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Modifying a structural equation model of child dietary intake using the Lagrange Multiplier test in SAS® PROC CALIS for MWSUG 2014

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ABSTRACT

Structural equation models (SEM) allow for simultaneous evaluation of causal pathways between multiple predictors and outcomes, including latent variables, given a hypothesized theoretical model. If an initial SEM does not meet appropriate fit criteria, model modification can achieved through utilization of statistical results from the Lagrange Multiplier (LM) test and theoretical knowledge.

To demonstrate modification of a SEM, baseline data from a garden-based nutrition intervention with elementary school children were examined using SAS® PROC CALIS. The two outcome variables of interest are fruit and vegetable intake (individually). Several measured determinants are hypothesized to predict these practices, including two latent variables, willingness to try fruit and vegetables, which have six indicator variables each.

Initial model fit was not acceptable (χ^2 : 407.9, 111 df, p<0.0001; RMSEA: 0.09, CFI: 0.89). From the LM statistic we identified several unaccounted for correlations between errors on indicators of willingness to try fruits and vegetables. These correlations are theoretically sensible given that items on these scales are worded identically, save for the words "fruit" and "vegetable", and these modifications were made. Once correlations between comparable indicators were added to the SEM, model fit was improved (χ^2 : 234.0, 105 df, p<0.0001; RMSEA: 0.06, CFI: 0.95). Additional relationships identified by the LM test were evaluated, but none were theoretically meaningful, and the second model was accepted as final.

Use of the LM test in PROC CALIS facilitates theoretically appropriate modification of an a priori SEM in order to improve overall model fit and produce more reliable parameter estimates.

INTRODUCTION

Structural equation modeling (SEM) is an analytical approach that allows for exploration of complex multivariate relationships. There are two primary benefits of this technique: 1) latent factors can be observed from knowledge of indicator variables, as with factor analysis, and 2) multiple relationships between independent and dependent variables can be simultaneously observed.

As with general linear regression, checks for model validity are an essential part of the modeling process. Unlike linear regression, statistical tests are performed to ensure the model meets criteria for acceptable fit. When overall model fit does not meet standards, the Lagrange Multiplier (LM) test can be employed to identify parameters that can be added to the model (either directional pathways or covariances between variables, factors or error terms) in order to improve fit. However, given its post-hoc nature, the LM test is a somewhat controversial approach, so must be used in conjunction with theoretical relevance.

This paper will demonstrate the use of the LM test to improve model fit in SEM using PROC CALIS. The specific application will be in exploring the determinants of fruit and vegetable intake and preferences in elementary school children, as part of a nutrition, cooking and gardening intervention for obesity prevention.^{1,2}

VARIABLES

One structural model will include variables specific to fruits, and variables specific to vegetables (all data are crosssectional). The model is structured in this way because fruits and vegetables have varying determinants,³ and thus the strength of relationships may differ between predictors of fruit versus vegetables intake and preferences. Also, there may be some effects of fruit –related variables on vegetable- related variables, and vice versa. The initial hypothesized model is included in **Figure 1**.



Figure 1: Initial hypothesized model

LIST OF VARIABLES

Fruit intake and **vegetable intake**: These variables are measured via the Block Kids Food Screener (last week version),⁴ and intake of each of these food types is provided in cup equivalents.

Fruit identification and **vegetable identification**: Students were given a list of 8 fruit items and 17 vegetable items, ⁵ and were asked if they knew these items, or not. A standardized sum (accounting for unanswered questions) was obtained, with a range of 6-8 items answered about fruits, and 11-17 items answered about vegetables.

Fruit preferences and **vegetable preferences**: For each item identified from the above question, students were asked if they liked this food "A lot", "A little", or "Not at all". Mean scores were used for fruit items and vegetable items (factor analysis was not used due to the large number of indicators).

Willingness to try fruit and **willingness to try vegetables**: The willingness to try scale⁶ has six items each for fruits and vegetables, with questions such as "How much do you like tasting new vegetables?" and responses ranging from "Not at all" to "A lot". The variables for the six items in each scale (specific to fruits or vegetables) served as indicators for a latent factor, either willingness to try fruits or willingness to try vegetables, respectively.

DATA ANALYSIS

DATA PREPARATION

Data appeared to be missing at random, so to account for missingness, the correlation matrix was analyzed instead of raw data. The following code was used to perform this task:

proc corr data=las2 out=las_corr nosimple noprob; var sex ethnicity bmiz age fruit_intake veg_intake fruit_id veg_id veg_willing_1 veg_willing_2 veg_willing_3 veg_willing_4 veg_willing_5 veg_willing_6 fruit_willing_1 fruit_willing_2 fruit_willing_3 fruit_willing_4 fruit_willing_5 fruit_willing_6 fruit_pref veg_pref;

run;

MODEL 1 SPECIFICATION

The code below was used to fit the initial hypothesized model. In the PROC line we specify that the data form is a correlation matrix, specify that we would like to use maximum likelihood estimation, and the 'mod' command indicates we would also like to output the results of the LM test.

Following the 'lineqs' statement, each equation represents a path we would like to include in the model. Terms such as 'b1' give a name to the parameter to reference in the output, and each equation also includes an error term. The terms 'f1' and 'f2' refer to our two latent factors, willingness to try fruits and willingness to try vegetables, and note

these must be indicated in this form (rather than creating a meaningful factor name, such as 'fruit_willingness_factor'; it is acceptable in CALIS to use original variable names rather than renaming as 'v1', etc. for those included in the database).

In the 'variance' statement (or 'std', which performs the same function), we indicate the parameters of which we want to estimate the variance. Similarly in the 'cov' statement, we indicate the relationships of which we would like to estimate the covariances, and we indicate a name for this parameter. We can name parameters in the 'variance' statement, as well, but this is not necessary. Note that for dependent variables (fruit and vegetable preferences and intake), we estimate the covariances of the error terms, not the variables themselves.

```
proc calis corr data=las corr method=ml mod;
```

```
lineqs
fruit pref = b1 fruit id + b2 veg id + b3 f1 + b4 f2 + e1,
veg pref = b5 fruit id + b6 veg id + b7 f1 + b8 f2 + e2,
fruit willing 1 = \overline{a1} f1 + e10,
fruit willing 2 = a2 f1 + e11,
fruit willing 3 = a3 f1 + e12,
fruit willing 4 = a4 f1 + e13,
fruit willing 5 = a5 f1 + e14,
fruit willing 6 = a6 f1 + e15,
veg willing 1 = a7 f2 + e16,
veg_willing_2 = a8 f2 + e17,
veg_willing_3 = a9 f2 + e18,
veg_willing_4 = a10 f2 + e19,
veg_willing_5 = a11 f2 + e20,
veg willing 6 = a12 f2 + e21,
fruit intake = b9 fruit id + b10 veg id + b11 f1 + b12 f2
      + b13 fruit pref + b14 veg pref + e3,
veg intake = b15 fruit id + b16 veg id + b17 f1 + b18 f2
      + b19 fruit pref + b20 veg pref + e4;
variance
e1, e2, e3, e4, e10, e11, e12, e13, e14,
e15, e16, e17, e18, e19, e20, e21;
COV
fruit_id veg_id= theta1,
f1 f2 = theta2,
fruit id fl = theta3,
veg id f2 = theta4,
fruit id f2 = theta5,
veg_id f1 = theta6,
e1 e2 = theta7,
e3 e4 = theta8;
```

run;

MODEL 1 OUTPUT

The following output summarizes the fit of this model. Fit of this model is not acceptable (we would like to see a value >0.95 for the Bentler Comparative Fit Index (CFI), and a value <0.05 for the Root Mean Square Error of Approximation (RMSEA) Index). Therefore, we look to the LM test results.

Fit Summary									
Modeling Info	350								
	N Variables	18							
	N Moments	171							
	N Parameters	60							
	N Active Constraints	0							
	Baseline Model Function Value	7.8962							
	Baseline Model Chi-Square	2755.7718							
	Baseline Model Chi-Square DF	153							
	Pr > Baseline Model Chi-Square	<.0001							
Absolute Index	Fit Function	1.1687							
	Chi-Square	407.8929							
	Chi-Square DF	111							
	Pr > Chi-Square	<.0001							
	Z-Test of Wilson & Hilferty	12.1838							
	Hoelter Critical N	117							
	Root Mean Square Residual (RMSR)	0.0500							
	Standardized RMSR (SRMSR)	0.0500							
	Goodness of Fit Index (GFI)	0.8678							
Parsimony Index	Adjusted GEI (AGEI)	0.7963							
	Parsimonious GFI	0.6296							
	RMSEA Estimate	0.0875							
	RMSEA Lower 90% Confidence Limit	0.0785							
	RMSEA Upper 90% Confidence Limit	0.0967							
	Probability of Close Fit	<.0001							
	ECVI Estimate	1.5324							
	ECVI Lower 90% Confidence Limit	1.3626							
	ECVI Upper 90% Confidence Limit	1.7252							
	Akaike Information Criterion	527.8929							
	Bozdogan CAIC	819.3689							
	Schwarz Bayesian Criterion	759.3689							
	McDonald Centrality	0.6543							
Incremental Index	Bentler Comparative Eit Index	0.8850							
Incremental index	Bentler Benett NEL	0.0000							
	Bentler-Bonett Non-normed Index	0.0020							
	Bollen Normed Index Rho1	0.7960							
	Bollen Non-normed Index Delta?	0.8877							
	James et al. Parsimonious NFI	0.6181							
	canno or an r aronnonous m r	0.0101							

Display 1: Selected output from first structural equation model (fit statistics)

Results of the LM test are included below. The first two tables indicate additional pathways that can be created between observed variables (distinguishing them such that the first table includes both 'predictor' and 'outcome' variables, and the second table only includes 'predictor' variables (ie, identification and willingness variables only); technically our willingness indicators are dependent on the willingness factor), and ranks them by magnitude of change to model fit. However, inclusion of any of these pathways indicated in the first two tables is not desirable. Since we hypothesize a relationship between the willingness to try factors and preferences and intake, it would not make sense to include pathways in the model directly linking the factor indicators with those outcome variables.

The third table shows additional covariances between errors that could be added to the model, again ranked by magnitude. From examining these, we see that some of these pathways make sense. For example, 'e21' and 'e15' both refer to the sixth item on the willingness to try scales, and these questions are worded identically, with the exception of the words 'fruits' or 'vegetables'. The same is true for 'e10' and 'e16', and 'e20' and 'e14'.

Rank Order of the 10 Largest LM Stat for Paths from Endogenous Variables											
То	From	LM Stat	Pr > ChiSq	Parm Change							
veg_willing_1	veg_pref	80.57136	<.0001	2.53750							
veg_willing_2	fruit_pref	56.52670	<.0001	-1.86176							
veg_willing_5	veg_intake	50.16653	<.0001	-0.60705							
veg_willing_1	fruit_pref	26.50015	<.0001	0.45302							
fruit_willing_1	fruit_pref	15.62408	<.0001	0.32701							
veg_willing_6	fruit_pref	5.09153	0.0240	0.23337							
veg_willing_6	veg_intake	3.14364	0.0762	0.13432							
veg_willing_5	fruit_intake	3.14335	0.0762	-0.06888							
fruit_willing_1	fruit_intake	1.90042	0.1680	0.06702							
veg_willing_3	veg_intake	1.88023	0.1703	0.08664							

Rank Order of the 7 Largest LM Stat for Paths from Exogenous Variables											
То	From	LM Stat	Pr > ChiSq	Parm Change							
fruit_willing_2	fruit_id	19.18740	<.0001	1.80624							
fruit_willing_4	fruit_id	5.41341	0.0200	-0.22117							
veg_willing_1	fruit_id	3.87754	0.0489	0.15839							
veg_willing_4	fruit_id	0.32662	0.5677	-0.03983							
veg_willing_2	fruit_id	0.17771	0.6733	0.08149							
fruit_willing_1	fruit_id	0.00521	0.9424	0.00511							
veg_willing_6	fruit_id	0.0003511	0.9851	-0.00284							

Rank Order of the 10 Largest LM Stat for Error Variances and Covariances									
Var1	Var2	LM Stat	Pr > ChiSq	Parm Change					
e12	e10	562.26681	<.0001	5.70035					
e21	e15	87.76411	<.0001	0.35950					
e16	e10	76.56286	<.0001	0.31856					
e14	e1	68.70051	<.0001	-0.69057					
e20	e12	66.66359	<.0001	-0.54926					
e14	e10	56.00892	<.0001	-0.65302					
e20	e14	51.67631	<.0001	0.33683					
e16	e14	37.46651	<.0001	-0.23992					
e19	e13	36.23283	<.0001	0.22154					
e13	e10	32,74577	< 0001	-0.31696					

Display 2: Selected output from first structural equation model (Lagrange Multiplier test)

MODEL 2 SPECIFICATION

To modify the model, we add six additional covariance parameters to the model, such that correlations are included between comparable items on the fruit and vegetable willingness to try subscales (**Figure 2**).



Figure 2: Modified structural model

The revised code is below, with the additional lines to the 'cov' statement being the only modifications made.

```
proc calis corr data=las corr method=ml mod;
lineqs
fruit pref = b1 fruit id + b2 veg id + b3 f1 + b4 f2 + e1,
veg pref = b5 fruit id + b6 veg id + b7 f1 + b8 f2 + e2,
fruit_willing_1 = a\overline{1} f1 + e10,
fruit_willing_2 = a2 f1 + e11,
fruit_willing_3 = a3 f1 + e12,
fruit_willing_4 = a4 f1 + e13,
fruit_willing_5 = a5 f1 + e14,
fruit_willing_6 = a6 f1 + e15,
veg willing 1 = a7 f2 + e16,
veg willing 2 = a8 f2 + e17,
veg_willing 3 = a9 f2 + e18,
veg willing 4 = a10 f2 + e19,
veg_willing_5 = all f2 + e20,
veg willing 6 = a12 f2 + e21,
fruit intake = b9 fruit id + b10 veg id + b11 f1
       + b12 f2 + b13 fruit_pref + b14 veg_pref + e3,
veg intake = b15 fruit id + b16 veg id + b17 f1
       + b18 f2 + b19 fruit_pref + b20 veg_pref + e4;
variance
e1, e2, e3, e4, e10, e11, e12, e13, e14,
e15, e16, e17, e18, e19, e20, e21;
COV
fruit_id veg_id= theta1,
f1 f2 = theta2,
fruit id f1 = theta3,
veg id f2 = theta4,
fruit id f2 = theta5,
veg id fl = theta6,
e1 e2 = theta7,
e3 e4 = theta8,
el0 el6 = theta9,
e15 e21 = theta10,
e13 \ e19 = theta11,
```

el1 e17 = theta12, e12 e18 = theta13, e14 e20 = theta14; run;

MODEL 2 OUTPUT

This code gives us the following output for fit statistics. We see that the model fit is improved, such that the CFI is within an acceptable range (0.95), and the RMSEA value is borderline (0.059). Also the Chi-square value is improved from 407.9 to 234.0 (with six fewer degrees of freedom), although still with a p-value >0.05. Results of the LM test were again evaluated, but none were theoretically meaningful, so we consider this model acceptable to move forward with hypothesis testing.

Fit Summary									
Modeling Info	N Observations	350							
	N Variables	18							
	N Moments	171							
	N Parameters	66							
	N Active Constraints	0							
	Baseline Model Function Value	7.8962							
	Baseline Model Chi-Square	2755.7718							
	Baseline Model Chi-Square DF	153							
	Pr > Baseline Model Chi-Square	<.0001							
Absolute Index	Fit Function	0.6705							
	Chi-Square	233.9922							
	Chi-Square DF	105							
	Pr > Chi-Square	<.0001							
	Z-Test of Wilson & Hilferty	6.7015							
	Hoelter Critical N	194							
	Root Mean Square Residual (RMSR)	0.0430							
	Standardized RMSR (SRMSR)	0.0430							
	Goodness of Fit Index (GFI)	0.9255							
Parsimony Index	Adjusted GFI (AGFI)	0.8786							
	Parsimonious GFI	0.6351							
	RMSEA Estimate	0.0593							
	RMSEA Lower 90% Confidence Limit	0.0491							
	RMSEA Upper 90% Confidence Limit	0.0695							
	Probability of Close Fit	0.0653							
	ECVI Estimate	1.0705							
	ECVI Lower 90% Confidence Limit	0.9526							
	ECVI Upper 90% Confidence Limit	1.2119							
	Akaike Information Criterion	365.9922							
	Bozdogan CAIC	686.6158							
	Schwarz Bayesian Criterion	620.6158							
	McDonald Centrality	0.8317							
Incremental Index	Bentler Comparative Fit Index	0.9504							
	Bentler-Bonett NFI	0.9151							
	Bentler-Bonett Non-normed Index	0.9278							
	Bollen Normed Index Rho1	0.8763							
	Bollen Non-normed Index Delta2	0.9513							
	James et al. Parsimonious NFI	0.6280							

Display 3: Selected output from second structural equation model (fit statistics)

The output below indicates the standardized estimates for the linear equations in the model, and the t-value indicates the significance of these associations. We see that fruit identification and willingness to try fruit are significant predictors of fruit preferences, and that willingness to try vegetables is a significant predictor of vegetable preferences. For fruit intake, only fruit identification is a significant predictor; and for vegetable intake, fruit identification, vegetables identification, and willingness to try vegetables are significant predictors.

Standardized Results for Linear Equations																												
fruit_pref	=	0.2258	*	fruit_id	+	-0.0948	*	veg_id	+	0.3290	*	f1	+	0.1392	*	f2	+	1.0000		e1							Τ	
Std Err		0.0524		b1		0.0542		b2		0.0746		b3		0.0720		b4												
t Value		4.3124				-1.7485				4.4085				1.9346														
veg_pref	=	-0.00427	*	fruit_id	+	-0.0459	*	veg_id	+	-0.1218	*	f1	+	0.5488	*	f2	+	1.0000		e2								
Std Err		0.0529		b5		0.0540		b6		0.0763		b7		0.0667		b8												
t Value		-0.0808				-0.8505				-1.5973				8.2286														
fruit_willing_1	=	0.5501	*	f1	+	1.0000		e10																				
Std Err		0.0417		a1																								
t Value		13.2048																										
fruit_willing_2	=	0.6278	*	f1	+	1.0000		e11																				
Std Err		0.0383		a2																								
t Value		16.3878																										
fruit_willing_3	=	0.7276	*	f1	+	1.0000		e12																				
Std Err		0.0326		a3																								
t Value		22.3219																										
fruit_willing_4	=	0.5811	*	f1	+	1.0000		e13																				
Std Err		0.0408		a4																								
t Value		14.2578																										
fruit_willing_5	=	0.6809	*	f1	+	1.0000		e14																				
Std Err		0.0350		a5																								
t Value		19.4423																										
fruit_willing_6	=	0.6417	*	f1	+	1.0000		e15																				
Std Err		0.0368		a6																								
t Value		17.4157																										
veg_willing_1	=	0.7614	*	f2	+	1.0000		e16																				
Std Err		0.0251		a7																								
t Value		30.3433																										
veg_willing_2	=	0.7760	*	f2	+	1.0000		e17																				
Std Err		0.0244		a8																								
t Value		31.8369																										
veg_willing_3	=	0.8244	*	f2	+	1.0000		e18																				
Std Err		0.0208		a9																								
t Value		39.6661																										
veg_willing_4	=	0.7181	*	f2	+	1.0000		e19																				
Std Err		0.0286		a10																								
t Value		25.0965																										
veg_willing_5	=	0.8085	*	f2	+	1.0000		e20																				
Std Err		0.0218		a11																								
t Value		37.0371																										
veg_willing_6	=	0.7652	*	f2	+	1.0000		e21																				
Std Err		0.0249		a12																								
t Value		30.6789																										
fruit_intake	=	-0.1302	*	fruit_id	+	0.1004	*	veg_id	+	0.0879	*	f1	+	0.0541	*	f2	+	0.0172	*	fruit_pref	+	0.0477	*	veg_pref	+	1.0000	e	.3
Std Err		0.0592		b9		0.0591		b10		0.0898		b11		0.0878		b12		0.0658		b13		0.0661		b14				
t Value		-2.1980				1.6991				0.9796				0.6161				0.2607				0.7217						
veg_intake	=	-0.2106	*	fruit_id	+	0.1868	*	veg_id	+	-0.0633	*	f1	+	0.2303	*	f2	+	0.0535	*	fruit_pref	+	0.0795	*	veg_pref	+	1.0000	e	.4
Std Err		0.0564		b15		0.0562		b16		0.0863		b17		0.0835		b18		0.0631		b19		0.0634		b20				
t Value		-3.7372				3.3219				-0.7333				2.7575				0.8479				1.2532						

Display 4: Selected output from second structural equation model (linear equations)

CONCLUSION

These results indicate that identification and willingness to try fruits and vegetables are predictors of preferences and intake, but preference is not predictive of intake, as has previously been demonstrated in the literature.⁷ A structural equation model is a useful way to examine these data because preferences can be evaluated as a mediator between intake and the predictors identification and willingness to try, but we see from these results that it is not one.

When specifying the model, the LM test is a helpful tool for model modification, to ensure that overall model fit is appropriate and that parameter estimates are reliable. However, prudence must be exercised when employing this approach so that models remain theoretically sound. Just as inappropriate model fit can diminish findings, so can too an illogical model.

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