Causal Mediation Analysis

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UK Stata Conference

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Causal Mediation Analysis

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Outline

- Introduction
- Overview of mediate
- Traditional mediation
- Causal inference
- Causal mediation analysis
- Examples

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Causal mediation analysis combines:

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Causal mediation analysis combines:

Causal inference

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Causal mediation analysis combines:

- Causal inference
- Mediation analysis

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Causal mediation analysis combines:

- Causal inference—What is the effect of a treatment on an outcome?
- Mediation analysis

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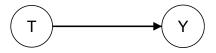
Causal mediation analysis combines:

- Causal inference—What is the effect of a treatment on an outcome?
- Mediation analysis—Can the total effect of a predictor on an outcome be decomposed into a direct effect and an indirect effect through a mediating variable?

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With causal inference, we want to answer questions about causality.

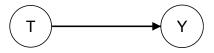
With causal inference, we want to answer questions about causality.



• What is the causal effect of T on Y?

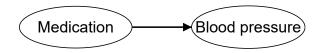
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With causal inference, we want to answer questions about causality.



- What is the causal effect of **T** on **Y**?
- What is the expected difference in **Y** if the treatment **T** is applied versus if the treatment is not applied?

What is the effect of a medication on blood pressure?



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What is the effect of a job-training program on probability of employment?



What is the effect of exercise on self-perceived well-being?

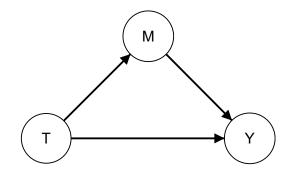


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With mediation analysis, we want to better understand the effect.

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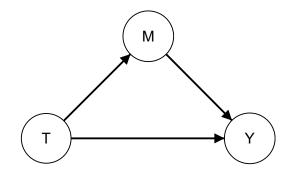
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• Why does **T** affect **Y**?

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With mediation analysis, we want to better understand the effect.

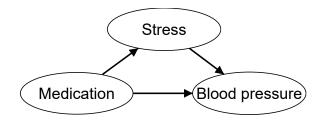


- Why does T affect Y?
- Can effect of **T** on **Y** be explained either completely or partially by a change in the mediator **M**?

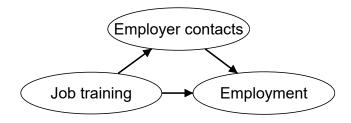
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Does the medication result in lower stress levels, which in turn, results in lower blood pressure?

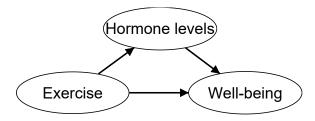


Does the job-training program put participants in contact with potential employers, which in turn, increases the probability of employment?



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Does exercise change levels of some hormones, which in turn, change self-perceived well-being?



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Causal mediation analysis

- With causal mediation analysis, we aim to draw causal inferences about the effect of a treatment on an outcome and to understand why the effect arises.
- To understand the why, we decompose the total effect into indirect effects through a mediator and direct effects.

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 In Stata 18, we introduced the mediate command to perform causal mediation analysis.

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mediate (ovar [omvarlist] [, omodel noconstant])
 (mvar [mmvarlist] [, mmodel noconsant])
 (tvar [, continuous(numlist)])
 [if] [in] [, stat options]

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• In Stata 18, we introduced the **mediate** command to perform causal mediation analysis.

mediate (ovar [omvarlist] [, omodel noconstant])
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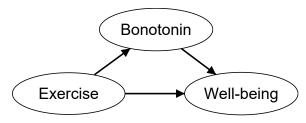
- ovar is a continuous, binary, or count outcome of interest.
- *mvar* is the mediator variable and may be continuous, binary, or count.
- *tvar* is the treatment variable and may be binary, multivalued, or continuous.

Mediator Outcome	linear	logit	probit	Poisson	exp. mean
	V	V	X	V	N N
linear	X	X	X	Х	X
logit		Х	Х	х	
probit	Х	Х	Х	х	Х
Poisson	Х	Х	Х	х	Х
exp. mean	Х	Х	Х	Х	Х

Note: X indicates a supported model combination

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For a simple example using the **mediate** command, we continue with our hypothesis that exercise affects well-being and that this may, at least in part, be because of a change in hormone levels. We will consider a fictional hormone **bonotonin**.



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```
. webuse wellbeing (Fictional well-being data)
```

```
. list wellbeing bonotonin exercise in 1/5, abbreviate(10)
```

	wellbeing	bonotonin	exercise
1.	71.73816	196.5467	Control
2.	68.66573	195.8572	Exercise
3. 4.	71.05155 69.44469	228.6035 206.6651	Exercise Exercise
4. 5.	75.62035	206.6651 261.6855	Exercise

- Both **wellbeing** and **bonotonin** are continuous, so we use the default linear model for the outcome and the mediator.
- **exercise** is a binary treatment variable with 0 representing the control group and 1 representing exercise group.

. mediate (wei Iteration 0:	EE criterion	= 5.104e-2	7			
Iteration 1: Causal mediat:		= 2.031e-2	8		Number of c	bs = 2,000
Outcome model Mediator mode Mediator varia Treatment type	l: Linear able: bonotoni	n				
wellbeing	Coefficient	Robust std. err.	Z	P> z	[95% conf.	interval]
NIE exercise (Exercise vs						
Control)	9.799821	.3943251	24.85	0.000	9.026958	10.57268
NDE exercise (Exercise vs Control)	2.891453	.2304278	12.55	0.000	2.439823	3.343083
TE exercise (Exercise vs						
Control)	12.69127	.4005941	31.68	0.000	11.90612	13.47642

Note: Outcome equation includes treatment-mediator interaction.

• We estimate the total effect of exercise on well-being is 12.7, with an indirect effect of 9.8 and a direct effect of 2.9.

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• What proportion of the total effect exercise on well-being is mediated through bonotonin levels?

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Proportion mediated

Number of obs = 2,000

wellbeing	Proportion	Robust std. err.	Z	P> z	[95% conf.	interval]
exercise (Exercise vs Control)	.77217	.0172979	44.64	0.000	.7382668	.8060732

Traditional mediation analysis: The formulation

 In traditional mediation analysis, we write models for the outcome and the mediator.

$$Y = \beta_0 + \beta_1 M + \beta_2 T + \epsilon$$
$$M = \alpha_0 + \alpha_1 T + \nu$$

• Then we define direct, indirect, and total effects as

$$\begin{aligned} \text{Direct} &= \beta_2 \\ \text{Indirect} &= \alpha_1 * \beta_1 \\ \text{Total} &= \beta_2 + \alpha_1 * \beta_1 \end{aligned}$$

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Traditional mediation analysis: Estimation

- We can fit the linear regression models using **regress** and then manually compute direct, indirect, and total effects and the corresponding standard errors.
- Alternatively, we can fit models simultaneously using, for instance, the sem command.

```
. sem (wellbeing <- bonotonin exercise)
      (bonotonin <- exercise)</pre>
```

After **sem**, we can use **estat teffects** to compute direct, indirect, and total effects and the corresponding standard errors.

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Traditional mediation analysis: Estimation

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Direct effects

	Coefficient	OIM std. err.	z	P> z	[95% conf.	interval]
Structural wellbeing						
 exercise	2.996658	.2119904	14.14	0.000	2.581164	3.412151

Indirect effects

	Coefficient	OIM std. err.	z	₽> z	[95% conf.	interval]
Structural wellbeing						
exercise	9.694617	.3717311	26.08	0.000	8.966037	10.4232

Total effects

Coeffici	OIM ient std.er	r. z	₽> z	[95% conf	. interval]
Structural wellbeing					
exercise 12.691	127 .3983777	7 31.86	0.000	11.91047	13.47208

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Traditional mediation analysis

- Traditional mediation analysis began with linear models for both the outcome and the mediator.
- The model for the outcome did not include a mediator by treatment interaction term.
- What can a causal inference approach add?

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Common steps in a causal inference approach:

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1 **Hypothetical modeling.** Researchers make assumptions about relationships among variables based on their understanding and expertise. These assumptions may be illustrated by using a causal diagram.

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Common steps in a causal inference approach:

- 1 **Hypothetical modeling.** Researchers make assumptions about relationships among variables based on their understanding and expertise. These assumptions may be illustrated by using a causal diagram.
- 2 **Causal effect identification.** Based on the assumptions made in the first phase, the researcher tries to determine whether the causal effect can be identified.

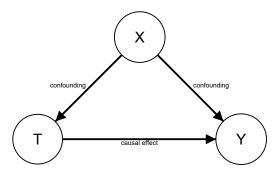
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Common steps in a causal inference approach:

- 1 **Hypothetical modeling.** Researchers make assumptions about relationships among variables based on their understanding and expertise. These assumptions may be illustrated by using a causal diagram.
- 2 **Causal effect identification.** Based on the assumptions made in the first phase, the researcher tries to determine whether the causal effect can be identified.
- 3 Parameter estimation. If the answer to the second phase is positive, the researcher can then to estimate the causal effect. Stata provides a variety of commands such as the teffects suite to estimate average treatment effects (ATEs), average treatment effects on the treated (ATETs), and other causal estimands of interest.

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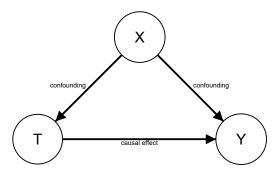
Causal diagram



 In this very simple causal diagram, we are interested in estimating the causal effect of treatment T on outcome Y, but we believe that X also affects both T and Y.

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Causal diagram



- In this very simple causal diagram, we are interested in estimating the causal effect of treatment T on outcome Y, but we believe that X also affects both T and Y.
- X is a counfounder, and we must somehow control for confounding to obtain an unbiased estimate of the causal effect.

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How do we define a causal effect?

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- A common causal-inference approach is based on the potential-outcomes framework.
- For a binary treatment *T*, we can define two potential outcomes.
 - Y(0) is the potential outcome that would have been observed if treatment T = 0 was assigned.
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- The individual treatment effect is the difference in the two potential outcomes, Y(1) - Y(0).
- The average treatment effect (ATE) is E[Y(1) Y(0)].

Fundamental problem of causal inference

Subject	Т	Y	Y(1)	Y(0)	Y(1) - Y(0)
1	0	2.1	?	2.1	?
2	1	3.7	3.7	?	?
3	1	4.2	4.2	?	?
4	0	6.2	?	6.2	?

For each individual, we can observe only one of Y(1) or Y(0).

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Can we estimate the ATE given that Y(1) - Y(0) is never observed?

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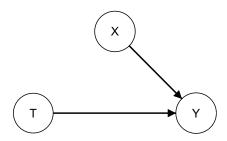
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• What if we have a randomized control trial (RCT)?

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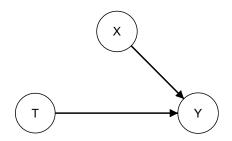
- What if we have a randomized control trial (RCT)?
- In an RCT, we randomize the treatment; therefore, T is independent of Y(0), Y(1), and X



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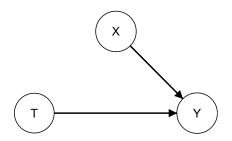


• In this case E[Y(0)] = E[Y|T = 0] and E[Y(1)] = E[Y|T = 1].

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- In this case E[Y(0)] = E[Y|T = 0] and E[Y(1)] = E[Y|T = 1].
- We can estimate the ATE as E[Y|T = 1] E[Y|T = 0].

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Can we estimate the ATE given that Y(1) - Y(0) is never observed?

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• What if we have observational data?

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 - Unconfoundedness or conditional independence. If we condition on confounders X, the treatment assignment is as good as random.
 - Stable unit treatment value assumption (SUTVA). The treatment of each individual is unrelated to the outcome of the treatment of all the other individuals in the population.

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 $E[Y(1)] - E[Y(0)] = E_x[E[Y|T = 1, X] - E[Y|T = 0, X]]$

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• We can use commands such as teffects to estimate the ATE.

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Estimating the causal effect

. teffects ra (wellbeing age) (exercise)			
Iteration 0: EE criterion = 1.261e-27			
Iteration 1: EE criterion = 1.707e-29			
Treatment-effects estimation	Number of obs	=	2,000
Estimator : regression adjustment			
Outcome model : linear			
Treatment model: none			

wellbeing	Coefficient	Robust std. err.	Z	P> z	[95% conf.	interval]
ATE exercise (Exercise vs Control)	12.76801	.3961873	32.23	0.000	11.9915	13.54452
POmean exercise Control	57.06904	.2738341	208.41	0.000	56.53234	57.60575

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Mediation analysis via potential outcomes

Kristin MacDonald (StataCorp LLC)

Causal Mediation Analysis

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Mediation analysis via potential outcomes

• We can extend the potential-outcomes framework to mediation analysis.

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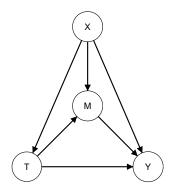
Mediation analysis via potential outcomes

- We can extend the potential-outcomes framework to mediation analysis.
- We can define a total average treatment effect as well as direct and indirect effects in terms of potential outcomes.

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Mediation analysis: Causal inference workflow

• A causal diagram



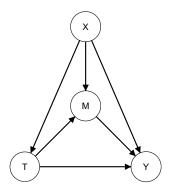
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Image: A matrix

Mediation analysis: Causal inference workflow

A causal diagram



• As before, we will make assumptions that allow us to get unbiased estimates of the causal effects, even in the presence of confounders.

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• We now have potential outcomes for the the mediator *M* and for the outcome *Y*.

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- We now have potential outcomes for the the mediator *M* and for the outcome *Y*.
- For the mediator, we have
 - ► M(0) is the potential outcome of the mediator that would have been observed if treatment T = 0 was assigned.
 - M(1) is the potential outcome of the mediator that would have been observed if treatment T = 1 was assigned.

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• Formally, let *t* be the treatment level with respect to the outcome, and let t' be the treatment level with respect to the mediator, the potential outcomes become Y[t, M(t')].

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- Formally, let *t* be the treatment level with respect to the outcome, and let t' be the treatment level with respect to the mediator, the potential outcomes become Y[t, M(t')].
- This leads to four types of potential outcomes:

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- Formally, let *t* be the treatment level with respect to the outcome, and let t' be the treatment level with respect to the mediator, the potential outcomes become Y[t, M(t')].
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- Formally, let t be the treatment level with respect to the outcome, and let t' be the treatment level with respect to the mediator, the potential outcomes become Y[t, M(t')].
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- Formally, let t be the treatment level with respect to the outcome, and let t' be the treatment level with respect to the mediator, the potential outcomes become Y[t, M(t')].
- This leads to four types of potential outcomes:
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 - Y[1, M(1)] is the potential outcome that would be observed if treatment T = 1 was assigned.
 - Y[1, M(0)] is the potential outcome that would be observed if treatment T = 1 was assigned, but the mediator is held at its value that would be observed if if T = 0 was assigned.

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- This leads to four types of potential outcomes:
 - Y[0, M(0)] is the potential outcome that would be observed if treatment T = 0 was assigned.
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 - Y[0, M(1)] is the potential outcome that would be observed if treatment T = 0 was assigned, but the mediator is held at its value that would be observed if if T = 1 was assigned.

Potential outcomes

. mediate (wellbeing) (bonotonin) (exercise), pomeans	
Iteration 0: EE criterion = $5.104e-27$	
Iteration 1: EE criterion = 2.023e-28	
Causal mediation analysis	Number of obs = $2,000$
Outcome model: Linear	
Mediator model: Linear	
Mediator variable: bonotonin	
Treatment type: Binary	

wellbeing	Coefficient	Robust std. err.	Z	₽> z	[95% conf.	interval]
POmeans YOMO Y1MO Y0M1	57.11317 60.00462 66.68199	.2753201 .3157888 .3258477	207.44 190.02 204.64	0.000 0.000 0.000	56.57355 59.38569 66.04334	57.65278 60.62356 67.32064
Y1M1	69.80444	.2898927	240.79	0.000	69.23626	70.37262

Note: Outcome equation includes treatment-mediator interaction.

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Kristin MacDonald (StataCorp LLC)

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• Average direct, indirect, and total treatment effects are contrasts between potential-outcome means.

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- Average direct, indirect, and total treatment effects are contrasts between potential-outcome means.
- The total effect is:

 $\tau \equiv E[Y(1)] - E[Y(0)] = E[Y(1, M(1))] - E[Y(0, M(0))]$

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 The effect of the treatment on the outcome through the mediator is the indirect effect:

$$\delta(t) \equiv E[Y(t, M(1))] - E[Y(t, M(0))], \quad t \in \{0, 1\}$$

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• The direct effect of the treatment is:

 $\zeta(t) \equiv E[Y(1, M(t))] - E[Y(0, M(t))], \quad t \in \{0, 1\}$

Notice that the total effect is the sum of direct and indirect effects

$$\tau = \delta(0) + \zeta(1)$$

$$\tau = \delta(1) + \zeta(0)$$

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Estimands

Denoting *E*[*Y*(*t*, *M*(*t'*))] as *Y*<sub>*tM*_{*t'*}, we define the following treatment effects of interest
</sub>

(Total) natural indirect effect (NIE)	$Y_{1M_1} - Y_{1M_0}$
(Pure) natural direct effect (NDE)	$Y_{1M_0} - Y_{0M_0}$
(Pure) natural indirect effect (PNIE)	$Y_{0M_1} - Y_{0M_0}$
(Total) natural direct effect (TNDE)	$Y_{1M_1} - Y_{0M_1}$
Total effect (TE)	$Y_{1M_1} - Y_{0M_0}$

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Decomposition 1

. mediate (wellbeing) (bonotonin) (exercise)
Iteration 0: EE criterion = 5.104e-27
Iteration 1: EE criterion = 2.031e-28
Causal mediation analysis Number of obs = 2,000
Outcome model: Linear
Mediator model: Linear
Mediator variable: bonotonin
Treatment type: Binary

wellbeing	Coefficient	Robust std. err.	Z	₽> z	[95% conf.	interval]
NIE exercise						
(Exercise vs Control)	9.799821	.3943251	24.85	0.000	9.026958	10.57268
NDE						
exercise (Exercise vs						
Control)	2.891453	.2304278	12.55	0.000	2.439823	3.343083
IE exercise (Exercise vs						
Control)	12.69127	.4005941	31.68	0.000	11.90612	13.47642
		1				

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Decomposition 2

. mediate (wellbeing) (bonotonin) (exercise), pnie tnde te

Iteration 0: EE criterion = 5.104e-27 Iteration 1: EE criterion = 3.672e-28

Causal mediation analysis

Outcome model:	Linear
Mediator model:	Linear
Mediator variable:	bonotonin
Treatment type:	Binary

Number of obs = 2,000

wellbeing	Coefficient	Robust std. err.	Z	₽> z	[95% conf.	interval]
PNIE						
exercise						
(Exercise						
VS						
Control)	9.568827	.3884522	24.63	0.000	8.807475	10.33018
INDE						
exercise						
(Exercise						
vs						
Control)	3.122447	.2418591	12.91	0.000	2.648412	3.596482
TE .						
exercise						
(Exercise						
VS					11.90612	
Control)	12.69127	.4005941	31.68	0.000		13.47642

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Causal Mediation Analysis

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Which decomposition do we want?

Nguyen, Schmid, and Stuart (2021) give suggestions for three scenarios:

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1 Want know whether there is a mediation effect? Use NIE and NDE.

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We are assuming there is some direct effect and we want to know whether any mediating effect also exists.

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We are assuming there is some direct effect and we want to know whether any mediating effect also exists.

2 Want to know whether any direct effect exists in addition to a mediation effect? Use PNIE and TNDE.

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We are assuming that some mediating effect exists and want to know whether there is an affect through any other mechanisms.

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We are assuming that some mediating effect exists and want to know whether there is an affect through any other mechanisms.

3 Have no prior assumption about whether direct or indirect effects exist? Use both decompositions.

We describe try to learn all we can from all decompositions.

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Behind the scenes

- mediate estimates all effects parameters, auxiliary parameters, and their variance-covariance matrix via generalized method of moments.
- We can specify **aequations** option to see estimated auxiliary parameters—the parameters estimated for the outcome and treatment models.

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Auxiliary parameter estimates

. mediate (wellbeing) (bonotonin) (exercise), aequations

Iteration 0: EE criterion = 5.104e-27 Iteration 1: EE criterion = 2.031e-28

Causal mediation analysis

Number of obs = 2,000

Outcome model:	Linear
Mediator model:	Linear
Mediator variable:	bonotonin
Treatment type:	Binary

wellbeing	Coefficient	Robust std. err.	Z	P> z	[95% conf.	interval]
NIE						
exercise						
(Exercise						
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Causal Mediation Analysis

Auxiliary parameter estimates

wellbeing exercise Exercise bonotonin	2.065871 .2130222	.8723559 .0034547	2.37 61.66	0.018	.3560846 .2062512	3.775657 .2197932
exercise# c.bonotonin Exercise	.0051424	.0046954	1.10	0.273	0040604	.0143452
cons	22.91374	.5633648	40.67	0.000	21.80956	24.01791
bonotonin exercise Exercise _cons	44.91939 160.544	1.641668 1.142508	27.36 140.52	0.000	41.70178 158.3047	48.137 162.7832

Note: Outcome equation includes treatment-mediator interaction.

• We defined our effects of interest in terms of potential-outcome means.

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- We need to consider what causal assumptions are required to identify those effects.

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- We defined our effects of interest in terms of potential-outcome means.
- We need to consider what causal assumptions are required to identify those effects.
- The potential-outcome means are the result of an integral of the conditional expectation of the outcome with respect to the conditional distribution of the mediator:

$$f[Y(t, M(t'))|\mathbf{X} = \mathbf{x}] = \int f[Y|\mathbf{M} = \mathbf{m}, T = t, \mathbf{X} = \mathbf{x}] dF[\mathbf{m}|T = t', \mathbf{X} = \mathbf{x}]$$

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- We need to consider what causal assumptions are required to identify those effects.
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$$f[Y(t, M(t'))|X = x] = \int f[Y|M = m, T = t, X = x] dF[m|T = t', X = x]$$

• This is sometimes referred to as the "mediation formula".

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- The potential-outcome means are the result of an integral of the conditional expectation of the outcome with respect to the conditional distribution of the mediator:

$$f[Y(t, M(t'))|X = x] = \int f[Y|M = m, T = t, X = x] dF[m|T = t', X = x]$$

- This is sometimes referred to as the "mediation formula".
- Notice that this is a nonparametric identification result.

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 In addition to assumptions for standard SUTVA and overlap assumptions, we need to make an assumption of sequential ignorability.

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- Sequential ignorability essentially means

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- Sequential ignorability essentially means
 - No unobserved confounding in the treatment-outcome relationship.
 - No unobserved confounding in the mediator-outcome relationship.
 - No unmeasured confounding in the treatment-mediator relationship.

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- In addition to assumptions for standard SUTVA and overlap assumptions, we need to make an assumption of sequential ignorability.
- Sequential ignorability essentially means
 - No unobserved confounding in the treatment-outcome relationship.
 - No unobserved confounding in the mediator-outcome relationship.
 - No unmeasured confounding in the treatment-mediator relationship.
 - There are no (observed) confounders in the mediator-outcome relationship that are caused by the treatment.

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- To meet these assumptions, we may need to add covariates to the model for the outcome, the model for the mediator, or both.
- Here, we adjust for **age** in both models before estimating potential-outcome means and the effects.

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Including covariates

. mediate (wellbeing age) (bonotonin age) (exercise)
Iteration 0: EE criterion = 6.163e-27
Iteration 1: EE criterion = 4.924e-29
Causal mediation analysis Number of obs = 2,000
Outcome model: Linear
Mediator model: Linear
Mediator variable: bonotonin
Treatment type: Binary

wellbeing	Coefficient	Robust std. err.	Z	P> z	[95% conf.	interval]
E exercise (Exercise vs Control)	9.851525	.3905733	25.22	0.000	9.086015	10.61703
E exercise (Exercise vs Control)	2.915712	.2327821	12.53	0.000	2.459468	3.371957
exercise (Exercise vs	2.913/12	.2327021	12.33		2.435400	

Kristin MacDonald (StataCorp LLC)

12 September 2024 48/62

Controlled direct effects

 What would be the causal effect if we could set the mediator to a specific value? To explore this, we estimate a controlled direct effect (CDE).

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- CDE(m) is then the average of the differences between potential outcomes.
- For binary treatment, CDE(m) is defined as Y(1|M = m) Y(0|M = m).
- Perhaps we want to know the effect of exercise on well-being if we had a medication that stabilized bonotonin levels at 200 for everyone in the population.

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. estat cde, mvalue(200) Controlled direct effect Mediator variable: bonotonin Mediator value = 200

Number of obs = 2,000

	CDE	Delta-method std. err.	Z	P> z	[95% conf.	interval]
exercise (Exercise vs Control)	3.121577	.2315869	13.48	0.000	2.667675	3.575479

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What if we have a different type of outcome, mediator, or treatment?

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What if we have a different type of outcome, mediator, or treatment?

• **mediate** allows a continuous, binary, or count outcome. You can specify a linear, logit, probit, Poisson, or exponential mean model for the outcome.

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- To demonstrate, we model a binary mediator, **bbonotonin**, which is an indicator for at least 10% increase in bonotonin over the baseline level.
- We also have a binary outcome, **bwellbeing**, which is an indicator for at least 10% improvement in well-being over the baseline value.
- We will fit a logit model for both the outcome and mediator and estimate the effects of interest using the same definitions based on potential-outcome means.

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> (bbonoto	ing basewell age, logit) nin basebono age, logit) e), nointeraction	
	riterion = 4.840e-18 riterion = 1.836e-33	
Causal mediation a	nalysis	Number of obs = $2,000$
Outcome model: Mediator model: Mediator variable: Treatment type:	Logit Logit bbonotonin Binary	

bwellbeing	Coefficient	Robust std. err.	Z	₽> z	[95% conf.	interval]
NIE						
exercise						
(Exercise vs						
Control)	.1110896	.0142334	7.80	0.000	.0831926	.1389866
NDE						
exercise						
(Exercise						
vs Control)	.146092	.0189224	7.72	0.000	.1090047	.1831792
CONCLOT)	.140092	.0109224	1.12	0.000	.1090047	.1031/92
TE						
exercise						
(Exercise						
VS	0574.046					
Control)	.2571816	.0143876	17.88	0.000	.2289824	.2853807

Note: Outcome equation does not include treatment-mediator interaction. (🗗) (🗄) ()

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• With a binary outcome, the effects are interpreted on a probability scale or as risk differences.

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- With a binary outcome, the effects are interpreted on a probability scale or as risk differences.
- We expect the probability of better well-being to be 0.26 higher if everyone in the population exercises than if no one exercises. Of that, the probability of better well-being is 0.11 higher because of an increase in bonotonin levels which and 0.15 higher because of other factors.

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- With a binary outcome, the effects are interpreted on a probability scale or as risk differences.
- We expect the probability of better well-being to be 0.26 higher if everyone in the population exercises than if no one exercises. Of that, the probability of better well-being is 0.11 higher because of an increase in bonotonin levels which and 0.15 higher because of other factors.
- We can use **estat rr** to report risk ratios or **estat or** to report odds ratios.

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Risk ratios

. estat rr

estat rr requires potential-outcome means; refitting model ...

Transformed treatment effects

Number of obs = 2,000

bwellbeing	Risk ratio	Robust std. err.	Z	₽> z	[95% conf.	interval]
NIE exercise (Exercise vs						
Control)	1.245647	.0392724	6.97	0.000	1.171004	1.325047
NDE exercise (Exercise vs Control)	1.477205	.0708189	8.14	0.000	1.344724	1.622738
TE exercise (Exercise vs Control)	1.840076	.0706258	15.89	0.000	1.70673	1.983839

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Odds ratios

. estat or

estat or requires potential-outcome means; refitting model ...

Transformed treatment effects

Number of obs = 2,000

bwellbeing	Odds ratio	Robust std. err.	Z	₽> z	[95% conf.	interval]
NIE exercise (Exercise vs						
Control)	1.562536	.0898293	7.76	0.000	1.396031	1.748901
NDE exercise (Exercise vs Control)	1.871182	.1490494	7.87	0.000	1.600713	2.187352
TE exercise (Exercise vs Control)	2.92379	.1841129	17.04	0.000	2.584315	3.307858

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Continuous treatment

- mediate supports binary, multivalued, and continuous treatments.
- When the treatment is continuous, we need to include the **continuous()** option in the treatment specification and define the values at which we want the potential-outcome means to be evaluated. The first value will be considered the control.

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Continuous treatment

. mediate (wellbeing) (bonotonin) (cexercise, continuous (30 60 90))

Iteration 0: EE criterion = 8.416e-28 Iteration 1: EE criterion = 8.416e-28 (backed up)

Causal mediation analysis

Number of obs = 2,000

Outcome model: Linear Mediator model: Linear Mediator variable: bonotonin Treatment type: Continuous

Continuous treatment levels:

```
0: cexercise = 30 (control)
1: cexercise = 60
2: cexercise = 90
```

wellbeing	Coefficient	Robust std. err.	z	P> z	[95% conf.	interval]
NIE						
cexercise						
(1 vs 0)	3.329037	.1613581	20.63	0.000	3.012781	3.645293
(2 vs 0)	6.630837	.3292353	20.14	0.000	5.985548	7.276127
NDE						
cexercise						
(1 vs 0)	.8769353	.0841601	10.42	0.000	.7119845	1.041886
(2 vs 0)	1.753871	.1683203	10.42	0.000	1.423969	2.083772
TE						
cexercise						
(1 vs 0)	4.205972	.1679266	25.05	0.000	3.876842	4.535103
(2 vs 0)	8.384708	.3394717	24.70	0.000	7.719356	9.05006

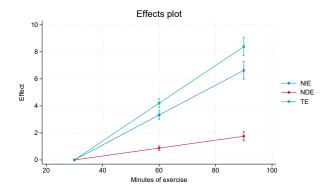
Note: Outcome equation includes treatment-mediator interaction.

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Graphing effects

 When we evaluate effects at multiple points, we can use estat effectsplot to easily compare the effects visually.



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Learn more: https://www.stata.com/manuals/causalmediate.pdf

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Thank you!

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References

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