'Translating back' common statistical test into their graphical causal language ancestors: adding practical statistical codes to the Jaccard & Jacoby 'tests-causal' model crosswalk

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Promisory note:

I will add open source R code in *daggity* and *lavaan* and *Onyx* to give practical (and testable) meaning to theoretical causal models behind common statistical analyses (as shown in Jaccard & Jacoby, 2009: 1.i. Appendix): t-test; [ANOVA; Pearson chi-square;] regression; mediation; [causal mediation, and more. We will tackle also how and where to 'add time' in causal models].

Actually: just 1 variable and 2 variable models today, sorry.

Refer for more to: Jaccard, J., & Jacoby, J. (2009). <u>Theory construction and model-building skills: A practical guide for social scientists</u>: Guilford Press.

1.i. APPENDIX: Inferring Theoretical Relationships from the Choice of Statistical

<u>Tests</u>

This is an applied worksession (partly), so:

For R codes and data for illustrations go to:

FIGURE 12.1. Causal models underlying statistical tests (text example on left, generic form on right). (a) Two Group/Condition *t*-Test; (b) One-Way Analysis of Variance; (c) Chi-Square Test of Independence and Test of Proportions; (d) Pearson Correlation/Linear Regression: Direct Cause Model;

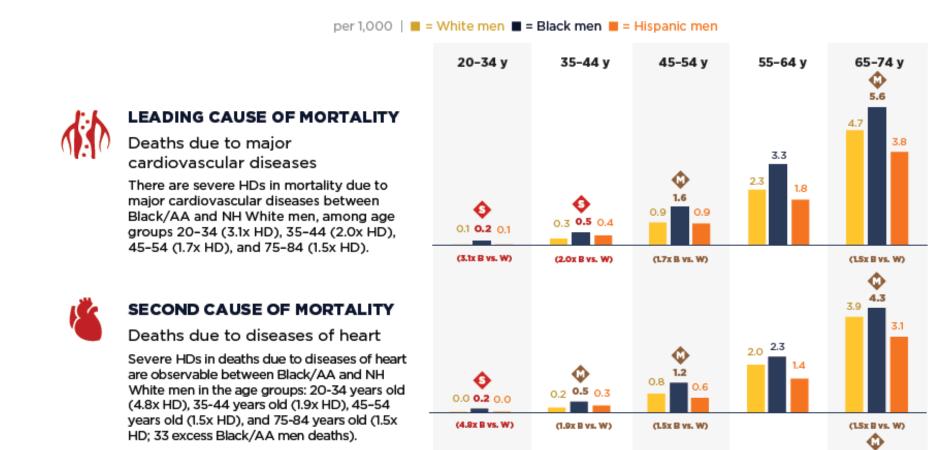
CONCLUDING ISSUES 348 Attitude Toward Gender Abortion --> contE binaryC Family Size b Religion Attitudes see above* С Political Party Gender Identification binaryC --> binary*E Money Donated Religiosity contC contE

Main points

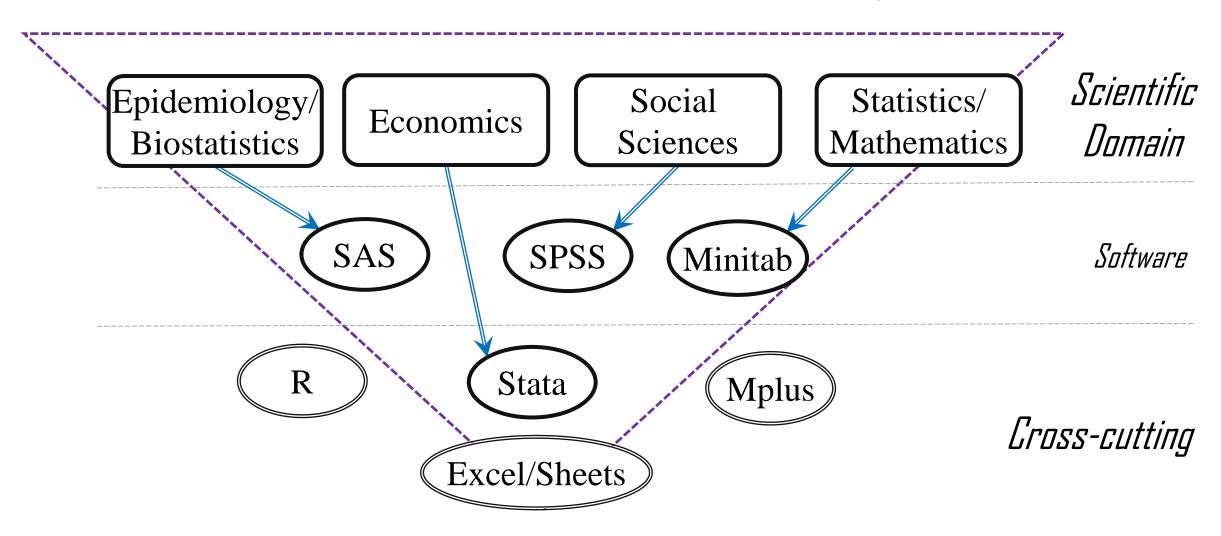
- 1. Statistical tests are *clueless* without a causal model;
 - Research questions need: info about research design and on a working model of how data is generated by nature (with researcher's or policy maker's input sometimes).
 - 2. In Null Hypothesis Significance Testing (NHST) framework: one needs a model for the 'null' state.
- 2. Instead of mere NHST from statistical tests, the focus shifts to causally well-informed models and tests of differences in fit between alternative/nested models.
- 3. The easiest way to model is graphically: will show how with *lavaan* and Ωnyx parametrically (and with data), and with *dagitty* and *MIIVsem* nonparametrically (data-less).

Who we are:

- -HDI: UConn Health Disparities Institute
- -BMoC: CT Report Card on Health Equity Among Boys And Men Of Color (BMoC) in CT: <u>full</u> <u>report</u>; <u>interview on WNPR</u> with Dr. Wizdom Powell
- -Biostatistics club UConn Health formerly CICATS, now Connecticut Convergence Institute for Translation in Regenerative Engineering



Statistical and software traditions - simplified



'Tracing rule' powers at work with 1 variable

We add σ 's, and run the 'tracing' tool along directed and bidirectional paths (except common effects along the way; and we can 'go back in time) + grab the cov/ariance of the variable at the 'root' on the pathway.



$$Cov(YY) = \sigma_{YY}$$

An <u>exogenous</u> variable (to the model!) is a 'big fat disturbance' or structural error: all of its variance is error, or unexplained. So:

$$Y = 1 \cdot u$$
 and therefore

$$\sigma_{YY} = \sigma_{uu}$$

"The correlation between two variables can be shown to equal the sum of the products of the chains of path coefficients along all of the paths by which the variables are connected." [:115]

Regression and how potential 'change' is deduced

Regression
$$\begin{array}{c}
U \\
& & \\
& & \\
X & & \\
\end{array}$$
 $X \longrightarrow \frac{\sigma_{XY}}{\sigma_{XY}} \longrightarrow Y$

With deviation scores one gets $\alpha_{Y} = 0$.

Notation: u is better here than ϵ because it represents 'ignored-for-now-other-causes', not just 'error'.

$$Y_i = \beta_{YX} \cdot X_i + 1 \cdot u_i$$
 [easier if $\alpha_Y = 0$]

Hence if one multiplies by X_i :

$$X_{i} \cdot Y_{i} = \beta_{YX} \cdot X_{i} \cdot X_{i} + \cdot X_{i} \cdot u_{i}$$

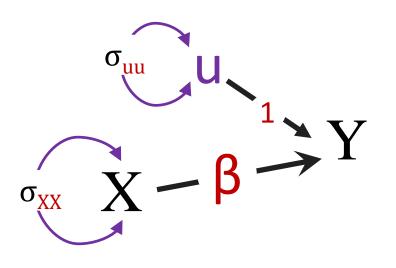
Sum across N (sample cases) & divide by N:

$$\frac{\sum_{i}^{N} X_{\mathbf{i}} \cdot Y_{\mathbf{i}}}{N} = \beta_{YX} \cdot \frac{\sum_{i}^{N} X_{\mathbf{i}} \cdot X_{\mathbf{i}}}{N} + \frac{\sum_{i}^{N} X_{\mathbf{i}} \cdot u_{\mathbf{i}}}{N} \qquad \qquad \text{Hence}:$$

$$\sigma_{vv} = \beta_{vv} \cdot \sigma_{vv} + \sigma_{vv}$$
 So:

$$\beta_{YX} = \frac{\sigma_{YX}}{\sigma_{XX}} - \sigma_{Xu} = \frac{Cov(Y,X)}{Cov(X,X)} - Cov(X,u)$$

'Tracing rule' powers at work



Cov(YX) is "sum of products path/structural coefficients, of all open pathways from X to Y":

$$\mathsf{Cov}(\mathsf{YX}) \overset{\textit{notation}}{\Longleftrightarrow} \sigma_{\underline{\mathsf{YX}}} \overset{\textit{Wright Tracing Rule}}{\Longleftrightarrow} \sigma_{\underline{\mathsf{XX}}} \cdot \beta$$

Hence:

$$\beta = \frac{\sigma_{XY}}{\sigma_{vv}}$$
 Simpler?

"The correlation between two variables can be shown to equal the sum of the products of the chains of path coefficients along all of the paths by which the variables are connected." [Wright:115]

'Tracing rule'

"Tracing rules (or Wright's rules) are simply a way to estimate the covariance between two variables by summing the appropriate connecting paths." [1:23]

- "Trace all paths between two variables (or a variable back to itself), multiplying all the coefficients along a given [open] path.
- i. You can start by going backwards along a single-headed arrow, but once you start going forward along these arrows you can no longer go backwards.
- ii. No loops! That is, you cannot go through the same variable more than once for a given path.
- iii. At maximum, there can be one double-headed arrow included in a path.
- iv. After tracing all the paths for a given relationship, sum all the paths." [1:24]
- +v. EC: A 'collider (z_1 --> w <-- z_2) is: not open/closed/disconnected
- Cannot go forward THEN backwards 'in time'

"The correlation between two variables can be shown to equal the sum of the products of the chains of path coefficients along all of the paths by which the variables are connected." [2:115]

^{1.} Beaujean, A. A. (2014). <u>Latent variable modeling using R</u>: A step-by-step guide: Routledge.

^{2.} Wright, S. (1921). Systems of mating. I. The biometric relations between parent and offspring. Genetics, 6(2), 111.

Original translational insight

Not clear which came first:

- 1. The decomposition idea
- 2. The regression->path conceptual leap.

kept unchanged, to the total standard deviation. A path coefficient differs from a coefficient of correlation in having direction.

The symbol $p_{X\cdot A}$ means the coefficient for the path of influence from A to X. In most cases in the present paper, however, it will be more convenient to represent the path coefficients by single letters.

It can be shown that the squares of the path coefficients measure the degree of determination by each cause. If the causes are independent of each other, the sum of the squared path coefficients is unity. If the causes are correlated, terms representing joint determination must be recognized. The complete determination of X in figure 1 by factor A and the correlated factors B and C can be expressed by the equation:

$$(1) a^2 + b^2 + c^2 + 2 bc r_{BC} = 1$$

The squared path coefficients and the expressions for joint determination measure the portion of the squared standard deviation of the effect due to the causes singly and jointly, respectively.

The correlation between two variables can be shown to equal the sum of the products of the chains of path coefficients along all of the paths by which the variables are connected. In figure 1, X and Y are connected by four paths.

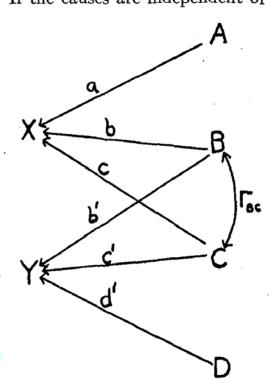


FIGURE 1.—A diagram illustrating the case of two variables (X and Y) determined in part by causes in common (B and C) which are correlated with each other.

$$r_{XY} = bb' + cc' + br_{BC}c' + cr_{BC}b'$$

First SEM model: 1920

"The path coefficient, measuring the importance of a given path of influence from cause to effect, is defined as the ratio of the variability of the effect to be found when all causes are constant except the one in question, the variability of which is kept unchanged, to the total variability.

Variability is measured by the standard deviation." [1]:329

Diagram illustrating the casual relations between litter mates (O, O') and between each of them and their parents. H, H', H'', H,''' represent the genetic constitutions of the four individuals, G, G', G'', and G''' that of four germ cells. E represents such environmental factors as are common to litter mates. D represents other factors, largely ontogenetic irregularity. The small letters stand for the various path coefficients.

•1. Wright, S. (1920). The relative importance of heredity and environment in determining the piebald pattern of guinea-pigs. *Proceedings of the National Academy of Sciences*, 6(6), 320-332.

Sire Chance Dam FIG. 5.

Original insight

The 'multiply- along-the-trek' rule may have come from the derivatives of composite functions formulas:

³Pearl (2013) calls them nested counterfactuals; the key insight Sewall Wright foresaw when proposing the path analytic method may have been that the change in Y in relation to the change in X (the slope $\delta Y/\delta X$), traced on the path through an intermediary M, is linked to the slopes $\delta M/\delta X$ and $\delta Y/\delta M$ following the composite function chain rule of derivatives: $\delta y/\delta x = \delta y/\delta M \cdot \delta m/\delta x$, which mirrors the Baron and Kenny $i = a \cdot b$. Adding the contributions of all such X-to-Y open paths yields the model predicted association between X and Y (see the "tracing rule," Loehlin, 2004).

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Origins of the Bayesian Networks

"Networks employing Directed Acyclic Graphs (DAGs) have a long and rich tradition, starting with the geneticist Wright (1921). He developed a method called path analysis [Wright, 1934] which later on, became an established representation of causal models in economics [Wold, 1964], sociology [Blalock, 1971] and psychology [Duncan, 1975]. Influence diagrams represent another application of DAG representation [Howard and Matheson, 1981], [Shachter, 1988] and [Smith, 1987]. These were developed for decision analysis and contain both chance nodes and decision nodes (our definition of causal models excludes decision nodes). Recursive models is the name given to such networks by statisticians seeking meaningful and effective decompositions of contingency tables (Lauritzen, 1982), (Wermuth & Lauritzen, 1983], [Kiiveri et al, 1984]. Bayesian Belief Networks (or Causal Networks) is the name adopted for describing networks that perform evidential reasoning ((Pearl, 1986a, 1988]). This paper establishes a clear semantics for these networks that might explain their wide usage as models for forecasting, decision analysis and evidential reasoning." [1]:136

Causal chains: early origins

"The ϕ 's may be thought of as quasiprobabilities of the events. They are defined successively as follows [2: 48]:

$$\Phi_0 = I$$
.

$$\Phi_1 = \pi_1.$$

$$\Phi_2 = I - (I - \pi_2) \cdot (I - \Phi_1 \cdot \pi_3)$$
.

$$\Phi_3 = I - (I - \Phi_1 \cdot \pi_4) \cdot (I - \Phi_2 \cdot \pi_5).$$

$$\Phi_4 = I - (I - \Phi_1 \cdot \pi_6) \cdot (I - \Phi_2 \cdot \pi_7) \cdot (I - \Phi_3 \cdot \pi_8)$$
.

$$S(\pi) = Q(\Phi_4, o) = -\log(I - \Phi_4).$$

^{1.} Good, I. J. (1961). A causal calculus (I). The British Journal for the Philosophy of Science, 11(44), 305-318.

^{2.} Good, I. J. (1961). A causal calculus (II). The British Journal for the Philosophy of Science, 12(45), 43-51.

Origins of the Bayesian Networks

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Probabilistic formulae of this kind are shorthand notation for the statement that for any instantiation i of the variables x_1, x_2, \ldots, x_n , the probability of the joint event $(x_1 = i) & (x_2 = i_2)$ &... & $(x_n = i_n)$ is equal to the product of the probabilities of the corresponding conditional events $(x_1 = i_1)$, $(x_2 = i_2 \text{ if } x_1 = i_1)$, $(x_3 = i_3 \text{ if } (x_2 = i_2 \&$ $x_1 = i_1$) For this expansion to be valid, we must require that P(E) > 0 for all conditioning events E." So, for example, the distribution corresponding to the graph of Fig. 1 can be written by inspection:

$$P(x_1, x_2, x_3, x_4, x_5, x_6) =$$

$$= P(x_6|x_5) \cdot P(x_6|x_5) \cdot P(x_5|x_2, x_3) \cdot$$

$$P(x_4|x_1, x_2) P(x_3|x_1) \cdot P(x_2|x_1) \cdot P(x_1) \text{ (Pearl 1986):244-5}$$

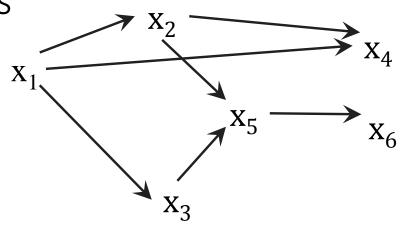


FIG. 1. A typical Bayesian network representing the distribution $P(x_1... x_6) = P(x_6|x_5) \cdot P(x_5|x_2, x_3) \cdot P(x_4|x_1, x_2) P(x_3|x_1) \cdot P(x_2|x_1) \cdot P(x_1)$

Modern Bayesian Networks

"In words, the qualitative statistical dependencies shown in this small Bayesian network can be described as follows:

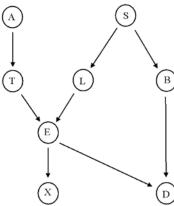
1. A recent trip to Asia (A) increases the chances of tuberculosis (T).

dyspnoea. " (Yedidia, Freeman et al. 2003):6

- 2. Smoking (S) is a risk factor for both lung cancer (L) and bronchitis (B).
- 3. The presence of either € tuberculosis or lung cancer can be detected by an X-ray result, but the X-ray alone cannot distinguish between them.
- 4. Dyspnoea (D) (shortness of breath) may be caused by bronchitis (B), or either € tuberculosis or lung cancer. Each node represents a variable that can be in a discrete number of possible states. We write x_i for the variable representing the different possible states of the node i. In addition to the qualitative dependencies described by the Bayesian network graph, there are quantitative statistical relationships that we assign to each arrow in the graph. Associated with each arrow is a conditional probability: for example, we write $p(x_L|x_S)$ for the conditional probability of a patient having lung cancer given that he does or does not smoke. For this link, we say that the S node is the 'parent' of the L node because x_L is conditionally dependent on x_S . Some nodes like the D node might have more than one parent; thus we write $p(x_D|x_S, x_S)$ for the conditional probability of having

Fig. 1 The fictional "Asia" Bayesian network, taken from (Lauritzen and Spiegelhalter 1988)
The over-all probability

$$p([x]) = p \; (x_A, \, x_S, \, x_T, \, x_L, \, x_B, \, x_E, \, x_X, \, x_D)$$
 that the patient has some combination of symptoms, test results, and diseases is just the product of all probabilities of the parent nodes and all the conditional probabilities':
$$p([x]) = p(x_A) \cdot p(x_S) \cdot p(x_T|x_A) \cdot p(x_L|x_S) \cdot p(x_B|x_S) \cdot p(x_E|x_L, \, x_T) \cdot p(x_D|x_B, \, x_E) \cdot p(x_X|x_E)$$
 (1: p. 6)



Origins of the Bayesian Networks

Figure 1 shows an example of such a diagram, derived from a prototype system MUNIN which forms part of a collaborative ESPRIT project to build an advisor in electromyography (EMG).

MU.LOSS Moderate Severe ATROPHY Total Other Atrophy Hormai **MU.STRUCTURE** MVA.RECRUITMENT Very small V.s.not stro. Reduced Discrete Sinot stro. No units Normal Inconclusive DISEASE Increased MVA.AMPLITUDE Very large Increased Chronic dist, axonal neu. Other Normal Reduced No units Moderate Inconclusive Severe TA.CONCLUSION Very small Normal Limb-girdle dystrophy MUP.CONCLUSION Increased Esriy Very small Large Late Very large Small Other Normal increased MUP.AMPLITUDE Large Mean amp. .6 SD 3,01 Very large Myotonic dystrophy MUP DURATION REGENERATION Subclinical Mean dur. .67 SD 3,04 MUP.POLYPHASIC Moderate Yes Congenit < 12% 12% - 24% POSTSYN.NEU.MUSC.TRANS > 24% Decrem. MUP.INSTABILITY Normal Normal/other Other Yes Normal Other MUP.SATELLITES PRESYN.NEU.MUSC.TRANS Increm Other REPSTIM.SLOW Decrement MYOTONIA Inconclusive REPSTIM.FAST Yes Decrement lamail. **FASCICULATIONS** Increment Inconclusive Fasc. SPONT.MYO.DIS Heumyo. DENERVATION SPONT.FASCICUI Moderate Fasciculation. Neumyo SPONT.DENV.POT Some Flenty p. 106

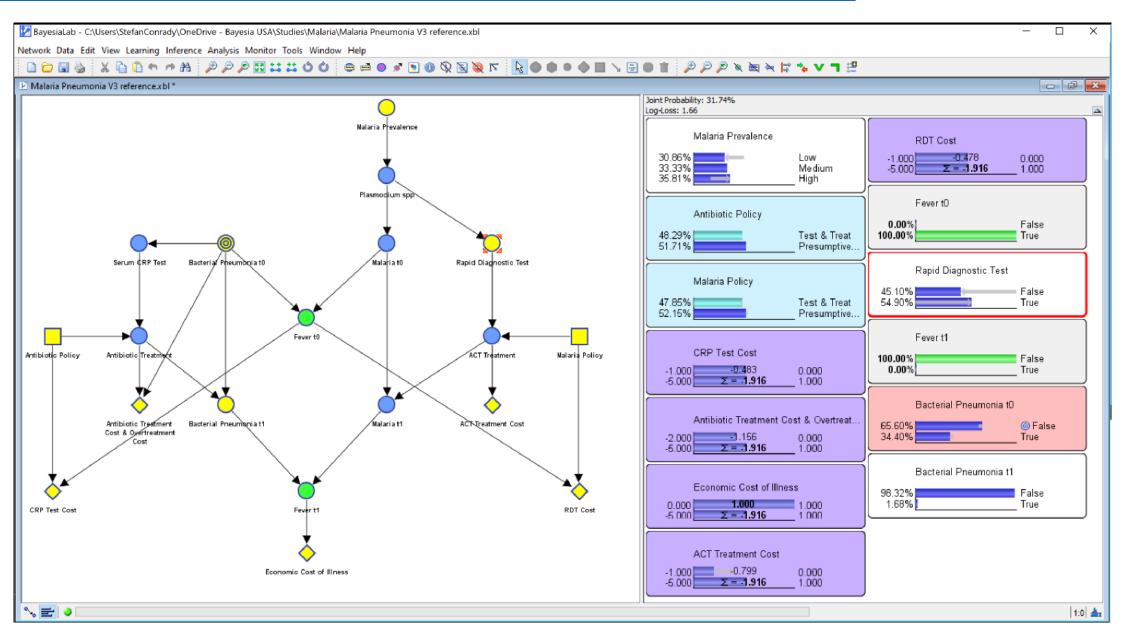
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Spiegelhalter, D. J., & Lauritzen, S. L. (1988). <u>Statistical reasoning and learning in knowledge-bases represented as causal networks</u> Expert Systems and Decision Support in Medicine (pp. 105-112): Springer.

Not unlike the BayesiaLab interface!

Bayesian Networks for Health Economics and Public Policy Research slide 139



BayesiaLab likely BayesiaLab likely

Epidemiology and Health 2011;33:e2011006

Nguefack-Tsague, G. (2011). Using Bayesian Networks to Model Hierarchical Relationships in Epidemiological Studies. Epidemiol Health, 33(0), e2011006-2011000. doi:10.4178/epih/e2011006

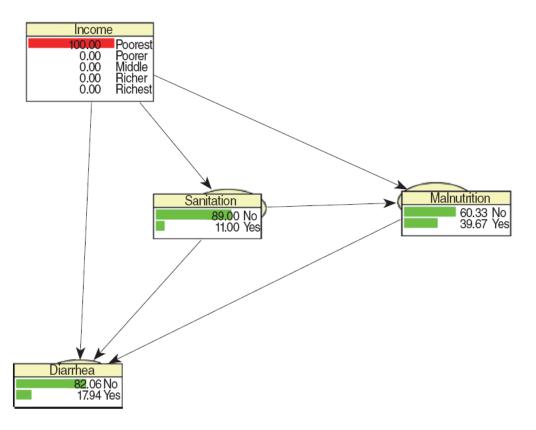


Figure 3. Frequency network showing posterior probabilities (%) when there is evidence that the child belongs to a family in the poorest quintile.

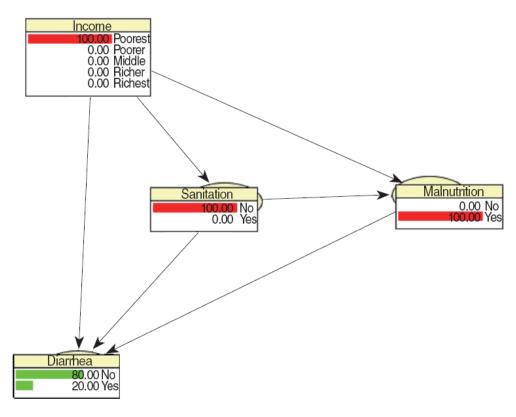


Figure 4. Frequency network showing posterior probabilities (%) of developing diarrhea when there is evidence that the child belongs to family in the poorest quintile, has poor sanitation conditions and is malnourished

How inventions 'happen'

"Jamie Robins (Figure 9.9), a pioneering statistician and epidemiologist at Harvard University who, together with Sander Greenland at the University of California, Los Angeles, is largely responsible for the widespread adoption of graphical models in epidemiology today. We collaborated for a couple of years, from 1993 to 1995, [and] he got me thinking about the problem of sequential intervention plans, which was one of his principal research interests.

After Jamie flew out to California to meet me on hearing about the "napkin problem" (Chapter 7), he was keenly interested in applying graphical methods to the sequential treatment plans that were his métier. Together we came up with a sequential back-door criterion for estimating the causal effect of such a treatment stream. I learned some important lessons from this collaboration. In particular, he showed me that two actions are sometimes easier to analyze than one because an action corresponds to erasing arrows on a graph, which makes it sparser." JP p. 328 & 330 "In 1976 Terry Speed invited me to Perth, Australia where he conducted a research seminar exploring relations between statistics and statistical physics. Among other things we studied the relation between the notion of interaction as used in contingency table analysis and in thermodynamics. To our delight they were formally the same [...]" (Lauritzen 1996)

Pearl, J., & Mackenzie, D. (2018). The Book of Why: The New Science of Cause and Effect: Hachette UK. My notes Lauritzen, S. L. (1996). Graphical models: Clarendon Press.

Ωnyx added value

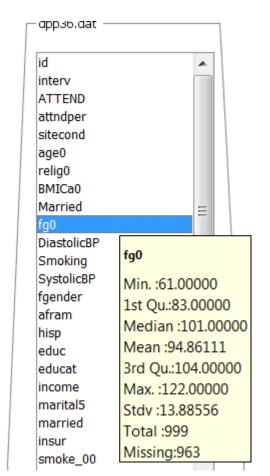
"Structural Equation Modeling is a frequently used multivariate analysis technique in the behavioral and social sciences. SEM are linear models of both observed and latent variables and their relationships. The maximum-likelihood-framework allows estimation of structural parameters even on the latent level by modeling the covariances and expectations of the observed variables.

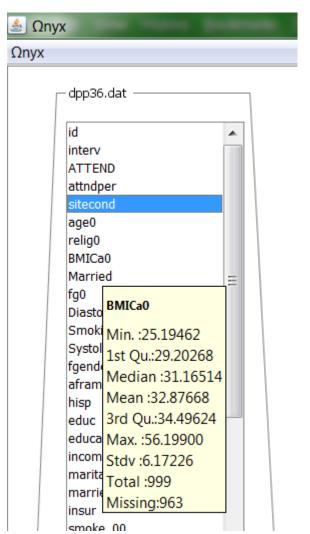
There are various text books that cover the essentials of SEM, for example, Bollen(1989). SEM can be conceived of as a unification of several multivariate analysis techniques under a single framework. Particularly, linear regression, ANOVA, correlation, path analysis, factor analysis, autoregression, and growth curve modeling can be considered special cases of SEM."

Ωnyx documentation from: http://onyx.brandmaier.de/documentation/

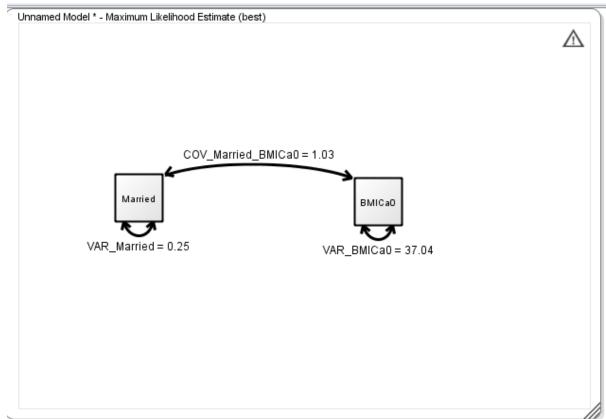
Robin Beaumont Ωnyx <u>Youtube</u>

Timo von Oertzen, Brandmaier, A. M., & Tsang, S. (2015). Structural Equation Modeling With Ωnyx. *Structural Equation Modeling: A Multidisciplinary Journal*, 22(1), 148-161. doi:10.1080/10705511.2014.935842





Robin Beaumont Ωnyx Youtube

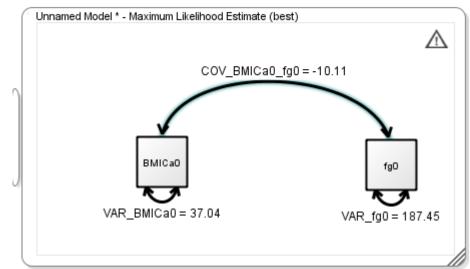


```
Estimate Summary
1st Qu.:0.00000
                 1st Qu.:29.20268
Median :1.00000
                 Median :31.16514
Mean :0.55556
                 Mean :32.87668
3rd Qu.:1.00000
                 3rd Qu.:34.49624
Max. :1.00000
                 Max. :56.19900
Stdv :0.50395
                 Stdv :6.17226
Total :36
                 Total :36
Missing:0
                 Missing:0
                              From / To|Estimate|Std.Error|lbound|rbound
        VAR Married|Married <-> Married| 0.24691| 0.05820|
         VAR BMICa0| BMICa0 <-> BMICa0|37.03850| 8.73006|
2|COV Married BMICa0| BMICa0 <-> Married| 1.02590| 0.53223|
Observed Statistics
                             : 3
Estimated Parameters
                             : 3
Non-Missing Ratio
                            : 0.036
Number of Observations
                            : 36
Minus Two Log Likelihood
                          : 279.602
Log Likelihood
                            : -139.801
Independent -2LL
                            : 284.004
Saturated -2LL
                            : 279.602
                             : 0.0
Restricted Degrees of Freedom: 0
```

R 2 variable 'model'

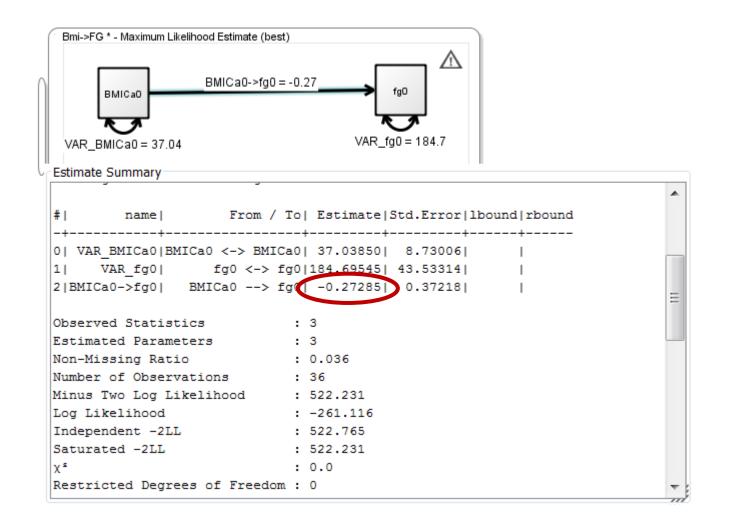
```
plot(fg0~BMICa0, data = dpp 36males Hartford fewer)
> with (dop 36males_Hartford_fewer, cov(fg0,BMICa0))
[1] (10.39484)
> mycorr <- with (dpp_36males_Hartford_fewer, cor(fg0,BMICa0, method="pearson"))</pre>
> mycorr
[1](-0.121286)
> rsquared <- mycorr^2</pre>
                                                                        0
> rsquared
[1] 0.0147103
                                                        100
                                                                                      0
                                                                                           0
                                                                  \infty
> mycorr <- with (dpp_36males_Hartford_fewer,</pre>
cor.test(fg0,BMICa0, alternative="two.sided",
                                                                                 0
                                                                       0
method="pearson"))
                                                                                                                 0
                                                        80
                                                                   0000
> mycorr
                                                                                              0
         Pearson's product-moment correlation
                                                                 0
data: fg0 and BMICa0
t = -0.71247, df = 34, p-value = 0.481
                                                        60
alternative hypothesis: true correlation is
not equal to 0
95 percent confidence interval:
                                                               25
                                                                       30
                                                                               35
                                                                                       40
                                                                                               45
                                                                                                       50
                                                                                                               55
 -0.4325846 0.2158507
sample estimates:
      cor
                                                                                        BMICa0
-0.121286
```

Robin Beaumont Ωnyx Youtube

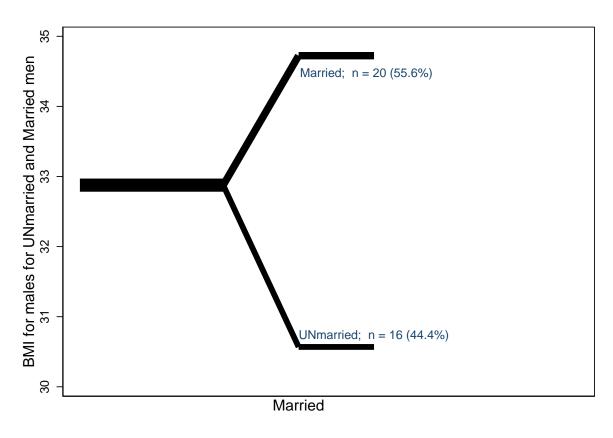


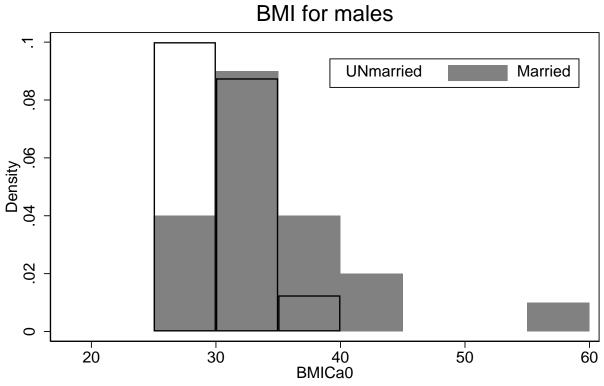
```
Estimate Summary
BMICa0
                   fg0
Min.
      :25.19462
                   Min.
                          :61.00000
1st Qu.:29.20268
                  1st Qu.:83.00000
Median :31.16514
                   Median :101.00000
     :32.87668
                        :94.86111
3rd Qu.:34.49624
                   3rd Qu.:104.00000
       :56.19900
                   Max. :122.00000
      :6.17226
                   Stdv :13.88556
Stdv
Total :36
                  Total :36
Missing:0
                  Missing:0
                         From / To| Estimate|Std.Error|lbound|rbound
            name
      VAR BMICa0|BMICa0 <-> BMICa0| 37.03850| 8.73006|
11
        VAR fg0|
                       fg0 <-> fg0|187.45293| 44.18308|
2|COV BMICa0 fg0|
                   fg0 <-> BMICa( -10.10610 | 13.98919 |
Observed Statistics
                              : 3
Estimated Parameters
                              : 3
Non-Missing Ratio
                              : 0.036
Number of Observations
                              : 36
Minus Two Log Likelihood
                              : 522.231
Log Likelihood
                              : -261.116
Independent -2LL
                              : 522.765
Saturated -2LL
                              : 522.231
                              : 0.0
```

Robin Beaumont Ωnyx <u>Youtube</u>



BMI of males: by marital status



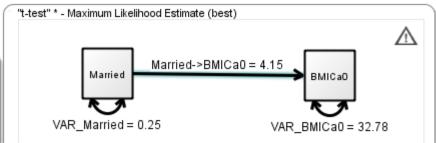


Obse	rap	Normal-based				
Į		Std. Err.		P> z	[95% Conf.	Interval]
	0148387			0.972	8369783	.8073009
kurtosis	1.515828	.410243	3.69	0.000	.7117664	2.319889

Observed Bootstrap			Normal-based				
	Coef.	Std. Err.	z	P> z	[95% Conf.	<pre>Interval]</pre>	
+							
1	.539829	.5361251	2.87	0.004	.4890433	2.590615	
sis 5	.160547	1.927927	2.68	0.007	1.381881	8.939214	
	 + 1	Coef. 	Coef. Std. Err.	Coef. Std. Err. z 1.539829 .5361251 2.87	Coef. Std. Err. z P> z 1.539829 .5361251 2.87 <u>0.004</u>	Coef. Std. Err. z P> z [95% Conf. 1.539829 .5361251 2.87 0.004 .4890433	

Marriage -> BMI of males

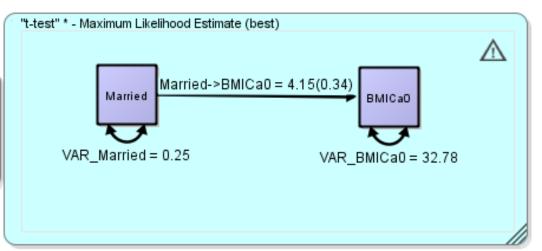
```
Z value for the Married->BMI is 4.15488/1.92024 = 2.16373
> zis<-4.15488/1.92024
Zis
[1] 2.16373
t.test
t = -2.1028, df = 34, p-value = 0.04296
sample estimates:
mean in group 0 mean in group 1
       30.56842
                 34,72329
Welch Two Sample t-test
t = -2.2389, df = 30.278, p-value = 0.03266
sample estimates:
mean in group 0 mean in group 1
       30.56842
                      34.72329
```



Estimate Summary From / TolEstimate|Std.Error| namel VAR BMICa0 | BMICa0 <-> BMICa0|32.77603| 7.72538| VAR Married | Married | 0.24691 | 0.05820 | 2|Married->BMICa0| Married --> BMICa0| 4.15488| 1.92024| Observed Statistics : 3 Estimated Parameters : 3 Non-Missing Ratio : 0.036 Number of Observations : 36 Minus Two Log Likelihood : 279.602 Log Likelihood : -139.801 Independent -2LL : 284.004 Saturated -2LL : 279.602 : 0.0 Restricted Degrees of Freedom : 0 AIC : 285.602 AICc : 285.602 BIC : 290.353 BIC (sample-size adjusted) : 290.353 Kulback-Leibler to Saturated : 0.0 xº from independent : 4.401 Degrees of Freedom (indep.) :1 RMSEA (df corrected) : 0.0 RMSEA (Kulback Leibler) : 0.0

RAM notation

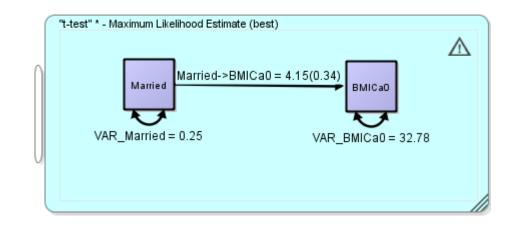
- 1. McArdle, J. J., & Boker, S. M. (1990). RAMpath: Path diagram software: Data Transforms.
- 2. McArdle, J. J. (2005). The development of the RAM rules for latent variable structural equation modeling. In A. Maydeu-Olivares & J.
- J. McArdle (Eds.), Contemporary psychometrics: A festschrift for Roderick P. McDonald (pp. 225-273).
- 3. Zhang, Z., Hamagami, F., Grimm, K. J., & McArdle, J. J. (2015). Using R Package RAMpath for Tracing SEM Path Diagrams and Conducting Complex Longitudinal Data Analysis. *Structural Equation Modeling: A Multidisciplinary Journal*, 22(1), 132-147.



RAM Matrices

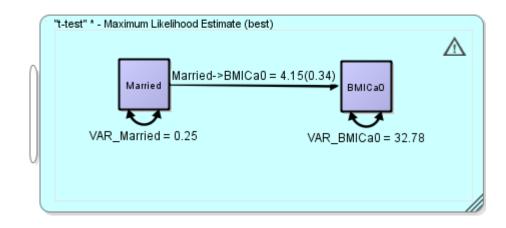
```
Variables: BMICa0, Married
                     0.00
      0.00
      0.00
                    Marr..0=4.15
       0.00
                     0.00
     VAR ..0=32.78 0.00
      0.00
                    VAR ..d=0.25
Model Covariance matrix = F(I-A)^{-1} S(I-A)^{-1} F^T
                        = F (I-A)^{-1} m
Model Mean Vector
```

Lavaan code for lazy folk: From Ωnix



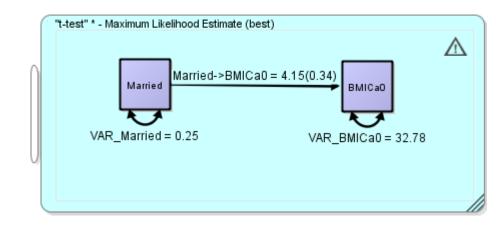
```
# This model specification was automatically generated by Onyx
library(lavaan);
modelData <- read.table(DATAFILENAME, header = TRUE) ;</pre>
 model<-"
! regressions
   BMICa0 ~ Married BMICa0*Married
! residuals, variances and covariances
   BMICa0 ~~ VAR BMICa0*BMICa0
   Married ~~ VAR Married*Married
! observed means
   BMICa0~1;
   Married~1;
II ;
result<-lavaan(model, data=modelData, fixed.x=FALSE, missing="FIML");
summary(result, fit.measures=TRUE);
```

sem code for lazy folk: From Ω nix



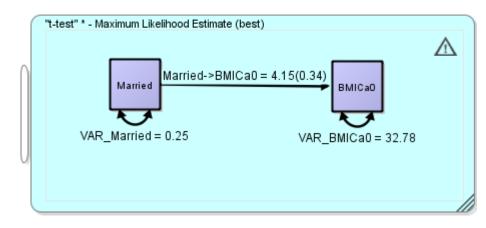
```
# This model specification was automatically generated by Onyx
#
require("sem");
modelData <- read.table(DATAFILENAME, header = TRUE)
paths <- c("BMICa0 <-> BMICa0", "Married <-> Married", "Married -> BMICa0")
parameter <- c("VAR_BMICa0", "VAR_Married", "Married_BMICa0")
values <- c("32.776030306921605", "0.24691358028835825", "4.154875879839016")
model <- array(c(paths, parameter, values), dim = c(3,3))
colnames(model) <- c("col1","col2","col3")
result <- sem(model = model, data = modelData)
summary(result)</pre>
```

Lavaan code for lazy folk: From Ωnix



```
# This model specification was automatically generated by Onyx
#
library(lavaan);
modelData <- read.table(DATAFILENAME, header = TRUE) ;</pre>
model<-"
! regressions
   BMICa0 ~ Married BMICa0*Married
! residuals, variances and covariances
   BMICa0 ~~ VAR BMICa0*BMICa0
  Married ~~ VAR Married*Married
! observed means
   BMICa0~1;
  Married~1;
· ;
result<-lavaan(model, data=modelData, fixed.x=FALSE, missing="FIML");
summary(result, fit.measures=TRUE);
```

OpenMx code for lazy folk: From Ωnix



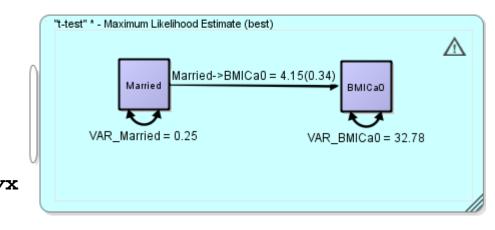
```
# This model specification was automatically generated by Onyx
require("OpenMx");
modelData <- read.table(DATAFILENAME, header = TRUE)</pre>
manifests<-c("BMICa0","Married")</pre>
latents<-c()</pre>
model <- mxModel(" t test ",</pre>
type="RAM",
manifestVars = manifests,
latentVars = latents,
mxPath(from="Married",to=c("BMICa0"), free=c(TRUE), value=c(1.0), arrows=1, label=c("Married BMICa0")
),
mxPath(from="BMICa0",to=c("BMICa0"), free=c(TRUE), value=c(1.0), arrows=2, label=c("VAR BMICa0")),
mxPath(from="Married", to=c("Married"), free=c(TRUE), value=c(1.0), arrows=2, label=c("VAR Married")),
mxPath(from="one",to=c("BMICa0","Married"), free=F, value=0, arrows=1),
mxData(modelData, type = "raw")
);
result <- mxRun(model)</pre>
summary(result)
```

Appendix: a 2 versions of t tests

```
> t.test(dpp 36males Hartford fewer$BMICa0~dpp 36males Hartford fewer$married,var.equal=TRUE)
        Two Sample t-test
data: dpp 36males Hartford fewer$BMICa0 by dpp 36males Hartford fewer$married
t = -2.1028, df = 34, p-value = 0.04296
alternative hypothesis: true difference in means is not equal to 0
95 percent confidence interval:
-8.1703995 -0.1393522
sample estimates:
mean in group 0 mean in group 1
       30.56842
                       34.72329
vs. Welch t-test
Welch Two Sample t-test
data: dpp 36males Hartford fewer$BMICa0 by dpp 36males Hartford fewer$married
t = -2.2389, df = 30.278, p-value = 0.03266
alternative hypothesis: true difference in means is not equal to 0
95 percent confidence interval:
-7.9434686 -0.3662831
sample estimates:
mean in group 0 mean in group 1
       30.56842
                      34,72329
```

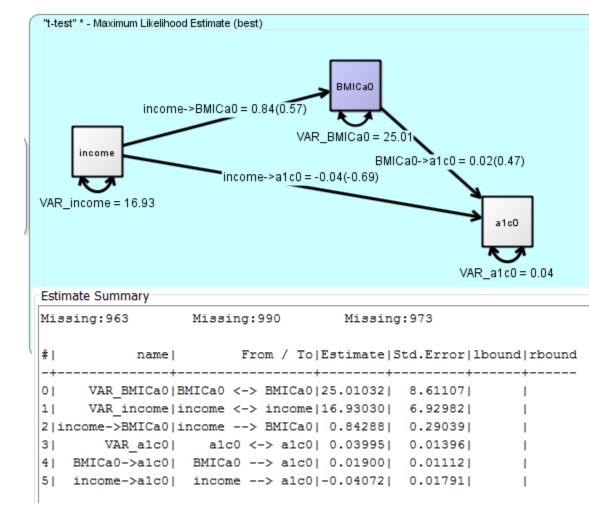
Mplus code for lazy folk: From Ω nix

```
Variables: BMICa0, Married
37.0385 1.0259
1.0259 0.2469
!This model specification was automatically created by Onyx
TITLE:
   "t-test"
DATA:
   FILE IS "DATAFILENAME";
VARIABLE: NAMES ARE BMICAO MARRIED ;
   USEVARIABLES ARE BMICAO MARRIED ;
MODEL:
! regressions of latents on manifest
! regressions of manifest on manifest
   BMICAO ON MARRIED*1.0;
! regressions of latents on latents or manifests
! residuals, variances and covariances
   BMICA0*1.0;
   MARRIED*1.0;
ANALYSIS:
   TYPE = general;
   ESTIMATOR = ml;
OUTPUT:
   sampstat;
```



A more informed model: Ωnix

There is likely an indirect effect too here.



```
fit <- sem(SEMJaccApp d, data = dpp 36males Hartford fewer)</pre>
> summary(fit, rsq = T)
lavaan (0.5-23.1097) converged normally after 17 iterations
 Number of observations
                                                    36
 Estimator
                                                    ML
 Minimum Function Test Statistic
                                                 0.000
 Degrees of freedom
                                                     0
Parameter Estimates:
  Information
                                              Expected
  Standard Errors
                                              Standard
Regressions:
                    Estimate Std.Err z-value P(>|z|) Std.lv Std.all
BMICa0 ~
   married
                      4.155
                               1.920
                                        2.164
                                                 0.030
                                                          4.155
                                                                   0.339
Variances:
                   Estimate Std.Err z-value P(>|z|)
                                                        Std.lv Std.all
                               7.725
                                      4.243
                                               0.000
   .BMICa0
                     32,776
                                                        32.776
                                                                   0.885
[only 1 variance estimated: but we have 2 groups...]
R-Square:
                   Estimate
    BMICa0
                      0.115
vs. Welch t-test
Welch Two Sample t-test
data: dpp 36males Hartford fewer$BMICa0 by dpp 36males Hartford fewer$married
t = -2.2389, df = 30.278, p-value = 0.03266
alternative hypothesis: true difference in means is not equal to 0
95 percent confidence interval:
-7.9434686 -0.3662831
sample estimates:
mean in group 0 mean in group 1
       30.56842
                       34.72329
```

Appendix: a

Appendix: a

```
SEMJaccApp a2 <- '
BMICa0~1
 fit <- sem(SEMJaccApp_a2,</pre>
             data = dpp36,
            group = "married")
Estimator
                                                   ML
  Model Fit Test Statistic
                                                  0.000
  Degrees of freedom
Number of observations per group
                                                     16
  0
Group 1 [0]:
Intercepts:
                   Estimate Std.Err z-value P(>|z|)
                                0.929
    BMICa0
                     30.568
                                        32.917
                                                  0.000
Variances:
                   Estimate Std.Err z-value P(>|z|)
    BMICa0
                     13.798
                                4.878
                                         2.828
                                                  0.005
vs. Welch t-test
Welch Two Sample t-test
t = -2.2389, df = 30.278, p-value = 0.03266
95 percent confidence interval:-7.9434686 -0.3662831
sample estimates:
mean in group 0 mean in group 1
       30.56842
                       34.72329
```

```
lavaan 0.6-3 ended normally after 15 iterations
 Optimization method
                                                NLMINB
  Number of free parameters
Chi-square for each group:
                                                  0.000
                                                  0.000
Parameter Estimates:
  Information
                                               Expected
  Information saturated (h1) model
                                             Structured
  Standard Errors
                                               Standard
Number of observations per group
                                                     20
Group 2 [1]:
Intercepts:
                             Std.Err z-value P(>|z|)
                   Estimate
                               1.549
                                        22,424
    BMICa0
                     34.723
                                                  0.000
Variances:
                   Estimate Std.Err z-value P(>|z|)
    BMICa0
                     47.958
                              15,166
                                         3.162
                                                  0.002
vs. Welch t-test
Welch Two Sample t-test
t = -2.2389, df = 30.278, p-value = 0.03266
alternative hypothesis: true difference in means is
not equal to 0
mean in group 0 mean in group 1
                       34.72329
       30.56842
```

Appendix: a

```
Df
                                                  AIC
                                     fitdif 0 236.19 239.36 0.0000
> fiteq <- sem(SEMJaccApp a2,</pre>
                                     fiteq
data = dpp36, group = "married",
group.equal = c("intercepts"))
lavaan 0.6-3 ended normally after 15 iterations
 Optimization method
                                                NLMINB
 Number of free parameters
 Estimator
                                                    MT.
 Model Fit Test Statistic
                                                  4.891
  Degrees of freedom
  P-value (Chi-square)
                                                  0.027
Number of observations per group
  0
                                                     16
Group 1 [0]:
Intercepts:
                   Estimate Std.Err z-value P(>|z|)
                               0.837
                                       37.703
                                                  0.000
    BMICa0
           (.p1.)
                     31.576
Variances:
                   Estimate Std.Err z-value P(>|z|)
                     14.813
                               5.237
                                        2.828
    BMICa0
                                                  0.005
vs. Welch t-test
Welch Two Sample t-test
t = -2.2389, df = 30.278, p-value = 0.03266
95 percent confidence interval:
 -7.9434686 -0.3662831
sample estimates:
mean in group 0 mean in group 1
       30.56842
                       34.72329
```

> anova(fitdif,fiteq)

```
Chi Square Difference Test
                   BIC Chisq Chisq diff Df diff Pr(>Chisq)
       1 232.45 237.20 4.8911 4.8911
                                                      0.027 *
                     Chi-square for each group:
                      0
                                                                     1.135
                                                                     3.756
                     Parameter Estimates:
                       Information
                                                                   Expected
                       Information saturated (h1) model
                                                                 Structured
                                                                   Standard
                       Standard Errors
                     Number of observations per group
                       1
                                                                         20
                     Group 2 [1]:
                     Intercepts:
                                        Estimate Std.Err z-value P(>|z|)
                                          31.576
                                                    0.837
                                                            37.703
                                                                      0.000
                         BMICa0
                                (.p1.)
                     Variances:
                                        Estimate Std.Err z-value P(>|z|)
                         BMICa0
                                          57.866
                                                  18,299
                                                             3.162
                                                                      0.002
                     vs. Welch t-test
                     Welch Two Sample t-test
                     t = -2.2389, df = 30.278, p-value = 0.03266
                     alternative hypothesis: true difference in means is
                     not equal to 0
                     sample estimates:
                     mean in group 0 mean in group 1
                            30.56842
                                            34.72329
```

Appendix a: Compare means in 2 independent groups

```
Classic
Model is... a t-test
Welch Two Sample t-test
t(df=1) = -2.2389, p = 0.033
```

Modern

Model is a 2 group 1 effect variable (no cause) [ContinuousEffect_{Group1} & ContinuousEffect_{Group2}]

```
Chi Square Difference Test \chi^2diff(df=1) = 4.8911 , p= 0.027
```

Pre-data: in DAG world

```
# 2 DAG
DagJaccApp_a2 <- dagitty('dag {</pre>
Married[pos="1,2"]
BMI [pos="2,2"]
Married-> BMI
}')
plot (DagJaccApp_a2)
#then can technically simulate data DID NOT WORK
DagJaccApp_a2 <- simulateSEM( g, .2, .3)</pre>
coef( summary( lm( BMI ~ Married, DagJaccApp_a2 ) ) )
```

Married > BMI

Pre-data: in MIIVsem world

```
##1 MIIVsem
model.MiivJaccApp_a2 <- '</pre>
BMI ~ Married
miivs(model.MiivJaccApp_a2)
Model Equation Information
 LHS RHS
             MIIVs
 BMI Married Married
```

Married > BMI

CRAN - Package MIIVsem

GitHub - zackfisher/MIIVsem

Bollen, K. A., & Fisher, Z. (2017). Model Implied Instrumental Variable (MIIV) Methods using MIIVsem: An R Package for Structural Equation Models (SEMs). Paper presented at the Modern Modeling Methods (M3) Conference, Storrs, CT.

https://miivs.shinyapps.io/miivs/

Bollen, K. A. (2018). Model Implied Instrumental Variables (MIIVs): An Alternative Orientation to Structural Equation Modeling. Multivariate Behavioral Research, 1-16.

Example 2: obesity paradox" Pre-data: in DAG world

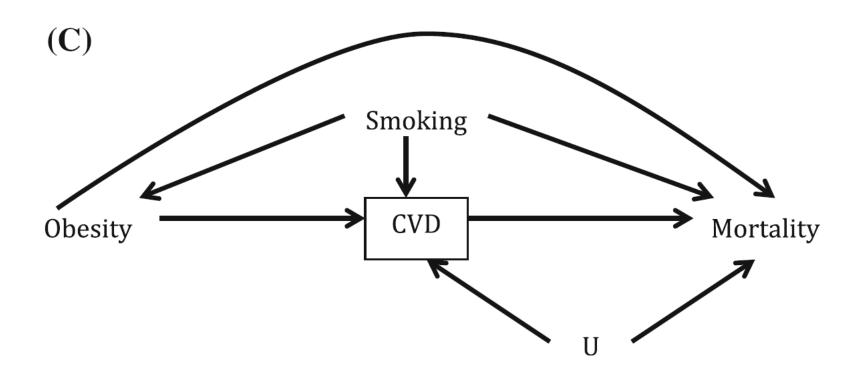


Fig. 1 a–c Directed acyclic graphs representing causal relations between obesity, cardiovascular disease (CVD), and mortality. Smoking and unmeasured factors (U) are included as confounders in (**b**, **c**), respectively

Obesity paradox" Pre-data: in DAG world

```
Smoking
g <- dagitty('dag {
Obesity [pos="0,1"]
CVD [pos="1,1"]
Mortality [pos="2,1"]
Smoking [pos="1,0"]
                                           Obesity
                                                       \geq CVD
                                                                    Mortality
MissedConf [pos="1.5,2"]
Obesity -> CVD -> Mortality <- MissedConf
Obesity <- Smoking -> CVD <- MissedConf
Smoking -> Mortality
Obesity-> Mortality
                                                             MissedConf
plot(g)
```

Johannes Textor, Andrew Forney, and Judea Pearl (2016). Causal Inference In Statistics: A Companion for R Users
Banack, H. R., & Kaufman, J. S. (2015). From bad to worse: collider stratification amplifies confounding bias in the "obesity paradox". *European journal of epidemiology, 1-4.*

Obesity paradox" Pre-data: in MIIVsem

```
model.obesparadox <- '
    Mortality ~ CVD +MissedConf + Smoking +Obesity
CVD ~ Obesity + MissedConf + Smoking
Obesity ~ Smoking
'
miivs(model.obesparadox)</pre>

(C)
    Hobesity
    Obesity
    Obes
```

```
LHS RHS
Mortality MissedConf, Smoking, CVD, Obesity CVD, Obesity, MissedConf, Smoking
CVD MissedConf, Smoking, Obesity Obesity, MissedConf, Smoking
Obesity Smoking MissedConf, Smoking
```

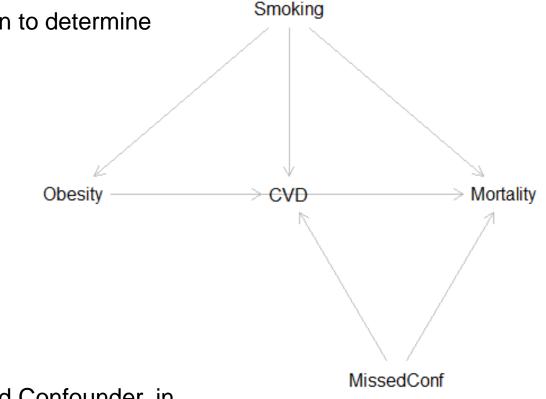
Model Equation Information

Obesity paradox" Which adjustments are required

"List all of the sets of variables that satisfy the backdoor criterion to determine the causal effect of Obesity on *Mortality*."

```
adj ustmentSets( g, "Obesity",
"Mortality", type="all" )

{ Smoking }
{ MissedConf, Smoking }
```



This is the cherry on the cake:

One **HAS TO to measure** smoking, OR smoking and the Missed Confounder, in order to identify the causal effect of Obesity on Mortality.

Why bother

Modern

- 1. It allows one to think in causal modeling manner
- 2. It is more flexible modeling: allows for fewer 'unrelaxable' assumptions, and can test assumptions along the way
- 3. The reality operates as different effects in different populations, e.g. health disparities: the effect in income on health differs for races/ethnic groups
- 4. It open the doors for formal causal inquiries: when and how can one recover causal effects with simple regressions, e.g.

What you may have learned

- 1. R is not that scary: nor is causal modeling
- 2. Causal calculus' has been implemented and is easy to use on dagitty
- 3. Several traditions have been merged in designing MIIVsem (economics, latent variables)
- 4. One can better teach/train by using graphical models: Onyx is there to help.
- 5. Answering 'what if' questions require solid causal footing:
- the science of cause and effect is here with us

Next:

The science of cause and effect is 'complete': Elias Bareinboim will come to Storrs to prove it.