevant for those who are retired, self-employed, or purchase insurance privately. In that sense, these two instruments could potentially be weak.

```

* Summarize available instruments
* summarize sratio lowincome multlc firmsz if linc!=.

```

Variable	Obs	Mean	Std. Dev.	Min	Max
<code>ssratio</code>	10089	.5365438	.3678175	0	9.25062
<code>lowincome</code>	10089	.1874319	.3902771	0	1
<code>multlc</code>	10089	.0620478	.2412543	0	1
<code>firmsz</code>	10089	.1405293	2.170389	0	50

We have four available instruments for one endogenous regressor. The obvious approach is to use all available instruments, because in theory this leads to the most efficient estimator. In practice, it may lead to larger small-sample bias because the small-sample biases of IV estimators increase with the number of instruments (Hahn and Hausman 2002).

At a minimum, it is informative to use `correlate` to view the gross correlation between endogenous variables and instruments and between instruments. When multiple instruments are available, as in the case of overidentified models, then it is actually the partial correlation after controlling for other available instruments that matters. This important step is deferred to sections 6.4.2 and 6.4.3.

6.3.4 IV estimation of an exactly identified model

We begin with IV regression of `ldrugexp` on the endogenous regressor `hi_empunio`, instrumented by the single instrument `ssiratio`, and several exogenous regressors.

We use `ivregress` with the `2sls` estimator and the options `vce(robust)` to control for heteroskedastic errors and `first` to provide output that additionally reports results from the first-stage regression. The output is in two parts:

```
. * IV estimation of a just-identified model with single endog regressor
. ivregress 2sls ldrugexp (hi_empunio = ssiratio) $x2list, vce(robust) first
```

First-stage regressions

	Coef.	Robust Std. Err.	t	P> t	[95% Conf. Interval]
hi_empunio					
totchr	.0127865	.0036655	3.49	0.000	.0056015 .0199716
age	-.0086323	.0007087	-12.18	0.000	-.0100216 -.0072431
female	-.07345	.0096392	-7.62	0.000	-.0923448 -.0545552
blhisp	-.06268	.0122742	-5.11	0.000	-.08674 -.0386201
linc	.0483937	.0066075	7.32	0.000	.0354417 .0613456
ssiratio	-.1916432	.0236326	-8.11	0.000	-.2379678 -.1453186
_cons	1.028981	.0581387	17.70	0.000	.9150172 1.142944

Number of obs = 10089
F(6, 10082) = 119.18
Prob > F = 0.000
R-squared = 0.0761
Adj R-squared = 0.0755
Root MSE = 0.4672

Instrumental variables (2SLS) regression

	Coef.	Robust Std. Err.	z	P> z	[95% Conf. Interval]
ldrugexp					
hi_empunio	-.8975913	.2211268	-4.06	0.000	-1.330992 -.4641908
totchr	.4502655	.0101969	44.16	0.000	.43028 .470251
age	-.0132176	.0029977	-4.41	0.000	-.0190931 -.0073421
female	-.020406	.0326114	-0.63	0.531	-.0843232 .0435113
blhisp	-.2174244	.0394944	-5.51	0.000	-.294832 -.1400167
linc	.0870018	.0226356	3.84	0.000	.0426368 .1313668
_cons	6.78717	.2688453	25.25	0.000	6.260243 7.314097

Number of obs = 10089
Wald chi2(6) = 2000.86
Prob > chi2 = 0.000
R-squared = 0.0640
Root MSE = 1.3177

Instrumented: hi_empunio
Instruments: totchr age female blhisp linc ssiratio

6.3.6 Testing for regressor endogeneity

The preceding analysis treats the insurance variable, `hi_empunio`, as endogenous. If instead the variable is exogenous, then the IV estimators (IV, 2SLS, or GMM) are still consistent, but they can be much less efficient than the OLS estimator.

The Hausman test principle provides a way to test whether a regressor is endogenous. If there is little difference between OLS and IV estimators, then there is no need to instrument, and we conclude that the regressor was exogenous. If instead there is considerable difference, then we needed to instrument and the regressor is endogenous. The test usually compares just the coefficients of the endogenous variables. In the case of just one potentially endogenous regressor with a coefficient denoted by β , the Hausman test statistic

$$T_H = \frac{(\hat{\beta}_{IV} - \hat{\beta}_{OLS})^2}{\hat{V}(\hat{\beta}_{IV} - \hat{\beta}_{OLS})}$$

is $\chi^2(1)$ distributed under the null hypothesis that the regressor is exogenous.

Before considering implementation of the test, we first obtain the OLS estimates to compare them with the earlier IV estimates. We have

```
. * Obtain OLS estimates to compare with preceding IV estimates
. regress ldrugexp hi_empunion $x2list, vce(robust) .

Linear regression                               Number of obs =   10089
                                                F(   6, 10082)    876.85
                                                Prob > F          0.0000
                                                R-squared         0.1770
                                                Root MSE         1.236
```

ldrugexp	Coef.	Robust Std. Err.	t	P> t	[95% Conf. Interval]	
hi_empunion	.0738788	.0259948	2.84	0.004	.0229435	.1248141
totchr	.4403807	.0093633	47.03	0.000	.4220268	.4587346
age	-.0035295	.001937	-1.82	0.068	-.0073264	.0002675
female	.0578055	.0253651	2.28	0.023	.0080848	.1075262
blhisp	-.1513068	.0341264	-4.43	0.000	-.2182013	-.0844122
linc	.0104815	.0137126	0.76	0.445	-.0163979	.037361
_cons	5.861131	.1571037	37.31	0.000	5.553176	6.169085

The OLS estimates differ substantially from the just-identified IV estimates given in section 6.3.4. The coefficient of `hi_empunion` has an OLS estimate of 0.074, greatly different from the IV estimate of -0.898 . This is strong evidence that `hi_empunion` is endogenous. Some coefficients of exogenous variables also change, notably, those for `age` and `female`. Note also the loss in precision in using IV. Most notably, the standard error of the instrumented regressor increases from 0.026 for OLS to 0.221 for IV, an eightfold increase, indicating the potential loss in efficiency due to IV estimation.

The `hausman` command can be used to compute T_H under the assumption that $\hat{V}(\hat{\beta}_{IV} - \hat{\beta}_{OLS}) = \hat{V}(\hat{\beta}_{IV}) - \hat{V}(\hat{\beta}_{OLS})$; see section 12.7.5. This greatly simplifies analysis because then all that is needed are coefficient estimates and standard errors from separate IV estimation (IV, 2SLS, or GMM) and OLS estimation. But this assumption is too strong. It is correct only if $\hat{\beta}_{OLS}$ is the fully efficient estimator under the null hypothesis of exogeneity, an assumption that is valid only under the very strong assumption that model errors are independent and homoskedastic. One possible variation is to perform an appropriate bootstrap; see section 13.4.6.

The postestimation `estat endogenous` command implements the related Durbin-Wu-Hausman (DWH) test. Because the DWH test uses the device of augmented regressors, it produces a robust test statistic (Davidson 2000). The essential idea is the following. Consider the model as specified in section 6.2.1. Rewrite the structural equation (6.2) with an additional variable, v_1 , that is the error from the first-stage equation (6.3) for y_2 . Then

$$y_{1i} = \beta_1 y_{2i} + x'_{1i} \beta_2 + \rho v_{1i} + u_i$$

Under the null hypothesis that y_2 is exogenous, $E(v_{1i} u_i | y_{2i}, x_{1i}) = 0$. If v_1 could be observed, then the test of exogeneity would be the test of $H_0: \rho = 0$ in the OLS regression of y_1 on y_2 , x_1 , and v_1 . Because v_1 is not directly observed, the fitted residual vector

\hat{v}_1 from the first-stage OLS regression (6.3) is instead substituted. For independent homoskedastic errors, this test is asymptotically equivalent to the earlier Hausman test. In the more realistic case of heteroskedastic errors, the test of $H_0: \rho = 0$ can still be implemented provided that we use robust variance estimates. This test can be extended to the multiple endogenous regressors case by including multiple residual vectors and testing separately for correlation of each with the error on the structural equation.

We apply the test to our example with one potentially endogenous regressor, `hi_empunio`, instrumented by `ssratio`. Then

```
. * Robust Durbin-Wu-Hausman test of endogeneity implemented by estat endogenous
. ivregress 2sls ldrugexp (hi_empunio = sratio) $x2list, vce(robust)
```

```
Instrumental variables (2SLS) regression                Number of obs =   10089
                                                         Wald chi2(6)      2009.86
                                                         Prob > chi2       0.0000
                                                         R-squared         0.0640
                                                         Root MSE        1.48177
```

ldrugexp	Robust		z	P> z	[95% Conf. Interval]	
	Coef.	Std. Err.				
hi_empunio	-.8975913	.2211268	-4.06	0.000	-1.330982	-.4641908
totchr	.4802655	.0101969	44.16	0.000	.43028	.470251
age	-.0132176	.0029977	-4.41	0.000	-.0190931	-.0073421
female	-.020406	.0326114	-0.63	0.531	-.0843232	.0435113
blhisp	-.2174244	.0394944	-5.51	0.000	-.294832	-.1400167
line	.0870018	.0226356	3.84	0.000	.0426368	.1313668
_cons	6.78717	.2688453	25.25	0.000	6.260243	7.314097

```
Instrumented: hi_empunio
Instruments: totchr age female blhisp line sratio
```

```
. estat endogenous
    Tests of endogeneity
    Ho: variables are exogenous

    Robust score chi2(1)      =   24.935   (p = 0.0000)
    Robust regression F(1,10081) =   26.4333   (p = 0.0000)
```

The last line of output is the robustified DWH test and leads to strong rejection of the null hypothesis that `hi_empunio` is exogenous. We conclude that it is endogenous.

We obtain exactly the same test statistic when we manually perform the robustified DWH test. We have

```
. * Robust Durbin-Wu-Hausman test of endogeneity implemented manually
. quietly regress hi_empunio sratio $x2list
. quietly predict vihat, resid
. quietly regress ldrugexp hi_empunio vihat $x2list, vce(robust)
. test vihat
( 1) vihat = 0
      F( 1, 10081) =   26.43
      Prob > F =   0.0000
```


3.7 Tests of overidentifying restrictions

The validity of an instrument cannot be tested in a just-identified model. But it is possible to test the validity of overidentifying instruments in an overidentified model provided that the parameters of the model are estimated using optimal GMM. The same test has several names, including overidentifying restrictions (OIR) test, overidentified (OID) test, Hansen's test, Sargan's test, and Hansen-Sargan test.

The starting point is the fitted value of the criterion function (6.8) after optimal GMM, i.e., $Q(\hat{\beta}) = \{(1/N)(y - X\hat{\beta})'Z\}\hat{S}^{-1}\{(1/N)Z'(y - X\hat{\beta})\}$. If the population moment conditions $E\{Z'(y - X\beta)\} = 0$ are correct, then $Z'(y - X\hat{\beta}) \simeq 0$, so $Q(\hat{\beta})$ should be close to zero. Under the null hypothesis that all instruments are valid, it can be shown that $Q(\hat{\beta})$ has an asymptotic chi-squared distribution with degrees of freedom equal to the number of overidentifying restrictions.

Large values of $Q(\hat{\beta})$ lead to rejection of $H_0: E\{Z'(y - X\beta)\} = 0$. Rejection is interpreted as indicating that at least one of the instruments is not valid. Tests can have power in other directions, however, as emphasized in section 3.5.5. It is possible that rejection of H_0 indicates that the model $X\beta$ for the conditional mean is misspecified. Going the other way, the test is only one of validity of the overidentifying instruments, so failure to reject H_0 does not guarantee that all the instruments are valid.

The test is implemented with the postestimation `estat overid` command following the `ivregress gmm` command for an overidentified model. We do so for the optimal GMM estimator with heteroskedastic errors and instruments, `ssratio` and `multic`. The example below implements `estat overid` under the overidentifying restriction.

```
. * Test of overidentifying restrictions following ivregress gmm
. quietly ivregress gmm ldrugexp (hi_empunio = sratio multic)
> $x2list, wmatrix(robust)

. estat overid
    Test of overidentifying restriction:
    Hansen's J chi2(1) = 1.04754 (p = 0.3061)
```

The test statistic is $\chi^2(1)$ distributed because the number of overidentifying restrictions equals $2 - 1 = 1$. Because $p > 0.05$, we do not reject the null hypothesis and conclude that the overidentifying restriction is valid.

A similar test using all four available instruments yields

```
. * Test of overidentifying restrictions following ivregress gmm
. ivregress gmm ldrugexp (hi_empunion = ssiratio lowincome multlc firmsz)
> $x2list, wmatrix(robust)
```

Instrumental variables (GMM) regression		Number of obs = 10089	
		Wald chi2(6) = 2042.12	
		Prob > chi2 = 0.0000	
		R-squared = 0.0829	
GMM weight matrix: Robust		Root MSE = 1.3043	

ldrugexp	Coef.	Robust Std. Err.	z	P> z	[95% Conf. Interval]	
hi_empunion	-.8124043	.1846433	-4.40	0.000	-1.174299	-.45051
totchr	.449488	.010047	44.74	0.000	.4297962	.4691799
age	-.0124598	.0027466	-4.54	0.000	-.0178432	-.0070765
female	-.0104528	.0306889	-0.34	0.733	-.0706019	.0496963
blhisp	-.2061018	.0382891	-5.38	0.000	-.2811471	-.1310566
linc	.0796532	.0203397	3.92	0.000	.0397882	.1195183
_cons	6.7126	.2425973	27.67	0.000	6.237118	7.188081

```
Instrumented: hi_empunion
Instruments: totchr age female blhisp linc ssiratio lowincome multlc firmsz

. estat overid

Test of overidentifying restriction:
Hansen's J chi2(3) = 11.5903 (p = 0.0089)
```

Now we reject the null hypothesis at level 0.05 and, barely, at level 0.01. Despite this rejection, the coefficient of the endogenous regressor `hi_empunion` is -0.812 , not all that different from the estimate when `ssiratio` is the only instrument.

4) (Демешев, Борзых, 18.1)

Величины X_i равномерны на отрезке $[-a; 3a]$ и независимы. Есть несколько наблюдений, $X_1 = 0.5$, $X_2 = 0.7$, $X_3 = -0.1$.

1. Найдите $\mathbb{E}(X_i)$ и $\mathbb{E}(|X_i|)$.
2. Постройте оценку метода моментов, используя $\mathbb{E}(X_i)$.
3. Постройте оценку метода моментов, используя $\mathbb{E}(|X_i|)$.
4. Постройте оценку обобщённого метода моментов используя моменты $\mathbb{E}(X_i)$, $\mathbb{E}(|X_i|)$ и взвешивающую матрицу.

$$W = \begin{pmatrix} 2 & 0 \\ 0 & 1 \end{pmatrix}$$

5. Найдите оптимальную теоретическую взвешивающую матрицу для обобщённого метода моментов
6. Постройте двухшаговую оценку обобщённого метода моментов, начав со взвешивающей матрицы W