

A Longitudinal Test of the Potential Reciprocal Effects between Self-rated Global Physical Health and Depression

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My Background

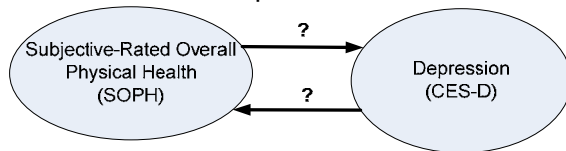
- I'm a methodologist with a focus on measurement and causal modeling
 - Especially via structural equation modeling (SEM)
- I have substantive training in social psychology and sociology of aging
- I'm currently interested in measurement models of physical health and psychological well-being and the causal linkages between these constructs
- Among other projects, I'm working with Neal Dawson and Eleanor Stoller on their Hepatitis C grant

Outline of Presentation

1. Obtain your prediction regarding the causal order between self-rated physical health and depression
2. Provide an overview of longitudinal analysis via growth curves vs. auto-regressive models vs. new “hybrid” models
3. Use an auto-regressive model to test for the potential reciprocal effects between self-rated health & depression

“Tease”

Predict Causal Link between SOPH and Depression



1. Does SOPH (e.g., self-rating based on “excellent,” “good,” “fair,” “poor”) cause depression?
 2. Does depression cause SOPH?
 3. Or is there a **reciprocal** association between them?
- Show of hands for 1 vs. 2 vs. 3?

Research Based on **Cross-Sectional** Survey Data

- A multitude of studies treat Physical Health (e.g., SOPH) as the predictor of Depression
- A multitude of studies treat Depression as the predictor of Physical Health
- Both of the above positions can't be right

Problems with Cross-Sectional Survey Data

- If the **causal order** researchers declare by “fiat” is **wrong**, their estimates of **effect size** will be **distorted**
- This bias occurs whether the hypothesized causal order is completely wrong, or the causal order is reciprocal

Benefits of Longitudinal (Panel) Survey Data

- Provides **empirical test** of causal order
- I'm assuming here that it is not feasible for one to use a true experimental design to test the causal order among variables whose causal order is in question

Problems with Longitudinal Analysis

- It is an order of magnitude more difficult than cross-sectional analysis
 - I'm assuming here that one is conducting longitudinal analysis with the more sophisticated (i.e., more appropriate) procedures
- Requirements
 - Extensive methods training
 - Extensive ("hands-on") experience with the procedures
 - Much more time to produce a final product
 - Enough "journal space" to adequately describe what one has done (especially given that it requires more "education" of the typical reader)

Overview of Longitudinal Analysis via Panel Designs

- Non-experimental research design
- Collects data on multiple individuals for at least two time points
- My focus here is on analytic tools for continuously-distributed outcome variables
 - Not survival (event history) analysis
- I would argue that SEM generally provides the best technique for analyzing these "repeated measure" designs based on interval-level outcome variables
 - Allows one to adjust for random and non-random measurement error
 - Is enormously flexible in the types of models it can test

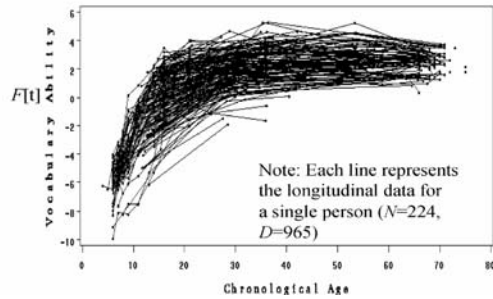
Two Major Types of Longitudinal Designs for Interval-Level Outcomes

- Auto-Regressive Models
 - Model "dynamic," "time-specific" causal processes
 - Used in current study to test potential reciprocal effects
 - "Standard" procedure until recently
- Growth Curve Models
 - Model "developmental" processes with "time-invariant" predictor variables (i.e., predictors that don't change over time)
 - Does not model reciprocal effects easily
 - Currently very "hot" technique that continues to grow in popularity

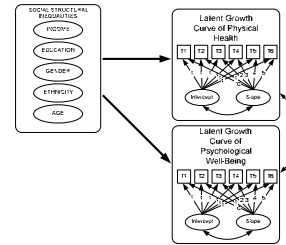
Brief Overview of Growth Curves

- Focus is on linking a person's score on a variable across all waves of data collection to create an individual "slope" (which can be linear or nonlinear)
- "Growth curve" is a bit of a misnomer, given that an individual's slope can be **negative** instead of **positive**

Fig. [7]: New longitudinal scores from the estimated IRT-scaled vocabulary (N=224)



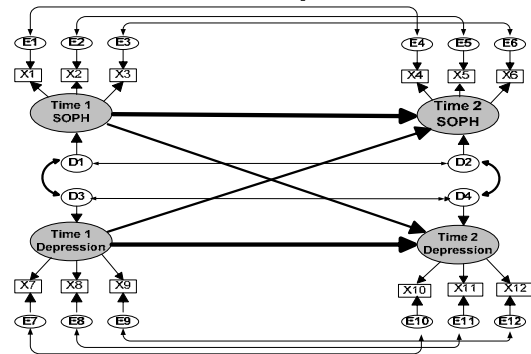
Social Status Characteristics as Time Invariant Predictors of the Latent Growth Curves of Physical Health and Psychological Well-being*
*Modified Diagram from Dissertation Perspectives of Chris Buhrm



Pros and Cons of Growth Curve Models

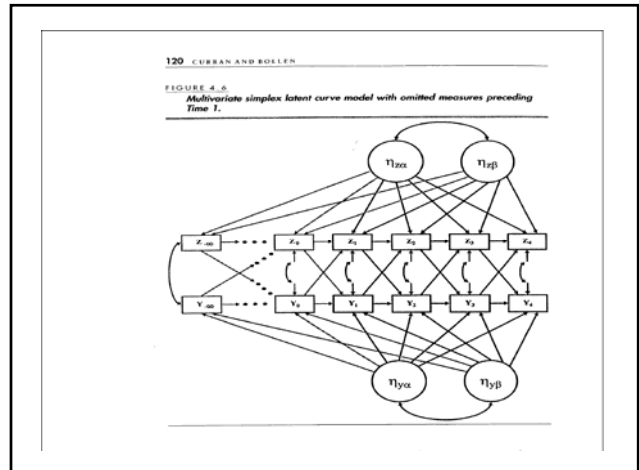
- Is ideal tool for modeling processes that may have an underlying “biological” component
 - Cognitive development
 - Anti-social behavior
 - Problem drinking, etc.
- But many (most?) variables do not have an underlying “developmental” trajectory
- Time-varying predictors generally cannot affect the growth process
- Does not easily incorporate reciprocal effects among variables

Auto-Regressive Model of SOPH and Depression



Pros and Cons of Auto-Regressive Models

- Readily incorporates time-varying predictors (time-specific, “dynamic” processes)
- Better design than growth curves for modeling reciprocal causal effects
- Cannot handle developmental processes



Pros and Cons of “Combo” Models

- Incorporates both dynamic and developmental processes
 - Controls for bias that can occur when modeling one process in the presence of the other process
 - Would appear to represent the current “state-of-the-art” for analyzing panel designs with interval-level outcome variables
- More complex than either auto-regressive or growth curve models
- Few researchers have any experience in using the procedure

References on Longitudinal Analysis

- Email me at kxk@case.edu
- I’ll send you a list with my recommendations on what order to read the material
- If your email system will handle large PDF files, I’ll send you the recommended articles via attached email

Theoretical Background for Proposed Reciprocal Effects between SOPH and Depression

- As noted earlier, my colleagues and I have found no research that tests for reciprocal effects between SOPH (or other dimensions of physical health) and depression
- However, there are various conceptual models researchers have proposed
- The next figure represents a summary of that conceptual background

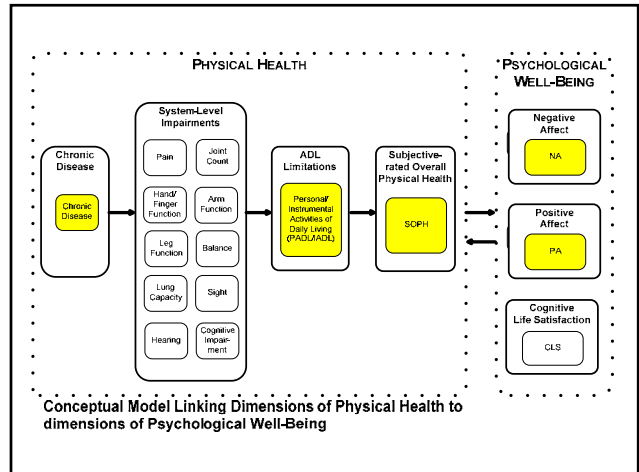
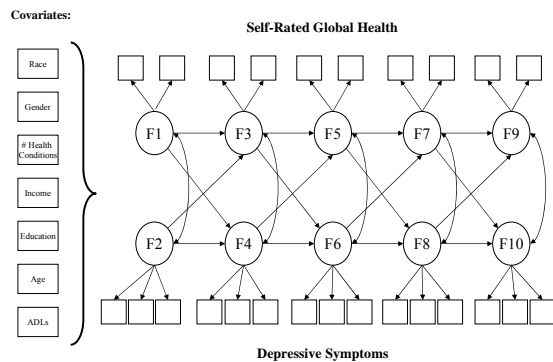


Figure 1
Auto-Regressive Cross-Lagged Model of Global Health and Depression ^a



Auto-Regressive Model (cont.)

- The previous causal model can incorporate equality constraints on factor loadings across waves
 - Important test of key assumption in longitudinal analysis
- Not depicted in previous diagram (to reduce clutter) are:
 - Auto-correlated disturbances
 - Auto-correlated measurement errors
- See prior – two-wave – model of SOPH & Depression for these additional auto-correlated paths

Sample

- Data taken from Rand version of Health & Retirement Survey (HRS)
- Panels collected in 92, 94, 96, 98, 2000
- Ages 51 to 61 in 1992
- Current study uses persons who completed all 5 waves
- N = 8,836

Measures

Depression (Short-version of CES-D)	<i>Response options collapsed into dichotomy:</i> 1 = Present all or most of time 0 = Absent all or most of time										
V1	Negative Affect (3-item composite): <ul style="list-style-type: none"> • I felt depressed • I felt lonely • I felt sad 										
V2	Positive Affect (2-item composite): <ul style="list-style-type: none"> • I was happy • I enjoy life 										
V3	Somatic Symptoms (3-item composite) <ul style="list-style-type: none"> • I felt everything I did was an effort • My sleep was restless • I could not get going 										
SOPH											
V4	Graded Response <i>Would you say your health is</i> <table style="margin-left: 20px;"> <tr> <td>1</td> <td>2</td> <td>3</td> <td>4</td> <td>5</td> </tr> <tr> <td>Poor</td> <td>Fair</td> <td>Good</td> <td>Very Good</td> <td>Excellent</td> </tr> </table>	1	2	3	4	5	Poor	Fair	Good	Very Good	Excellent
1	2	3	4	5							
Poor	Fair	Good	Very Good	Excellent							
V5	Work-Related Health <i>How I want to ask how your health affects paid work activities. Do you have any impairment or health problem that limits the kind or amount of paid work activities?</i> 1 = No 0 = Yes										

Measures (cont.)

Chronic Illness V6	<i>Has a doctor ever told you had.....(Seven-item composite with range 0 to 7)</i> <ul style="list-style-type: none"> • High blood pressure • Diabetes • Cancer • Lung disease • Heart disease • Stroke • Arthritis
Functional Disability V7	Activities of Daily Living (five-item composite): <i>Response options collapsed into dichotomous indicators:</i> 1 = Any difficulty 0 = No difficulty <ul style="list-style-type: none"> • Bathing • Eating • Dressing • Walking across a room • Getting in/out of bed Cronbach's alpha = .70
Education V8	<i>Five-point scale ranging from (1) high school dropout to (5) college graduate and above.</i>
Income V9	<i>Thirty-point scale ranging from no income to \$145,000 and above in \$5,000 increments.</i>
Gender V10	1 = Females 0 = Males
Race V11	1 = African-American 0 = Caucasian

Table 1
Means and Standard Deviations of Variables
(n=8836)

Variables	Wave 1	Wave 2	Wave 3	Wave 4	Wave 5
Race (0=White 1=Black)	.16 (.36)	-	-	-	-
Gender (0=male 1=female)	.59 (.49)	-	-	-	-
# of Chronic Conditions (0 to 7)	.95 (.99)	-	-	-	-
Income (0=no income to 30=\$145,000 or more, \$5,000 increments)	10.7 (7.8)	-	-	-	-
Education (0= less than high school to 5=college grad or beyond)	3.1 (1.4)	-	-	-	-
Age (actual age)	54.9 (5.5)	-	-	-	-
Functional Impairment (0= none to 5=high impairment)	.06 (.35)	-	-	-	-
Self-Assessed Health (1=poor to 5=excellent)	3.6 (1.1)	3.5 (1.1)	3.5 (1.1)	3.2 (1.1)	3.3 (1.1)
Health Limits Work (0=yes 1=no)	.82 (.39)	.78 (.41)	.76 (.43)	.75 (.43)	.73 (.44)
Positive Affect (0=high to 2=low)	.31 (.63)	.18 (.50)	.18 (.49)	.21 (.53)	.20 (.51)
Somatic Complaints (0=low to 3=high)	.30 (.65)	.66 (.94)	.63 (.91)	.79 (.95)	.75 (.94)
Depressive Affect (0=low to 3=high)	.12 (.47)	.41 (.87)	.43 (.85)	.51 (.91)	.52 (.92)

Table 2
Correlations Among Exogenous Variables
(n=8836)

Variables	Gender	Chronic Conditions	Income	Education	Age	ADL Impairment
Race (0=White 1=Black)	.06*	.11*	-.14*	-.16*	-.01	.07*
Gender (0=male 1=female)		.01	-.09*	-.06*	-.27*	.03*
# of Chronic Conditions (0 to 7)			-.16*	-.15*	.18*	.20*
Income (0=no income to 30=\$145,000 or more)				.38*	-.12*	-.13*
Education (0= less than high school to 5=college grad or beyond)					-.05*	-.11*
Age (actual age)						.02
ADL Impairment (0 = no impairment to 5=high impairment)						—

* p < .05

Table 3
Effects of Covariates on Endogenous Variables^a
(n=8836)

Variables		Race	Gender	Chronic Conditions	Income	Education	Age	ADL
Self-Assessed Health	Time 1	-.19*	-.02*	-.61*	.38*	.40*	-.11*	-.39*
	Time 2	-.02	-.01	.02	.01	.03*	-.01	.03*
	Time 3	.01	.01	-.04*	-.01	.02*	.01	-.01
	Time 4	-.02	.02	-.03*	.01	.02*	.01	.01
	Time 5	-.01	-.01	-.01	.01	.02*	.01	-.02
Depression	Time 1	.09*	.10*	.20*	-.21*	-.21*	-.06*	.31*
	Time 2	.01	.09*	-.06*	-.01	-.08*	-.04*	.01
	Time 3	.01	.02	.01	-.01	-.05*	-.05*	.03*
	Time 4	.02	.06*	-.03*	-.01	-.04*	.01	-.01
	Time 5	-.01	.08*	-.01	.01	-.01	-.01	-.01

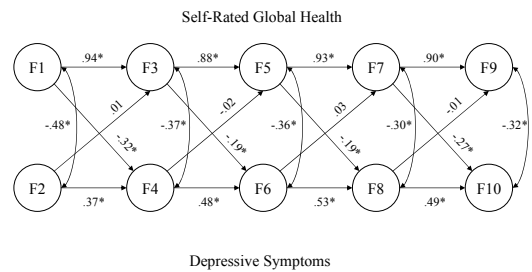
^a Values at Time 1 are correlations; all other values are standardized regression coefficients
* p < .05

Overall Model Fit

With unstandardized factor loadings constrained equal across waves:

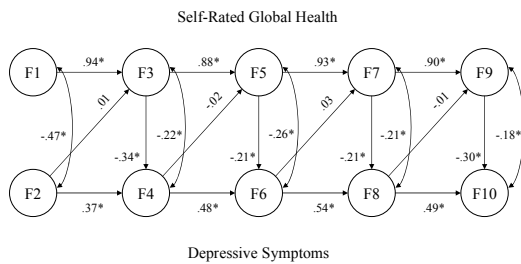
- CFI = .95
- Tucker-Lewis = .93
- RMSEA = .04
- SRMR = .06

Figure 2
Estimated Auto-Regressive Cross-Lag Model^{a,b}
(n=8,836)



^a To simplify presentation, correlated measurement errors and auto-correlated disturbance terms are not presented
^b Standardized coefficients
* p < .05

Figure 3
Auto-regressive Cross-Lag Model with Contemporaneous Effect
of Health on Depression^{a,b}
(n=8,836)



^a To simplify presentation, correlated measurement errors and auto-correlated disturbance terms are not presented
^b Standardized coefficients
* p < .05

Conclusions

- SOPH affects Depression
- Depression does not affect SOPH
- Accordingly, our study finds no evidence of reciprocal effects
- Cross-sectional studies should be cautious in their claims that depression affects physical health

Future Research

- Replicate study using:
 - More complete measures of SOPH and Depression
 - Better second indicator for SOPH
 - Full CES-D with non-dichotomous response options
 - More complete set of measures for other physical health dimensions that we would treat as time-varying rather than time-invariant predictors (as in present study)
 - A sample of old-old persons for which SOPH will vary more (i.e., a sample that is not predominantly healthy)
 - More variation in SOPH would allow us to
 - Model *shorter lag* periods (e.g., 1-year rather than 2-year)
 - Model *contemporaneous reciprocal* effects more completely
 - “Hybrid” models that allow simultaneous assessment of developmental (growth curve) and dynamic (auto-regressive) processes