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# Introduction

# **1.1** Observed and latent variables

Observed variables are those that can be directly measured, such as systolic blood pressure, diastolic blood pressure, waist-hip ratio, body mass index, and heart rate. Measurements from observed variables provide data as the basic source of information for statistical analysis. In medical, social, and psychological research, it is common to encounter latent constructs that cannot be directly measured by a single observed variable. Simple examples are intelligence, health condition, obesity, and blood pressure. To assess the nature of a latent construct, a combination of several observed variables is needed. For example, systolic blood pressure and diastolic blood pressure should be combined to evaluate blood pressure; and waist-hip ratio and body mass index should be combined to evaluate obesity. In statistical inference, a latent construct is analyzed through a latent variable which is appropriately defined by a combination of several observed variables.

For practical research in social and biomedical sciences, it is often necessary to examine the relationships among the variables of interest. For example, in a study that focuses on kidney disease of type 2 diabetic patients (see Appendix 1.1), we have data from the following observed key variables: plasma creatine (PCr), urinary albumin creatinine ratio (ACR), systolic blood pressure (SBP), diastolic blood pressure (DBP), body mass index (BMI), waist–hip ratio (WHR), glycated hemoglobin (HbAlc), and fasting plasma glucose (FPG). From the basic medical knowledge about kidney disease, we know that the severity of this disease is reflected by both PCr and ACR. In order to understand the effects of the explanatory (independent) variables such as SBP and BMI on kidney disease, one possible approach is to apply the well-known regression model by treating PCr and ACR as outcome (dependent) variables and regressing them on the observed explanatory (independent) variables as follows:

 $PCr = \alpha_1 SBP + \alpha_2 DBP + \alpha_3 BMI + \alpha_4 WHR + \alpha_5 HbA1c + \alpha_6 FPG + \epsilon_1, \quad (1.1)$ 

$$ACR = \beta_1 SBP + \beta_2 DBP + \beta_3 BMI + \beta_4 WHR + \beta_5 HbA1c + \beta_6 FPG + \epsilon_2.$$
(1.2)

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From the estimates of the  $\alpha$ s and  $\beta$ s, we can assess the effects of the explanatory variables on PCr and ACR. For example, based on the estimates of  $\alpha_1$  and  $\beta_1$ , we can evaluate the effects of SBP on PCr and ACR, respectively. However, this result cannot provide a clear and direct answer to the question about the effect of SBP on kidney disease. Similarly, the effects of other observed explanatory variables on kidney disease cannot be directly assessed from results obtained from regression analysis of equations (1.1) and (1.2). The deficiency of the regression model when applied to this study is due to the fact that kidney disease is a latent variable (construct) rather than an observed variable. A better approach is to appropriately combine PCr and ACR into a latent variable 'kidney disease (KD)' and regress this latent variable on the explanatory variables. Moreover, one may be interested in the effect of blood pressure rather than in the separate effects of SBP and DBP. Although the estimates of  $\alpha_1$  and  $\alpha_2$  can be used to examine the respective effects of SBP and DBP on PCr, they cannot provide a direct and clear assessment on the effect of blood pressure on PCr. Hence, it is desirable to group SBP and DBP together to form a latent variable that can be interpreted as 'blood pressure (BP)', and then use BP as an explanatory variable. Based on similar reasoning, {BMI, WHR} and {HbA1c, FPG} are appropriately grouped together to form latent variables that can be interpreted as 'obesity (OB)' and 'glycemic control (GC)', respectively. To study the effects of blood pressure, obesity, and glycemic control on kidney disease, we consider the following simple regression equation with latent variables:

$$KD = \gamma_1 BP + \gamma_2 OB + \gamma_3 GC + \delta.$$
(1.3)

This simple regression equation can be generalized to the multiple regression equation with product terms. For example, the following regression model can be used to assess the additional interactive effects among blood pressure, obesity, and glycemic control on kidney disease:

$$KD = \gamma_1 BP + \gamma_2 OB + \gamma_3 GC + \gamma_4 (BP \times OB) + \gamma_5 (BP \times GC) + \gamma_6 (OB \times GC) + \delta.$$
(1.4)

Note that studying these interactive effects by using the regression equations with the observed variables (see (1.1) and (1.2)) is extremely tedious.

It is obvious from the above simple example that incorporating latent variables in developing models for practical research is advantageous. First, it can reduce the number of variables in the key regression equation. Comparing equation (1.3) with (1.1) and (1.2), the number of explanatory variables is reduced from six to three. Second, as highly correlated observed variables are grouped into latent variables, the problem induced by multicollinearity is alleviated. For example, the multicollinearity induced by the highly correlated variables SBP and DBP in analyzing regression equation (1.1) or (1.2) does not exist in regression equation (1.3). Third, it gives better assessments on the interrelationships of latent constructs. For instance, direct and interactive effects among the latent constructs blood pressure, obesity, and glycemic control can be assessed through the regression model (1.4). Hence, it is important to have a statistical method that simultaneously groups highly correlated observed variables into latent variables and assesses interrelationships among latent variables through a regression model of latent variables. This strong demand is the motivation for the development of structural equation models.

#### OBJECTIVES OF THE BOOK 3

# **1.2** Structural equation model

The structural equation model (SEM) is a powerful multivariate tool for studying interrelationships among observed and latent variables. This statistical method is very popular in behavioral, educational, psychological, and social research. Recently, it has also received a great deal of attention in biomedical research; see, for example, Bentler and Stein (1992) and Pugesek *et al.* (2003).

The basic SEM, for example, the widely used LISREL model (Jöreskogand Sörbom, 1996), consists of two components. The first component is a confirmatory factor analysis (CFA) model which groups the highly correlated observed variables into latent variables and takes the measurement error into account. This component can be regarded as a regression model which regresses the observed variables on a smaller number of latent variables. As the covariance matrix of the latent variables is allowed to be nondiagonal, the correlations/covariances of the latent variables can be evaluated. However, various effects of the explanatory latent variables on the key outcome latent variables of interest cannot be assessed by the CFA model of the first component. Hence, a second component is needed. This component is again a regression type model, in which the outcome latent variables are regressed on the explanatory latent variables. As a result, the SEM is conceptually formulated by the familiar regression type model. However, as latent variables in the model are random, the standard technique in regression analysis cannot be applied to analyze SEMs.

It is often important in substantive research to develop an appropriate model to evaluate a series of simultaneous hypotheses on the impacts of some explanatory observed and latent variables on the key outcome variables. Based on its particular formulation, the SEM is very useful for achieving the above objective. Furthermore, it is easy to appreciate the key idea of the SEM, and to apply it to substantive research; one only needs to understand the basic concepts of latent variables and the familiar regression model. As a result, this model has been extensively applied to behavioral, educational, psychological, and social research. Due to the strong demand, more than a dozen user-friendly SEM software packages have been developed; typical examples are AMOS, EQS6, LISREL, and Mplus. Recently, the SEM has become a popular statistical tool for biomedical and environmental research. For instance, it has been applied to the analysis of the effects of *in utero* methylmercury exposure on neurodevelopment (Sánchez *et al.*, 2005), to the study of ecological and evolutionary biology (Pugesek *et al.*, 2003), and to the evaluation of the interrelationships among latent domains in quality of life (e.g. Lee *et al.*, 2005).

## **1.3** Objectives of the book

Like most other statistical methods, the methodological developments of standard SEMs depend on crucial assumptions. More specifically, the most basic assumptions are as follows: (i) The regression model in the second component is based on a simple linear regression equation in which higher-order product terms (such as quadratic terms or interaction terms) cannot be assessed. (ii) The observed random variables are assumed to be continuous, and independently and identically normally distributed. As these assumptions may not be valid in substantive research, they induce limitations in applying SEMs to the analysis of real data in relation to complex situations. Motivated by the need to overcome these limitations, the growth of SEMs has been very rapid in recent years. New models and statistical methods have been

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developed to relax various aspects of the crucial assumptions for better analyses of complex data structure in practical research. These include, but are not limited to: nonlinear SEMs with covariates (e.g. Schumacker and Marcoulides, 1998; Lee and Song, 2003a); SEMs with mixed continuous, ordered and/or unordered categorical variables (e.g. Shi and Lee, 2000; Moustaki, 2003; Song and Lee, 2004; Song et al., 2007); multilevel SEMs (e.g. Lee and Shi, 2001; Rabe-Hesketh et al., 2004; Song and Lee, 2004; Lee and Song, 2005); mixture SEMs (e.g. Dolan and van der Maas, 1998; Zhu and Lee, 2001; Lee and Song, 2003b); SEMs with missing data (e.g. Jamshidian and Bentler, 1999; Lee and Tang, 2006; Song and Lee, 2006); SEMs with variables from exponential family distributions (e.g. Wedel and Kamakura, 2001; Song and Lee, 2007); longitudinal SEMs (Dunson, 2003; Song et al., 2008); semiparametric SEMs (Lee et al., 2008; Song et al., 2009; Yang and Dunson, 2010; Song and Lu, 2010); and transformation SEMs (van Montfort *et al.*, 2009; Song and Lu, 2012). As the existing software packages in SEMs are developed on the basis of the covariance structure approach, and their primary goal is to analyze the standard SEM under usual assumptions, they cannot be effectively and efficiently applied to the analysis of the more complex models and/or data structures mentioned above. Blindly applying these software packages to complex situations has a very high chance of producing questionable results and drawing misleading conclusions.

In substantive research, data obtained for evaluating hypotheses of complex diseases are usually very complicated. In analyzing these complicated data, more subtle models and rigorous statistically methods are important for providing correct conclusions. In view of this, there is an urgent need to introduce into applied research statistically sound methods that have recently been developed. This is the main objective in writing this book. As we write, there has only been a limited amount of work on SEM. Bollen (1989) was devoted to standard SEMs and focused on the covariance structure approach. Compared to Bollen (1989), this book introduces more advanced SEMs and emphasizes the Bayesian approach which is more flexible than the covariance structure approach in handling complex data and models. Lee (2007) provides a Bayesian approach for analyzing the standard and more subtle SEMs. Compared to Lee (2007), the first four chapters of this book provide less technical discussions and explanations of the basic ideas in addition to the more involved, theoretical developments of the statistical methods, so that they can be understood without much difficulty by applied researchers. Another objective of this book is to introduce important models that have recently been developed and were not covered by Lee (2007), including innovative growth curve models and longitudinal SEMs for analyzing longitudinal data and for studying the dynamic changes of characteristics with respect to time; semiparametric SEMs for relaxing the normality assumption and for assessing the true distributions of explanatory latent variables; SEMs with a nonparametric structural equation for capturing the true general relationships among latent variables, and transformation SEMs for analyzing highly nonnormal data. We believe that these advanced SEMs are very useful in substantive research.

## **1.4 The Bayesian approach**

A traditional method in analyzing SEMs is the covariance structure approach which focuses on fitting the covariance structure under the proposed model to the sample covariance matrix computed from the observed data. For simple SEMs, when the underlying distribution of the observed data is normal, this approach works fine with reasonably large sample sizes. However, some serious difficulties may be encountered in many complex situations in which

#### REAL DATA SETS AND NOTATION 5

deriving the covariance structure or obtaining an appropriate sample covariance matrix for statistical inferences is difficult.

Thanks to recent advances in statistical computing, such as the development of various efficient Markov chain Monte Carlo (MCMC) algorithms, the Bayesian approach has been extensively applied to analyze many complex statistical models. Inspired by its wide applications in statistics, we will use the Bayesian approach to analyze the advanced SEMs that are useful for medical and social-psychological research. Moreover, in formulating and fitting the model, we emphasize the raw individual random observations rather than the sample covariance matrix. The Bayesian approach coupled with the formulation based on raw individual observations has several advantages. First, the development of statistical methods is based on the first moment properties of the raw individual observations which are simpler than the second moment properties of the sample covariance matrix. Hence, it has the potential to be applied to more complex situations. Second, it produces a direct estimation of latent variables, which cannot be obtained with classical methods. Third, it directly models observed variables with their latent variables through the familiar regression equations; hence, it gives a more direct interpretation and can utilize the common techniques in regression such as outlier and residual analyses in conducting statistical analysis. Fourth, in addition to the information that is available in the observed data, the Bayesian approach allows the use of genuine prior information for producing better results. Fifth, the Bayesian approach provides more easily assessable statistics for goodness of fit and model comparison, and also other useful statistics such as the mean and percentiles of the posterior distribution. Sixth, it can give more reliable results for small samples (see Dunson, 2000; Lee and Song, 2004). For methodological researchers in SEMs, technical details that are necessary in developing the theory and the MCMC methods are given in the appendices to the chapters. Applied researchers who are not interested in the methodological developments can skip those appendices. For convenience, we will introduce the freely available software WinBUGS (Spiegelhalter, et al., 2003) through analyses of simulated and real data sets. This software is able to produce reliable Bayesian statistics including the Bayesian estimates and their standard error estimates for a wide range of statistical models (Congdon, 2003) and for SEMs (Lee, 2007).

## **1.5** Real data sets and notation

We will use several real data sets for the purpose of motivating the models and illustrating the proposed Bayesian methodologies. These data sets are respectively related to the studies about: (i) job and life satisfaction, work attitude, and other related social-political issues; (ii) effects of some phenotype and genotype explanatory latent variables on kidney disease for type 2 diabetic patients; (iii) quality of life for residents of several countries, and for stroke patients; (iv) the development of and findings from an AIDS preventative intervention for Filipina commercial sex workers; (v) the longitudinal characteristics of cocaine and polydrug use; (vi) the functional relationships between bone mineral density (BMD) and its observed and latent determinants for old men; and (vii) academic achievement and its influential factors for American youth. Some information on these data sets is given in Appendix 1.1.

In the discussion of various models and their associated statistical methods, we will encounter different types of observations in relation to observable continuous and discrete variables or covariates; unobservable measurements in relation to missing data or continuous measurements underlying the discrete data; latent variables; as well as different types of

Table 1	.1 Ty	pical	notation.
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Symbol	Meaning
ω	Latent vector in the measurement equation
η	Outcome (dependent) latent vector in the structural equation
ξ	Explanatory (independent) latent vector in the structural equation
ε, δ	Random vectors of measurement errors
Λ	Factor loading matrix in the measurement equation
<b>B</b> , <b>Π</b> , <b>Γ</b> , $\Lambda_{\omega}$	Matrices of regression coefficients in the structural equation
Φ	Covariance matrix of explanatory latent variables
$\Psi_\epsilon, \Psi_\delta$	Diagonal covariance matrices of measurement errors, with diagonal elements $\psi_{\epsilon k}$ and $\psi_{\delta k}$ , respectively
$lpha_{0\epsilon k},eta_{0\epsilon k},\ lpha_{0\delta k},eta_{0\delta k}$	Hyperparameters in the gamma distributions of $\psi_{\epsilon k}$ and $\psi_{\delta k}$
$\mathbf{R}_0, \rho_0$	Hyperparameters in the Wishart distribution related to the prior distribution of $\Phi$
$\mathbf{\Lambda}_{0k}, \mathbf{H}_{0yk}$	Hyperparameters in the multivariate normal distribution related to the prior distribution of the <i>k</i> th row of $\Lambda$ in the measurement equation
$\mathbf{\Lambda}_{0\omega k}, \mathbf{H}_{0\omega k}$	Hyperparameters in the multivariate normal distribution related to the prior distribution of the <i>k</i> th row of $\Gamma$ in the structural equation
$\mathbf{I}_q$	A $q \times q$ identity matrix; sometimes we just use I to denote an identity matrix if its dimension is clear.

parameters, such as thresholds, structural parameters in the model, and hyperparameters in the prior distributions. Hence, we have a shortage of symbols. If the context is clear, some Greek letters may serve different purposes. For example,  $\alpha$  has been used to denote an unknown threshold in defining an ordered categorical variable, and to denote a hyperparameter in some prior distributions. Nevertheless, some general notation is given in Table 1.1.

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# **Appendix 1.1** Information on real data sets

# Inter-university Consortium for Political and Social Research (ICPSR) data

The ICPSR data set was collected in the World Values Survey 1981–1984 and 1990–1993 project (World Values Study Group, ICPSR Version). The whole data set consists of answers to a questionnaire survey about work attitude, job and family life, religious belief, interest in politics, attitude towards competition, etc. The items that have been used in the illustrative examples in this book are given below.

Thinking about your reasons for doing voluntary work, please use the following five-point scale to indicate how important each of the reasons below have been in your own case (1 is unimportant and 5 is very important).

V 62 Religious beliefs 1 2 3 4 5

During the past few weeks, did you ever feel ... (Yes: 1; No: 2)

V 89	Bored	1	2	2									
V 91	Depres	sed of	r very	unha	прру	1	2						
V 93	Upset b	becaus	se son	neboo	dy criti	icized :	you	1	2				
V 96	All thin	ngs co	nside	red, ł	now sa	tisfied	are you	u with	your	life as a w	hole th	nese days	s?
	1	2	3	4	5	6	7	8	9	10			
	Dissatis	fied								Satisfied			

Here are some aspects of a job that people say are important. Please look at them and tell me which ones you personally think are important in a job. (Mentioned: 1; Not Mentioned: 2)

V 99	Good	Pay	1	2							
V 100	Pleas	sant pe	ople to	o wor	k witl	h 1		2			
V 102	Good	l job s	ecurity	/	1	2					
V 103	Good	l chan	ces for	pron	notior	n 1	2	2			
V 111	A res	sponsil	ole job	)	1	2					
V 115 How much pride, if any, do you take in the work that you do?											
	1	A gr	eat de	al	2	Som	e	3	Little	4	None
V 116	Over								vith you	r job?	
	1	2	3	4	5	6	7	8	9	10	
]	Dissati	sfied								Satisf	ied
V 117	How	free a	re you	to m	ake de	ecision	s in y	our jo	b?		
	1	2	3	4	5	6	7	8	9	10	
I	Not at	all							А	great	deal
V 129	Whe	n jobs	are sca	arce,	peopl	e shoul	d be f	forced	l to retir	e early	ζ,
	1	Agre	ee,	2	Nei	ther,	3	D	isagree		
V 132	How	satisfi	ed are	you	with t	he fina	ncial	situat	ion of ye	our ho	usehold?
	1	2	3	4	5	6	7	8	9	10	
1	Dissati	sfied								Satisf	ied

V 176	76 How important is God in your life? 10 means very important and											
	1 me	eans n	ot at al	ll imp	ortant							
	1	2	3	4	5	6	7	8	9	10		
V 179	How	often	do you	u pray	to Go	od outs	ide of	religio	ous ser	vices?	Would you say	
	1	Ofte	en		2	Some	times					
	3	Har	dly eve	er	4	Only	in tim	es of c	risis	5	Never	
V 180	Over	all, ho	ow sati	sfied o	or diss	atisfie	d are y	ou wit	h you	r home	life?	
	1	2	3	4	5	6	7	8	9	10		
Ι	Dissati	sfied								Satisfi	ied	
V 241	How	intere	ested w	vould	you sa	iy you	are in	politic	s?			
	1	Very	y intere	ested		2	Som	lewhat	intere	sted		
	3	Not	very in	nteres	ted	4	Not	Not at all interested				

Now I'd like you to tell me your views on various issues. How would you place your views on this scale? 1 means you agree completely with the statement on the left, 10 means you agree completely with the statement on the right, or you can choose any number in between.

V 252										
	1	2	3	4	5	6	7	8	9	10
Individu more res providin	sponsi	bility	for							The state should take more responsibility to ensure that everyone
P	8									is provided for.
V 253										-
	1	2	3	4	5	6	7	8	9	10
People w should ha available unemplo	ave to or los	take a se thei	iny jol r					sł	nould	e who are unemployed I have the right to refuse hey do not want.
V 254										
	1	2	-	4	5	6	7	8	9	10
Competi		-							-	etition is harmful. It
stimulate	-	-						br	ings	out the worst in people.
hard and	l deve	lop ne	w idea	as.						
V 255										
	1	2	3	4	5	6	7	8	9	10
In the lo usually b	-							b	ring	work doesn't generally success – it's more a r of luck and connections.

Please tell me for each of the following statements whether you think it can always be justified, never be justified, or something in between.

V 296	Clai	ming	govern	ment	benefits	whi	ch you	are no	ot enti	tled to	
	1	2	3	4	5	6	7	8	9	10	
	Never									Always	

APPENDIX 9

V 297	Avo	iding a	fare	on pub	lic tra	insport					
	1	2	3	4	5	6	7	8	9	10	
Ν	Vever									Always	
V 298	Chea	ating o	n tax	if you	have	the cha	nce				
	1	2	3	4	5	6	7	8	9	10	
Ν	Vever									Always	
V 314 Failing to report damage you've done accidentally to a parked vehicle											
	1	2	3	4	5	6	7	8	9	10	
Ν	Jever									Always	
I am going to read out some statements about the government and the economy. For each one,											
could you t	ell me	e how	much	you ag	gree o	r disag	ree?				
V 336	Our	govern	nment	should	l be n	nade m	uch m	ore op	en to	the public	
	1	2	3	4	5	6					
Ag	gree C	Comple	tely		Γ	Disagree	e Com	pletely	/		
V 337	We a	are mo	re like	ely to h	ave a	health	y econ	iomy i	f the	government allows	
	mor	e freed	lom fo	or indiv	vidual	s to do	as the	y wish	l I		
	1	2	3	4	5	6					
Ag	gree C	Comple	tely		Γ	Disagree	e Com	pletely	/		
V 339	Poli	tical re	form	in this	count	try is m	noving	too ra	pidly	,	
	1	2	3	4	5	6					
Ag	gree C	Comple	etely		Γ	Disagree	e Com	pletely	/		

## Type 2 diabetic patients data

The data set was collected from an applied genomics program conducted by the Institute of Diabetes, the Chinese University of Hong Kong. It aims to examine the clinical and molecular epidemiology of type 2 diabetes in Hong Kong Chinese, with particular emphasis on diabetic nephropathy. A consecutive cohort of 1188 type 2 diabetic patients was enrolled into the Hong Kong Diabetes Registry. All patients underwent a structured 4-hour clinical and biochemical assessment including renal function measured by plasma creatine (PCr) and urinary albumin creatinine ratio (ACR); continuous phenotype variables such as systolic blood pressure (SBP), diastolic blood pressure (DBP), body mass index (BMI), waist–hip ratio (WHR), glycated hemoglobin (HbA1c), fasting plasma glucose (FPG), non-high-density lipoprotein cholesterol (non-HDL-C), lower-density lipoprotein cholesterol (LDL-C), plasma triglyceride (TG); and multinomial genotype variables such as beta-3 adrenergic receptor (ADR $\beta$ 3), beta-2 adrenergic receptor SNP1 (ADR $\beta$ 21), beta-2 adrenergic receptor SNP2 (ADR $\beta$ 22), angiotensin converting enzyme (DCP1 intro 16 del/ins (DCP1)), and angiotensin II receptor type 1 AgtR1 A1166C (AGTR1).

## WHOQOL-BREF quality of life assessment data

The WHOQOL-100 assessment was developed by the WHOQOL group in 15 international field centers for the assessment of quality of life (QOL). The WHOQOL-BREF instrument is a short version of WHOQOL-100 consisting of 24 ordinal categorical items selected from the 100 items. This instrument was established to evaluate four domains: physical health, mental health, social relationships, and environment. The instrument also includes two ordinal categorical items for overall QOL and health-related QOL, giving a total of

26 items. All of the items are measured on a 5-point scale (1 = 'not at all/very dissatisfied'; 2 = 'a little/dissatisfied'; 3 = 'wery much/satisfied'; 5 = 'extremely/very satisfied'). The frequencies of the ordinal scores of the items are as follows:

		O	Number of			
WHOQOL item	1	2	3	4	5	incomplete obs.
Q1 Overall QOL	3	41	90	233	107	1
Q2 Overall health	32	127	104	154	58	0
Q3 Pain and discomfort	21	65	105	156	127	1
Q4 Medical treatment dependence	21	57	73	83	239	2
Q5 Energy and fatigue	15	57	166	111	118	8
Q6 Mobility	16	36	58	120	243	2
Q7 Sleep and rest	28	87	95	182	83	0
Q8 Daily activities	7	73	70	224	100	1
Q9 Work capacity	19	83	88	191	91	3
Q10 Positive feelings	2	30	141	241	59	2
Q11 Spirituality/personal beliefs	13	45	149	203	61	4
Q12 Memory and concentration	4	40	222	184	21	4
Q13 Bodily image and appearance	9	46	175	137	106	2
Q14 Self-esteem	13	72	130	210	50	0
Q15 Negative feelings	4	54	137	239	39	2
Q16 Personal relationships	8	46	68	218	134	1
Q17 Sexual activity	25	55	137	149	76	33
Q18 Social support	2	23	84	228	136	2
Q19 Physical safety and security	2	25	193	191	62	2
Q20 Physical environment	4	29	187	206	43	6
Q21 Financial resources	27	56	231	105	54	2
Q22 Daily life information	5	27	176	194	70	3
Q23 Participation in leisure activity	10	99	156	163	47	0
Q24 Living conditions	9	27	53	235	151	0
Q25 Health accessibility and quality	0	17	75	321	61	1
Q26 Transportation	8	38	61	253	113	2

## **AIDS preventative intervention data**

The data set was collected from female commercial sex workers (CSWs) in 95 establishments (bars, night clubs, karaoke TV and massage parlours) in cities in the Philippines. The whole questionnaire consists of 134 items on areas of demographic knowledge, attitudes, beliefs, behaviors, self-efficacy for condom use, and social desirability. The primary concern is finding an AIDS preventative intervention for Filipina CSWs. Questions are as follows:

(1) How much of a threat do you think AIDS is to the health of people? no threat at all/very small/moderate/strong/very great

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- (2) What are the chances that you yourself might get AIDS? none/very small/moderate/great/very great
- (3) How worried are you about getting AIDS? not worried/slightly/moderate/very/extremely

How great is the risk of getting AIDS or the AIDS virus from sexual intercourse with someone:

- (4) Who has the AIDS virus using a condom? none/very small/moderate/great/very great
- (5) Whom you don't know very well without using a condom? none/very small/moderate/great/very great
- (6) Who injects drugs? none/very small/moderate/great/very great
- (7) How often did you perform vaginal sex in the last 7 days?
- (8) How often did you perform manual sex in the last 7 days?
- (9) How often did you perform oral sex in the last 7 days?
- (10) Have you ever used a condom? Yes/No
- (11) Did you use a condom the last time you had sex? Yes/No
- (12) Have you ever put a condom on a customer? Yes/No
- (13) Do you agree or disagree that condoms make sex less enjoyable? strongly agree/agree/neutral/disagree/strongly disagree
- (14) Do you agree or disagree that condoms cause a man to lose his erection? strongly agree/agree/neutral/disagree/strongly disagree
- (15) Do you agree or disagree that condoms cause pain or discomfort? strongly agree/agree/neutral/disagree/strongly disagree
- (16) Are condoms available at your establishment for the workers who work there? Yes/No
- (17) How much do you think you know the disease called AIDS? nothing/a little/somewhat/moderate/a great deal
- (18) Have you ever had an AIDS test? Yes/No

#### Polydrug use and treatment retention data

This is a longitudinal study of polydrug use initiated by California voters and conducted in five California counties in 2004. Proposition 36 directs drug offenders to a communitybased drug treatment to reduce drug abuse using proven and effective treatment strategies. One of the objectives of the study is to examine why court-mandated offenders drop out of the drug treatment and to compare the characteristics, treatment experiences, perceptions, and outcomes of treatment completers (see Evans *et al.*, 2009). Data were collected from self-reported and administrative questionnaires about the retention of drug treatment (i.e.

the days of stay in treatment), drug use history, drug-related crime history, and service and test received for 1588 participants at intake, 3-month, and 12-month follow-up interviews. In addition, variables about treatment motivation (Mtsum01, Mtsum02, and Mtsum03) were collected at intake. Variables include:

- (1) Drgplm30: Drug problems in past 30 days at intake, which ranges from 0 to 30.
- (2) Drgday30: Drug use in past 30 days at intake, which ranges from 0 to 30.
- (3) DrgN30: The number of kinds of drugs used in past 30 days at intake, which ranges from 1 to 8.
- (4) Incar: The number of incarcerations in lifetime at intake, which ranges from 0 to 216.
- (5) ArrN: The number of arrests in lifetime at intake, which ranges from 1 to 115.
- (6) Agefirstarrest: The age of first arrest, which ranges from 6 to 57.
- (7) Retent: Days of stay in treatment or retention, which ranges from 0 to 365.
- (8) M12drg30: Primary drug use in past 30 days at 12-month interview, which ranges from 1 to 5.
- (9) Servicem: Services received in past 3 months at TSI 3-month interview.
- (10) DrugtestTX: The number of drug tests by TX in past 3 months at TSI 3-month interview, which ranges from 0 to 36.
- (11) DrugtestCJ: The number of drug tests by criminal justice in past 3 months at TSI 3-month interview, which ranges from 0 to 12.
- (12) Mtm01: Motivation subscale 1 at intake, which ranges from 1 to 5.
- (13) Mtm02: Motivation subscale 2 at intake, which ranges from 1 to 5.
- (14) Mtm03: Motivation subscale 3 at intake, which ranges from 1 to 5.

## Quality of life for stroke survivors data

The setting for this study was the Prince of Wales Hospital (PWH) in Hong Kong which is a regional university hospital with 1500 beds serving a population of 0.7 million people. Patients with acute stroke within 2 days of admission were identified and followed up at 3, 6, and 12 months post stroke. All patients included in the study were ethnic Chinese. As the aim was to study those with a first disabling stroke, patients were excluded if they had moderate or severe premorbid handicap level (Rankin Scale score greater than 2). Outcome measures are obtained from questionnaires, which respectively measure respondents' functional status, depression, health-related quality of life, and handicap situation, including (1) modified Barthel Index (MBI) score, (2) Geriatric Depression Scale (GDS) score, (3) Chinese Mini-Mental State Examination (MMSE) score, (4) World Health Organization Quality of Life measure (abbreviated Hong Kong version) (WHOQOL BREF (HK)) score, and (5) the London Handicap Scale (LHS) score.

APPENDIX 13

## Cocaine use data

This data set was obtained from a longitudinal study about cocaine use conducted at the UCLA Center for Advancing Longitudinal Drug Abuse Research. The UCLA Center collected various measures from patients admitted in 1988–1989 to the West Los Angeles Veterans Affairs Medical Center and met the DSM III-R criteria for cocaine dependence. The cocaine-dependent patients were assessed at baseline, 1 year after treatment, 2 years after treatment, and 12 years after treatment in 2002. Measures at each time point include the following:

- cocaine use (CC), an ordered categorical variable coded 1 to 5 to denote days of cocaine use per month that are fewer than 2 days, 2–7 days, 8–14 days, 15–25 days, and more than 25 days, respectively;
- (2) Beck inventory (BI), an ordered categorical variable coded 1 to 5 to denote scores that are less than 3.0, between 3.0 and 8.0, between 9.0 and 20.0, between 21 and 30, and greater than 30;
- (3) depression (DEP), an ordered categorical variable based on the Hopkins Symptom Checklist-58 scores, coded 1 to 5 to denote scores that are less than 1.1, between 1.1 and 1.4, between 1.4 and 1.8, between 1.8 and 2.5, and greater than 2.5;
- (4) number of friends (NF), an ordered categorical variable coded 1 to 5 to denote no friend, 1 friend, 2–4 friends, 5–8 friends, and more than 9 friends;
- (5) 'have someone to talk to about problem (TP)', {0, 1} for {No, Yes};
- (6) 'currently employed (EMP)', {0, 1} for {No, Yes};
- (7) 'alcohol dependence (AD) at baseline', {0, 1} for {No, Yes}.

## Bone mineral density data

This data set was collected from a partial study on osteoporosis prevention and control. The study concerned the influence of serum concentration of sex hormones, their precursors and metabolites on bone mineral density (BMD) in older men. It was part of a multicenter prospective cohort study of risk factors of osteoporotic fractures in older people. A total of 1446 Chinese men aged 65 years and older were recruited using a combination of private solicitation and public advertising from community centers and public housing estates.

The observed variables include: spine BMD, hip BMD, estrone (E1), estrone sulfate (E1-S), estradiol (E2), testosterone (TESTO), 5-androstenediol (5-DIOL), dihydrotestosterone (DHT), androstenedione (4-DIONE), dehydroepiandrosterone (DHEA), DHEA sulfate (DHEA-S), androsterone (ADT), ADT glucuronide (ADT-G),  $3\alpha$ -diol-3G (3G), and  $3\alpha$ -diol-17G (17G). Weight and age were also measured.

## National longitudinal surveys of youth (NLSY) data

The four-decade-long NLSY is one of the most comprehensive longitudinal studies of youths conducted in North America. The NLSY data include a nationally representative sample of youths who were 14–21 years old in 1979 and 29–36 years old in 1994.

The data set derived for the illustrative examples in this book includes 1660 observations and the following measures: the Peabody Individual Achievement Tests (PIAT) with continuous scales in the three domains of math, reading recognition, and reading comprehension; the Behavior Problem Index (BPI) with an ordinal scale in the five domains of anti-social, anxious, dependent, headstrong, and hyperactive behavior; home environment in the three domains of cognitive stimulation, emotional support, and household conditions; and friendship in the two domains of the number of boyfriends and the number of girlfriends. The instruments for measuring these constructs were taken from a short form of Home Observation for Measurement of the Environment (HOME) Inventory.

## References

- Bentler, P. M. and Stein, J. A. (1992) Structural equation models in medical research. *Statistical Methods in Medical Research*, 1, 159–181.
- Bollen, K. A. (1989) Structural Equations with Latent Variables. New York: John Wiley & Sons, Inc.
- Congdon, P. (2003) Applied Bayesian Modelling. Chichester: John Wiley & Sons, Ltd.
- Dolan, C. V. and van der Maas, H. L. J. (1998) Fitting multivariage normal finite mixtures subject to structural equation modeling. *Psychometrika*, **63**, 227–253.
- Dunson, D. B. (2000) Bayesian latent variable models for clustered mixed outcomes. Journal of the Royal Statistical Society, Series B, 62, 355–366.
- Dunson, D. B. (2003) Dynamic latent trait models for multidimensional longitudinal data. *Journal of the American Statistical Association*, 98, 555–563.
- Evans, E., Li, L. and Hser, Y. I. (2009) Client and program factors associated with dropout from court mandated drug treatment. *Evaluation and Program Planning*, **32**, 204–212.
- Jöreskog, K. G. and Sörbom, D. (1996) LISREL 8: Structural Equation Modeling with the SIMPLIS Command Language. Hove: Scientific Software International.
- Jamshidian, M. and Bentler, P. M. (1999) ML estimation of mean and covariance structures with missing data using complete data routines. *Journal of Educational and Behavioral Statistics*, **24**, 21–41.
- Lee, S. Y. (2007) *Structural Equation Modeling: A Bayesian Approach*. Chichester: John Wiley & Sons, Ltd.
- Lee, S. Y. and Shi, J. Q. (2001) Maximum likelihood estimation of two-level latent variable models with mixed continuous and polytomous data. *Biometrics*, **57**, 787–794.
- Lee, S. Y. and Song X. Y. (2003a) Model comparison of nonlinear structural equation models with fixed covariates. *Psychometrika*, **68**, 27–47.
- Lee, S. Y. and Song, X. Y. (2003b) Maximum likelihood estimation and model comparison for mixtures of structural equation models with ignorable missing data. *Journal of Classification*, 20, 221–255.
- Lee, S. Y. and Song, X. Y. (2004) Evaluation of the Bayesian and maximum likelihood approaches in analyzing structural equation models with small sample sizes. *Multivariate Behavioral Research*, 39, 653–686.
- Lee, S. Y. and Song, X. Y. (2005) Maximum likelihood analysis of a two-level nonlinear structural equation model with fixed covariates. *Journal of Educational and Behavioral Statistics*, **30**, 1–26.
- Lee, S. Y. and Tang, N. S. (2006) Bayesian analysis of nonlinear structural equation models with nonignorable missing data. *Psychometrika*, **71**, 541–564.
- Lee, S. Y., Song, X. Y., Skevington, S. and Hao, Y. T. (2005) Application of structural equation models to quality of life. *Structural Equation Modeling A Multidisciplinary Journal*, **12**, 435–453.

- Lee, S. Y., Lu, B. and Song, X. Y. (2008) Semiparametric Bayesian analysis of structural equation models with fixed covariates. *Statistics in Medicine*, 27, 2341–2360.
- Moustaki, I. (2003) A general class of latent variable models for ordinal manifest variables with covariate effects on the manifest and latent variables. *British Journal of Mathematical and Statistical Psychology*, **56**, 337–357.
- Pugesek, B. H., Tomer, A. and von Eye, A. (2003) Structural Equation Modeling: Applications in Ecological and Evolutionary Biology, Cambridge: Cambridge University Press.
- Rabe-Hesketh, S., Skrondal, A. and Pickles, A. (2004) Generalized multilevel structural equation modeling. *Psychometrika*, 69, 167–190.
- Sánchez, B. N., Budtz-Jorgensen, E., Ryan L, M. and Hu, H. (2005) Structural equation models: a review with applications to environmental epidemiology. *Journal of the American Statistical Association*, 100, 1443–1455.
- Schumacker, R. E. and Marcoulides, G. A. (1998) Interaction and Nonlinear Effects in Structural Equation Modeling. Mahwah, NJ: Lawrence Erlbaum Associates.
- Shi, J. Q. and Lee, S. Y. (2000) Latent variable models with mixed continuous and polytomous data. *Journal of the Royal Statistical Society, Series B*, 62, 77–87.
- Song, X. Y. and Lee, S. Y. (2004) Bayesian analysis of two-level nonlinear structural equation models with continuous and polytomous data. *British Journal of Mathematical and Statistical Psychology*, 57, 29–52.
- Song, X. Y. and Lee, S. Y. (2006) A maximum likelihood approach for multisample nonlinear structural equation models with missing continuous and dichotomous data. *Structural Equation Modeling – A Multidisciplinary Journal*, 13, 325–351.
- Song, X. Y. and Lee, S. Y. (2007) Bayesian analysis of latent variable models with non-ignorable missing outcomes from exponential family. *Statistics in Medicine*, 26, 681–693.
- Song, X. Y. and Lu, Z. H. (2010) Semiparametric latent variable models with Bayesian P-splines. *Journal of Computational and Graphical Statistics*, 19, 590–608.
- Song, X. Y. and Lu, Z. H. (2012) Semiparametric transformation models with Bayesian P-splines. *Statistics and Computing*, to appear.
- Song, X. Y., Lee, S. Y., Ng, M. C. Y., So, W. Y. and Chan, J. C. N. (2007) Bayesian analysis of structural equation models with multinomial variables and an application to type 2 diabetic nephropathy. *Statistics in Medicine*, 26, 2348–2369.
- Song, X. Y., Lee, S. Y. and Hser, Y. I. (2008) A two-level structural equation model approach for analyzing multivariate longitudinal responses. *Statistics in Medicine*, 27, 3017–3041.
- Song, X. Y., Xia, Y. M. and Lee, S. Y. (2009) Bayesian semiparametric analysis of structural equation models with mixed continuous and unordered categorical variables. *Statistics in Medicine*, 28, 2253–2276.
- Spiegelhalter, D. J., Thomas, A., Best, N. G. and Lunn, D. (2003) *WinBugs User Manual. Version 1.4*. Cambridge: MRC Biostatistics Unit.
- van Montfort, K., Mooijaart, A. and Meijerink, F. (2009) Estimating structural equation models with non-normal variables by using transformations. *Statistica Neerlandica*, **63**, 213–226.
- Wedel, M. and Kamakura, W. A. (2001) Factor analysis with (mixed) observed and latent variables in the exponential family. *Psychometrika*, **66**, 515–530.
- Yang, M. G. and Dunson, D. B. (2010) Bayesian semiparametric structural equation models with latent variables. *Psychometrika*, **75**, 675–693.
- Zhu, H. T. and Lee, S. Y. (2001) A Bayesian analysis of finite mixtures in the LISREL model. *Psychometrika*, **66**, 133–152.