Psychology 454: Latent Variable Modeling Review and final comments

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Outline

Conceptual overview

A theory of data

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Regression

Multivariate Regression and Partial Correlation

Goodness of fit measures

Absolute and Relative Fit measures

Absolute fit indices

Incremental or relative fit indices

Practical Advice

Measuring change and factorial invariance

Factorial Invariance

Measuring change

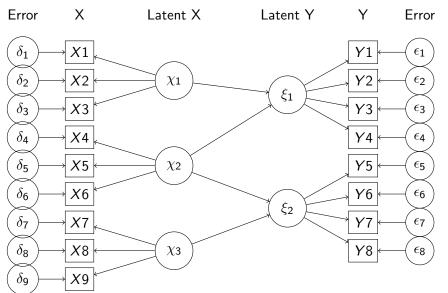
Regression without correcting for reliability

Model misspecification

Mediation

Latent model approach
Problems with SEM

Latent Variable Modeling: A conceptual Syllabus

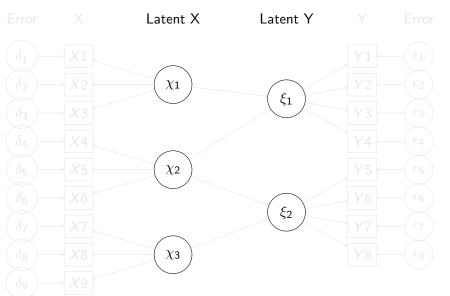


Observed Variables

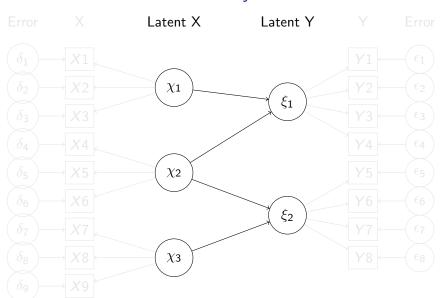
Χ		Υ	
X1		Y1	
X2		Y2	
X3		Y3	
X4		Y4	
X5		Y5	
X6		Y6	
X7		Y7	
X8		Y8	
<i>X</i> 9			



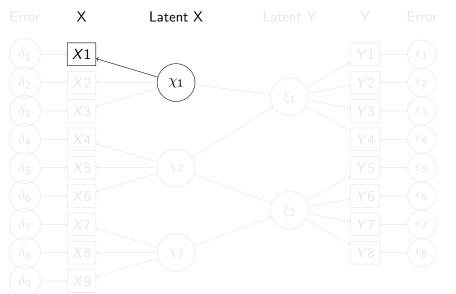
Latent Variables



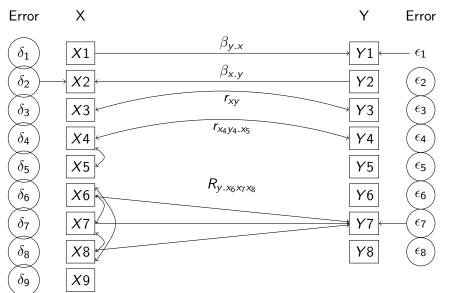
Theory



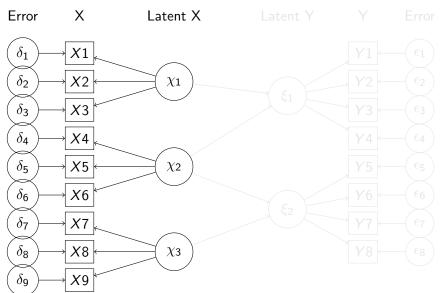
A theory of data and fundamentals of scaling



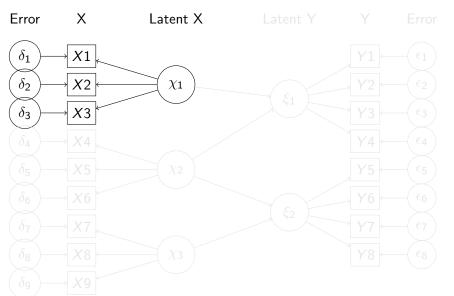
Correlation, Regression, Partial Correlation, Multiple Regression



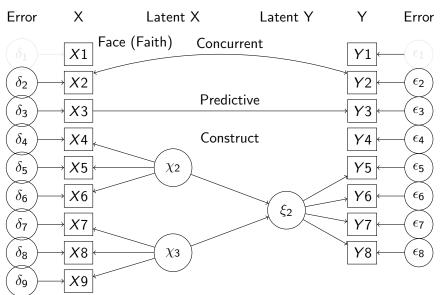
Measurement: A latent variable approach.



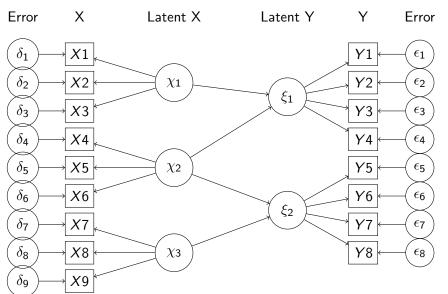
Reliability: How well does a test reflect one latent trait?



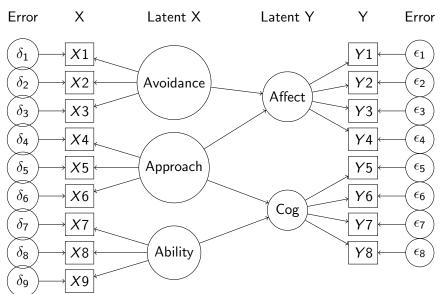
Face, Concurrent, Predictive, Consruct



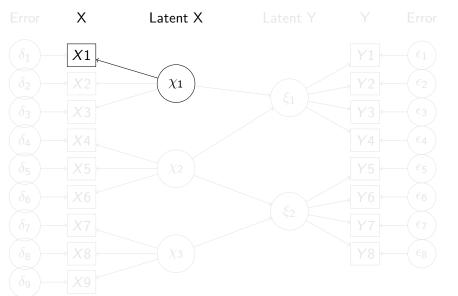
Psychometric Theory: Data, Measurement, Theory



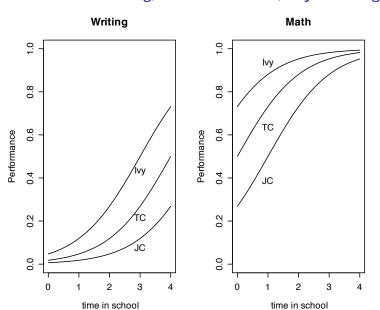
Psychometric Theory: Data, Measurement, Theory



A theory of data and fundamentals of scaling

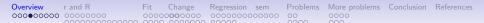


Effect of teaching, effect of students, or just scaling?

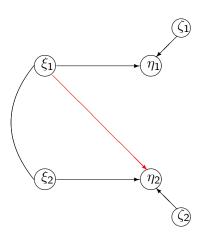


The best scale is the one that works best

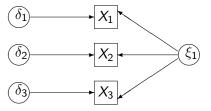
- 1. Money is linear but negatively accelerated with utility.
- 2. Perceived intensity is a log function of physical intensity.
- 3. Probability of being correct is a logistic or cumulative normal function of ability.
- 4. Energy used to heat a house is linear function of outdoor temperature.
- 5. Time to fall a particular distance varies as the square root of the distance.
- 6. Gravitational attraction varies as $1/distance^2$
- 7. Hull speed of sailboat varies as square root of length of boat.
- 8. Sound intensity in db is log(observed/reference)
- 9. pH of solutions is -log(concentration of hydrogen ions)



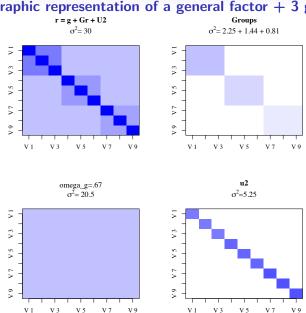
Theory: A regression model of latent variables n



A measurement model for X Reliability



A graphic representation of a general factor + 3 group factors



Factor analytic definition of a general factor

Let F = [g + G] be a column wise concatenation of a general factor and a set of group factors:

Variable	g	Group 1	Group 2	Group 3	
	g	F1	F2	F3	
V1	g1	a1	0	0	
V2	g2	a2	0	0	
V3	g3	a3	0	0	
V4	g4	0	b4	0	
V5	g5	0	b5	0	
V6	g6	0	b6	0	
V7	g7	0	0	c7	
V8	g8	0	0	с8	
V9	g9	0	0	с9	

Variance decomposition of R

$$R = FF' + U^2$$
 where $F = [g + G]$

For orthogonal F and G, the correlation matrix is a function of the general loadings as well as the group loadings:

$$R = gg' + GG' + U^2$$

The amount of variance attributable to the general factor is just $\omega_{\rm g}$ (McDonald, 1999) where

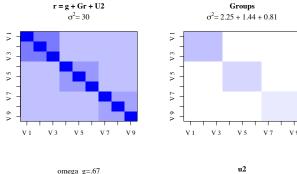
$$\omega_{\mathsf{g}} = \frac{1'\mathsf{g}\mathsf{g}'1}{1'\mathsf{R}1}$$

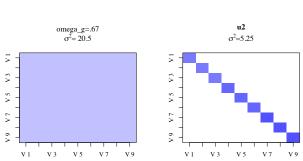
The total amount of reliable variance (that which is attributable to general + groups) is ω_t

$$\omega_t = \frac{1'gg'1 + 1'GG'1}{1'R1}$$

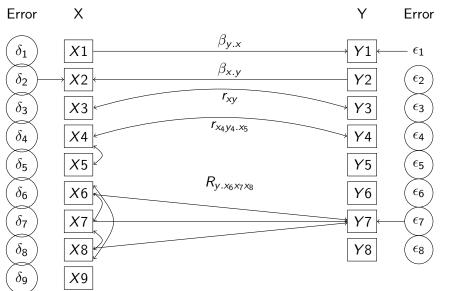
The problem then is how to find ω_{g} .





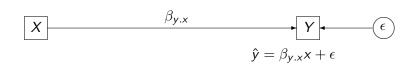


Correlation, Regression, Partial Correlation, Multiple Regression



Bivariate Regression

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$$\beta_{y.x} = \frac{\sigma_{xy}}{\sigma_x^2}$$

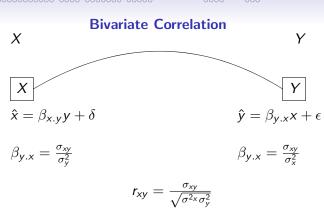


$$\hat{\mathbf{x}} = \beta_{\mathbf{x}.\mathbf{y}}\mathbf{y} + \delta$$

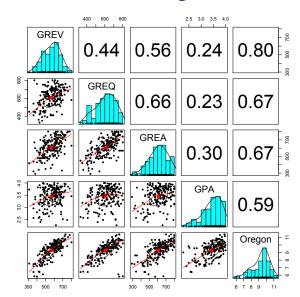
$$\beta_{y.x} = \frac{\sigma_{xy}}{\sigma_y^2}$$

X

δ



Scatter Plot Matrix showing correlation and LOESS regression

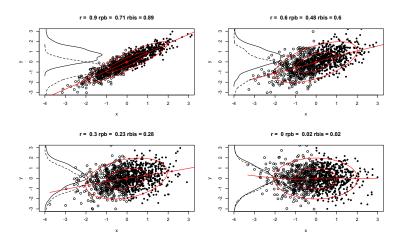


Alternative versions of the correlation coefficient

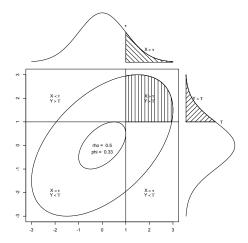
Table: A number of correlations are Pearson r in different forms, or with particular assumptions. If $r = \frac{\sum x_i y_i}{\sqrt{\sum x_i^2 \sum y_i^2}}$, then depending upon the type of data being analyzed, a variety of correlations are found.

Coefficient	symbol	X	Υ	Assumptions
Pearson	r	continuous	continuous	
Spearman	rho (ho)	ranks	ranks	
Point bi-serial	r_{pb}	dichotomous	continuous	
Phi	$\dot{\phi}$	dichotomous	dichotomous	
Bi-serial	r _{bis}	dichotomous	continuous	normality
Tetrachoric	r_{tet}	dichotomous	dichotomous	bivariate normality
Polychoric	r_{pc}	categorical	categorical	bivariate normality

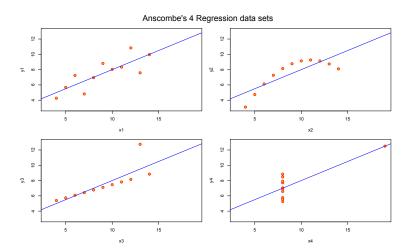
The biserial correlation estimates the latent correlation



The tetrachoric correlation estimates the latent correlation



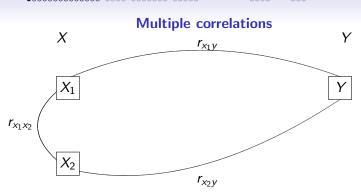
Cautions about correlations: Anscombe data set



The ubiquitous correlation coefficient

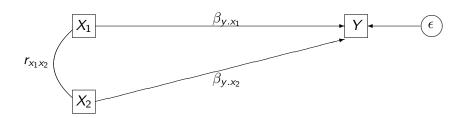
Table: Alternative Estimates of effect size. Using the correlation as a scale free estimate of effect size allows for combining experimental and correlational data in a metric that is directly interpretable as the effect of a standardized unit change in \times leads to r change in standardized y.

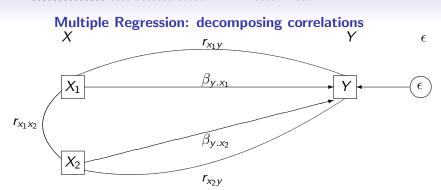
Statistic	Estimate	r equivalent	as a function of r
Pearson correlation	$r_{xy} = \frac{C_{xy}}{\sigma_x \sigma_y}$	r _{xy}	
Regression	$b_{y.x} = \frac{Cxy}{\sigma_z^2}$	$r = b_{y.x} \frac{\sigma_y}{\sigma_x}$	$b_{y.x} = r \frac{\sigma_x}{\sigma_y}$
Cohen's d	$d=rac{X_1-\hat{X}_2}{\sigma_{\times}}$	$r = \frac{d}{\sqrt{d^2 + 4}}$	$d = \frac{2r}{\sqrt{1-r^2}}$
Hedge's g	$g=\frac{X_1-X_2}{s_x}$	$r = \frac{g}{\sqrt{g^2 + 4(df/N)}}$	$g = \frac{2r\sqrt{df/N}}{\sqrt{1-r^2}}$
t - test	$t = \frac{d\sqrt{df}}{2}$	$r = \sqrt{t^2/(t^2 + df)}$	$t = \sqrt{\frac{r^2 df}{1 - r^2}}$
F-test	$F = \frac{d^2df}{4}$	$r = \sqrt{F/(F + df)}$	$F = \frac{r^2 df}{1 - r^2}$
Chi Square		$r = \sqrt{\chi^2/n}$	$\chi^2 = r^2 n$
Odds ratio	$d = \frac{\ln(OR)}{1.81}$	$r = \frac{\ln(OR)}{1.81\sqrt{(\ln(OR)/1.81)^2 + 4}}$	$In(OR) = \frac{3.62r}{\sqrt{1-r^2}}$
r _{equivalent}	r with probability p	$r = r_{equivalent}$	·

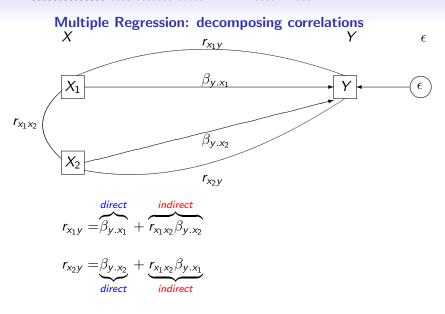


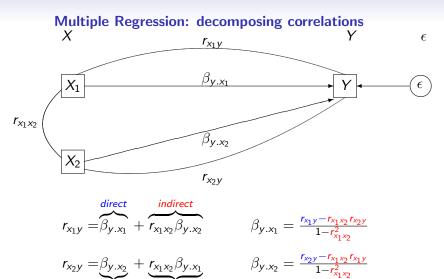
Multiple Regression



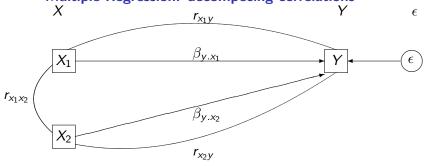










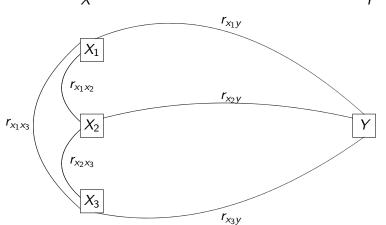


$$r_{x_{1}y} = \beta_{y.x_{1}} + r_{x_{1}x_{2}}\beta_{y.x_{2}} \qquad \beta_{y.x_{1}} = \frac{r_{x_{1}y} - r_{x_{1}x_{2}}r_{x_{2}y}}{1 - r_{x_{1}x_{2}}^{2}}$$

$$r_{x_{2}y} = \beta_{y.x_{2}} + r_{x_{1}x_{2}}\beta_{y.x_{1}} \qquad \beta_{y.x_{2}} = \frac{r_{x_{2}y} - r_{x_{1}x_{2}}r_{x_{1}y}}{1 - r_{x_{1}x_{2}}^{2}}$$

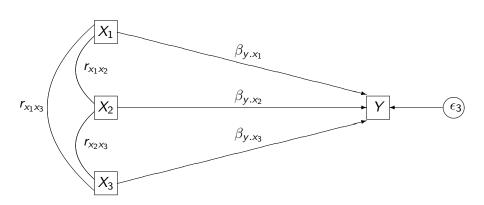
 $R^2 = r_{x_1 y} \beta_{y.x_1} + r_{x_2 y} \beta_{y.x_2}$

What happens with 3 predictors? The correlations

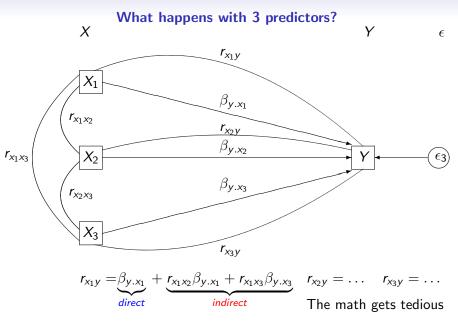


What happens with 3 predictors? β weights

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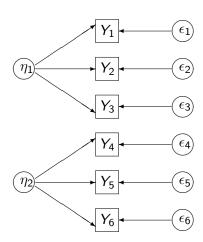




Multiple regression and matrix algebra

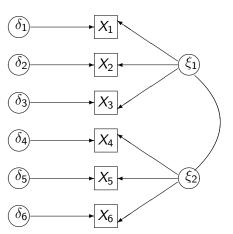
- Multiple regression requires solving multiple, simultaneous equations to estimate the direct and indirect effects.
 - Each equation is expressed as a r_{x_iy} in terms of direct and indirect effects.
 - Direct effect is $\beta_{v.x_i}$
 - Indirect effect is $\sum_{j\neq i} beta_{y.x_j} r_{x_j y}$
- How to solve these equations?
- Tediously, or just use matrix algebra

A measurement model for Y - uncorrelated factors ${\boldsymbol \eta}$

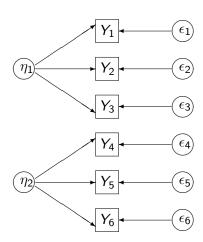


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A measurement model for X – Correlated factors X



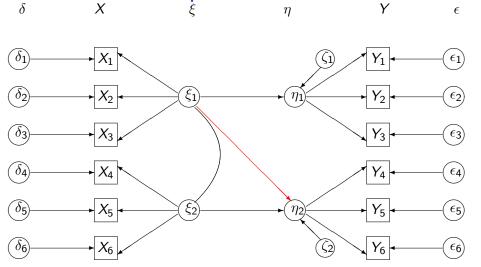
A measurement model for Y - uncorrelated factors ${\boldsymbol \eta}$



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Exploratory versus confirmatory

- The real power of confirmatory analysis is the ability to test paths and models.
- EFA fits $\mathbf{R} = \mathbf{FF'} + \mathbf{U^2}$
 - for k variables and nf factors
 - The number of parameters to estimate is k * nf (nf * (nf 1)/2)
 - The number of observed correlations is k * (k-1)/2
 - degrees of freedom = k * (k-1)/2 k * nf (nf * (nf 1)/2)
 - all parameters are allowed to vary
- Confirmatory analysis fixes some parameters to zero, doesn't estimate others
 - This makes a more constrained model, with more degrees of freedom
- Compare the EFA and CFA of the Thurstone data sets (12 versus 24 df)
- Fixing parameters to specified values is also possible

A number of tests of fit taken from Marsh et al. (2005)

- 1. Marsh, Hau & Grayson (2005) lists 40 different proposed measures of goodness of fit
- Measures of absolute fit.
 - $I_o = \text{index of fit for original or baseline model}$
 - $I_t = \text{index of fit for target or "true" model}$
- 3. Measures of incremental fit Type I
 - $\frac{|I_t I_o|}{Max(I_o I_o)}$ which is either

 - $\frac{I_o I_t}{I_o}$ or $\frac{I_t I_o}{I_o}$
- 4. Measures of incremental fit Type II
 - $\frac{|I_t I_o|}{E(I_t I_o)}$ which is either

 - $\frac{I_o I_t}{I_o E(I_t)}$ or $\frac{I_t I_o}{E(I_t) I_o}$

Fit functions from Jöreskog

- 1. Ordinary least squares $F = \frac{1}{2}tr(S \Sigma)^2$
 - The squared difference between the observed (S) and model (Σ) covariance matrices
 - tr means trace of the sum of the diagonal values of the matrix of squared deviations
- 2. Generalized least squares $F = \frac{1}{2}tr(I S^{-1}\Sigma)^2$
 - I is the identity matrix
 - if the model = data, then $S^{-1}\Sigma$ should be I
 - weight the fit by the inverse of the observed covariances
- 3. Maximum Likelihood $F = log|\Sigma| + tr(S\Sigma^{-1}) log|S| p$
 - weight the fit by the inverse of the modeled covariance
 - p is the number of variables
 - tr (I) = p, and thus the ML should be 0 if the model fits the data

Fit-function based indices

- 1. Fit Function Minimum fit function (FF)
 - $FF = \frac{\chi^2}{(N-1)}$
- 2. Likelihood ratio $LHR = e^{\frac{-1}{2}FF}$
- 3. χ^2 (minimum fit function chi square)

•
$$\chi^2 = tr(\Sigma^{-1}S - I) - log|\Sigma^{-1}S| = (N-1)FF$$

- 4. $p(\chi^2)$ probability of observing a χ^2 this larger or larger given that the model fits
- 5. $\frac{\chi^2}{df}$ has expected value of 1

Non-centrality based indices

- 1. Non-centrality parameter
 - $NCP = \chi^2 df$
- Dk: Rescaled noncentrality paramter (McDonald & Marsh, 1990)

•
$$Dk = FF - \frac{df}{N-1} = \frac{\chi^2}{(N-1)} - \frac{df}{N-1} = \frac{\chi^2 - df}{N-1}$$

- 3. PDF (population discrepancy function = DK normed to be non-negative)
 - $PDF = max(\frac{\chi^2 df}{N 1}, 0)$
- Mc: Measure of centrality (CENTRA, MacDonald Fit Index (MFI)
 - $Mc = e^{\frac{-(\chi^2 df)}{2(N-1)}}$

Error of approximation indices

How large are the residuals, estimated several different ways

- 1. RMSEA (root mean square error of approximation)
 - $RMSEA = \sqrt{PDF/df} = \sqrt{\frac{\max(\frac{\chi^2 df}{N 1}, 0)}{df}}$
 - based upon the non-central χ^2 distribution to find the error of fit
- MSEA (mean square error of approximation unnormed version of RMSEA)
 - $MSEA = \frac{Dk}{df} = \frac{\chi^2 df}{(N-1)df}$
- RMSEAP (root mean square error of approximation of close fit)
 - RMSEA < .05
- 4. RMR Root mean square residual (or, if S and Σ are standardized, the SRMR). Just
 - · square root of the average squared residual
 - $\sqrt{\frac{2\sum(S-\Sigma)^2}{p*(p+1)}}$

Information indices

Compare the information of a model with the number of parameters used for the model. These allow for comparisons of different models with different degrees of freedom.

- AIC (Akaike Information Criterion for a model penalizes for using up df)
 - $AIC = \chi^2 + p * (p+1) 2df = \chi^2 + 2K$
 - where $K = \frac{p*(p+1)}{2} df$
- 2. Baysian Information Criterion
 - $-2Log(L) + plog(N) = \chi^2 Klog(N(.5(p(p+1))))$

Goodness of fit indices

1. GFI from LISREL

•
$$GFI = 1 - \frac{tr(\Sigma^{-1}S - I)^2}{tr(\Sigma^{-1}S)^2}$$

2. Adjusted Goodness of Fit (Lisrel)

•
$$AGFI = 1 - \frac{p(p+1)}{2df}(1 - GFI)$$

3. Unbiased GFI (from Steiger)

• GFI =
$$\frac{p}{2\frac{(\chi^2-df)}{(N-1)}+p}$$

Comparing solutions to solutions

- 1. Incremental fit indices without correction for model complexity
 - RNI (relative non-centrality) McDonald and Marsh
 - CFI Comparative fit index (normed version of RNI) Bentler
 - Normed Fit index (Bentler and Bonett)
- 2. Incremental fit indices correcting for model complexity
 - Tucker Lewis Index
 - Normed Tucker Lewis
 - Incremental Fit index
 - Relative Fit Index
- 3. Parsimony indices

Incremental fit indices without correction for model complexity

- 1. RNI (relative non-centrality) McDonald and Marsh
 - $RNI = 1 \frac{Dk_t}{Dk_n}$
 - where $DK = \frac{\chi^2 df}{N 1}$ for either the null or the tested model
- 2. CFI Comparative fit index (normed version of RNI) Bentler
 - Just norm the RNI to be greater than 0.
 - $CFI = 1 \frac{MAX(NCP_t, 0)}{MAX(NCP_n, 0)}$
- 3. Normed Fit index (Bentler and Bonett)

Fitting functions from Loehlin

- 1. Let S be the "strung out" data matrix
- 2. Let Σ be the "strung out" model matrix
- 3. $Fit = (S \Sigma)'W^{-1}(S \Sigma)$
- 4. Where W =
 - Ordinary Least Squares W = I
 - Generalized Least Squares W = SS'
 - Maximum likelihood $W = \Sigma \Sigma'$

Practical advice

- 1. Taken from Kenny
 - http://davidkenny.net/cf/fit.htm
- 2. Bentler and Bonnet Normed Fit Index
 - $\frac{\chi_{Null}^2 \chi_{Model}^2}{\chi_{Null}^2}$
 - Between .90 and .95 is acceptable
 - > .95 is "good"
- 3. RMSEA
 - if $\chi^2 < df$, then RMSEA = 0
 - "good" models have RMSEA < .05
 - "poor" models have RMSEA > .10
- 4. p of close fit
 - Null hypothesis is that RMSEA is .05
 - test if RMSEA > .05
 - Claim good fit if p(RMSEA > .05) < .05

Considering rules of thumb and fit

- 1. Fit functions have distributions and thus are susceptible to problems of type I and type II error.
 - Compare the fits for correct model as well as those for a simpler incorrect (?) model
- Should we just use chi square and reject models that don't fit, or should we reason about why they don't fit.
- 3. All models are wrong, some are useful.
- "The epistemological basis of statistics has moved away from being a set of procedures, applied mechanistically, and moved toward building and evaluating statistical and scientific models." (Rodgers, 2010)

What does it mean if the model does not fit

- 1. Model is wrong
- 2. Measurement is wrong
- 3. Structure is wrong
- 4. Assumptions are wrong
- 5. at least one of above, but which one?

Specification & Respecification

- 1. Is the measurement model consistent
 - revise it
 - evaluate loadings
 - evaluate error variances
 - more or fewer factors
 - correlated errors?
- 2. Structural model:
 - adjust paths
 - drop paths
 - add paths
- 3. Equivalent models
 - What models are equivalent
 - Do they make equally good sense

Factorial invariance: Does a test measure the same thing in different groups?

- 1. Across groups
 - Different schools
 - Different groups (e.g., ethnicity, age, gender)
- 2. Across time
 - Is todays' measure the same as next year's measure?
- 3. Types of invariance
 - Configural: Are the arrows the same
 - Weak invariance: Are the loadings the same across groups
 - Strong invariance: Loadings and intercepts are equal across groups
 - Super strong: Loadigs, intercepts and means are equal across groups

Measuring structure at two (or more) time points

- 1. Is the structure the same
 - Structural Invariance (is the graph the same)
 - Measurement invariance (are the loadings the same)
 - Strong measurement invariance (are the item intercepts the same?)
 - Measuring change
- 2. Do the means change (is there growth)
 - This is the means of the latent trait, not the means of the items
- 3. Do the latent traits correlate across two or more occasions?
 - Just two occasions, can not separate trait from state effects
 - With > 2 occasions, can examine trait and state effects
- 4. Compare several different simulations

Observed scores over 4 time points

Do they differ in means? Do they measure the same thing?

 X_1

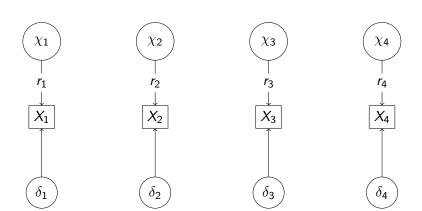
 X_2

*X*₃

 X_4

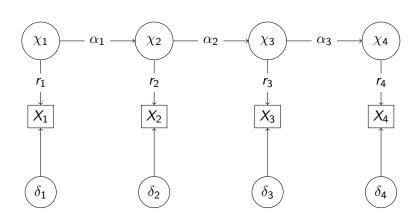
Modeled 4 time points

Are the measures measures of the same construct? Are the measures invariant?

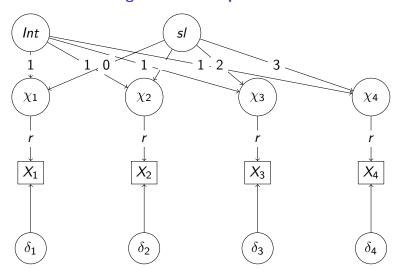


Model state change over 4 time points: simplex but with too many parameters

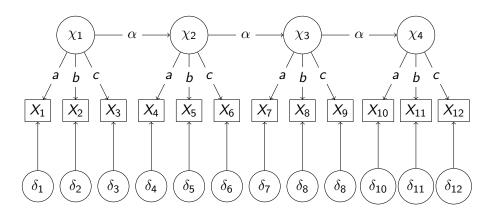
 $\sqrt{r_i}$ is reliability, α_i is autocorrelation.



Model change over 4 time points: traits are stable



Model change over 4 time points: states with measurement



State growth

models <- 'L1 = a*x1 +b *x2 + c* x3 L2 = a* x4 + b * x5 + c* x6 L3 = a*x7 + b*x8 +c*x9 L4 = a*x10 + b*x11 + c*x12

> L2 ~ alpha* L1 L3 ~ alpha * L2 L4 ~ alpha * L3'

fits <- growth(models,data=simp4\$observed)
summary(fits)</pre>

lavaan (0.5-19) converged normally after 32 iterations

Number of observations 1000

 Estimator
 ML

 Minimum Function Test Statistic
 84.661

 Degrees of freedom
 67

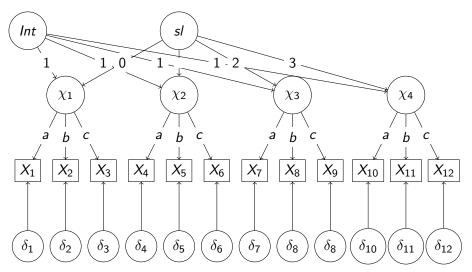
 P-value (Chi-square)
 0.071

Parameter Estimates:

Information Expected
Standard Errors Standard

Latent Variables:

Model change over 4 time points: traits with measurement



Model misspecification: failure to include variables

A classic problem in statements of causal structure is the failure to include appropriate variables. Such model misspecification is the bane of using correlations to infer anything about causality, for there is always the lurking third variable that could explain the relationship.

- In an attempt to demonstrate this effect, consider the correlation between three variables at time 1 as predictors of an important outcome at time 2.
- 2. The measured variables at time 1 are
 - Yellow Fingers
 - Yellow Teeth
 - Bad Breath.
- 3. The outcome variable is probability of Lung Cancer (rescored with a logistic transformation to be a continuous variable ranging from -3 to 3.) ¹

¹As I hope is obvious, this is an artificial example. It was inspired, in part, by the webpage on causal and statistical reasoning at Carnegie Mellon

For the purposes of this demonstration, we create an artificial correlation matrix of these four variables by defining a latent variable, θ , with factor loadings theta. The product of $\theta\theta^T$ is the observed correlation matrix:

```
theta <- matrix(c(0.8, 0.7, 0.6, 0.5), nrow = 4)
observed <- theta %*% t(theta)
diag(observed) <- 1
rownames(observed) <- colnames(observed) <- c("breath", "teeth",
    "fingers", "cancer")
observed
```

```
breath teeth fingers cancer
         1.00 0.56
                       0.48
breath
                             0.40
teeth
         0.56 1.00
                       0.42
                             0.35
         0.48 0.42
                             0.30
fingers
                       1.00
         0.40 0.35
                       0.30
                             1.00
cancer
```

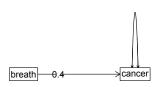
Misspecified linear regression: A series of models

```
R code setCor(y="cancer",x="breath",data=observed)
```

```
Call: setCor(y = "cancer", x = "breath", data = observed)
Multiple Regression from matrix input
Beta weights
       cancer
          0.4
breath
Multiple R
       cancer
          0.4
cancer
multiple R2
       cancer
         0.16
cancer
Unweighted multiple R
cancer
   0.4
Unweighted multiple R2
cancer
 0.16
>
```

1 predictor

Regression Models



Misspecified linear regression: A series of models

R code

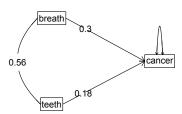
setCor(y="cancer",x=c("breath","teeth"),data=observed)

```
Call: setCor(y = "cancer", x = c("breath", "teeth"), data = observed)
Multiple Regression from matrix input
Beta weights
       cancer
         0.30
breath
teeth
         0.18
Multiple R
       cancer
cancer
        0.43
multiple R2
       cancer
         0.18
cancer
Unweighted multiple R
cancer
 0.42
Unweighted multiple R2
cancer
```

0.18

2 predictors

Regression Models



Misspecified linear regression: A series of models

R code

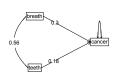
setCor(y="cancer",x=c("breath","teeth","fingers"),data=observed)

```
Call: setCor(v = "cancer", x = c("breath", "teeth", "fingers"), data = observed)
Multiple Regression from matrix input
Beta weights
        cancer
breath
          0.26
teeth
          0.16
fingers 0.11
Multiple R
       cancer
        0.44
cancer
multiple R2
       cancer
cancer 0.19
 Unweighted multiple R
cancer
 0.43
 Unweighted multiple R2
cancer
 0.19
```

2 and 3 predictors

Regression Models

Regression Models



0.56 0.48 (teeth -0.16 cancer)

unweighted matrix correlation = 0.42

unweighted matrix correlation = 0.43

Add in a perfectly causal variable

```
theta <- matrix(c(1, 0.8, 0.7, 0.6, 0.5), nrow = 5)
observed <- theta %*% t(theta)
diag(observed) <- 1
rownames(observed) <- colnames(observed) <- c("smoking", "breath",
    "teeth", "fingers", "cancer")
observed
```

observed

	smoking	breath	teeth	fingers	cancer
smoking	1.0	0.80	0.70	0.60	0.50
breath	0.8	1.00	0.56	0.48	0.40
teeth	0.7	0.56	1.00	0.42	0.35
fingers	0.6	0.48	0.42	1.00	0.30
cancer	0.5	0.40	0.35	0.30	1.00

Try the regression with all 4 predictors

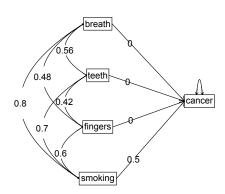
```
R code
```

```
setCor(y="cancer",x=c("breath","teeth","fingers","smoking"),data=observed)
```

```
Call: setCor(v = "cancer", x = c("breath", "teeth", "fingers", "smoking"),
    data = observed)
Multiple Regression from matrix input
Beta weights
        cancer
breath
           0.0
teeth
           0.0
fingers
         0.0
smoking
           0.5
Multiple R
       cancer
          0.5
cancer
multiple R2
       cancer
         0.25
cancer
Unweighted multiple R
cancer
 0.46
Unweighted multiple R2
cancer
  0.22
```

A perfect predictor

Regression Models



But what if the causal variable is not a perfect measure?

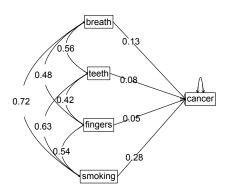
```
Add in some error to smoking
```

```
theta <- matrix(c(.9, 0.8, 0.7, 0.6, 0.5), nrow = 5)
observed <- theta %*% t(theta)
diag(observed) <- 1
rownames(observed) <- colnames(observed) <- c("smoking", "breath",
    "teeth", "fingers", "cancer")
observed</pre>
```

	smoking	breath	teeth	fingers	cancer
smoking	1.00	0.72	0.63	0.54	0.45
breath	0.72	1.00	0.56	0.48	0.40
teeth	0.63	0.56	1.00	0.42	0.35
fingers	0.54	0.48	0.42	1.00	0.30
cancer	0.45	0.40	0.35	0.30	1.00

An imperfect predictor

Regression Models



Improper specification of a mediation model will produce incorrect results

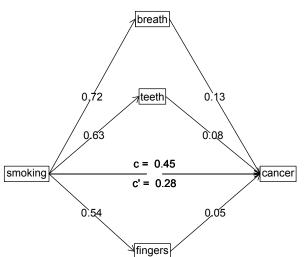
- 1. Although mediation models are popular ways of organizing regression models, they are not guaranteed to be corrrect.
- 2. Even with time lags built into the model, the relationships are not necessarily correct.
- 3. Consider our smoking cancer example

Does hygiene mediate smoking - cancer relationship?

R code mediate(y="cancer",m=c("breath","teeth","fingers"),x="smoking",data=observed) Mediation analysis Call: mediate(y = "cancer", x = "smoking", m = c("breath", "teeth", "fingers"), data = observed) The DV (Y) was cancer. The IV (X) was smoking. The mediating variable(s) = breath teeth fingers. Total Direct effect(c) of smoking on cancer = 0.45 S.E. = 0.03 t direct = 15.92 with probability = 0 Direct effect (c') of smoking on cancer removing breath teeth fingers = 0.28 S.E. = 0.05 t direct = 6.08 with probability = 1.8e-09Indirect effect (ab) of smoking on cancer through breath teeth fingers = 0.17 Mean bootstrapped indirect effect = 0.15 with standard error = 0.03 Lower CI = 0.08 Upper CI = 0.22 Summary of a, b, and ab estimates and ab confidence intervals a cancer ab mean.ab ci.ablower ci.abupper breath 0.72 0.13 0.09 0.00 0.05 0.11 0.63 0.08 0.05 0.04 0.00 0.09 teeth fingers 0.54 0.05 0.03 0.06 0.02 0.09

Smoking effect on cancer is partially mediated by hygiene

Mediation model



Try it the other way: smoking is the mediator

Call: mediate(v = "cancer", x = c("breath", "teeth", "fingers"), m = "smoking",

0.04

0.5 0.5 0.5

Mediation analysis

data = observed)

fingers 0.20 0.28 0.05 0.06

ratio of indirect to total effect=

ratio of indirect to direct effect= 0.99 0.61 0.42

mediate(y="cancer",x=c("breath","teeth","fingers"),m="smoking",data=observed)

The DV (Y) was cancer. The IV (X) was breath teeth fingers. The mediating variable(s) = smoking.

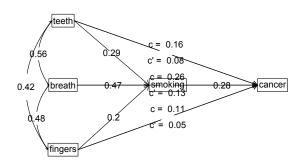
```
Total Direct effect(c) of breath on cancer = 0.26 S.E. = 0.03 t direct = 8.62 with p = 0
Direct effect (c') of breath on cancer removing smoking = 0.13 S.E. = 0.04 t direct = 3.12
Indirect effect (ab) of breath on cancer through smoking = 0.13
Mean bootstrapped indirect effect = 0.13 with standard error = 0.02 Lower CI = 0.08
                                                                                  Upper CI =
Total Direct effect(c) of teeth on cancer = 0.16 S.E. = 0.03 t direct = 5.19 p = 2.6e-07
Direct effect (c') of teeth on cancer removing smoking = 0.08 S.E. = 0.04 t direct = 2.17
Indirect effect (ab) of teeth on cancer through smoking = 0.08
Mean bootstrapped indirect effect = 0.07 with standard error = 0.01 Lower CI = 0.05
                                                                                  Upper CJ =
Total Direct effect(c) of fingers on cancer = 0.11 S.E. = 0.03 t direct = 3.5 p= 0.00048
Direct effect (c') of fingers on cancer removing smoking = 0.05 S.E. = 0.03 t direct = 1.62
Indirect effect (ab) of fingers on cancer through smoking = 0.05
Mean bootstrapped indirect effect = 0.06 with standard error = 0.01 Lower CI = 0.04 Upper CI =
Summary of a, b, and ab estimates and ab confidence intervals
         a cancer ab mean.ab ci.ablower ci.abupper
breath 0.47 0.28 0.13 0.13
                                 0.08
                                            0.17
teeth 0.29 0.28 0.08 0.07
                                 0.05
                                            0.10
```

0.08



Smoking partially mediates hygiene effects

Mediation model



Latent variable modeling can correct for unreliability of measurement

- 1. Part of the problem is regression is measures are imperfect
- 2. We can not see that in the raw correlations
- 3. Can either estimate reliabilities (somehow) or can form latent variables

```
library(lavaan)
model1 <- 'theta =" breath + teeth + fingers
```

fit1 <- cfa(model=model1,sample.cov=observed,sample.nobs=1000,std.lv=TRUE) summary(fit1)

lavaan.diagram(fit1)

teeth

c ·

```
avaan (0.5-18) converged normally after 14 iterations
  Number of observations
                                                   1000
  Estimator
                                                     MI.
  Minimum Function Test Statistic
                                                  0.000
  Degrees of freedom
  Minimum Function Value
                                       0.0000000000000
Parameter estimates:
  Information
                                               Expected
                                               Standard
  Standard Errors
                   Estimate Std.err Z-value P(>|z|)
Latent variables:
  theta =~
                      0.800
                               0.035
                                        22.813
                                                  0.000
    breath
    teeth
                      0.700
                               0.034
                                        20.413
                                                  0.000
    fingers
                      0.600
                               0.034
                                      17.855
                                                  0.000
Variances:
                      0.360
                               0.041
    breath
                      0.509
                               0.037
```

0 000

0 000

Add in the Cancer variable

```
model2 <- 'theta = breath + teeth + fingers
cancer theta

fit2 <- cfa(model=model2,sample.cov=observed,sample.nobs=1000,std.lv=TRUE)
summary(fit2)
lavaan.diagram(fit2)
```

```
lavaan (0.5-18) converged normally after 15 iterations
  Number of observations
                                                    1000
  Estimator
                                                     MI.
  Minimum Function Test Statistic
                                                   0.000
  Degrees of freedom
  P-value (Chi-square)
                                                   1.000
Parameter estimates:
  Information
                                               Expected
  Standard Errors
                                               Standard
                   Estimate Std.err Z-value P(>|z|)
Latent variables:
  theta =~
                                0.032
                                        24.874
                                                   0.000
    breath
                      0.800
    teeth
                      0.700
                                0.032
                                        21.645
                                                   0.000
    fingers
                      0.600
                                0.033
                                        18.319
                                                   0.000
Regressions:
  cancer ~
    theta
                       0.500
                                0.034
                                        14.897
                                                   0.000
```

Add in smoking

```
model3 <- 'theta = smoking + breath + teeth + fingers
cancer theta

fit3 <- cfa(model=model3,sample.cov=observed,sample.nobs=1000,std.lv=TRUE)
summary(fit3)
lavaan.diagram(fit3)
```

```
lavaan (0.5-18) converged normally after 16 iterations
  Number of observations
                                                   1000
  Estimator
                                                      ML
  Minimum Function Test Statistic
                                                   0.000
  Degrees of freedom
  P-value (Chi-square)
                                                   1.000
Parameter estimates:
  Information
                                               Expected
  Standard Errors
                                               Standard
                   Estimate Std.err Z-value P(>|z|)
Latent variables:
  theta =~
    smoking
                      0.900
                                0.026
                                        34.113
                                                  0.000
    breath
                      0.800
                                0.028
                                        28.838
                                                  0.000
    teeth
                      0.700
                                0.029
                                        24.109
                                                  0.000
    fingers
                      0.600
                                0.030
                                        19.824
                                                  0.000
Regressions:
  cancer
    theta
                      0.500
                                0.031
                                        15.975
                                                  0.000
```

44 ways to fool yourself with SEM

Adapted from Rex Kline; Principals and Practice of Structural Equation Modeling, 2005

- 1. Specification
- 2. Data
- 3. Analysis and Respecification
- 4. Interpretation

Specification errors

- 1. Specifying the model after the data are collected.
 - Particularly a problem when using archival data.
- 2. Are key variables omitted?
- 3. Is the model identifiable?
- Omitting causes that are correlated with other variables in the structural model.
- Failure to have sufficient number of indicators of latent variables.
 - "Two might be fine, three is better, four is best, anything more is gravy" (Kenny, 1979)
- 6. Failure to give careful consideration to directionality.
 - Path techniques are good for estimating strengths if we know the underlying model, but are not good for determining the model (Meehl and Walker, 2002)

Specification errors (continued)

- 7. Specifying feedback loops ("non recursive models") as a way to mask uncertainty
- 8. Overfit the model, ignoring parsimony
- 9. Add disturbances ("measurement error correlations" aka "correlated residuals") with substantive reason
- Specifying indicators that are multivocal without substantive reason

Data Errors

- 1. Failure to check the accuracy of data input or coding
 - Missing data codes (use a clear missing value)
 - Misytyped, mis-scanned data matrices
 - Improperly reversed items
 - Let the computer do it for you
 - Why reverse an item when a negative sign will do it for you?
- Ignoring the pattern of missing data, is it random or systematic.
- 3. Failure to examine distributional characteristics
 - Weird data -> weird results
- 4. Failure to screen for outliers
 - Outliers due to mistakes
 - Outliers due to systematic differences

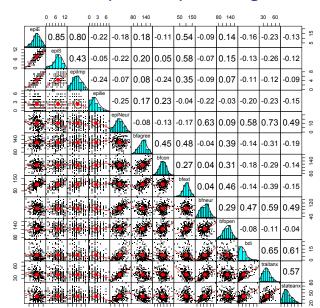
Describe the data

```
> describe(epi.bfi)
```

pairs.panels(epi.bfi,pch=".",gap=0) #mind the gap

	var	n	mean	sd	median	trimmed	mad	\min	max	range	skew	kurtosis	se
epiE	1	231	13.33	4.14	14	13.49	4.45	1	22	21	-0.33	-0.06	0.27
epiS	2	231	7.58	2.69	8	7.77	2.97	0	13	13	-0.57	-0.02	0.18
epiImp	3	231	4.37	1.88	4	4.36	1.48	0	9	9	0.06	-0.62	0.12
epilie	4	231	2.38	1.50	2	2.27	1.48	0	7	7	0.66	0.24	0.10
epiNeur	5	231	10.41	4.90	10	10.39	4.45	0	23	23	0.06	-0.50	0.32
bfagree	6	231	125.00	18.14	126	125.26	17.79	74	167	93	-0.21	-0.27	1.19
bfcon	7	231	113.25	21.88	114	113.42	22.24	53	178	125	-0.02	0.23	1.44
bfext	8	231	102.18	26.45	104	102.99	22.24	8	168	160	-0.41	0.51	1.74
bfneur	9	231	87.97	23.34	90	87.70	23.72	34	152	118	0.07	-0.55	1.54
bfopen	10	231	123.43	20.51	125	123.78	20.76	73	173	100	-0.16	-0.16	1.35
bdi	11	231	6.78	5.78	6	5.97	4.45	0	27	27	1.29	1.50	0.38
traitanx	12	231	39.01	9.52	38	38.36	8.90	22	71	49	0.67	0.47	0.63
stateany	1.3	231	39.85	11.48	38	38 92	10 38	21	79	58	0.72	-0.01	0.76

Graphic descriptions using SPLOMs



Data errors (continued)

- 5. Assuming all relationships are linear without checking
 - graphical techniques are helpful for non-linearities
 - Simple graphical techniques do not help for interactions
- 6. Ignoring lack of independence among observations
 - Nesting of subjects within pairs, within classrooms, with managers
 - Can we model the nesting?

Errors of analysis and respecification

- 1. Failure to check the accuracy of computer syntax
 - · Direction of effects
 - Error specifications
 - Omitted paths
- 2. Respecifying the model based entirely on statistical criteria
 - Just because it does not fit does not mean it should be fixed
- 3. Failure to check for admissible solutions
 - Are some of the paths impossible?
 - Do some of the variables have negative variances?
- 4. Reporting only standardized estimates
 - These are sample based estimates and reflect variances (errorful) and covariances (supposedly error free)
- 5. Analyzing a correlation matrix when the covariance matrix is more appropriate
 - Anything that has growth or change component must be done with covariances

Errors of Analysis and respecification (continued)

- 6. Analyzing a data set with extremely high correlations
 - solution will either be unstable or will not work if variables are too "colinear"
- 7. Not enough subjects for complexity of the data
 - This is ambiguous what is enough?
 - Remember, the standard error of a correlation reflects sample size $se_r = \frac{1}{\sqrt{(1-r^2)(n-2)}}$
 - And thus, the t value associated with any correlation is $\frac{r}{\sqrt{(1-r^2)(n-2)}}$

Errors of Analysis and respecification (continued)

- 8. Setting scales of latent variables inappropriately.
 - particularly a problem with multiple group comparisons
- 9. Ignoring the start values or giving bad ones.
 - Supplying reasonable start values helps a great deal
- 10. Do different start values lead to different solutions?
- 11. Failure to recognize empirical underidentification
 - for some data sets, the model is underidentified even though there are enough parameters
 - Failure to separate measurement from structural portion of model
 - Use the two or four step procedure

Errors of Analysis and respecification (continued)

- 12. Estimating means and intercepts without showing measurement invariance
- 13. Analyzing parcels without checking if parcels are in fact factorially homogeneous.
 - Factorial Homogeneous Item Domains (FHID)
 - Homogenous Item Composites (HIC)
 - (but consider contradictory advice on parcels)

Errors of Interpretation

- 1. Looking only at indexes of overall fit
 - need to examine the residuals to see where there is misfit, even though overall model is fine
- 2. Interpreting good fit as meaning model is "proved".
 - consider alternative models
 - better able to reject alternatives
- 3. Interpreting good fit as meaning that the endogenous variables are strongly predicted.
 - What is predicted is the covariance of the variables, not the variables
 - Are the residual covariances small, not whether the error variance is small
- 4. Relying solely on statistical criterion in model evaluation
 - What can the model not explain
 - What are alternative models
 - What constraints does the model imply

Errors of interpretation (continued)

- 5. Relying too much on statistical tests
 - significance of particular path coefficients does not imply effect size or importance
 - Overall statistical fit (χ^2) is sensitive to model misfit as f(N)
- Misinterpreting the standardized solution in multiple group problems
- 7. Failure to consider equivalent models
 - Why is this model better than equivalent models?
- 8. Failure to consider non-equivalent models
 - · Why is this model better than other, non-equivalent models?
- 9. Reifying the latent variables
 - · Latent variables are just models of observed data
 - "Factors are fictions"
- 10. Believing that naming a factor means it is understood

Errors of interpretation (continued)

- 11. Believing that a strong analytical method like SEM can overcome poor theory or poor design.
- 12. Failure to report enough so that you can be replicated
- Interpreting estimates of large effects as evidence for "causality"

Final Comments

1. Theory First

- What are the alternative theories?
- Are there specific differences in the theories that are testable?

Measurement Model

- Comparison of measurement models?
- How many latent variables? How do you know?
- Measurement Invariance?

3. Structural Model

- · Comparison of multiple models?
- What happens if the arrows are reversed?

4. Theory Last

- What do we know now that we did not know before?
- What do we have shown is not correct?

Conclusion

- 1. Latent variable models are a powerful theoretical aid but do not replace theory.
- Nor do latent modeling algorithms replace the need for good scale development.
- 3. Latent variable models are a supplement to the conventional regression models of observed scores.
- 4. Other latent models (not fully considered) include
 - Item Response Theory
 - Latent Class Analysis
 - Latent Growth Curve analysis

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