

Fitting fixed and random effects meta-analysis models using structural equation models

Tom M. Palmer Jonathan A. C. Sterne

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Mathematics
& Statistics

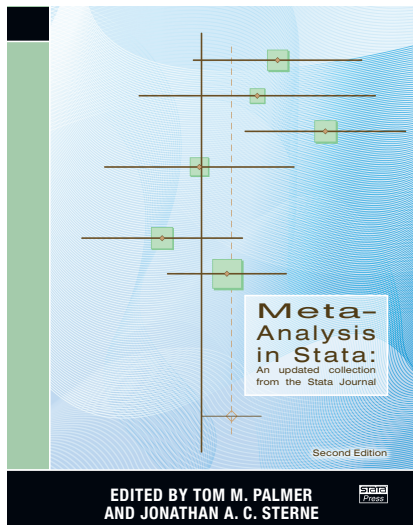
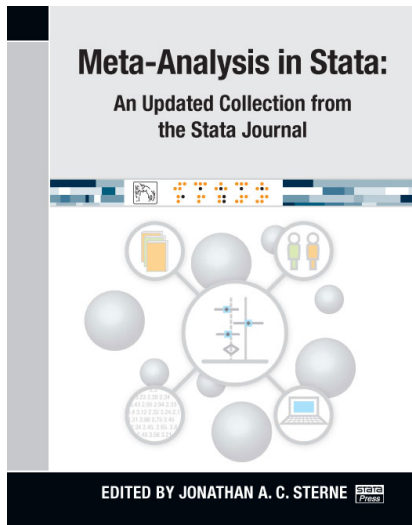
Lancaster
University



- ▶ Introduction
- 1. Univariate fixed effect meta-analysis
- 2. Univariate random effects meta-analysis
- 3. Multivariate meta-analysis with non-zero within study covariances
- ▶ Summary

Introduction I

- ▶ This talk focuses on the use of Stata and follows Palmer & Sterne (Stata Journal, forthcoming)



- ▶ 27 Stata Journal articles, 11 new since 1st ed. (3 forthcoming)

Introduction II

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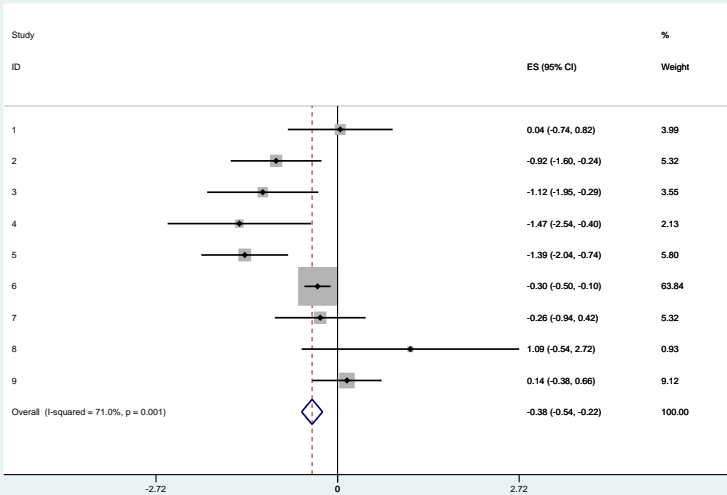
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- ▶ User specifies constraints on variances of the error terms of the outcome (i.e. variance of the residuals).
- ▶ Using SEM for meta-analysis well developed in Psychology
- ▶ Discussed in articles (and a new book) by Cheung (2008, 2010, 2013a, 2013b, 2013c, 2015)
- ▶ `metaSEM` package in R (automates use of `OpenMx`) also by Cheung

1. Univariate outcome meta-analysis models: fixed effect

$$y_i \sim N(\theta, \sigma_i^2)$$

i.e. y_i and σ_i^2 estimated in each study.

- ▶ Example, Turner et al. (2000)
- ▶ 9 trials investigating effect of taking diuretics during pregnancy on risk of pre-eclampsia
- ▶ log odds ratios for association between pre-eclampsia and diuretics from each study and SE



Pooled OR: 0.68 (95% CI 0.58, 0.80) – lower risk of pre-eclampsia for diuretic group

Syntax 1

To fit the model in sem we generate a weighting variable of inverse variances:

- ▶ $Y \sim N(X\theta, \sigma^2 W^{-1})$
- ▶ WLS estimate:
$$\hat{\theta} = (X'WX)^{-1}X'WY = (\sum_{i=1}^N w_i y_i) / (\sum_{i=1}^N w_i)$$
- ▶ Variance of WLS estimate = $\sigma^2 (X'WX)^{-1}$
- ▶ But we require the pooled variance to be:
$$1 / \sum_{i=1}^N w_i = (X'WX)^{-1}$$
- ▶ Hence we constrain $\sigma^2 = 1$.

Syntax 1

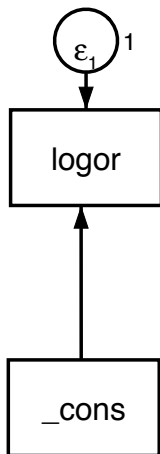
```
. gen double weight = 1/varlogor  
. sem (logor <- ) [iw=weight], var(e.logor@1) nodescribe nocnsreport nolog
```

```
Structural equation model           Number of obs   =           9  
Estimation method   = ml  
Log likelihood      = -157.71614
```

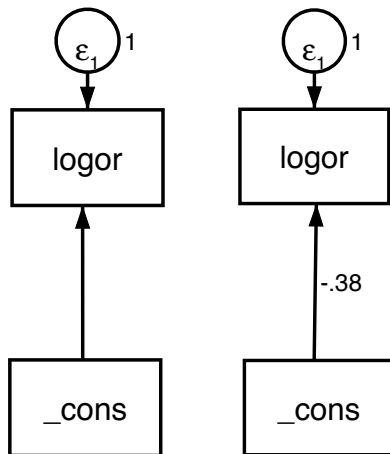
```
-----+-----  
                |               OIM  
                |      Coef.   Std. Err.   z   P>|z|   [95% Conf. Interval]  
-----+-----  
Structural      |  
  logor <-     |  
    _cons      |  -.3815467   .0799025   -4.78   0.000   -.5381527   -.2249406  
-----+-----  
var(e.logor)   |           1 (constrained)  
-----+-----
```

```
LR test of model vs. saturated: chi2(1)   =   143.07, Prob > chi2 = 0.0000
```

Stata SEM builder path diagrams I



Stata SEM builder path diagrams I



After fitting the estimated coefficient is shown

Syntax 2

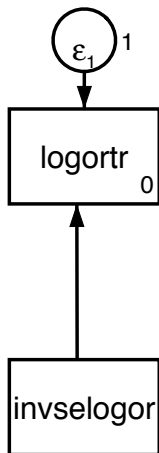
- ▶ Fit the same model by scaling all the variables by 1/SEs
- ▶ Scale the vector of 1's for the **intercept**
- ▶ Constrain $\sigma^2 = 1$

```
. gen double invselogor = 1/selogor
. gen double logortr = logor*invselogor
. sem (logortr <- invselogor, nocons), noheader nodeline nocnsreport nolog var(e.logortr@1)
```

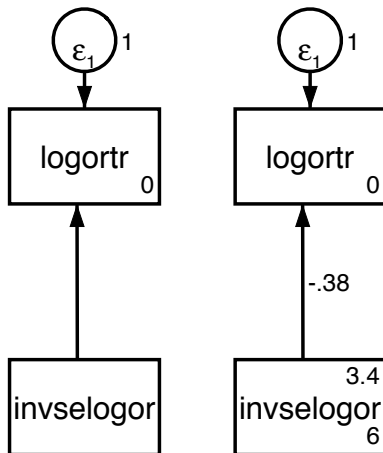
```
-----
                |                OIM
                |                Coef.   Std. Err.   z   P>|z|   [95% Conf. Interval]
-----+-----
Structural      |
logortr <-      |
invselogor     |   -.3815467   .0799025   -4.78   0.000   -.5381527   -.2249406
  _cons         |                0   (constrained)
-----+-----
var(e.logortr) |                1   (constrained)
-----
```

```
LR test of model vs. saturated: chi2(2)   =   9.10, Prob > chi2 = 0.0106
```

Stata SEM builder path diagrams II



Stata SEM builder path diagrams II



After fitting the estimated coefficient is shown, along with mean and variance of covariate (scaled intercept)

Heterogeneity test

- ▶ Remove constraint from σ^2 .

```
. sem (logortr <- invselogor, nocons), noheader nodescribe nocnsreport nolog
-----+-----
                |               OIM
                |      Coef.   Std. Err.   z   P>|z|   [95% Conf. Interval]
-----+-----
Structural      |
logortr <-      |
  invselogor    | -0.3815467   .1398305   -2.73   0.006   -0.6556094   -0.107484
  _cons         |          0 (constrained)
-----+-----
var(e.logortr) |  3.062546   1.443698                1.215689   7.715122
-----+-----
LR test of model vs. saturated: chi2(1)   =      0.61, Prob > chi2 = 0.4344
```

- ▶ $Q = \widehat{\sigma^2} N = 3.016 \times 9 = 27.56$, $P = 0.00056$
- ▶ $I^2 = \frac{Q - df}{Q} = (27.56 - 8) / 27.56 = 0.71$

2. Univariate outcome random effects meta-analysis

$$y_i \sim N(\theta + \nu_i, \sigma_i^2)$$
$$\nu_i \sim N(0, \tau^2)$$

- ▶ Syntax 1: 9 studies – 9 random effects
- ▶ Syntax 2: interact 1 random effect with standard errors (untransformed variables)
- ▶ Syntax 3: interact 1 random effect with the inverse standard error transformed variables
- ▶ Same example meta-analysis
- ▶ metan RE DL pooled log OR: -0.516 (95% CI -0.908, -0.124)
- ▶ $Q = 27.56$ ($p=0.001$), $I^2 = 71\%$, $\tau^2 = 0.2185$

Syntax 3 – use 1/SE transformed variables

- ▶ Constrain coefficient of interaction of inverse SEs and RE to 1.
- ▶ Constrain variance of residuals to 1.
- ▶ Variance of RE, $\text{var}(M)$, is estimate of τ^2 .

```
. gsem (logotr <- invselogor c.invselogor#c.M@1, nocons), ///  
> var(e.logotr@1) latent(M) nolog nocnsreport
```

```
Generalized structural equation model          Number of obs   =           9  
Log likelihood =  -18.8726
```

	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
logotr <-						
invselogor	-.5166151	.2059448	-2.51	0.012	-.9202594	-.1129708
c.invselogor#c.M	1	(constrained)				
_cons	0	(omitted)				
var(M)	.2377469	.1950926			.0476023	1.187413
var(e.logotr)	1	(constrained)				

Syntax 3 – use 1/SE transformed variables

- ▶ Constrain coefficient of interaction of inverse SEs and RE to 1.
- ▶ Constrain variance of residuals to 1.
- ▶ Variance of RE, $\text{var}(M)$, is estimate of τ^2 .

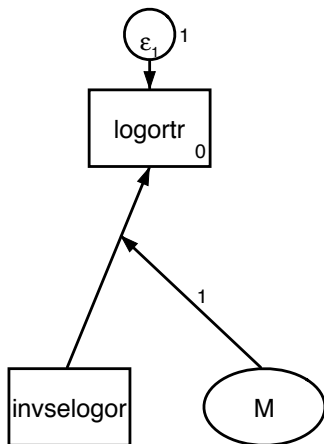
```
. gsem (logotr <- invselogor c.invselogor#c.M@1, nocons), ///  
> var(e.logotr@1) latent(M) nolog nocnsreport
```

```
Generalized structural equation model          Number of obs   =           9  
Log likelihood =  -18.8726
```

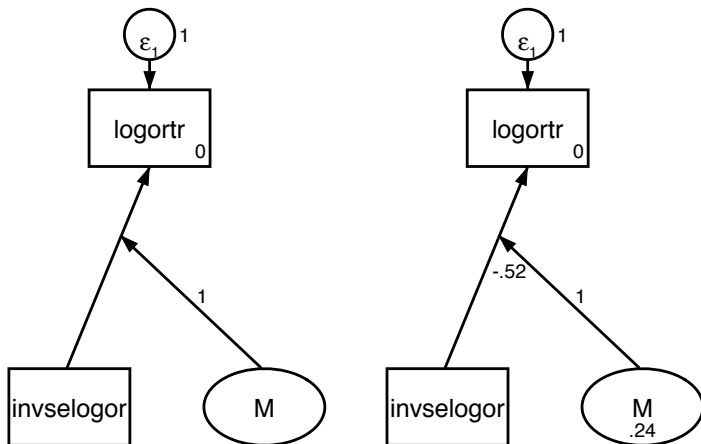
	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
logotr <-						
invselogor	-.5166151	.2059448	-2.51	0.012	-.9202594	-.1129708
c.invselogor#c.M	1	(constrained)				
_cons	0	(omitted)				
var(M)	.2377469	.1950926			.0476023	1.187413
var(e.logotr)	1	(constrained)				

- ▶ gsem can't do REML estimation of τ^2 (metaSEM in R can).
- ▶ can derive a prediction interval for pooled estimate

Stata SEM builder syntax 3 path diagram



Stata SEM builder syntax 3 path diagram



3.1 Multivariate fixed effect meta-analysis with non-zero within study covariances

$$\blacktriangleright Y = \begin{bmatrix} y_{11} \\ y_{12} \\ \dots \\ y_{N1} \\ y_{N2} \end{bmatrix}, \theta = \begin{bmatrix} \theta_1 \\ \theta_2 \end{bmatrix}, V_i = \begin{bmatrix} \sigma_{i,11}^2 & \sigma_{i,12} \\ \sigma_{i,12} & \sigma_{i,22}^2 \end{bmatrix}$$

$$\blacktriangleright \Sigma = \begin{bmatrix} V_1 & 0 & 0 \\ 0 & \dots & 0 \\ 0 & 0 & V_N \end{bmatrix} \quad Y \sim \text{MVN}(\theta, \Sigma)$$

- \blacktriangleright Transformation multivariate equivalent of 1/SE scaling – Cholesky decomposition of inverse of within study covariance matrix, i.e. $W_i^{1/2} = V_i^{-1/2}$
- \blacktriangleright $W^{1/2}Y \sim \text{MVN}(W^{1/2}X\theta, W^{1/2}\Sigma(W^{1/2})')$
- \blacktriangleright Fibrinogen Studies Collaboration (2004): incidence of CHD (log hazard ratio), 31 studies using 2 outcomes

Multivariate fixed effect meta-analysis with non-zero within study covariances

```
. use FSCstage1, clear

. * code to generate transformed outcome and outcome indicator variables

. sem (ystarstack <- xstarstack1 xstarstack2, nocons), ///
>     var(e.ystarstack@1) nocapslatent nolog nocnsr nodescribe

Structural equation model           Number of obs       =           62
Estimation method   = ml
Log likelihood      = -384.49772
```

```
-----+-----
                |              OIM
                |      Coef.   Std. Err.   z   P>|z|   [95% Conf. Interval]
-----+-----
Structural      |
  ystarstack <- |
    xstarstack1 |   .2042387   .0529888    3.85  0.000   .1003826   .3080947
    xstarstack2 |   .8639001   .0536208   16.11  0.000   .7588052   .968995
      _cons      |           0 (constrained)
-----+-----
var(e.ystarstack) |           1 (constrained)
-----+-----
LR test of model vs. saturated: chi2(2)   =       15.87, Prob > chi2 = 0.0004
```

Heterogeneity test for both outcomes jointly

► Again remove constraint from variance of residuals

```
. quietly sem (ystarstack <- xstarstack1 xstarstack2, nocons), nocapslatent

. di "var(e.ystarstack) = " _b[var(e.ystarstack):_cons]
var(e.ystarstack) = 1.8483607

. local Q = _b[var(e.ystarstack):_cons]*e(N)

. local df = e(N) - 2

. di "Het. test statistic = " 'Q'
Het. test statistic = 114.59836

. di "Het. test p-value = " chi2tail('df', 'Q')
Het. test p-value = .00002803
```

Decompose the heterogeneity test for each outcome

- ▶ reshape data to wide format
- ▶ Specify model using 2 equations – 1 for each outcome; each has a residual variance

```
qui sem (ystarstack1 <- xstarstack11 xstarstack21@c1) ///  
>         (ystarstack2 <- xstarstack22@c1), nocons ///  
>         nocaps nolog nocnsr nodescribe
```

Outcome	Approach	Q	P	I ² (95% CI)
1	Multivariate	48.12	P=0.019 (sem)	18 (mvmeta)
1	Univariate	36.74	P=0.185	18 (0, 48)
2	Multivariate	66.50	P<0.0001 (sem)	55 (mvmeta)
2	Univariate	66.19	P<0.0001	55 (32, 70)

3.2 Random effects multivariate meta-analysis with non-zero within study covariance

▶ $Y \sim \text{MVN}(\theta + \nu, \Sigma)$

▶ $\nu \sim N(\mathbf{0}, \mathbf{T}^2)$, for a 2 outcome model $\mathbf{T}^2 = \begin{bmatrix} \tau_1^2 & \tau_{12} \\ \tau_{12} & \tau_2^2 \end{bmatrix}$

▶ long format data – specify **study** level random effects

```
. gsem (ystarstack <- c.xstarstack1#c.M1[study]@1 c.xstarstack2#c.M2[study]@1 ///
> xstarstack1 xstarstack2, nocons), ///
> latent(M1 M2) nocnsreport nolog ///
> cov(e.ystarstack@1 (M1[study]*M2[study]))
```

Generalized structural equation model Number of obs = 62
Log likelihood = -101.66433

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

ystarstack <-						
xstarstack1	.1875603	.0690866	2.71	0.007	.0521531	.3229675
xstarstack2	.8585811	.0887304	9.68	0.000	.6846728	1.032489
...						
var(M1[study])	.0221546	.0324089			.0012597	.3896245
var(M2[study])	.0945799	.0614174			.0264883	.3377098
cov(M2[study],M1[study])	.0272542	.0382754	0.71	0.476	-.0477642	.1022726
var(e.ystarstack)		1	(constrained)			

Equivalent model for wide format data (2 equations).

```
. gsem (ystarstack1 <- c.xstarstack1#c.M1@1 c.xstarstack21#c.M2@1 ///
>         xstarstack11 xstarstack21@c1, nocons) ///
>         (ystarstack2 <- c.xstarstack22#c.M2@1 xstarstack22@c1, nocons), ///
>         cov(e.ystarstack1@1 e.ystarstack2@1) latent(M1 M2) ///
>         collinear nocnsreport nolog
```

Generalized structural equation model Number of obs = 31
 Log likelihood = -101.66433

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

ystarstack1 <-					
xstarstack11	.1875603	.0690866	2.71	0.007	.0521531 .3229675
xstarstack21	.8585811	.0887304	9.68	0.000	.6846728 1.032489
...					

ystarstack2 <-					
xstarstack22	.8585811	.0887304	9.68	0.000	.6846728 1.032489
...					

var(M1)	.0221546	.0324089			.0012597 .3896245
var(M2)	.0945799	.0614174			.0264883 .3377098

cov(M2,M1)	.0272542	.0382754	0.71	0.476	-.0477642 .1022726

var(e.ystarstack1)	1	(constrained)			
var(e.ystarstack2)	1	(constrained)			

```
. di "corr(M1,M2)=", _b[cov(M2,M1):_cons]/sqrt(_b[var(M1):_cons]*_b[var(M2):_cons])
```

corr(M1,M2)= .59539071

Summary

- ▶ Can fit these models using `metan`; `metareg`; `mvmeta` (White, 2009, 2011)
- ▶ Fixed effect meta-analysis – 2 syntaxes
- ▶ Random effect meta-analysis – 3 syntaxes
- ▶ Meta-regression – FE and RE
- ▶ Multivariate outcome – FE and RE – with zero and non-zero within study covariances
- ▶ (and by extension) Multivariate meta-regression
- ▶ For RE models `gsem` cannot perform REML estimation – `metaSEM` in R can.
- ▶ Cochran heterogeneity test after FE models (joint test and test for each multivariate outcome)

References

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Acknowledgements

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- ▶ Mike Cheung (National University of Singapore)
- ▶ Rebecca Pope, Stephanie White, and the development team of the `gsem` command at StataCorp
- ▶ Medical and Pharmaceutical Statistics (MPS) Research Unit, Lancaster University

- ▶ Constrain coefficients of study and RE interactions to 1.
- ▶ Constrain the studies to be independent with variance as estimated in each study.
- ▶ Variance of residuals $\text{var}(e.\text{logor})$ is estimate of τ^2

Syntax 1: 9 random effects

```
. mkmat varlogor, mat(f)

. mat f = diag(f)

. qui tabulate trial, gen(tr)

. gsem (logor <- M1#c.tr1@1 M2#c.tr2@1 M3#c.tr3@1 ///
>       M4#c.tr4@1 M5#c.tr5@1 M6#c.tr6@1 ///
>       M7#c.tr7@1 M8#c.tr8@1 M9#c.tr9@1) ///
>       , covstructure(_LEx, fixed(f)) intmethod(laplace) nocnsreport nolog
```

```
Generalized structural equation model          Number of obs   =           9
Log likelihood = -9.4552759
```

	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
logor <-						
c.tr1#c.M1	1	(constrained)				
...						
_cons	-.5166151	.2059448	-2.51	0.012	-.9202594	-.1129707
var(M1)	.16	(constrained)				
var(M2)	.12	(constrained)				
var(M3)	.18	(constrained)				
var(M4)	.3	(constrained)				
var(M5)	.11	(constrained)				
var(M6)	.01	(constrained)				
var(M7)	.12	(constrained)				
var(M8)	.69	(constrained)				
var(M9)	.07	(constrained)				
var(e.logor)	.2377469	.1950926			.0476023	1.187413

► Can derive 95% prediction interval for pooled effect

```
. local settotal = sqrt(_se[logor:_cons]^2 + _b[var(e.logor):_cons])  
. local pilow = _b[logor:_cons] - invt(e(N) - 2, .975)*'settotal'  
. local piupp = _b[logor:_cons] + invt(e(N) - 2, .975)*'settotal'  
. di "95% Prediction interval:", 'pilow', 'piupp'  
95% Prediction interval: -1.7682144 .73498424
```

Syntax 2: 1 random effect interacted with SEs

- ▶ Constrain interaction coefficients to 1.
- ▶ Constrain variance of REs to 1.
- ▶ Variance of residuals $\text{var}(e.\text{logor})$ is estimate of τ^2 .

```
. gsem (logor <- ibn.trial#c.selogor#c.M01), var(M01) nolog nocnsreport
```

```
Generalized structural equation model          Number of obs   =           9  
Log likelihood = -9.4552759
```

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

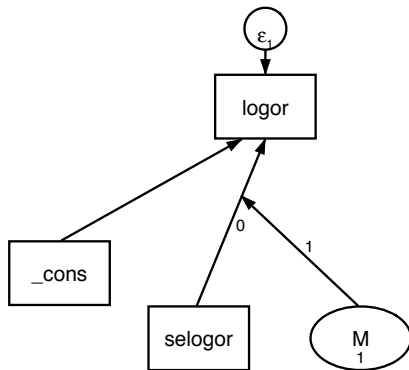
logor <-						
trial#c.selogor#c.M						
1	1	(constrained)				
2	1	(constrained)				
3	1	(constrained)				
4	1	(constrained)				
5	1	(constrained)				
6	1	(constrained)				
7	1	(constrained)				
8	1	(constrained)				
9	1	(constrained)				

_cons	-.5166151	.2059448	-2.51	0.012	-.9202594	-.1129708

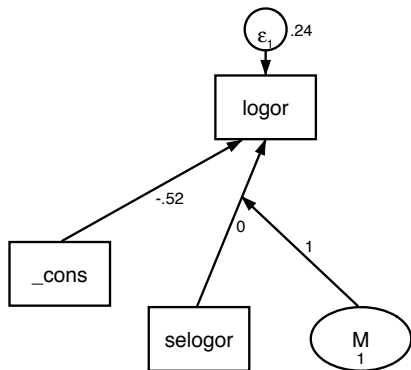
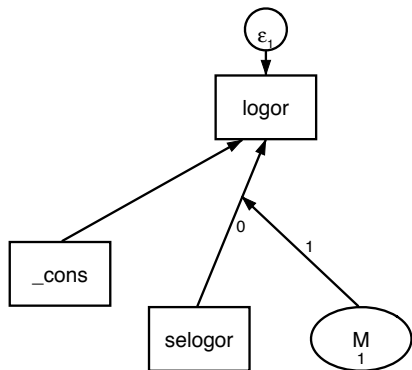
var(M)	1	(constrained)				

var(e.logor)	.2377469	.1950926			.0476023	1.187413

Stata SEM builder random effects syntax 2 path diagram



Stata SEM builder random effects syntax 2 path diagram



2.1 Fixed effect meta-regression

- ▶ $y_i \sim N(X_i\theta, \sigma_i^2)$
- ▶ Not recommend – assumes het. explained by covariates
- ▶ Tends to give too small SEs with moderate/large heterogeneity
- ▶ We need to fit it to obtain heterogeneity test
- ▶ Example data (Thompson & Sharp 1999) 28 RCTs of cholesterol lowering interventions for reducing risk of IHD.
- ▶ Each study reports log odds ratio and its SE, and a variable summarising the cholesterol reduction in each trial.

Fixed effect meta-regression

```
. use cholesterol, clear
(Serum cholesterol reduction & IHD)

. gen double invselogor = 1/sqrt(varlogor)

. gen double logortr = logor*invselogor

. gen double cholreductr = cholreduc*invselogor

. sem (logortr <- cholreductr invselogor, nocons), ///
>      noddescribe nolog nocnsreport var(e.logortr@1)
```

```
Structural equation model          Number of obs    =          28
Estimation method = ml
Log likelihood      = -165.21497
```

```
-----+-----
          |              OIM
          |      Coef.   Std. Err.   z   P>|z|   [95% Conf. Interval]
-----+-----
Structural |
logortr <- |
cholreductr |  -.4752451   .1382083   -3.44   0.001   -.7461284   -.2043617
invselogor |   .1207613   .0972033    1.24   0.214   -.0697538    .3112763
   _cons |              0 (constrained)
-----+-----
var(e.logortr)|              1 (constrained)
-----+-----
```

```
LR test of model vs. saturated: chi2(2) = 1.42, Prob > chi2 = 0.4907
```

Heterogeneity test for meta-regression

► Remove constraint from variance of residuals

```
. quietly sem (logortr <- cholreductr invselogor, nocons), ///  
>           nodeline nolog nocnsreport  
  
. local Q = _b[var(e.logortr):_cons]*e(N)  
  
. local df = e(N) - 2  
  
. di "Het. test statistic = " 'Q'  
Het. test statistic = 37.866258  
  
. di "Het. test p-value = " chi2tail('df', 'Q')  
Het. test p-value = .06231403
```

3.1 Fixed effect multivariate MA with zero within study covariances

$$\blacktriangleright Y = \begin{bmatrix} y_{11} \\ y_{12} \\ \dots \\ y_{N1} \\ y_{N2} \end{bmatrix}, \theta = \begin{bmatrix} \theta_1 \\ \theta_2 \end{bmatrix}, V_i = \begin{bmatrix} \sigma_{i,11}^2 & 0 \\ 0 & \sigma_{i,22}^2 \end{bmatrix}$$

$$\blacktriangleright \Sigma = \begin{bmatrix} V_1 & 0 & 0 \\ 0 & \dots & 0 \\ 0 & 0 & V_N \end{bmatrix}$$

$$\blacktriangleright Y \sim \text{MVN}(\theta, \Sigma)$$

- ▶ Example meta-analysis (Riley et al. 2007) 10 studies, diagnostic accuracy of tumour marker for bladder cancer, each report logit of sensitivity and specificity

```
. use telomerase, clear
(Riley's telomerase data)
```

```
. reshape long y s, i(study) j(outcome)
(note: j = 1 2)
```

Data	wide	->	long
Number of obs.	10	->	20
Number of variables	5	->	4
j variable (2 values)		->	outcome
xij variables:			
	y1 y2	->	y
	s1 s2	->	s

```
. gen byte y2cons = (outcome == 2)
```

```
. gen double invse = 1/s
```

```
. gen double ytr = y*invse
```

```
. gen double y2constr = y2cons*invse
```

```
. sem (ytr <- y2constr invse, nocons), nocaps nodelscribe nolog nocnsr var(e.ytr@1)
```

```
Structural equation model          Number of obs    =          20
Estimation method = ml
Log likelihood      = -116.12748
```

	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
Structural						
ytr <-						
y2constr	.0834338	.2104572	0.40	0.692	-.3290547	.4959223
invse	1.126318	.1177527	9.57	0.000	.8955267	1.357109
_cons	0	(constrained)				
var(e.ytr)	1	(constrained)				

```
LR test of model vs. saturated: chi2(2) = 47.82, Prob > chi2 = 0.0000
```

```
. lincom [ytr]invse + [ytr]y2constr
```

```
( 1) [ytr]y2constr + [ytr]invse = 0
```

	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
(1)	1.209751	.174432	6.94	0.000	.867871	1.551632

- ▶ Remove constraint from variance of residuals

```
. quietly sem (ytr <- y2constr invse, nocons), nocaps nodeline nolog nocnsr
. local Q = _b[var(e.ytr):_cons]*e(N)
. local df = e(N) - 2
. di "Het. test statistic = " 'Q'
Het. test statistic = 90.865377
. di "Het. test p-value = " chi2tail('df', 'Q')
Het. test p-value = 1.009e-11
```

3.2 Random effects multivariate outcomes with zero within study covariances

▶ $Y \sim \text{MVN}(\boldsymbol{\theta} + \boldsymbol{\nu}, \Sigma)$

▶ $\boldsymbol{\nu} \sim N(\mathbf{0}, \mathbf{T}^2)$, for a 2 outcome model $\mathbf{T}^2 = \begin{bmatrix} \tau_1^2 & \tau_{12} \\ \tau_{12} & \tau_2^2 \end{bmatrix}$

```
. use telomerase, clear
. gen double y1tr = y1/s1
. gen double invs1 = 1/s1
. gen double y2tr = y2/s2
. gen double invs2 = 1/s2
```



```

. gsem (y1tr <- c.invs1#c.M1@1 invs1, nocons) ///
> (y2tr <- c.invs2#c.M2@1 invs2, nocons), ///
> cov(e.y1tr@1 e.y2tr@1 e.y1tr*e.y2tr@0) ///
> latent(M1 M2) nolog nocnsreport

```

Generalized structural equation model Number of obs = 10
Log likelihood = -37.273657

	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]	
y1tr <-						
invs1	1.158561	.1616837	7.17	0.000	.8416669	1.475455
...						
y2tr <-						
invs2	2.00511	.4581216	4.38	0.000	1.107208	2.903012
...						
var(M1)	.1179669	.0000813			.1178077	.1181264
var(M2)	1.628624	.0018461			1.62501	1.632246
cov(M2,M1)	-.4383192	.0001342	-3265.57	0.000	-.4385823	-.4380561
var(e.y1tr)	1	(constrained)				
var(e.y2tr)	1	(constrained)				