


```

+ "Psychological_factor", "Pain_intensity"), c("Movement",
+
+ "Psychological_factor", "Pain_intensity"))), `Grotle2004B` =
structure(c(1,-0.11,-0.23,-0.11,1,0.21,-0.23,0.21,1), .Dim = c(3L, 3L),
.Dimnames = list(c("Movement",
+
+ "Psychological_factor", "Pain_intensity"), c("Movement",
+
+ "Psychological_factor", "Pain_intensity"))), `Kim2017` =
structure(c(1,-0.03,-0.02,-0.03,1,0.25,-0.02,0.25,1), .Dim = c(3L, 3L),
.Dimnames = list(c("Movement",
+
+ "Psychological_factor", "Pain_intensity"), c("Movement",
+
+ "Psychological_factor", "Pain_intensity"))), .Names = c("Trost2012",
"Thomas2008A", "Thomas2008B", "Thomas2008C", "Watson1997A",
"Watson1997B", "Alschuler2009", "Grotle2004A", "Grotle2004B",
"Kim2017")), n = c(51, 36, 36, 36, 36, 36, 76, 123, 233, 30),
Questionnaire = c("PASS", "PASS", "TSK", "PCS", "FABQ", "self_efficacy",
"TSK", "FABQ", "FABQ", "FABQ")), .Names = c("data", "n",
+
+ "Questionnaire"))
>

```

```
> ## show the studies ##
```

```
>
> head(Studiestest10$data)
$`Trost2012`
```

	Movement	Psychological_factor	Pain_intensity
Movement	1.00	-0.38	-0.11
Psychological_factor	-0.38	1.00	0.14
Pain_intensity	-0.11	0.14	1.00

```
$Thomas2008A
```

	Movement	Psychological_factor	Pain_intensity
Movement	1.00	-0.54	-0.24
Psychological_factor	-0.54	1.00	0.28
Pain_intensity	-0.24	0.28	1.00

```
$Thomas2008B
```

	Movement	Psychological_factor	Pain_intensity
Movement	1.00	-0.41	-0.24
Psychological_factor	-0.41	1.00	0.28
Pain_intensity	-0.24	0.28	1.00

```
$Thomas2008C
```

	Movement	Psychological_factor	Pain_intensity
Movement	1.00	-0.35	-0.24
Psychological_factor	-0.35	1.00	0.28
Pain_intensity	-0.24	0.28	1.00

```
$Watson1997A
```

	Movement	Psychological_factor	Pain_intensity
Movement	1.00	-0.28	-0.20
Psychological_factor	-0.28	1.00	0.28
Pain_intensity	-0.20	0.28	1.00

```
$Watson1997B
```

	Movement	Psychological_factor	Pain_intensity
--	----------	----------------------	----------------

```

Movement                1.00                -0.27                -0.20
Psychological_factor    -0.27                1.00                0.28
Pain_intensity          -0.20                0.28                1.00

>
> ## Display the sample sizes
> Studiestest10$n
[1] 51 36 36 36 36 36 76 123 233 30
>
> ## Variables used in the analysis
> var.names <- c("Mov", "Psy", "Pai")

> ### Stage 1 from Hagger (give the same results as version from Cheung)
> random1 <- tssem1(Studiestest10$data, Studiestest10$n, method="REM",
RE.type="Diag", acov="weighted")
> summary(random1)

```

Call:

```

meta(y = ES, v = acovR, RE.constraints = Diag(paste0(RE.startvalues,
"*Tau2_", 1:no.es, "_", 1:no.es)), RE.lbound = RE.lbound,
I2 = I2, model.name = model.name, suppressWarnings = TRUE,
silent = silent, run = run)

```

95% confidence intervals: z statistic approximation

Coefficients:

	Estimate	Std.Error	lbound	ubound	z value
Pr(> z)					
Intercept1	-2.6959e-01	5.5461e-02	-3.7830e-01	-1.6089e-01	-4.8610
	1.168e-06	***			
Intercept2	-1.9026e-01	1.7026e-02	-2.2363e-01	-1.5689e-01	-11.1744 <
	2.2e-16	***			
Intercept3	2.1824e-01	3.4400e-02	1.5082e-01	2.8566e-01	6.3442
	2.236e-10	***			
Tau2_1_1	1.3085e-02	1.0824e-02	-8.1289e-03	3.4299e-02	1.2089
	0.2267				
Tau2_2_2	1.0000e-10	NA	NA	NA	NA
	NA				
Tau2_3_3	1.0000e-10	NA	NA	NA	NA
	NA				

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Q statistic on the homogeneity of effect sizes: 25.07557

Degrees of freedom of the Q statistic: 27

P value of the Q statistic: 0.570227

Heterogeneity indices (based on the estimated Tau2):

	Estimate
Intercept1: I2 (Q statistic)	0.4793
Intercept2: I2 (Q statistic)	0.0000
Intercept3: I2 (Q statistic)	0.0000

Number of studies (or clusters): 10

Number of observed statistics: 30

Number of estimated parameters: 6

Degrees of freedom: 24

-2 log likelihood: -47.77288

OpenMx status1: 5 ("0" or "1": The optimization is considered fine.

```

Other values may indicate problems.)
>
> ## Extract the fixed-effects estimates
> (est_fixed <- coef(random1, select="fixed"))
Intercept1 Intercept2 Intercept3
-0.2695942 -0.1902597  0.2182395
>
> ## Convert the estimated vector to a symmetrical matrix
> ## where the diagonals are fixed at 1 (for a correlation matrix)
> vec2symMat(est_fixed, diag=FALSE)
      [,1]      [,2]      [,3]
[1,] 1.0000000 -0.2695942 -0.1902597
[2,] -0.2695942 1.0000000  0.2182395
[3,] -0.1902597  0.2182395  1.0000000

> ## stage 2
> ## Regression coefficients --> why 0.2 (same as Hagger) ???
> A1 <- create.mxMatrix(c(0, "0.2*PsytoMov", "0.2*PaitoMov",
+ 0, 0, "0.2*PaitoPsy",
+ 0, 0, 0),
+ type="Full", byrow=TRUE, ncol=3, nrow=3,
+ as.mxMatrix=FALSE)
>
> ## This step is not necessary but it is useful for inspecting the
model.
> dimnames(A1)[[1]] <- dimnames(A1)[[2]] <- c("Mov", "Psy", "Pai")
> A1
      Mov Psy      Pai
Mov "0" "0.2*PsytoMov" "0.2*PaitoMov"
Psy "0" "0"           "0.2*PaitoPsy"
Pai "0" "0"           "0"

>
> ## Covariance matrix among the variables (CAUTION: NOT SURE FOR THIS
STEP --> why 0.2 ???)
> S1 <- create.mxMatrix(c("0.2*e_Mov", 0, 0,
+ 0, "0.2*e_Psy", 0,
+ 0, 0, "0.2*e_Pai"),
+ type="Full", byrow=TRUE, ncol=3, nrow=3, as.mxMatrix=FALSE)
>
> ## This step is not necessary but it is useful for inspecting the
model.
> dimnames(S1)[[1]] <- dimnames(S1)[[2]] <- c("Mov", "Psy", "Pai")
> S1
      Mov      Psy      Pai
Mov "0.2*e_Mov" "0"      "0"
Psy "0"         "0.2*e_Psy" "0"
Pai "0"         "0"      "0.2*e_Pai"

> ## Stage 2 analysis: different option are in the website from Cheung
and in its paper
> ## option 3 (Cheung/Hagger) --> works like option 1; but cannot have
the indirect effect
>
> random2 <- tssem2(random1, Amatrix=A1, Smatrix=S1,
diag.constraints=TRUE,
+ intervals.type="LB", mx.algebras( Ind=mxAlgebra(PaitoPsy*PaitoMov,
name="Ind")))
+

```

```

+
+ ## Refit again to have all the LBCI
+ random2 <- rerun(random2)
Erreur : unexpected symbol in:
"## Refit again to have all the LBCI
random2"
> summary(random2)

Call:
wls(Cov = pooledS, aCov = aCov, n = tssem1.obj$total.n, Amatrix =
Amatrix,
     Smatrix = Smatrix, Fmatrix = Fmatrix, diag.constraints =
diag.constraints,
     cor.analysis = cor.analysis, intervals.type = intervals.type,
     mx.algebras = mx.algebras, model.name = model.name, suppressWarnings
= suppressWarnings,
     silent = silent, run = run)

```

95% confidence intervals: Likelihood-based statistic
Coefficients:

	Estimate	Std.Error	lbound	ubound	z value	Pr(> z)
PaitoMov	-0.137996	NA	-0.177185	-0.098776	NA	NA
PsytoMov	-0.239478	NA	-0.353150	-0.125946	NA	NA
PsytoPai	0.218240	NA	0.150818	0.285661	NA	NA

Goodness-of-fit indices:

	Value
Sample size	693.00
Chi-square of target model	0.00
DF of target model	0.00
p value of target model	0.00
Number of constraints imposed on "Smatrix"	0.00
DF manually adjusted	0.00
Chi-square of independence model	141.58
DF of independence model	3.00
RMSEA	0.00
RMSEA lower 95% CI	0.00
RMSEA upper 95% CI	0.00
SRMR	0.00
TLI	-Inf
CFI	1.00
AIC	0.00
BIC	0.00

OpenMx status1: 0 ("0" or "1": The optimization is considered fine.
Other values indicate problems.)

```

>
> ## Display the A matrix
> mxEval(Amatrix, random2$mx.fit)
      Mov      Psy      Pai
Mov    0 -0.2394781 -0.137996
Psy    0  0.0000000  0.000000
Pai    0  0.2182395  0.000000
>
>
> ## Draw the output
> library(semPlot)
>
> ## Convert the model to semPlotModel object

```

```
> my.plot <- meta2semPlot(random2)
>
> ## Plot the parameter estimates
> semPaths(my.plot, whatLabels="est", nCharNodes=10, color="yellow",
+         layout="circle2", edge.label.cex = 0.8)
>
```