# **Sensitivity Analyses for Decision Modeling**

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# Content

## • Why sensitivity analyses?

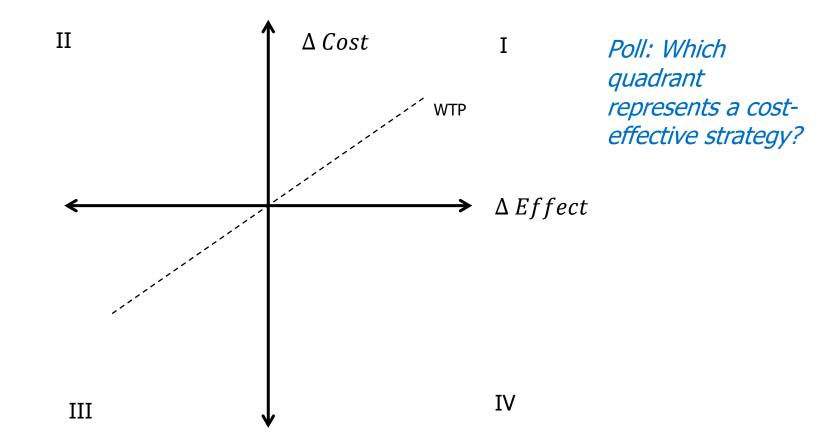
# Types of Sensitivity Analyses

- One-way sensitivity Analyses
- Tornado Diagrams
- Scenario Analyses
- Probabilistic Sensitivity Analyses

# **Output of a Decision Model**

Type of Model	Output	
Budget Impact Model	Cost per strategy	
Cost Benefit Model	Net social benefit = Incremental Benefit (cost) – Incremental Costs	Point
Cost-Effectiveness Model	$ICER = \frac{\Delta cost}{\Delta health effect}$	Estimates
Cost-utility Model	$ICER = \frac{\Delta \cos t}{\Delta QALYs}$	

### **Cost-effectiveness Model quadrants**



# **Cost-effectiveness Model quadrants**

### <u>Quadrant I</u>:

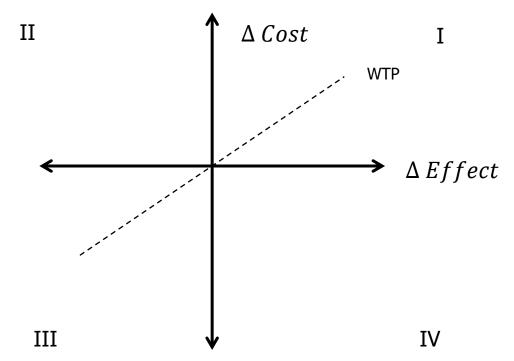
 More costly and more effective (if below WTP)

### Quadrant II:

More costly and less effective
 (No)

### Quadrant III:

 Less costly and less effective (If below WTP)

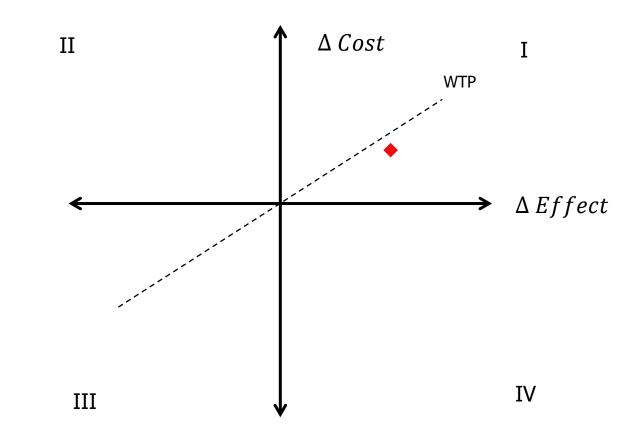


### Quadrant IV:

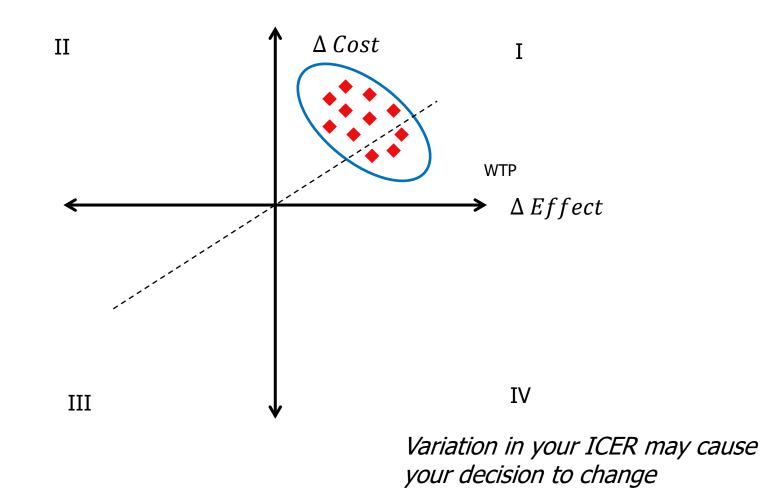
Less costly and more effective (Yes!)

## Poll 2

Would you recommend to adopt a new technology, based on this ICER result?



### **Cost-effectiveness Model output**



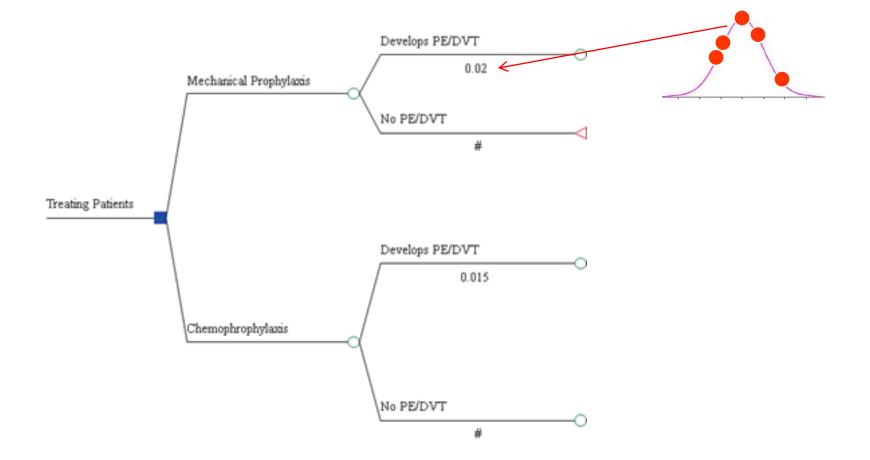
# Why sensitivity analysis?

Evaluate how uncertainty/variation in model <u>inputs</u> affects the model <u>outputs</u>

- Base-case model  $\rightarrow$  ICERs
- Sensitivity Analyses  $\rightarrow$  Variation in ICER

Statistical Analysis	<b>Cost-Effectiveness Analysis</b>		
Mean	ICER (Base-Case)		
Variation around Mean	Variation around ICER		

### Varying point estimates (TreeAge model)



# **General Approach, Sensitivity Analysis**

- 1. Change model input
- 2. Recalculate ICER
- If new ICER is substantially different from old ICER → model is sensitive to that parameter
   *In this case, it is very important to be accurate about this parameter!*

# **Types of inputs**

### Cost

### Health Effect

- Life Years Saved
- Utilities
- Cases of Disease Avoided
- Infections Cured
- Probabilities

### Discount Rate

# **Types of Uncertainty**

Term
Stochastic Uncertainty
Parameter Uncertainty
Heterogeneity

Briggs et al. 2012 Model Parameter Estimation and Uncertainty: A Report of the ISPOR-SMDM Modeling Good Research Practices Task Force – 6. *Value in Health*, 15: 835-842.

# **Types of Uncertainty**

Term	Models	How to handle in a decision model	Analagous term in regression	Example
Stochastic Uncertainty	Variation between identical patients	microsimulation	Error term	19% of Medicare beneficiaries readmitted to the hospital within 30 days. Person 1 = readmitted, Persons 2, 3, 4, 5 = not readmitted
Parameter Uncertainty	Uncertainty in estimation of parameter of interest	Probalistic sensitivity analysis (PSA)	Standard Error of the estimate	Toss a fair coin 100 times. You get 55 "heads" and 45 "tails"
Heterogeneity	Differences in patient characteristics	Scenario Analysis	Beta-coefficients/test of sig. amongst different levels of a covariate	Drug is effective for people with mild/moderate disease; it is not effective for people with severe disease

Briggs et al. 2012 Model Parameter Estimation and Uncertainty: A Report of the ISPOR-SMDM Modeling Good Research Practices Task Force – 6. *Value in Health*, 15: 835-842.

# Types of Sensitivity Analyses





# **Types of Sensitivity Analyses**

- One-way sensitivity Analyses
- Tornado Diagrams
- Scenario Analyses
- Probabilistic Sensitivity Analyses

Often Deterministic

# **Types of Sensitivity Analyses**

### Deterministic (DSA)

 model input is specified as <u>multiple point estimates</u> (sequentially) and varied manually

### Probabilistic (PSA)

model inputs are specified as a <u>distribution</u> and varied

# **DSA versus PSA**

#### Example: Cost input, cost of outpatient visit

	DSA	PSA
Base case	\$100	\$100
Input	\$80, \$90, \$110, \$120	
Results	ICER A (when cost is \$80) ICER B (when cost is \$90) ICER C (when cost is \$110) ICER D (when cost is \$120)	The mean ICER when we vary the base-case using a normal distribution with a mean of \$100 and standard deviation of \$10 is X, using 1000 iterations

# **DSA, PSA and Model structure**

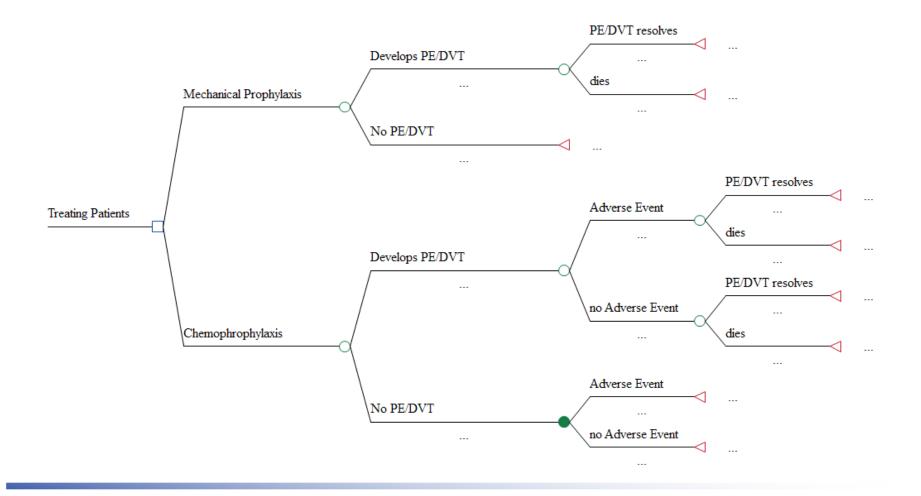
	DSA	PSA
Markov Cohort	Х	Х
Individual-level Markov Model	Х	Х
<b>Discrete-Event Simulation</b>	X	X

# Sensitivity Analyses in TreeAge

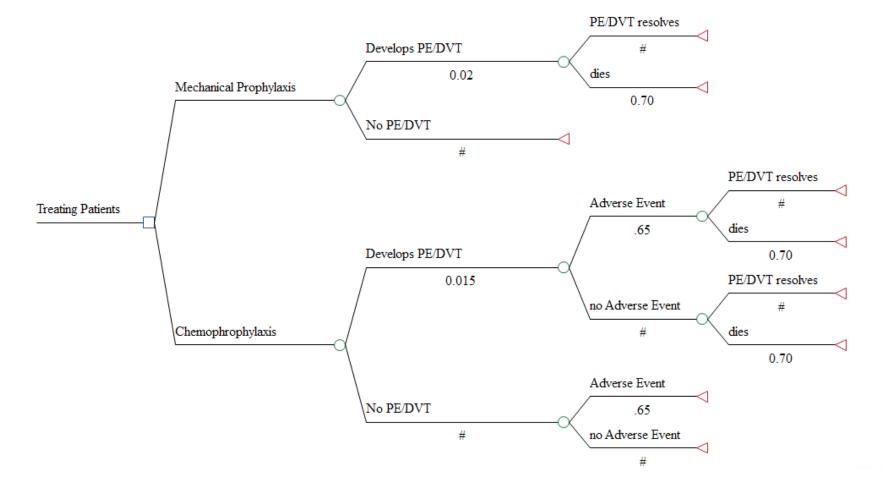




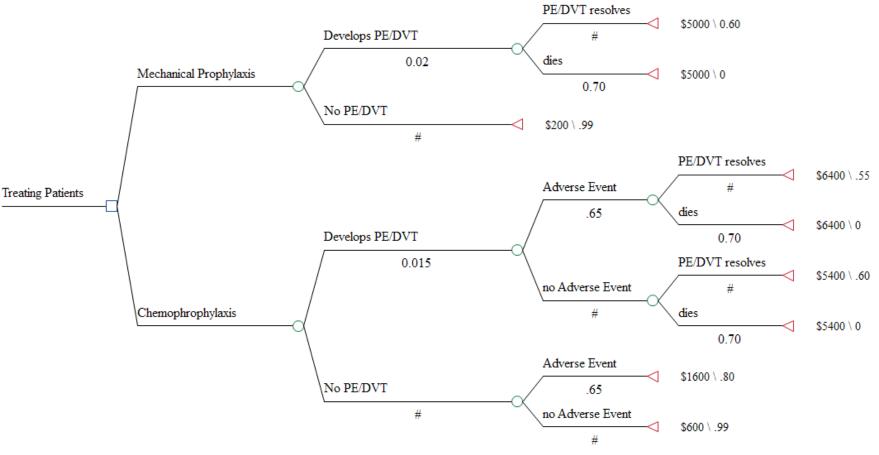
# **PE/DVT example**



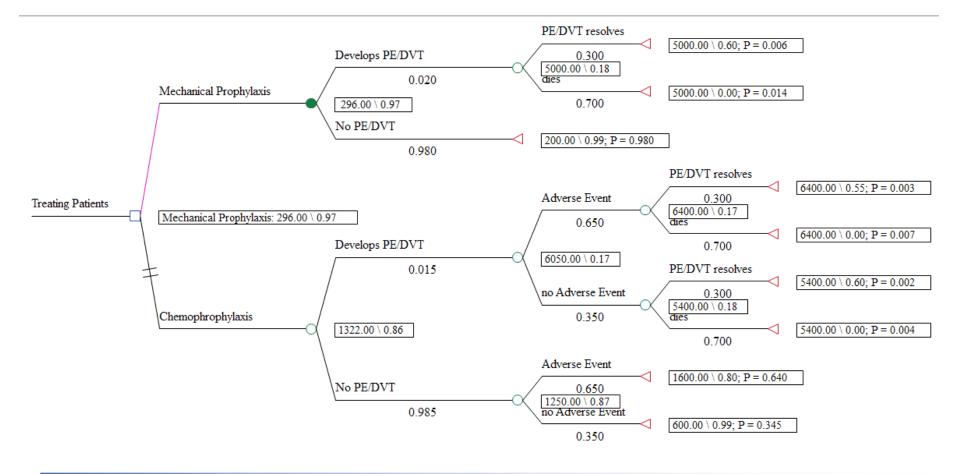
# PE/DVT example – Hypothetical Probabilities



# PE/DVT example – Hypothetical full inputs



# Model results, with point estimates



### One-Way Sensitivity Analyses





# **One-way sensitivity analysis**

- Vary one input (parameter) at a time, and see how model results are affected
- Deterministic Example: probability of AE\_chemo
  - Base-case: 0.02
  - Sensitivity analysis: range from 1-8%
    - Run 8 models, each with the following input: 0.01, 0.02, 0.03, 0.04, 0.05, 0.06, 0.07, 0.08
- Probabilistic Example
  - Base-case: 0.02
  - Sensitivity analysis: insert a *distribution*, each iteration selects a single value from this distribution to be used as the Prob of AE\_chemo

# **Inputting variables to run a sensitivity analysis: best Practices**

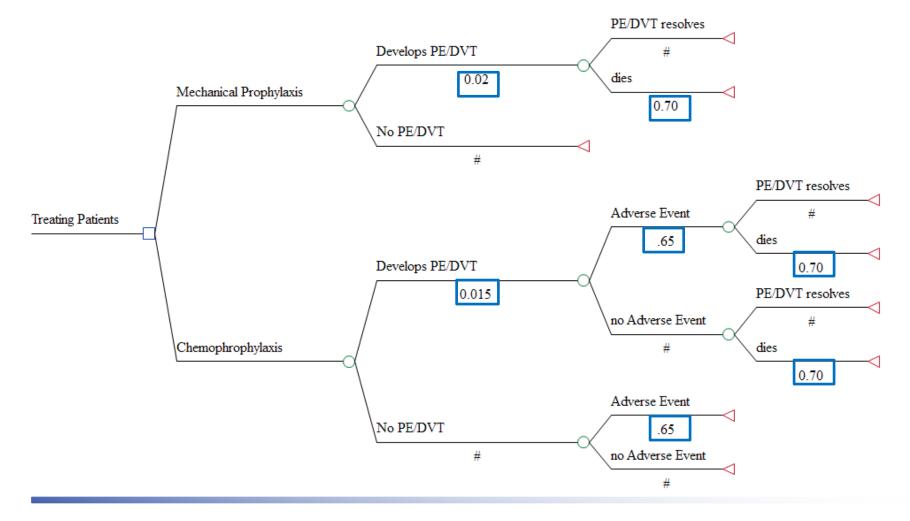
### 1. Insert variables, not point estimates

- Example: probability of PE, mechanical prophylaxis
  - "0.02" (Point estimate)
  - "p\_PEDVT\_mechan" (Variable)

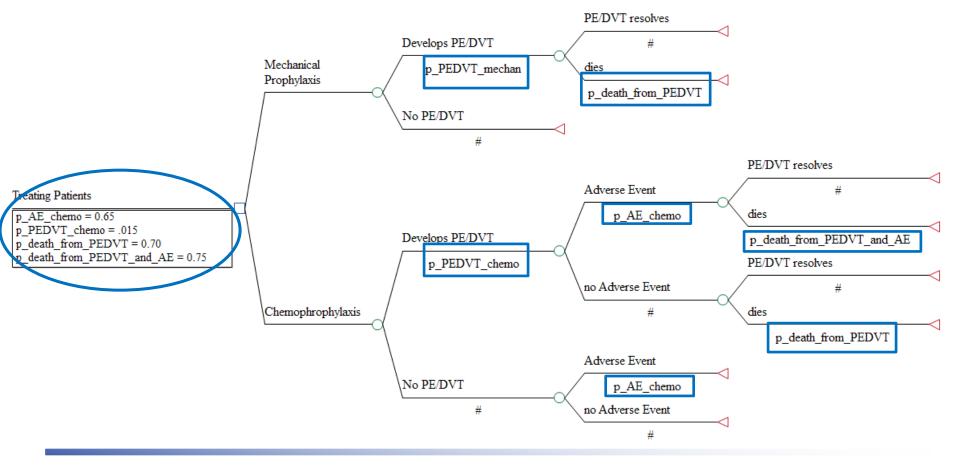
### 2. Then, define variables as:

- Point estimates (DSA) or
- Distributions (PSA)
- Example: definition of probability of PE/DVT, mechanical
  - Defining variable as a point estimate: "p\_PEDVT\_mechan = 0.02"
  - Defining variable as a distribution: "p\_PEDVT\_mechan = dist\_PEDVT\_mechan"

# PE/DVT example – Probabilities as Point Estimates



# **PE/DVT example – Probabilities as Variables and Variables defined as Point Estimates**



# **One-way sensitivity analyses**

### Define your range

#### 🌾 One-Way Sensitivity Analysis Setup

Variable	Low value	High value	Intervals	Definitions	Correlations
p_AE_chemo	0.4	0.8	4	[Treating Patients: 0	

X

# **Output, one-way sensitivity analyses**

#### Sensitivity Cost Effectiveness Analysis

p_AE_chemo	Strategy	Cost	Incr cost	Eff	Incr Eff	C/E	Incr C/E (ICER)	Dominance
<b>⊡</b> 4								
	Mechanical Prophylaxis	296.00	0.00	0.97	0.00	303.96	0.00	
	Chemophrophylaxis	1072.00	776.00	0.90	-0.07	1187.50	-10919.58	(Dominated)
ė.5								
	Mechanical Prophylaxis	296.00	0.00	0.97	0.00	303.96	0.00	
	Chemophrophylaxis	1172.00	876.00	0.88	-0.09	1325.86	-9750.26	(Dominated)
<b>⊡</b> 0.6								
	Mechanical Prophylaxis	296.00	0.00	0.97	0.00	303.96	0.00	
	Chemophrophylaxis	1272.00	976.00	0.87	-0.11	1470.22	-8985.25	(Dominated)
ė. <b>0.7</b>								
	Mechanical Prophylaxis	296.00	0.00	0.97	0.00	303.96	0.00	
	Chemophrophylaxis	1372.00	1076.00	0.85	-0.13	1620.99	-8445.76	(Dominated)
<b>⊡</b> 0.8								
	Mechanical Prophylaxis	296.00	0.00	0.97	0.00	303.96	0.00	
	Chemophrophylaxis	1472.00	1176.00	0.83	-0.15	1778.59	-8044.88	(Dominated)

# Inputs for a one-way sensitivity analysis

- Range from reported 95% Confidence Interval
- Varying a parameter an arbitrary range, such as ± 50% -- not a great practice
  - This will demonstrate model sensitivity, but does not reflect uncertainty
- Expert Opinion

# Series of One-way Sensitivity Analyses

### 1) Vary probability of chemoprophylaxisrelated adverse event

a. Compare these ICERs to base-case ICER

### 2) Vary cost of treating adverse event

a. Compare these ICERs to base-case ICER

### 3) Vary probability of death from PE/DVT

a. Compare these ICERs to base-case ICER

4) Etc.

# Caution

- Generally, a series of one-way sensitivity analyses will underestimate uncertainty in a cost-effectiveness ratio:
  - The ICER is based off of multiple parameters, not just one
  - Here, you are assuming that uncertainty exists only in one parameter
  - Solution: Probabilistic Sensitivity Analyses!

# But...

You should still do one-way sensitivity analyses!

Easy way to understand which parameters matter

# **Tornado diagrams**

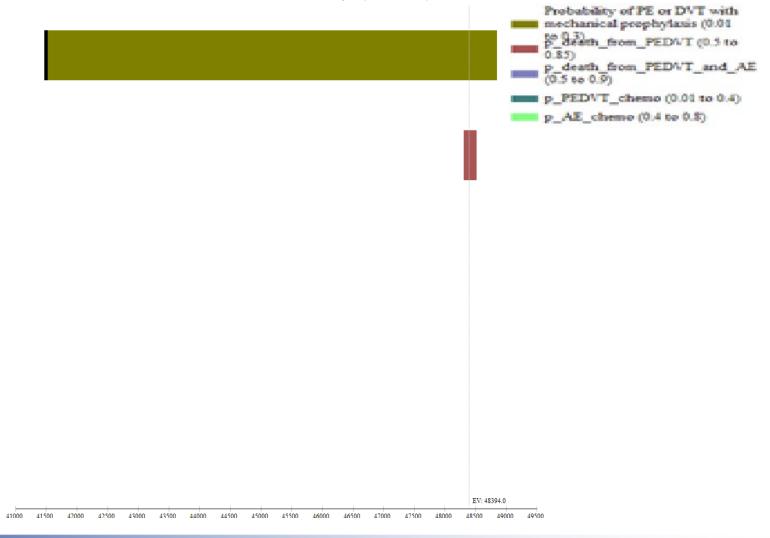
- Tell you which of your one-way sensitivity analyses had the greatest impact on model results
- Bar: a one-way sensitivity analysis
- Width of bar represents impact on model results

# **Conducting a tornado diagram**

Variable  Low    p_PEDVT_mechan  0.0    p_PEDVT_chemo  0.0    p_death_from_PEDVT  0.5    p_death_from_PEDVT  0.5	-	High value	Intervals		Add Remove 🗠 v				
p_PEDVT_mechan  0.0    p_PEDVT_chemo  0.0    p_death_from_PEDVT  0.5	)1		Intervals						
p_PEDVT_chemo 0.0 p_death_from_PEDVT 0.5	-	~ ~		Definitions	Correlations				
p_death_from_PEDVT 0.5	11	0.3	4	[Treating Patients: .02]					
	1	0.4	4	[Treating Patients: .0					
p death from PEDVT 0.5	i	0.85	4	[Treating Patients: 0					
	i	0.9	4	[Treating Patients: 0					
p_AE_chemo 0.4	ł	0.8	4	[Treating Patients: 0					
Check coherence									
Extend bars using threshold	l info								
Willingness-to-pay									
50000									
Calculation type									
Net monetary benefits									
C Net health benefits									
					OK Cancel				

# **Tornado Diagram (Net Benefits)**

Tornado Analysis (Net Benefits)



# Tornado Results (ICER) – recommended graph to view

Probability of PE or DVT with mechanical prophylaxis (0.01 Stath from PEDVT (0.5 to 0.85% p death from PEDVT and AE (0.5 to 0.9) p\_PEDVT\_chemo (0.01 to 0.4) p\_AE\_chemo (0.4 to 0.8) EV: -8694.04032 -45000 -40000 -25000 Ó 5000 -35000 -30000 -20000 -15000 -10000 -5000

# Tornado diagram, text report

#### **Tornado Sensitivity Analysis - ICER Report**

VARIABLE_NAME	VARIABLE_RANGE	LOW_VALUE		SPREAD	SPREAD_SQR	RISK_PCT	CUMUL_PCT
p_PEDVT_mechan	0.01 to 0.3	-43639.51223	599.24346	44238.75569	1957067504.59758	35.90785	35.90785
p_AE_chemo	0.4 to 0.8	-10919.58067	-8044.87618	2874.70449	8263925.87916	0.15162	36.09902
p_PEDVT_chemo	0.01 to 0.4	-8755.5842	-7313.90762	1441.67658	2078431.34776	0.03813	35.94598
p_death_from_PEDVT	0.5 to 0.85	-8792.95107	-8565.56971	227.38136	51702.28401	0.00095	35.94693
p_death_from_PEDVT_and_AE	0.5 to 0.9	-8793.94024	-8635.18248	158.75776	25204.02665	0.00046	35.94739

- The high value for p\_PEDVT\_mechan results in chemoprophylaxis now being the preferred strategy
- Tells us we need to be more precise with our estimate of PE/DVT associated with mechanical prophylaxis

# **Limitations of Tornado diagrams**

 Just a series of one-way sensitivity analyses, with results presented on top of one another

There is not just uncertainty in one parameter – there is uncertainty in most, if not all, parameters

#### Scenario Analyses





### **Scenario analyses**

- Interested in subgroups
  - Cost-effectiveness of chemical versus mechanical prophylaxis in 85+ only
    - Change risk of PE/DVT, risk of AE, risk of death from PE/DVT/AE
- Changes the <u>point estimate</u> of multiple parameters
- Do not incorporate uncertainty !

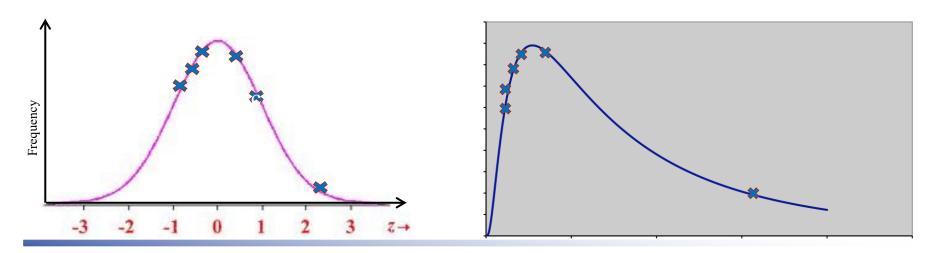
#### Probabilistic Sensitivity Analyses





# **Probabilistic sensitivity analysis**

- Vary multiple parameters simultaneously
- Each variable comes from a *distribution*
- Model is run many times (1,000, 10,000, etc.)
  - Each model iteration plucks a value from that distribution and uses it as the model input





- Values are sampled with replacement!
- Values sampled based on their likelihood of occurrence
- **Results** (comparing strategy A to B):
  - Mean Cost<sub>A</sub> & variation in Cost<sub>A</sub>
  - Mean Cost<sub>B</sub> & variation in Cost<sub>B</sub>
  - Mean Health Effect<sub>A</sub> & variation in Health Effect<sub>A</sub>
  - Mean Health Effect<sub>B</sub> & variation in Health Effect<sub>B</sub>

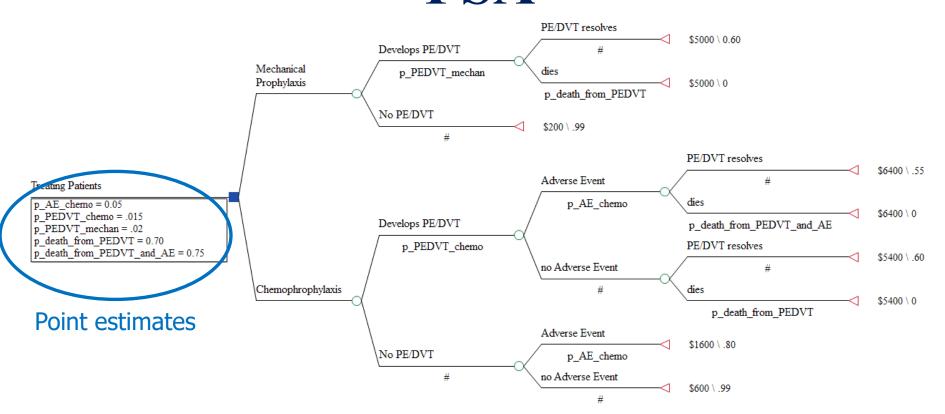
**Choosing distributions for your PSA – general guidance** 

Costs: log-normal, normal

Probabilities: beta

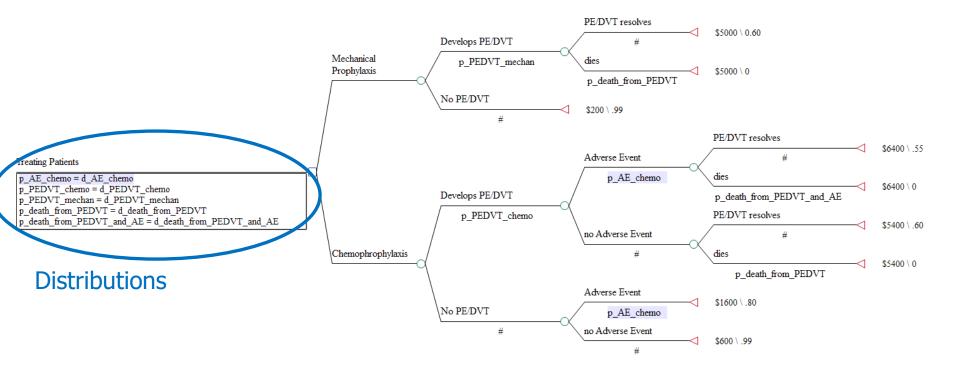
Utilities: beta

#### **Inputting variables into your PSA**



 Need to define variables in terms of distributions, rather than point estimates

### **Defining distributions in a PSA**



# Creating distribution-based definitions

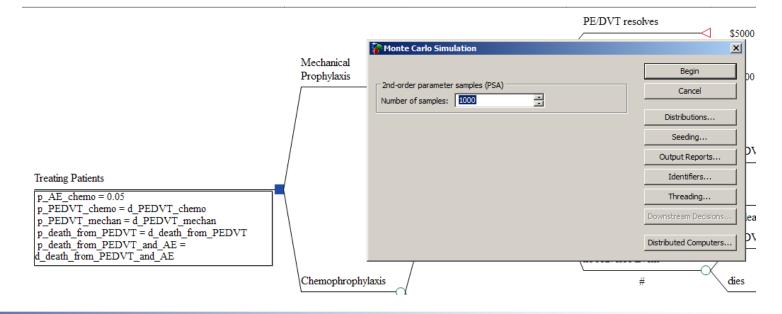
- 1. Create the distribution: d\_AE\_chemoprophyalxis
  - Define the distribution in terms of its shape
    - normal, beta, etc
  - Define the parameters for that distribution
    - mean/variance, alpha/beta, etc.

#### 2. Assign the distribution to a variable: prob\_AE\_chemoprophylaxis = d\_AE\_chemoprophylaxis

# **Running a PSA**

#### Define all variables (model inputs) as distributions

Determine your number of iterations



# Ways to show uncertainty in the ICER

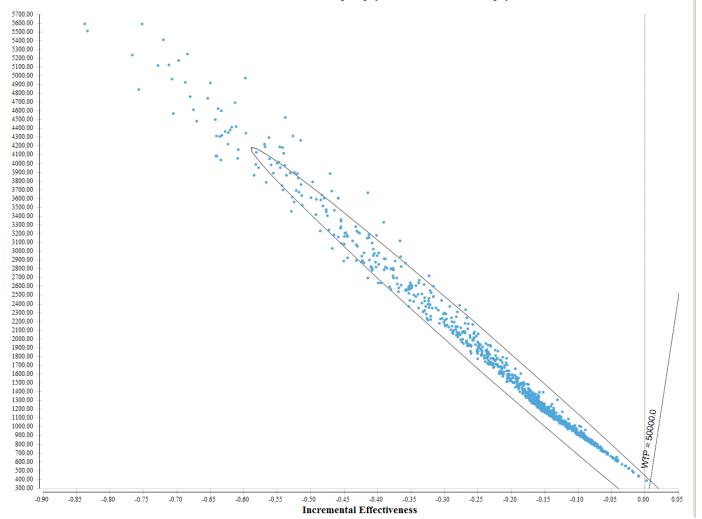
Cost-effectiveness planes (CE scatterplot)

Cost-effectiveness acceptability curve

Net benefits

#### **CE Scatter Plot**





## "ICE Report"

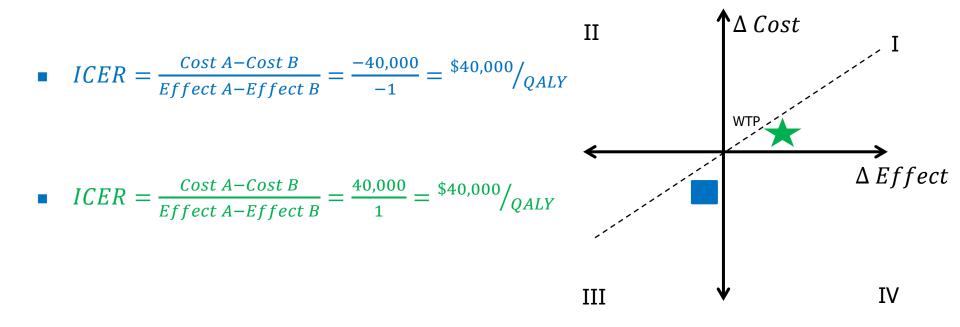
Incremental CE Plot Report Chemophrophylaxis v. Mechanical Prophylaxis						
COMPONENT	QUADRANT	INCREFF	INCRCOST	INCRCE	FREQUENCY	PROPORTION
C1	IV	IE>0	IC<0	Superior	0	0
C2	I	IE>0	IC>0	ICER<50000.0	0	0
C3	III	IE<0	IC<0	ICER>50000.0	0	0
C4	I	IE>0	IC>0	ICER>50000.0	1	0.001
C5	III	IE<0	IC<0	ICER<50000.0	0	0
C6	п	IE<0	IC>0	Inferior	999	0.999
Indiff	origin	IE=0	IC=0	0/0	0	0

In this hypothetical example (with entirely made-up data) Mechanical Prophylaxis is cost-effective compared to Chemo Prophylaxis 99.9% of the time

- Costs less AND provides more health benefit

# Ways one should <u>not</u> show uncertainty in the ICER

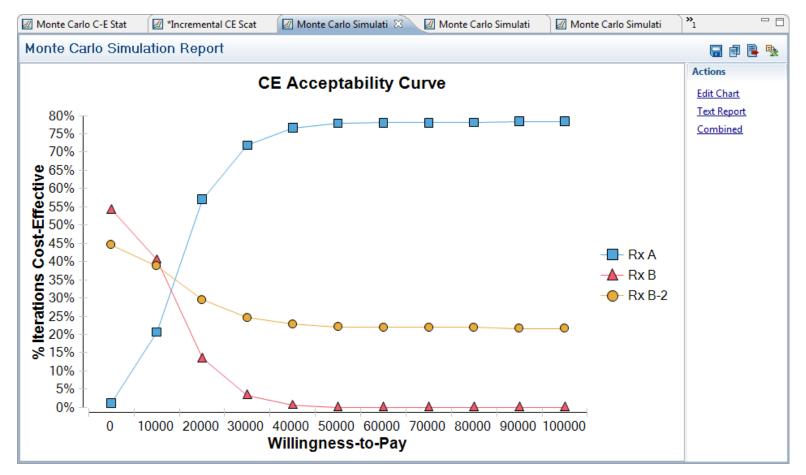
- Show only the numeric value of the ICER and Confidence Interval



# Willingness to pay (WTP)

- Previously, I had to specify my WTP
- What if you don't know what that is?
  Or different decision makers have different WTP?
- Use a <u>Cost-Effectiveness Acceptability Curve</u>
  Percentage of iterations that favor each strategy, over a range of WTP

# Cost-effectiveness acceptability curves – hypothetical



# How many iterations in a PSA?

- More distributions = more iterations
- Stop when the simulations generate mean values (without seeding) that are very similar

#### **Monte Carlo C-E Statistics**

Attribute	Statistic	Mechanical Prophylaxis	Chemophrophyl	Attribute	Statistic
⊡ Cost				- Cost	
	Mean	295.98	1371.17		Mean
	Std Deviation	14.14	966.99		Std Devia
	Minimum	258.19	614.93		Minimum
	2.5%	270.26	625.63		2.5%
	10%	278.24	645.27		10%
	Median	295.36	944.17		Median
	90%	315.24	2839.58		90%
	97.5%	325.44	4053.16		97.5%
	Maximum	338.22	5235.56		Maximum
	Size (n)	1000.00	1000.00		Size (n)
	Variance	199.99	935077.00		Variance
	Variance/Size	0.20	935.08		Variance/
	SQRT[Varianc	0.45	30.58		SQRT[Va
⊡Eff	Mean	0.97	0.86	<b>⊡</b> ∙Eff	Mean

#### **Monte Carlo C-E Statistics**

All de la	Obe Kelle	Mashanial Prophylauia	Channelburghad
Attribute	Statistic	Mechanical Prophylaxis	Chemophrophyl
Cost	-		
	Mean	295.92	1351.17
	Std Deviation	13.87	900.21
	Minimum	258.06	613.43
	2.5%	270.30	631.42
	10%	277.89	651.39
	Median	294.83	950.08
	90%	313.93	2682.31
	97.5%	322.97	3850.64
	Maximum	347.62	5115.89
	Size (n)	1000.00	1000.00
	Variance	192.33	810375.85
	Variance/Size	0.19	810.38
	SQRT[Varianc	0.44	28.47
Ė Eff	_		
	Mean	0.97	0.86

#### **100 iterations**

#### **Monte Carlo C-E Statistics**

Attribute	Statistic	Mechanical Prophylaxis	Chemophrophyl
⊡ Cost			
	Mean	297.80	1413.88
	Std Deviation	13.17	919.06
	Minimum	269.18	613.56
	2.5%	278.24	620.09
	10%	281.11	654.41
	Median	295.40	1056.64
	90%	315.54	2697.37
	97.5%	324.32	3593.22
	Maximum	336.49	5047.80
	Size (n)	100.00	100.00
	Variance	173.49	844673.03
	Variance/Size	1.73	8446.73
	SQRT[Varianc	1.32	91.91
⊨ Eff			
	Mean	0.97	0.85

#### **Monte Carlo C-E Statistics**

Attribute	Statistic	Mechanical Prophylaxis	Chemophrophyl
⊡. Cost			
	Mean	296.30	1274.05
	Std Deviation	14.44	891.76
	Minimum	260.79	614.87
	2.5%	261.01	626.80
	10%	280.79	641.58
	Median	296.48	929.81
	90%	315.42	2678.31
	97.5%	322.91	3994.27
	Maximum	335.50	4528.79
	Size (n)	100.00	100.00
	Variance	208.37	795237.48
	Variance/Size	2.08	7952.37
	SQRT[Varianc	1.44	89.18
⊨ • Eff	_		
	Mean	0.97	0.88

### **PSA Summary**

- Looks at model results when multiple sources of uncertainty are evaluated simultaneously
- Results presented in terms of:
   C-E planes (quadrants)
   C-E acceptability curves
- Required in order to publish in a peerreviewed journal!

#### Joint Parameter Uncertainty

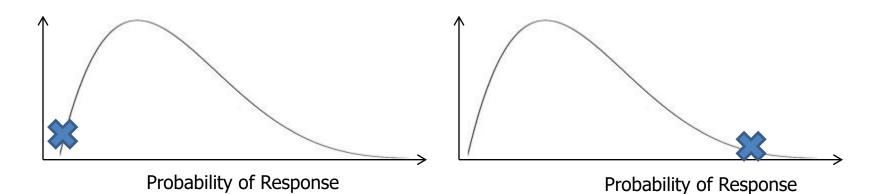




Joint Parameter uncertainty The model will assume no covariance between parameters unless you specify otherwise

Probability of response at 26 weeks

Probability of response at 52 weeks



# Accommodating Joint Parameter uncertainty

Define one variable in terms of the other

X = Y + (Y \* 0.2)

Use a table to link variables, have PSA identify Index

Variable X = if(PSA = 1; Table 1[Index; 1]; 0.55) -

Variable Y = if(PSA = 1; Table 1[Index; 2]; 0.65)

Index	X	Y
1	0.60	0.67
2	0.480	0.89
3	0.89	0.93

- If the PSA indicator is turned on:
  - go to Table 1, choose the row (Index) corresponding with the model cycle we are in and use the value in column 1
- otherwise, use a value of 0.55

#### SUMMARY

#### Summary

- All model inputs have variation/uncertainty
- Test how variation/uncertainty affects model results
  Do so by varying model inputs
- Tornado diagrams: first-pass understanding of the most important variables in your model
- Need to run a PSA in order to fully evaluate the combination of variation/uncertainty in all/most model inputs on robustness of model results

– Be careful to accommodate joint parameter variation

### References

#### General Overview:

 Hunink M, Glasziou P, Siegel J, et al. "Chapter 11: Variability and Uncertainty" in <u>Decision Making in Health</u> <u>and Medicine: Integrating Evidence and Values</u>. Cambridge, UK: Cambridge Press, 2004. 339-363.

#### Best Practices:

 Briggs et al. Model Parameter Estimation and Uncertainty: A Report of the ISPOR-SMDM Modeling Good Research Practices Task Force – 6. *Value in Health*, 2012, 15: 835-842.

#### **QUESTIONS?**