How to measure what people prefer: Health preference research to optimize health-related interventions

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Session 1 Welcome and introductions



Head of Countries



JASON J. ONG — AUSTRALIA



NITTAYA PHANUPHAK — THAILAND



NINA T. CASTILLO-CARANDANG — PHILIPPINES

HOPE Project Manager



WARITTHA TIEOSAPJAROEN — AUSTRALIA

 Aim to reduce HIV transmission in East Asia by optimizing the roll-out, implementation, and real-world effectiveness of pre-exposure prophylaxis (PrEP) in Australia, Thailand, and the Philippines









 Establish a network of regional experts from research, communities and policymakers, we will co-design intervention strategies with key populations to improve PrEP care adherence and persistence among key populations.





- Ηόρε
- These strategies will be informed by socially innovative methods (e.g., crowdsourcing, discrete choice experiments) and implemented in each country site.
- (2024 economic evaluations alongside trials)





Overall aims of the workshop

Introduce health preference research with a focus on DCE
 Describe the steps of conducting a DCE
 Share examples of DCEs in health research

Program

- Each session
 - ~30 minutes of lectures
 - ~30 minutes of group work

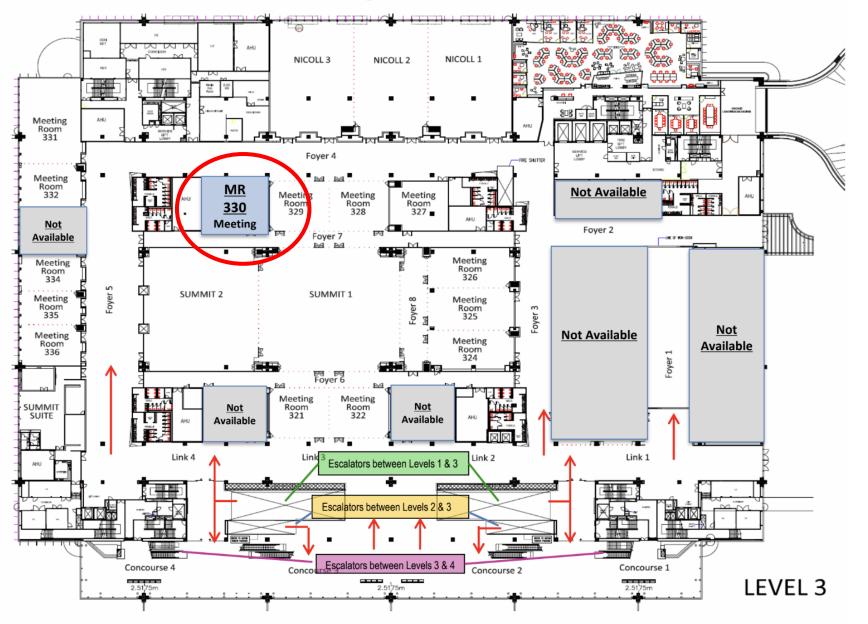


Logistics

- Toilets
- Food
- Emergency Exit

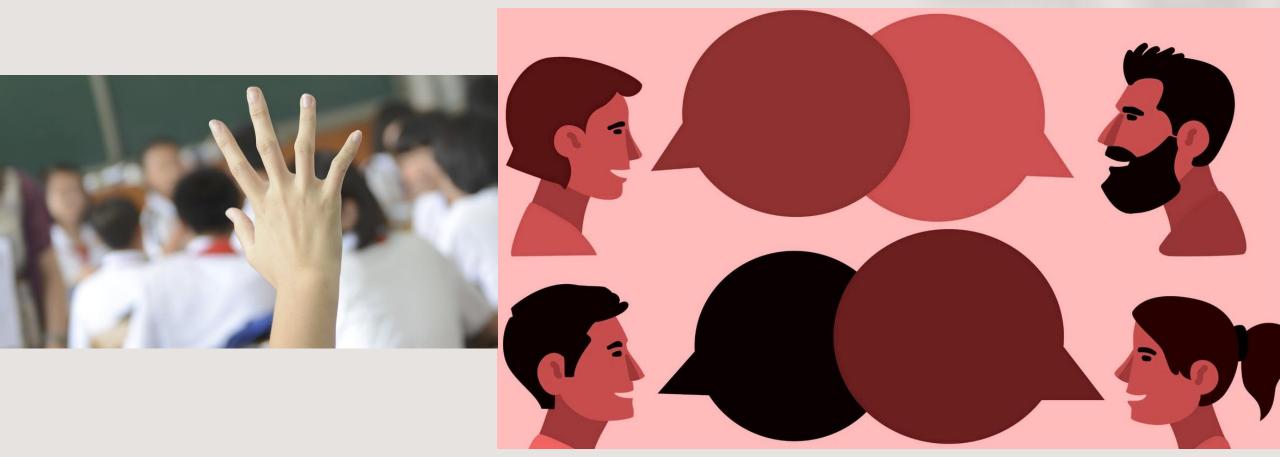


Suntec Singapore - Level 3



- Seek first to understand, then to be understood
- Maintain curiosity
- Share the floor and let others speak
- Respect different views, needs and priorities
- Use "I" statements own your experience and opinions
- Maintain confidentiality de-identify stories where necessary
- Assume good intentions
- Be supportive and collaborative
- Enjoy the process

Learn from one another





Who am I?



Sexual health physician

- Melbourne Sexual Health Centre
- Clients with HIV/STI

Researcher that models

- Understanding epidemics and human behaviors/preferences to improve health services (and outcomes)
 - Infectious disease and statistical models
 - Decision modelling



Join at **slido.com #8942 967**



https://admin.sli.do/event/1zLjSiS7yF eFaBoBCJDo8T/polls

Session 2. Overview of how to conduct discrete choice experiments

Objectives

- Describe the motivation for conducting DCEs
- Describe limitations of DCEs
- Present the steps for conducting a DCE

Motivation

Outline

What is a DCE?

Steps of conducting a DCE

Motivation

Motivation

- A one-size-fits-all models does not work for all people.
- People-centred care = Gold standard
 - Precision public health
 - Differentiated service delivery
 - responsive, client-centred approach that simplifies and adapts services to better serve individual needs and reduce unnecessary burdens on the health system.

Preference sensitive care

Incorporate patient preferences and values – "patient voice"



Monthly Every 2 months Every 3 months Every 6 months



Physician Clinical officer Nurse Pharmacist Community health worker Patient / peer / family Client



HIV clinic / hospital Primary care clinic Drop-in centre Community Home

WHAT

ART initiation / refills Clinical monitoring Adherence support Laboratory tests Opportunistic infections treatment Psychosocial support

YOU CAN AFFORD ANYTH NG. YOU JUST CAN'T AFFORD EVERYTHING.





How do YOU evaluate people's values and preferences?

How do we make choices?

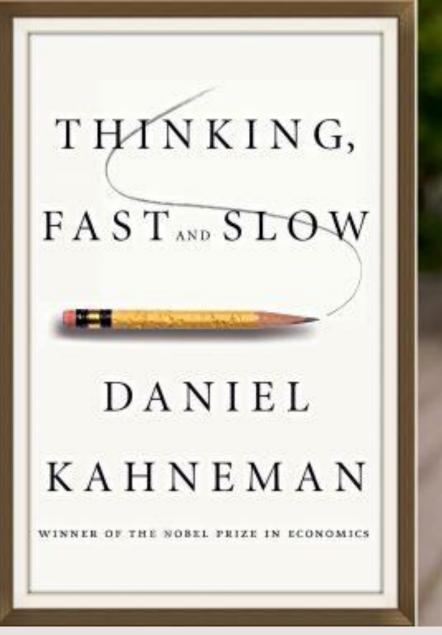


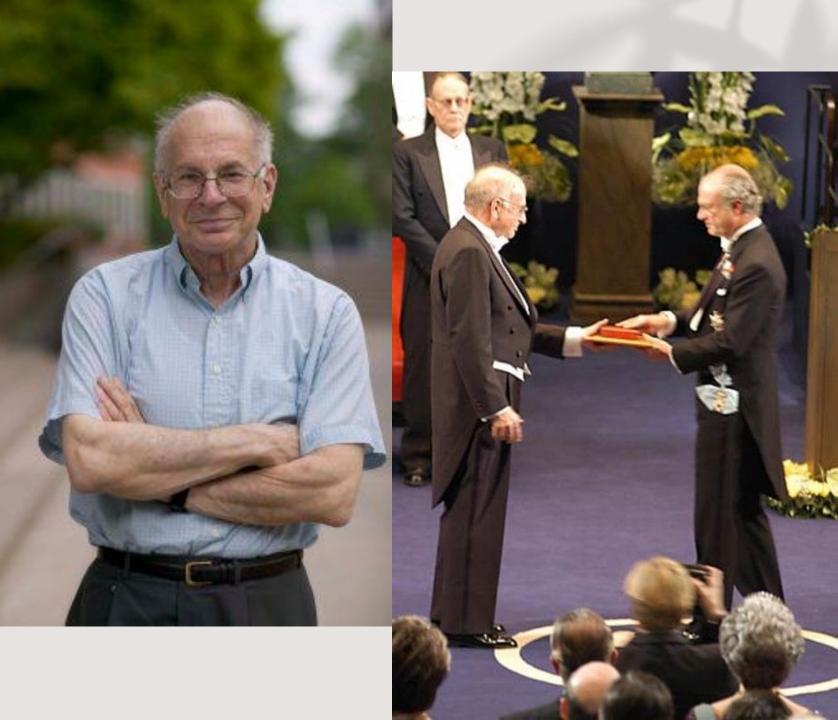


Economic theory: humans make rational decisions









System 1 Intuitive and instinctive

Unconscious

Fast

Associative

Automatic - you have no control over it



System 2 Rational and logical

Effortful

Slow

Logical

Deliberate - you control yourself

Daniel McFadden's Nobel prize speech

- <u>https://www.nobelprize.org/prizes/economic-</u> <u>sciences/2000/mcfadden/lecture/</u>
 - Discrete choice modeling



Health preference research

- "Choice defines value".
- Determine the type of goods and services most valued by a target population
- With a better understanding of what patients want, providers, regulators, and policy makers can better meet the patient's needs
- Resources are allocated efficiently and to where they are most valued and needed.

Health preference research

Revealed preference

- Observe choices of consumers in the market
- No information on potential demand for new products



Health preference research

Stated preference

- Hypothetical choices
- When market information is not available (e.g. new products)
 - Potential impact of proposed changes or design of products, policies or programs



Health preference research

Systematically identify what people want so that providers, regulators, policy makers can better meet their needs.

"Preference sensitive" goods and services





What is a DCE?

Discrete choice experiment?

Choice set?

Attributes?

Multinomial logit models?

What is a DCE?

- Stated preference method
 - Quantitative measurement of relative preference
 - How do people trade-off between attributes?
- Goods/Services described as combination of attributes
- People rationally choose the combination that maximizes their utility

What is a DCE?

"Choice set"

 Choices made from a set of mutually exclusive and collectively exhaustive alternatives

Choice modelling

- Parameters are estimated from a sample of observed choices made by decision makers when confronted with a choice situation
- Identifying most influential parameters that led to observed choices and accounting for sources of unobserved influence

Choice set



If these were your only options, which of the following laptops would you choose to purchase?

Brand	Microsoft	Apple	Google	
Operating System	Windows 10	OS X	Chrome OS	
Screen Size	13.5″	13″	12″	
Battery Life	12 hours	10 hours	12 hours	
Front Camera	Yes	Yes	No	
Rear Camera	Yes	No	No	
Stylus	Yes	No	No	I would not
Removable Keyboard	Yes	No	No	choose any o
Price	\$1,499	\$1,299	\$999	these.
	0	0	0	0

Alternatives

If these were your only options, which of the following laptops would you choose to purchase?

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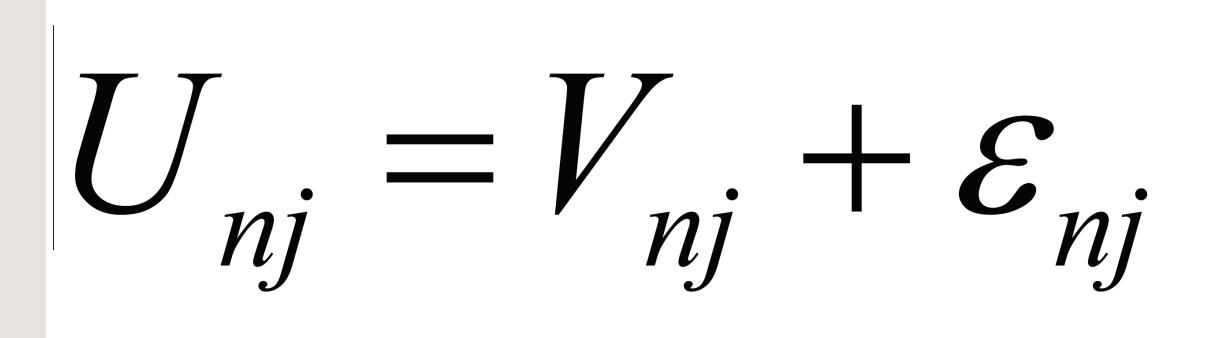
Levels

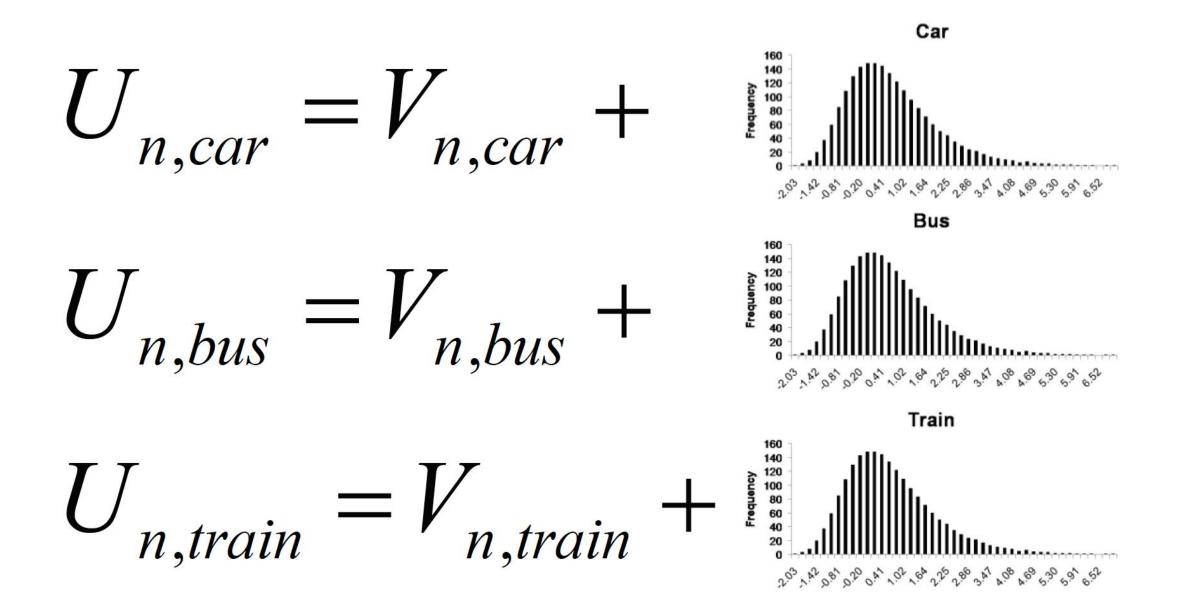
Choice modelling

Consumer theory

- People choose goods/services to maximise their utility
- Utility
 - Satisfaction, value
- Assume goods/services can be described by series of attributes and respondent will consider all attributes before choosing one that gives the best utility

Utility (U) = observed component (V) + unobserved component (ϵ)





- − U*(choice 1) > U*(choice 0) \rightarrow Choose 1
- − U*(choice 1) ≤ U*(choice 0) → Choose 0



Used in many disciplines





Lays

Masala

28 g - Rs 10.00

Evening

1

Standard Delivery: Tomorrow

The Express Delivery: Out of stock

ADD 📷

MRP: Rs 10

Bingo Yumitos Potato Chips - Original Style, Chilli

28 g - Rs 9.50 MRP:Rs 10 Rs 9.50 Standard Delivery: Tomorrow Evening The Express Delivery: Out of stock

ADD 📷



Potato Chips - Calm Cream & **Onion Flavour**

30 g Pouch - Rs 10.00 MRP: Rs 10

bbstar Price Rs 9.70 🕐

Standard Delivery: Tomorrow Evening

Express Delivery: Out of stock Qty 1 ADD 📾



Potato Chips - Flirty Tomato Tango

30 g Pouch - Rs 10.00 MRP: Rs 10 Standard Delivery: Tomorrow Evening





Attribute	Situation A		Situation B	
Re-opening schools	In 4 weeks		Immediately	
Re-opening restaurants and bars	Immediately	101	In 4 weeks	
Tracing app	Voluntary		Mandatory	§ 🖻
Quarantine for persons above 70 years	No	ijŶ	Yes	
Available ICU capacities	Sufficient) (M	Temporarily overloaded	!
Unemployment rate	10 %	₩	5 %	İ
I decide for		1]

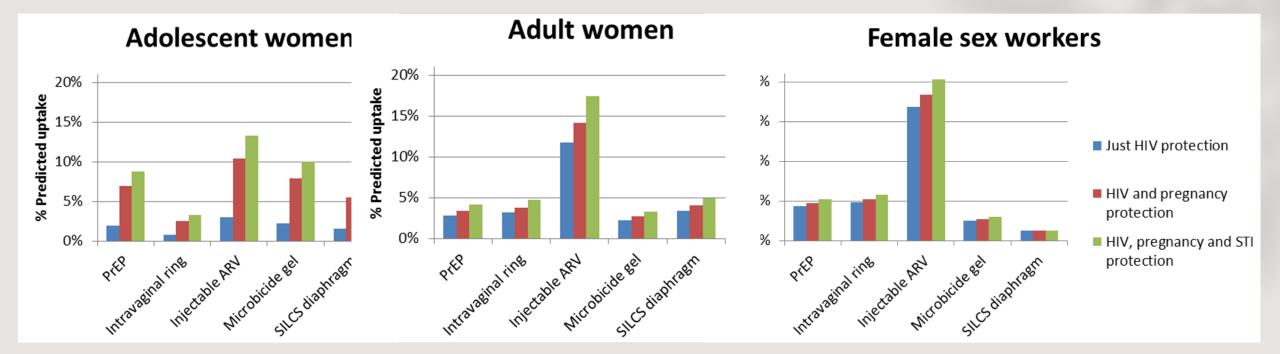
Useful outputs

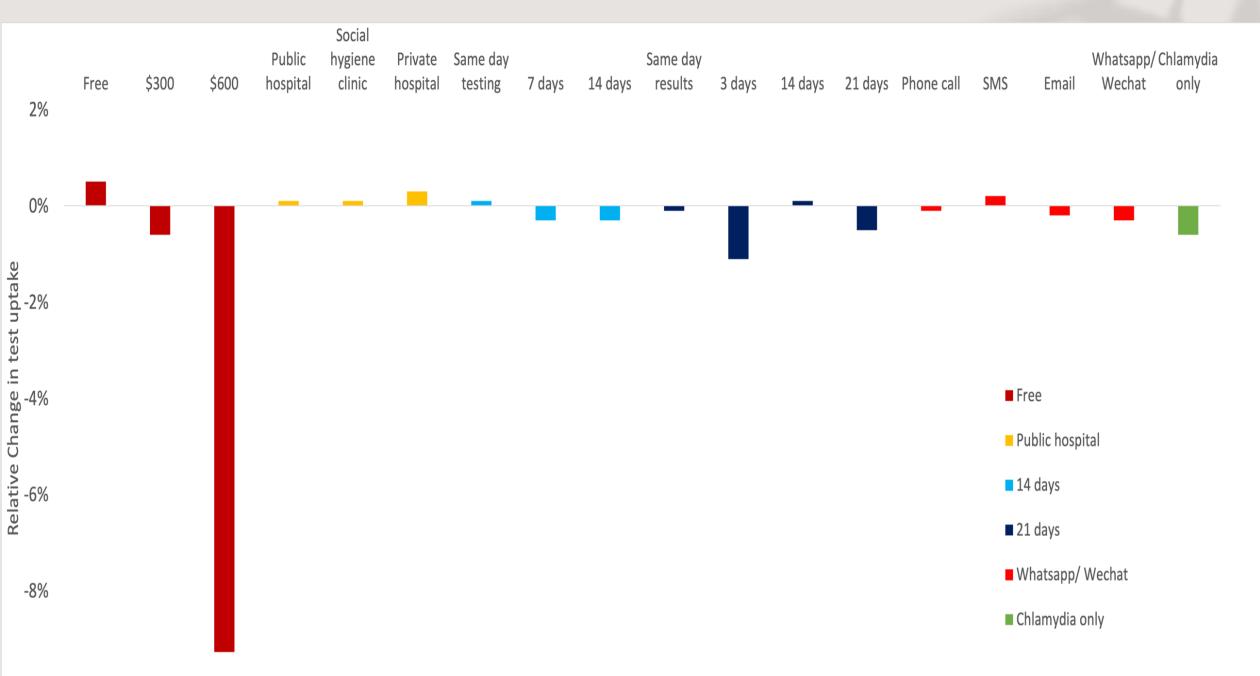
Valuation of individual components and overall valuation

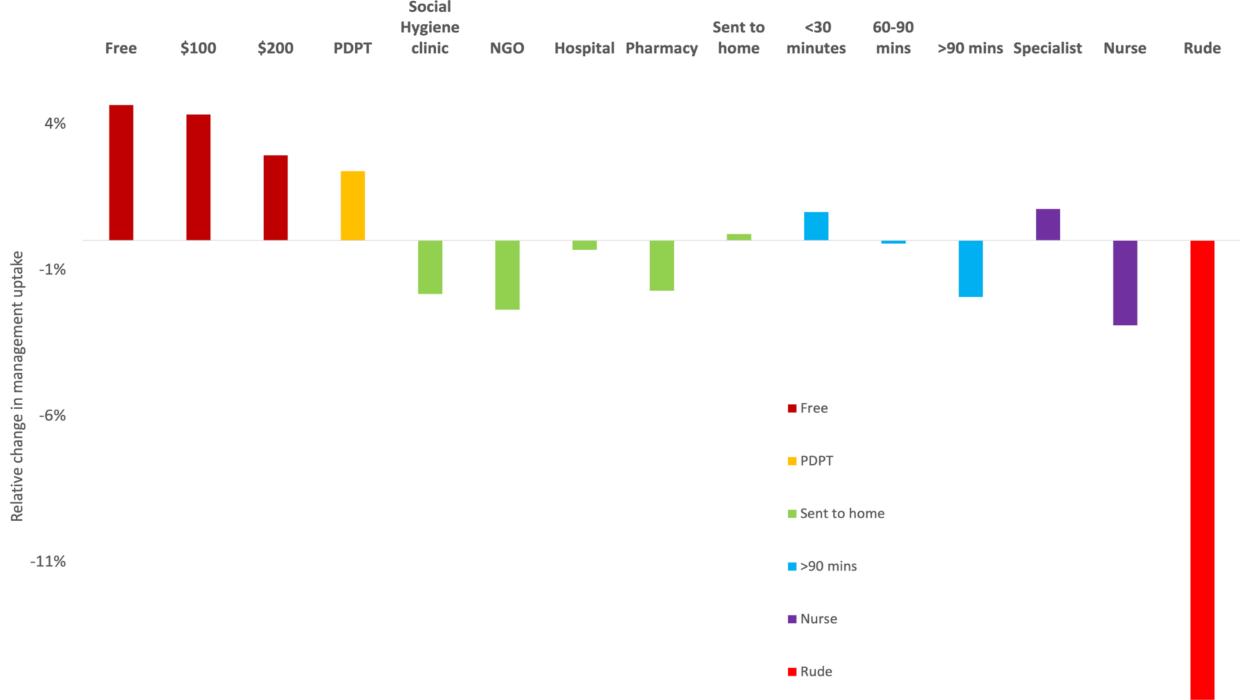
- Exploring heterogeneity
 - Subpopulations
- Willingness to pay

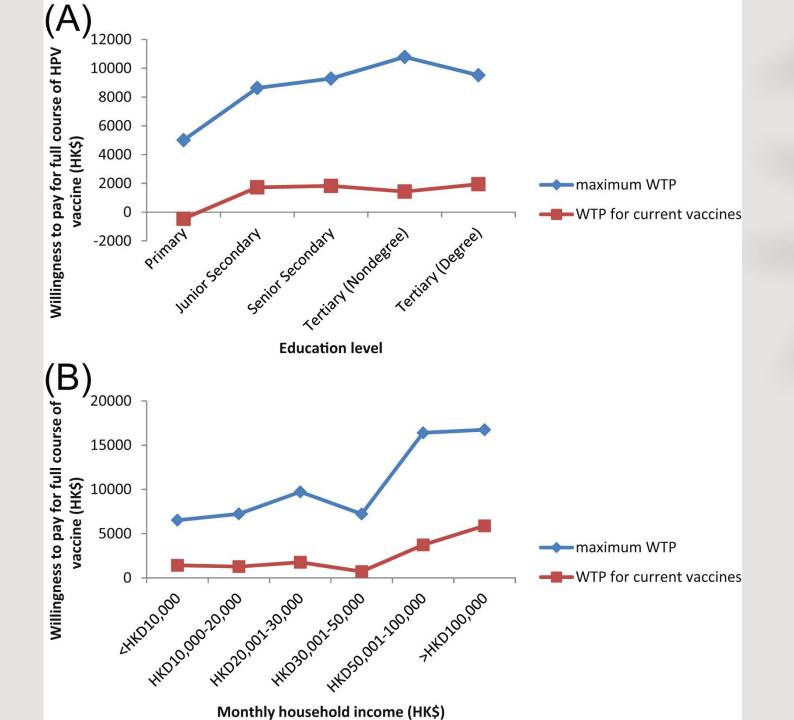
- Forecasting of choices/demand in specific scenarios
 - Predicted market shares

Uptake Predictions: scenario 5 all products









Limitations

- Hypothetical bias
 - Framing
 - Providing enough information
- External validity
 - Capturing right attributes/levels need good inputs
 - Stability of preferences over time
- Assumption of utility maximization
 - Reality can be more complicated ... emotions, social norms (?)
 - Predicting how you will act ...

But do DCEs generate valid uptake predictions?

0

0

0.2

0.4

0.6

Specificity

0.8

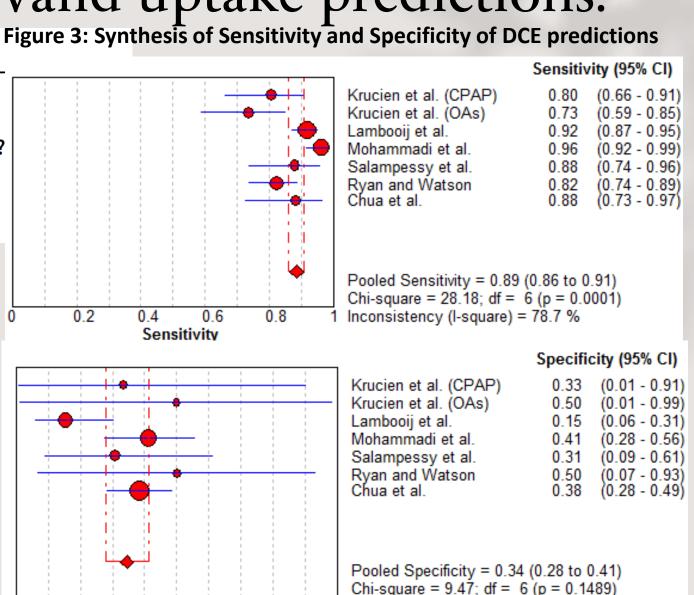
The European Journal of Health Economics https://doi.org/10.1007/s10198-018-0954-6

ORIGINAL PAPER

How well do discrete choice experiments predict health choices? A systematic review and meta-analysis of external validity

Matthew Ouaife¹ · Fern Terris-Prestholt¹ · Gian Luca Di Tanna² · Peter Vickerman³

- Few studies estimate DCE prediction validity (7)
- Pooled sensitivity and specificity estimates were 88% and 34%, respectively.
- DCEs are better at predicting who would opt-in to a health-related decision rather than who would not"



Inconsistency (I-square) = 36.6 %



- Approach to quantitatively estimate preferences for product/service characteristics
- Choice sets built from qualitative phase
- Survey with repeated scenario responses with varying characteristics
- Force trade offs between attributes allowing quantitative estimation relative values (utilities).
- Can also be used to value health states, estimate societal preferences and WTP



- A.Discrete choices
- B. Choice sets
- C.Alternatives defined by combination of attributes
- D.Survey with multiple choice sets
- E. Respondent characteristics

Steps for conducting a DCE

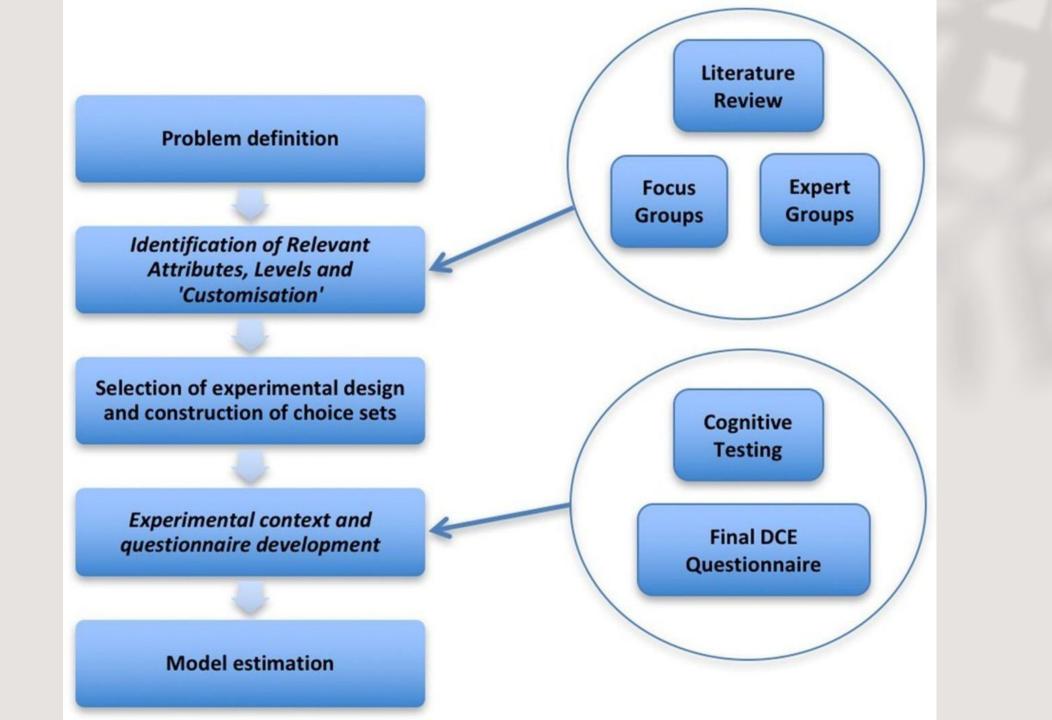


Steps of a DCE

USER GUIDE WITH CASE STUDIES

How to Conduct a Discrete Choice Experiment for Health Workforce Recruitment and Retention in Remote and Rural Areas







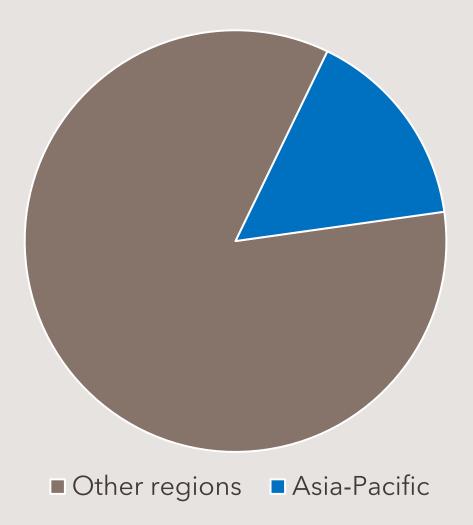
Questions?

Session 2 – small group discussion

Introduction to the scenario of HIV in East Asia

Tasks

- Name your group
- Which population(s) do you want to focus on?
- How do you identify relevant attributes / levels?
 - What methods can you use?
- List all likely attributes that would influence someone using PrEP



38.4 million people living with HIV in 2021

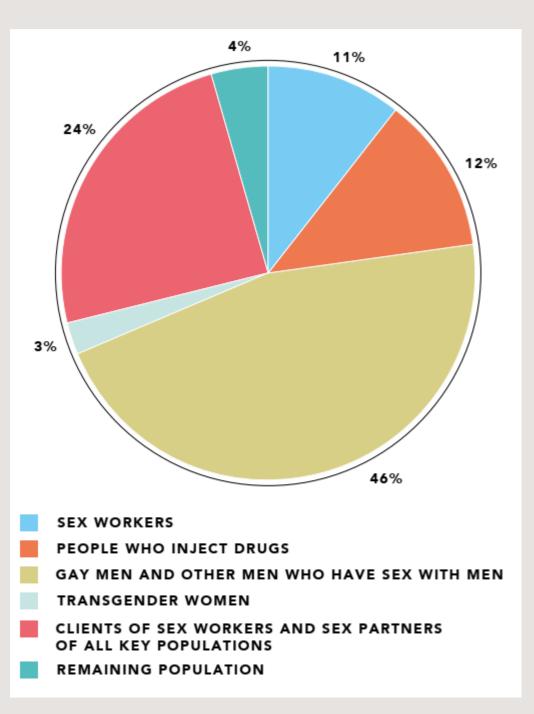
15.6

U/n

or about 6 million PLWH lived in the Asia-Pacific region ~140 k

AIDS-related deaths in the Asia-Pacific region

UNAIDS DATA BOOK 2022



Among 260k new HIV infections in the Asia-Pacific region in 2021,

46%

Gay men and other men who have sex with men

UNAIDS DATA BOOK 2022

UNAIDSFast-Track Targets500 000

New HIV infections or fewer

90-90-90

by 2030 **95-95-95** HIV treatment

200 000 New HIV infections or fewer

New HIV infections or fewe

ZERO

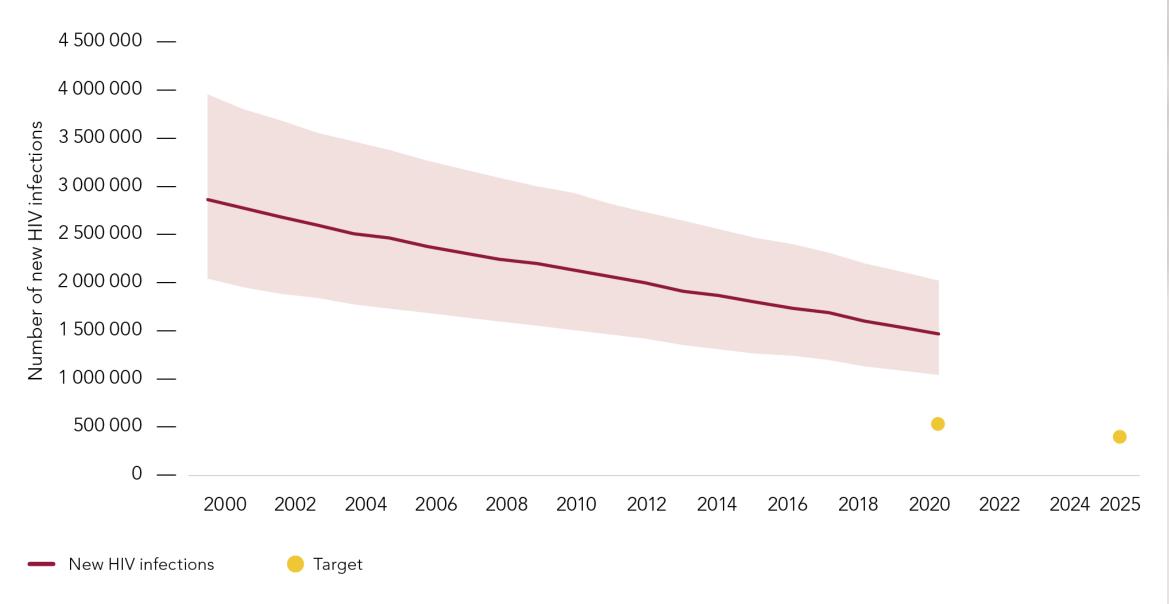
Discrimination

by 2020

ZERO

Discrimination

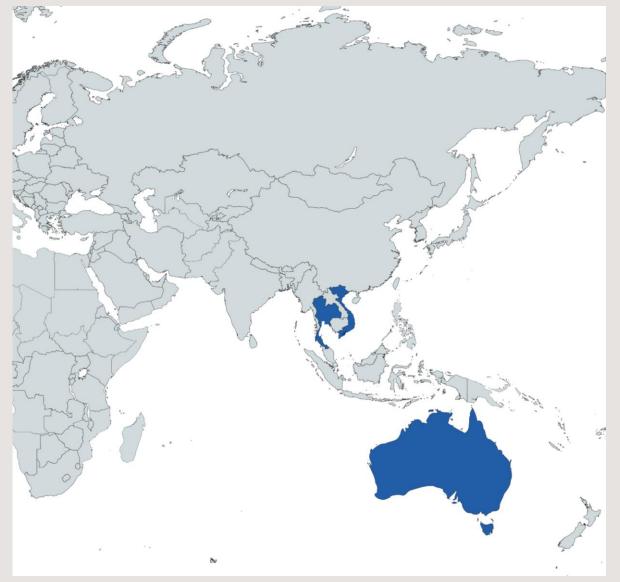
New HIV infections, global, 2000–2025, and 2020 and 2025 targets



Source: UNAIDS epidemiological estimates, 2021 (https://aidsinfo.unaids.org/).







PrEP is included in the national HIV respon

- Thailand
- Vietnam
- Australia

Global Advocacy for HIV Prevention (AVAC)

FINANCIAL CONSTRAINTS

STIGMA

ACCESS PrEP

STRUCTURAL BARRIERS

Prep Preference

Tiffany 2022, Budi 2022



Monthly Every 2 months Every 3 months Every 6 months



Physician Clinical officer Nurse Pharmacist Community health worker Patient / peer / family Client



HIV clinic / hospital Primary care clinic Drop-in centre Community Home

WHAT

ART initiation / refills Clinical monitoring Adherence support Laboratory tests Opportunistic infections treatment Psychosocial support

Tips about attributes

- Should be independent of one another
 - Gender of provider
 - Cost of consultation
- Objective, unambiguous, precise
 - "Cost" vs. "monthly cost of medication"
- One concept
 - Try to avoid compounded descriptions
 - "Injection that requires you to visit every 2 months"

Make sure everyone understands the terms in the same way

• "Nausea"

Mild to moderate nausea, diarrhea, and/or vomiting 3-4 times / week ¹	3 Days of nausea a month for first 3 months ²	Nausea ³	Risk of GI problems ⁴
 None Resolves after taking medicine for 2 weeks Continues as long as patient takes medicine 	NoneMildModerate	 None 30 min/d 90 min/d 	 0% (no risk of GI problems) 100 of 1000 people (10%) have GI problems 200 of 1000 people (20%) have GI problems 300 of 1000 people (30%) have GI problems

Small group discussion

Tasks

- Name your group
- Which population(s) do you want to focus on?
- How do you identify relevant attributes / levels?
 - What methods can you use?
- List all likely attributes that would influence someone using PrEP

Session 3. How to identify the right attributes / levels Not as easy as you think!

Objectives

- Discuss the various methods to choose attributes and levels
- Describe ways to prioritize attributes



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Stylus	Yes	No	No	I would not
Removable Keyboard	Yes	No	No	choose any o
Price	\$1,499	\$1,299	\$999	these.
	0	0	0	0

Attributes

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	0	0	0	0

Levels

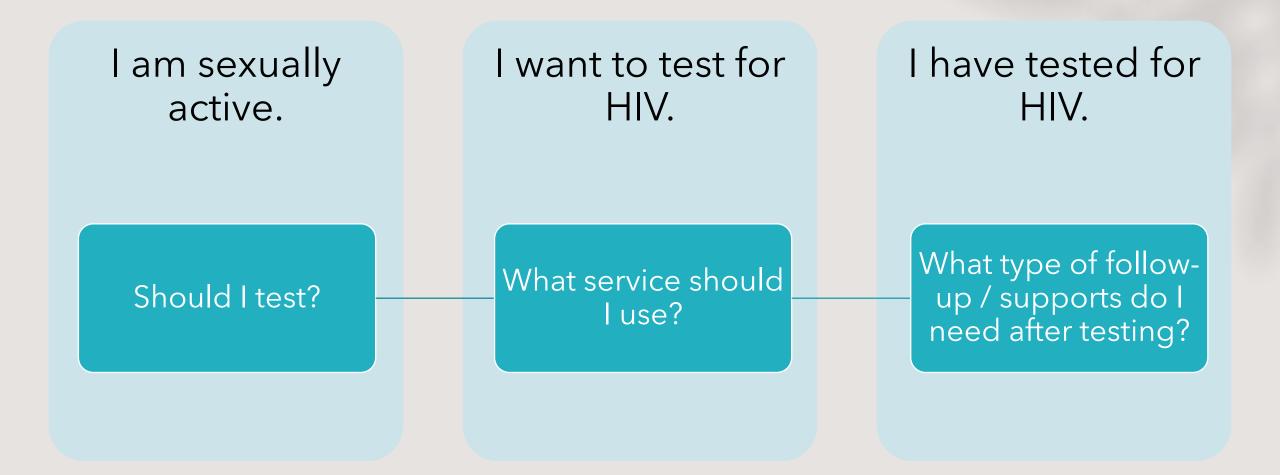
Attributes	Attribute Levels
PrEP modality	 Daily oral pill (1)
	 On-demand oral pill (2)
	 Long-acting injectable PrEP (3)
	PrEP Implant (4)
PrEP prescription	 Same-day prescription (1)
	• 2-visit prescription (2)
	 Telehealth prescription (3)
Medication pick-up	Clinics/hospitals (1)
	 Primary community health center (2)
	 Pharmacy (3)
	 MSM-focus CBOs (4)
	Home delivery (5)
Enhance support (for	 Physical or e-calendar (1)
medical adherence and	 Smartphone application (2)
follow-up visit reminder)	• Text reminder (3)
	 Anonymous peer support group (4)
Cost (out-of-pocket)	• Free (1)
	• 65% off (350 RMB) <mark>(2)</mark>
	• 30% off (700 RMB) (3)
	 Full price (1000 RMB) (4)

How to choose your attributes/levels?

- 1. Specify the choice scenario
- 2. What are drivers of choice?
- 3. Prioritise what to include in your choice set

1) Specify your choice scenario VERY CAREFULLY!!!

What decision point are you interested in?



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	0	0	0	0

Choice scenario – Any problems with these statements?

"Choose the service you prefer" "Imagine you want to see a doctor. Which service would you prefer to attend?" "Imagine you have a flu-like illness. Which service would you prefer to attend?"

"Ideal" number of attributes / levels?

How many things can you hold in your head at once?



Imagine you have a minor symptom, such as getting a cold, coughing and so on...

There are two facilities, each with different characteristics. If you have a choice, which option would you

choose from below?

Characteristics	Facility A	Facility B
Type of service	General service	Specialized service
Treatment measures	Traditional Chinese Medicine	Modern Medicine
Cost (CNY)	300CNY	100CNY
Travel time	>30mins	≤30mins
Care provider	Senior medical practitioner	Junior medical practitioner
Which facility would you choose? (Please		
tick only one box at the right)		

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Which facility would you choose?	(Please	
tick only one box at the right)		

3.1 You have been diagnosed with diabetes type 2. Your doctor asks you to decide between therapy A and therapy B. Which therapy would you choose?

Attribute	Therapy A	Therapy B
Additional <u>healthy</u> life years	+3 Years	+1 Years
Risk of urinary tract infection	Low 5 out of 100 (5%)	High 10 out of 100 (10%)
Risk of gastrointestinal problems	Low 4 out of 100 (4%)	High 12 out of 100 (12%)
Adjustment of long term blood glucose level (HbA1c)	Good (7,0 - 7,5%)	Very good (6,5 - 7,0%)
Risk of genital infection	Low 10 out of 100 (10%)	High 20 out of 100 (20%)
Possible weight change	+4 kg	+/- 0 kg
Possible hypoglycemia	Severe (severe symptoms)	Mild (without symptoms)
	0	0
	© Axel C. Mühlbacher	

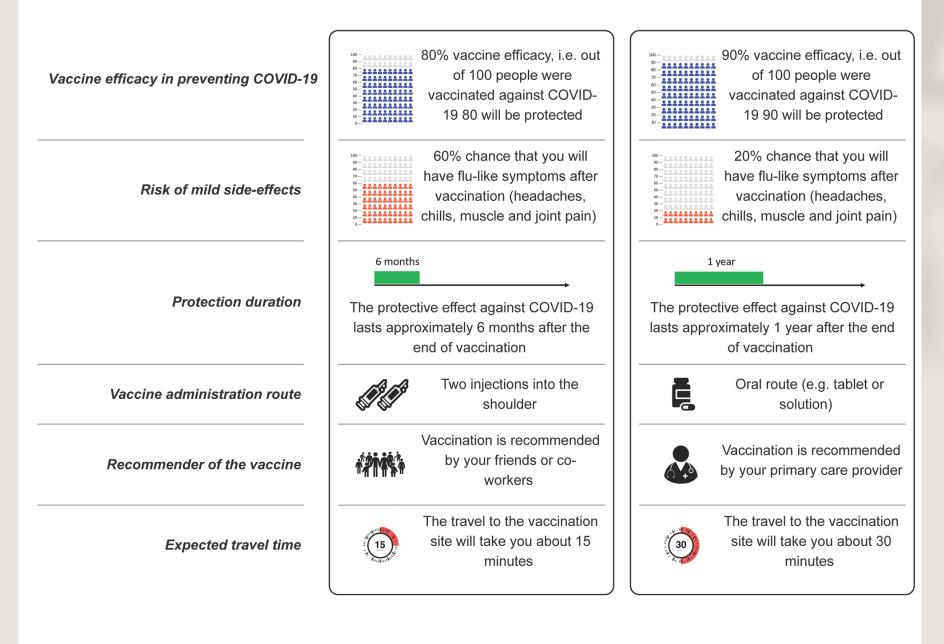
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Possible hypoglycemia	Severe (severe symptoms)	Mid (without symptoms)		
	0	0		
© Axel C. Möhlbacher				

3.1 You have been diagnosed with diabetes type 2. Your doctor asks you to decide between therapy A and therapy B. Which therapy would you choose?

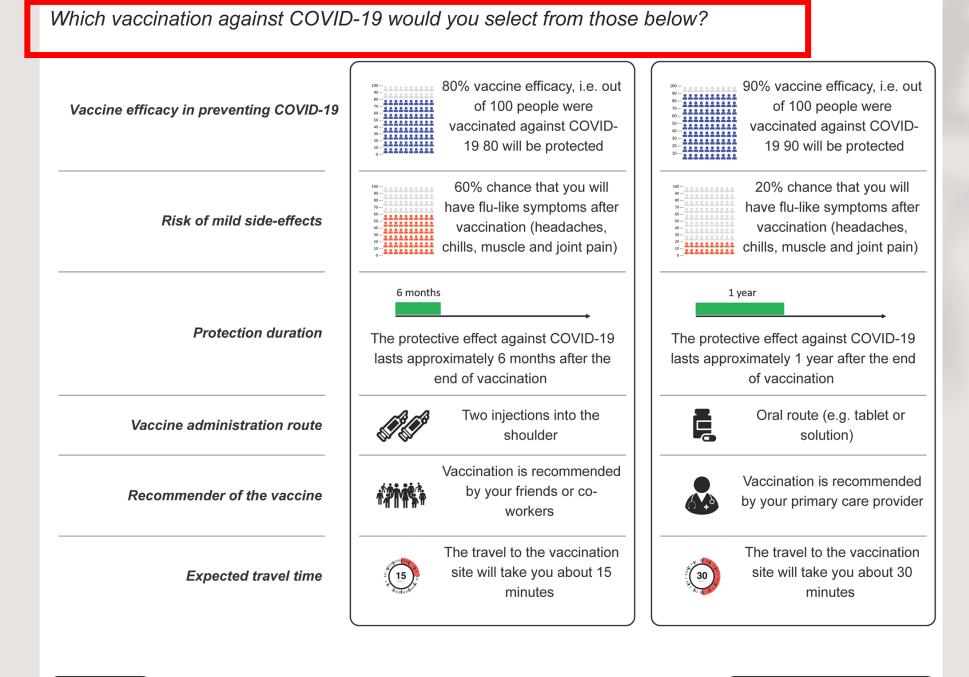
Attribute		Therapy A	Therapy B	
Additional <u>healthy</u> life years		+3 Years	+1 Years	
Risk of urinary tract infection	5	Low	High 10 out of 100 (10%)	
Risk of gastrointestinal problems	4	Low	High 12 out of 100 (12%)	
Adjustment of long term blood glucose level (HbA1c)		Good (7,0 - 7,5%)	Very good (6,5 - 7,0%)	
Risk of genital infection		Low	High 20 out of 100 (20%)	
Possible weight change		+4 kg	+/- 0 kg	
Possible hypoglycemia		Severe (severe mptoms)	Mild (without symptoms)	
		0	0	
© Axel C. Mohlbacher				

Which vaccination against COVID-19 would you select from those below?



Go back

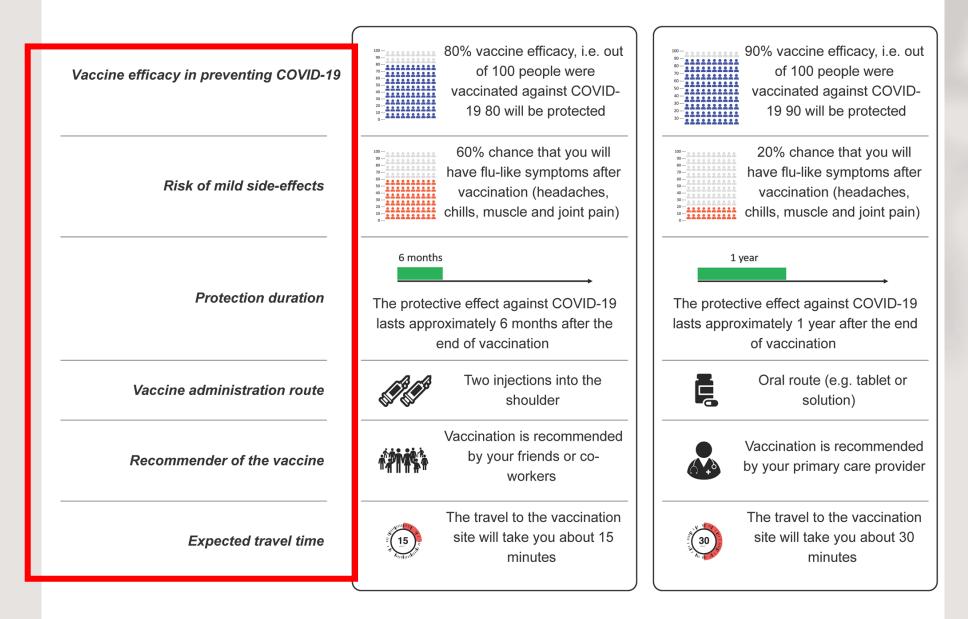
I choose neither



Go back

I choose neither

Which vaccination against COVID-19 would you select from those below?



I choose neither

2) What are the drivers of choice?

Methods

- Can use more than one
- Qualitative methods
 - Focus group discussions
 - One-on-one interviews
 - Consultation with stakeholders
- Literature review
- "Experts" choosing attributes and levels
- Pre-determined by specific scenario e.g. pipeline of new HIV drugs
- Pilot studies
 - Ranking of attributes

Health Economics

Research Article 🔂 Full Access

Using qualitative methods for attribute development for discrete choice experiments: issues and recommendations

Joanna Coast 🔀, Hareth Al-Janabi, Eileen J. Sutton, Susan A. Horrocks, A. Jane Vosper, Dawn R. Swancutt, Terry N. Flynn

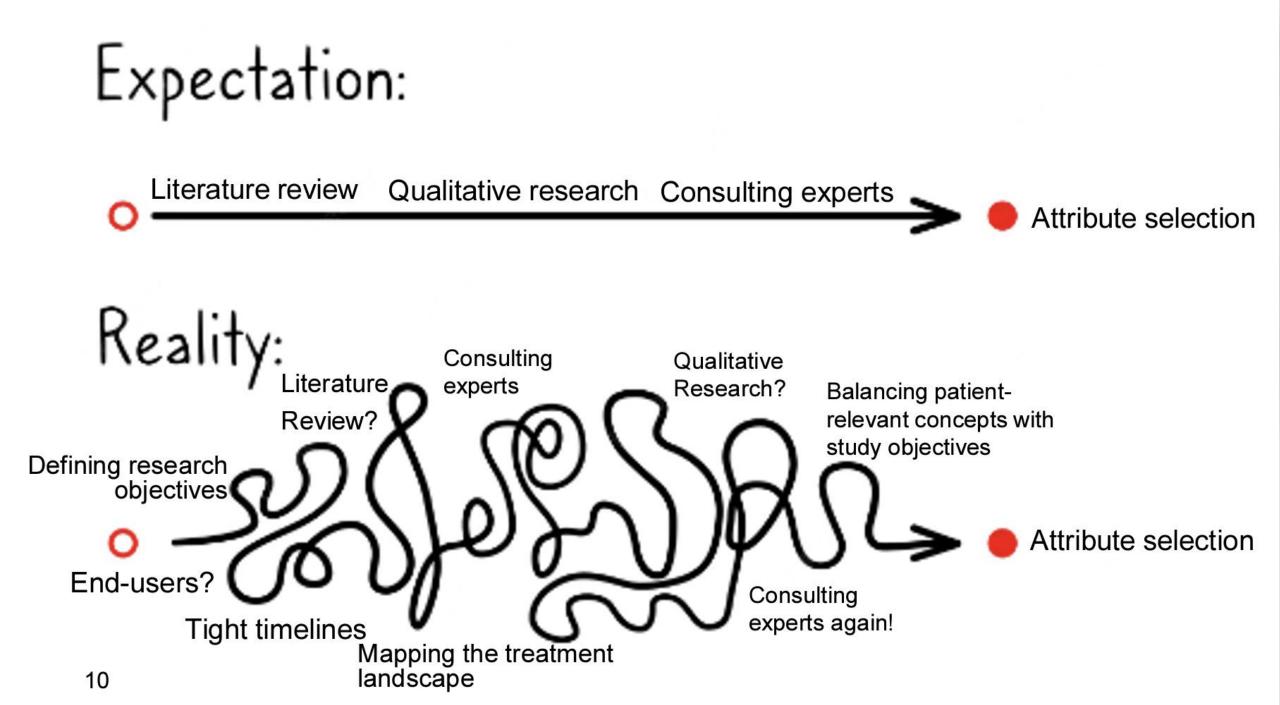
First published: 06 May 2011 | https://doi.org/10.1002/hec.1739 | Citations: 321

Key points

- 2 stage process
 - Conceptual development
 - Refinement of language to convey the intended meaning
 - Pilot!

Key points

- Choose attributes
 - Not too close to latent construct "utility"
 - No attribute labelled "utility" or that expresses overall happiness with the alternative
 - Important to decision-maker ('drivers of choice')
 - Capable of being traded
 - Not too dominant that you have no one trading
 - Able to be 'manipulated'



Science

- Literature review
- Qualitative research
- Potentially ranking/rating methods to refine

Attribute selection

Art

- Research question
- Business question
- Consider end-use and end-user/key stakeholders
- Stage of product development
- Level of rigour required
- Timelines and budgets

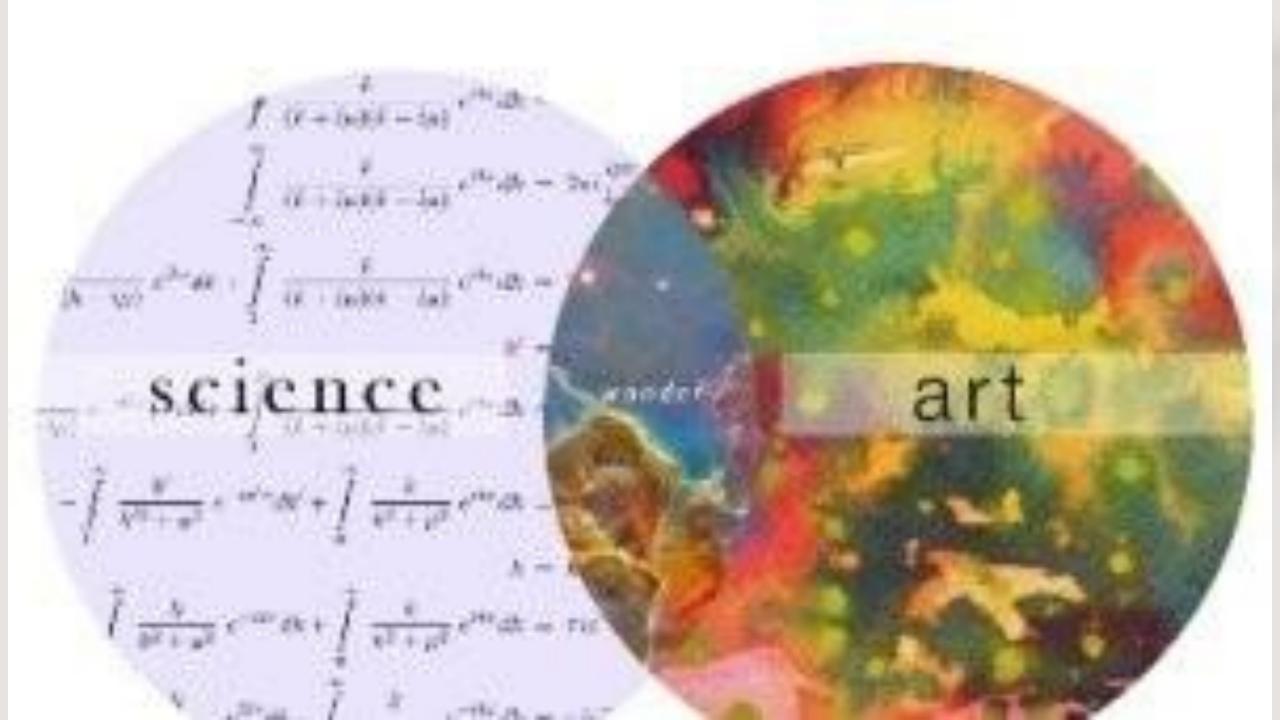
~20 potential attributes

3) How do you prioritize which attributes to present?

~4-6 final attributes

Possible activities

- Ranking exercise
- Utilise qualitative studies
 - Show choice sets (at the end of interview)
 - Ask if anything important missing
- Discussions with 'experts'
- Model data from first 10% of total sample





Questions?

Session 3 – small group discussion

Session 3 – discussion

- What other questions would you include in your survey?
- How will you collect survey responses
 - discuss pros/cons of online vs. paper vs. interviewer-assisted?
- How will you recruit?

Session 4. Experimental design and presentation of choice sets

Objectives

- Understand different types of experimental designs
 - Demonstrate NGENE and Qualtrics
- Understand different ways choice sets can be presented
- Describe what survey platforms are compatible for DCE surveys

1) Experimental designs

Experimental design

- Determines what combinations of levels are shown in a choice set?
- Why does it matter?
 - Maximise amount of information from each choice made
 - => reduce sample size

- Full factorial design
 - Every possible combination of attribute levels
 - 2 alternatives, 3 attributes with 4 levels
 - Total combination = ?

- Full factorial design
 - Every possible combination of attribute levels
 - 2 alternatives, 3 attributes with 4 levels
 - Total combination = (4 X 4 X 4) x (4 X 4 X 4) = 4,096!

- Full factorial design
 - Every possible combination of attribute levels
 - 2 alternatives, 3 attributes with 4 levels
 - Total combination = (4 X 4 X 4) x (4 X 4 X 4) = 4,096!



Slides from

 https://www.sydney.edu.au/business/our-research/institute-oftransport-and-logistics-studies/courses/discrete-choiceanalysis.html



Discrete Choice Analysis

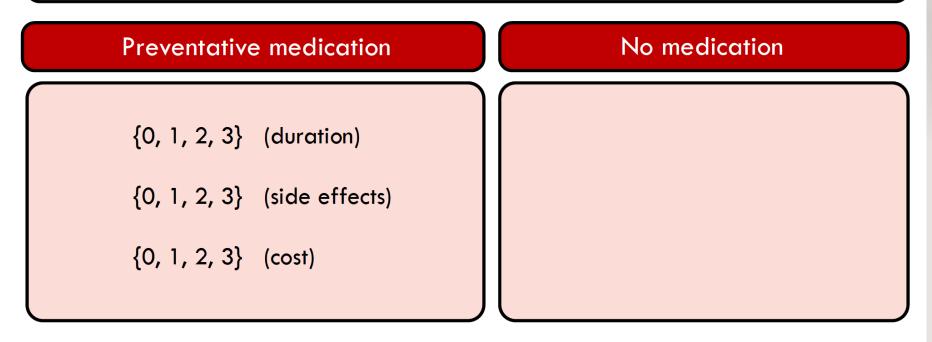
Example

Consider a one week safari trip to **Kruger Park**. Which malaria prevention option would you prefer, knowing that medication **reduces risk with 90 per cent**?

Preventative medication	No medication
{2, 4, 6, 8} weeks duration {no, mild, moderate, severe} side effects {\$10, \$40, \$70, \$100} cost	

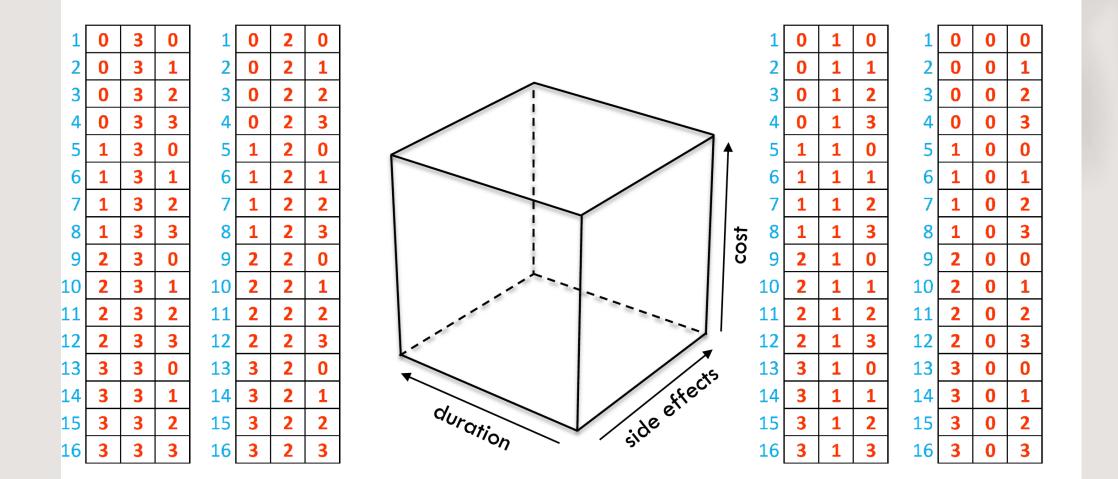
Example (design coding)

Consider a one week safari trip to **Kruger Park**. Which malaria prevention option would you prefer, knowing that medication **reduces risk with 90 per cent**?





64 possible choice tasks (full factorial)



64 possible choice tasks (full factorial)

0	3	0	1	(
0		1	2	(
0	3	2	3	(
0	3	З	4	(
1	3	0	5	1
1	3	1	6	
1	3	2	7	1
1	3	3	8	1
2	3	0	9	
2	3	1	10	1
2	3	2	11	1
2	3	3	12	1
3	3	0	13	
3	3	1	14	
3	3	2	15	
3	3	3	16	
	0 0 1 1 1 1 2 2 2 2 2 3 3 3 3 3 3 3	 0 3 0 3 1 3 1 3 1 3 3 2 3 4 4<	031032033130131132133231232232233330331332	0 3 1 2 0 3 2 3 0 3 3 4 1 3 0 5 1 3 1 6 1 3 2 7 1 3 2 7 1 3 2 7 1 3 3 8 2 3 1 10 2 3 1 10 2 3 2 11 2 3 3 12 3 3 0 13 3 3 1 14 3 3 2 15

0_0_0

0	1	0	
		1	
0	1	2	
0	1	3	
1	1	0	
1	1	1	
1	1	2	
1	1	3	
2	1	0	
2	1	1	
2	1	2	
2	1	3	
3	1	0	
3	1	1	
3	1	2	
3	1	3	
	0 0 1 1 1 1 2 2 2 2 2 3 3 3 3 3 3	0 1 0 1	0 1 1 0 1 2

1	0	0	0
2	0	0	0 1 2
3	0 0	0 0	2
4	0	0 0 0	3
5	0 1	0	0
6	1	0	1
7	1	0	2
1 2 3 4 5 6 7 8 9	1	0 0 0	3
	2	0	0
10	1 1 2 2 2 2	0 0 0	0 1 2 3 0 1 2 3
11	2	0	2
11 12 13	2	0	3
13	3	0	0
14	3	0	0 1 2
14 15	3 3	0 0	2
16	3	0	3

64 possible choice tasks (full factorial)

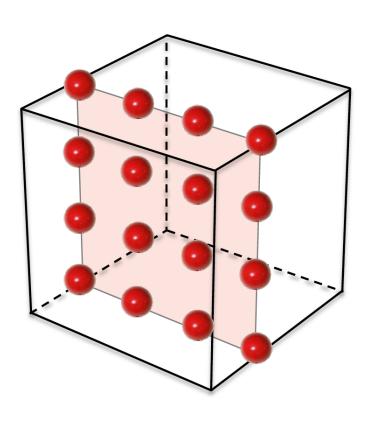
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3	0	3 3 3 3 3	0 1 2 3 0	1 2 3 4 5 6 7 8	0
4	0	3	3	4	0
5	1	3	0	5	1
6	1	3	1	6	1
7	1	3	2	7	1
8	1 1 1	3 3 3	1 2 3	8	1 1 1
9	2 2 2 2 3	3 3 3 3 3	0 1 2 3 0	9	2 2 2 2 3
10	2	3	1	10	2
11	2	3	2	11	2
12	2	3	3	12	2
1 3	3	3	0	12 13 14	3
14	3	3	1	14	3
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16	3	3 3 3	1 2 3	15 16	3 3
16	3	3	3	16	3

1	0	1	0	
2	0	1		
3	0	1	1 2 3	
4	0 0 0	1	3	
5	1	1	0	
6	1	1 1	1	
1 2 3 4 5 6 7 8	1	1 1	1 2 3	
8	1	1	3	
9	2	1	0	
10	1 1 1 2 2 2 2 2	1	1	
11	2	1	2	
12	2	1	3	
12 13	3	1	0	
14	3 3 3 3	1 1 1 1 1 1 1 1	1 2 3 0 1 2 3	
15	3	1	2	
16	3	1	3	

1	0	0	0
2	0	0 0	1
3	0	0	2
4	0	0	
1 2 3 4 5 6 7 8	0 0 1	0	3 0 1 2
6	1	0	1
7	1	0	2
8	1	0	3 0
9	2	0	0
10	1 1 2 2 2 2 2	0	1
11	2	0	2
12 13	2	0	3
13	3	0	0
14	3 3 3	0	1 2 3 0 1 2
14 15	3	0 0 0 0 0 0 0 0 0 0 0 0 0	
16	3	0	3

64 possible choice tasks (full factorial)

1	0	3	0	1	0	2
2	0	3	1	2	0	2
2 3	0	3	2	3	0	2
4	0	3	3	4	0	2
5 6	1	3	0	5 6	1	2
6	1	3	1		1	2
7	1	3	2	7	1	2
8	1	3	3	8	1	2
9	2	3	0	9	2	2
10	2	3	1	10	2	2
	2	3	2	11	2	2
11 12 13	2	3	3	12	2	2 2
13	3	3	0	13	3	2
14	3	3	1	11 12 13 14 15	3	2
15	3	3	2	15	3	2
16	3	3	3	16	3	2

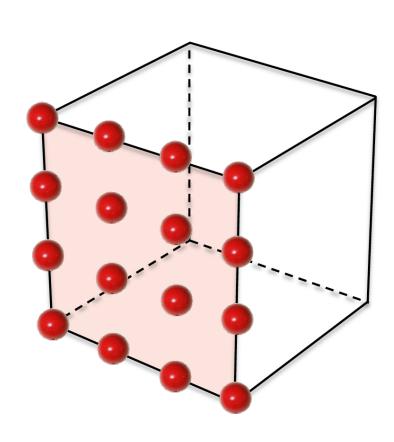


1	0	1	0	
2	0	1	1	
3	0	1	2	
1 2 3 4 5	0	1	0 1 2 3	
5	1	1	0	
6	1	1	1	
6 7 8 9	1	1	2	
8	1	1	3	
9	2	1	0	
10	0 0 1 1 1 2 2 2 2 2	1	0 1 2 3 0 1 2 3	
11	2	1	2	
11 12	2	1	3	
13		1	0	
14 15	3 3 3 3	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	0 1 2	
15	3	1	2	
16	3	1	3	

1	0	0	0
2	0	0	1
3	0	0	2
1 2 3 4 5 6 7 8	0	0	3
5	1	0	0
6	1	0	1
7	1	0	2
8	1	0	3
9	2	0	0
10	2	0	1
11 12	2	0	2
12	2	0	3
13	3	0	0
14	3	0	1
13 14 15 16	0 0 1 1 1 1 2 2 2 2 3 3 3 3 3 3	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 1 2 3 0 1 2 3 0 1 2 3 0 1 2 3 0 1 2 3 3
16	3	0	3

64 possible choice tasks (full factorial)

1	0	3	0	1
2	0	3	1	2
3	0 0	3	1 2 3	3
4	0	3		4
5	1 1 1	3	0 1 2	5
6	1	3	1	6
7	1	3	2	7
1 2 3 4 5 6 7 8 9	1 2 2 2 2 3	3	3	1 2 3 4 5 6 7 8 9
9	2	3	0	9
10	2	3	1 2 3	10
11	2	3	2	11
12	2	3		12
13	3	3	0	13
14	3	3	1 2	14
12 13 14 15	3 3	3		15
16	3	3	3	16



				_
1	0	1	0	
2	0	1	1	
3	0	1	2	
1 2 3 4 5 6 7 8 9 10	0 0 1 1 1 1 2 2 2 2 2	1	3	
5	1	1	0	
6	1	1	1	
7	1	1	2	
8	1	1	3	
9	2	1	0	
10	2	1	1	
11	2	1	2	
12	2	1	3	
13	3	1	0	
11 12 13 14 15	3 3	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	0 1 2 3 0 1 2 3 0 1 2 3 0 1 2 3 3 3	
15	3 3	1	2	
16	3	1	3	

1	0	
2	0	
2 3 4 5 6 7 8	0 0 0 0	
4	0	
5	1 1 1 2 2 2 3 3 3	
6	1	
7	1	
8	1	
9	2	
9 10	2	
11	2	
12	2	
13	3	
14	3	
15	3	
16	3	

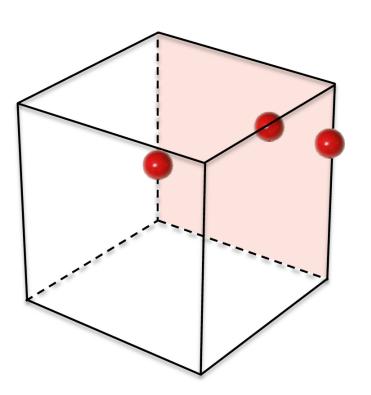
64 possible choice tasks (full factorial)

- Do we really need all 64 choice tasks? Or can we make a sub-selection?

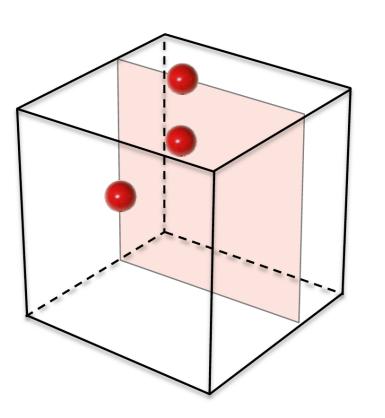
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2	0	3	1	2	0	2	1	2	0	1	1	2	0	0	1
3	0	3	2	3	0	2	2		0	1	2	3	0	0	2
4	0	3	3	4	0	2	3		0	1	3	4	0	0	3
5	1	3	0	5	1	2	0		1	1	0	5	1	0	0
6	1	3	1	6	1	2	1		1	1	1	6	1	0	1
7	1	3	2	7	1	2	2		1	1	2	7	1	0	2
8	1	3	3	8	1	2	3		1	1	3	8	1	0	3
9	2	3	0	9	2	2	0		2	1	0	9	2	0	0
10	2	3	1	10	2	2	1		2	1	1	10	2	0	1
11	2	3	2	11	2	2	2		2	1	2	11	2	0	2
12	2	3	3	12	2	2	3		2	1	3	12	2	0	3
13	3	3	0	13	3	2	0	13	3	1	0	13	3	0	0
14	3	3	1	14	3	2	1		3	1	1	14	3	0	1
15	3	3	2	15	3	2	2	15	3	1	2	15	3	0	2
16	3	3	3	16	3	2	3	16	3	1	3	16	3	0	3

- Fractional factorial design
 - Subset of total possible combinations
 - How do you choose which subset?

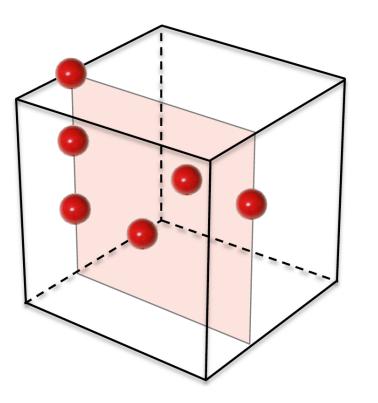
- Fractional factorial design
 - Subset of total possible combinations
 - How do you choose which subset?
 - Random
 - Give 1st subset to 1st respondent, 2nd subset to 2nd respondent, and so on...
 - Risk of seeing only certain levels of attributes



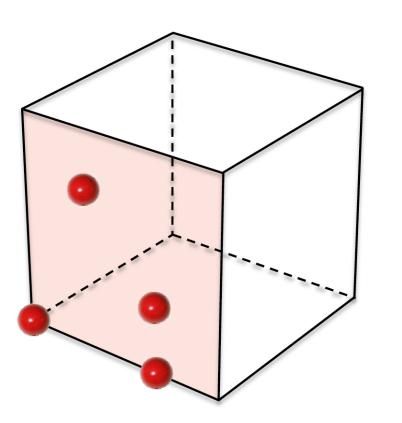
1	0	3	1
2	2	2	-
2	2	3	2
3	3	3	2
4	0	2	1
5	1	2	2
6	1	2	3
1 2 3 4 5 6 7 8 9 10	0	1	1
8	0	1	2
9	0	1	3
10	1	1	1
11	2	1	2
12	3	1	2
11 12 13 14 15	0 2 3 0 1 1 0 0 1 2 3 0 1 2 2 2	3 3 2 2 1 1 1 1 1 1 1 0 0	1 2 1 2 3 1 2 3 1 2 2 0 2 0 2 0
14	1	0	2
15	2	0	0
16	2	0	1



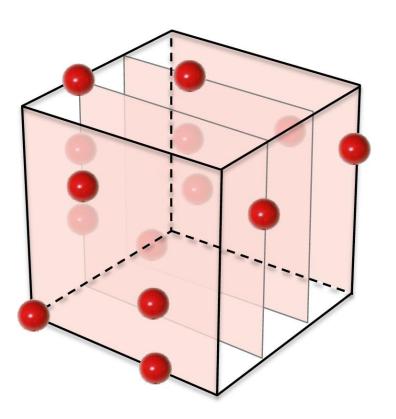
1	0	3	1
2	2	3 3	2
3	3	3	2
4	0	2	1
5	1	2	2
6	1	3 2 2 2	1 2 1 2 3
1 2 3 4 5 6 7 8 9	0	1	1
8	0	1	2
9	0	1	3
10	1	1 1 1 1	1 2 3 1 2 2 0 2 0 2 0
11	2	1	2
11 12	3	1	2
13 14	0	0	0
14	1	0	2
15	0 2 3 0 1 2 0 1 2 3 0 1 2 2 2	1 1 0 0	0
16	2	0	1



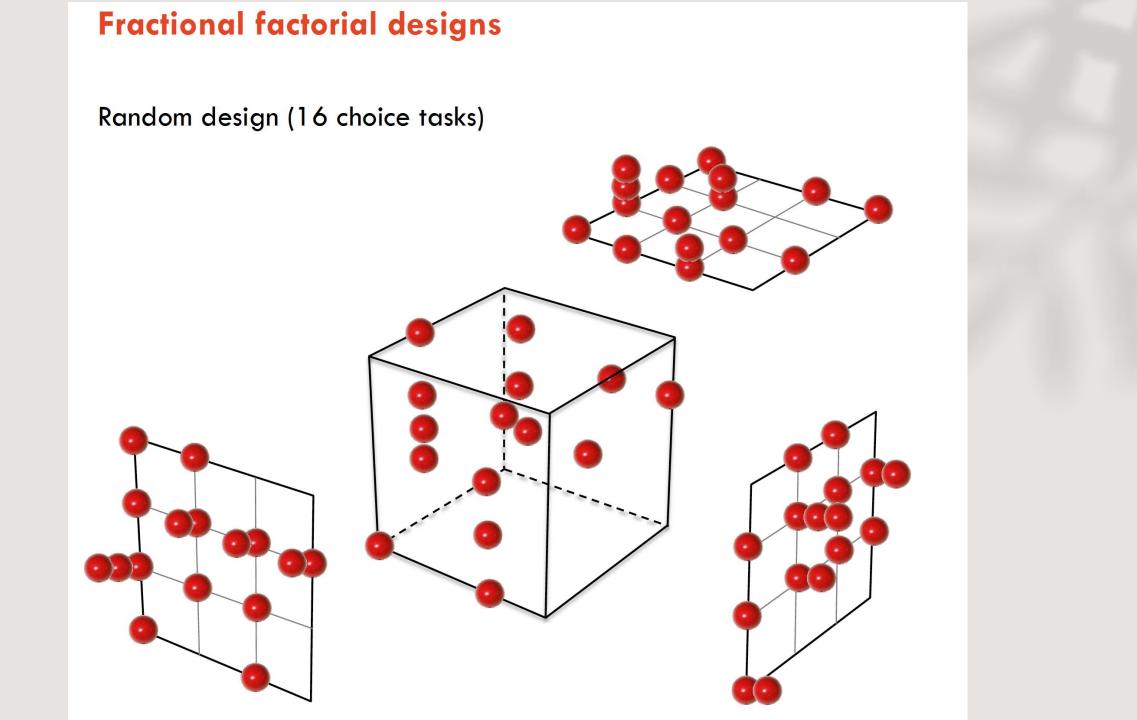
0	3	1
2	3	2
3	3	2
0	2	1
1	2	2
1	2	1 2 1 2 3
0		1
0	1	2
0	1	3
1	1	1
2	1	2
3	1	2
0	0	1 2 3 1 2 2 0 2
1	0	2
2	0	0
2	0	1
	0 2 3 0 1 1 0 0 1 2 3 0 1 2 2	0 1 0 1



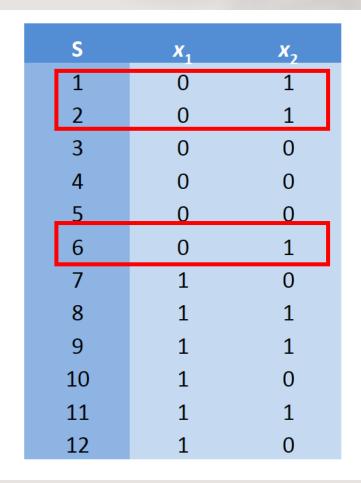
1	0	3	1
2	2	3	2
3	3	3	2
1 2 3 4 5 6 7 8 9	0	2	1
5	1	2	2
6	1	2	3
7	0	1	1
8	0	1	2
9	0	3 3 2 2 2 1 1 1 1 1 1 1 1 0 0	1 2 1 2 3 1 2 3 1 2 2 2 0 2 2
10	1	1	1
11	2	1	2
12	3	1	2
13	0	0	0
14	1	0	2
14 15	0 2 3 0 1 1 0 0 1 2 3 0 1 2 2 2	0	0
16	2	0	1



1	0	3	1
2	2	3	2
3	3	3	2
4	0	2	1
5	1	2	2
6	1	2	3
7	0	1	1
8	0	1	2
1 2 4 5 6 7 8 9 10	0	1	3
10	1	1	1
11 12	2	1	2
12	3	1	2
13	0	0	0
14	1	0	2
13 14 15	0 2 3 0 1 1 0 0 0 1 2 3 0 1 2 2	3 3 2 2 1 1 1 1 1 1 1 0 0 0 0	1 2 1 2 3 1 2 3 1 2 3 1 2 0 2 0 2 0 0
16	2	0	1



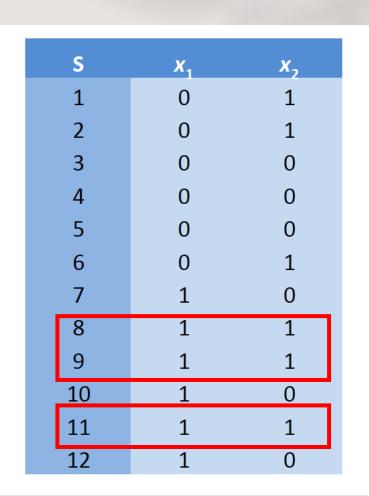
- Fractional factorial design
 - Orthogonal designs
 - Uncorrelated attribute levels
 - Every pair of levels occurs equally (rows)



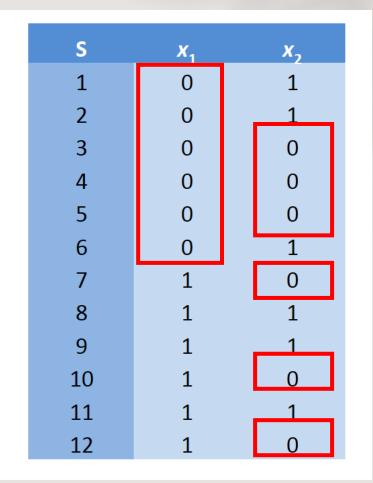
- Fractional factorial design
 - Orthogonal designs
 - Uncorrelated attribute levels
 - Every pair of levels occurs equally (rows)

S	x ₁	x ₂
1	0	1
2	0	1
3	0	0
4 5	0	0
5	0	0
6	0	1
7	1	0
8	1	1
9	1	1
10	1	0
11	1	1
12	1	0

- Fractional factorial design
 - Orthogonal designs
 - Uncorrelated attribute levels
 - Every pair of levels occurs equally (rows)



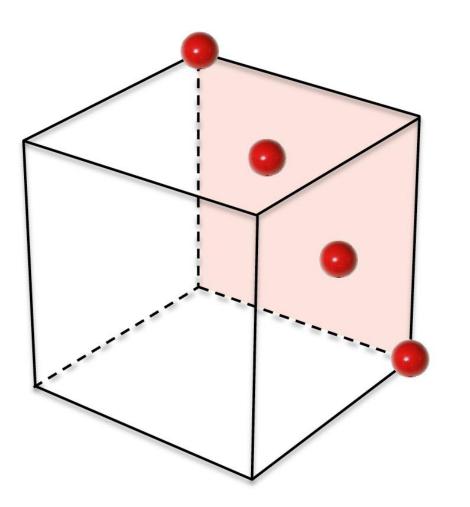
- Fractional factorial design
 - Orthogonal designs
 - Uncorrelated attribute levels
 - Every pair of levels occurs equally (rows)
 - Each level appears equal number of times for each attribute (columns)



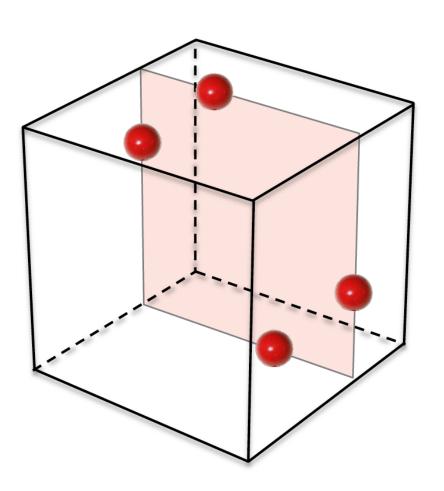
- Fractional factorial design
 - Orthogonal designs
 - Almost impossible to manually generate orthogonal arrays
 - Design libraries
 - http://support.sas.com/techsup/technote/ts723_Designs.txt
 - <u>http://neilsloane.com/oadir/</u>
 - Software (e.g. NGENE, SAS, SPSS)

Types of experimental designs

- Fractional factorial design
 - Orthogonal designs
 - Design may be orthogonal but data used in estimation is often not orthogonal
 - Respondents not completing all choice sets assigned
 - Blocks of choice sets not equally distributed
 - Undesirable combinations
 - Dominant
 - Impossible

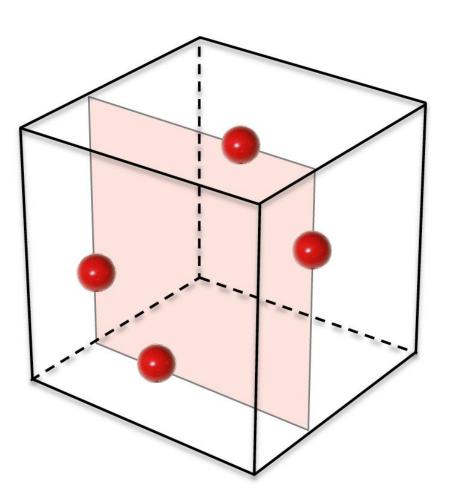


1	0	3	3
2	1	3	2
3	1 2	3	1
1 2 3 4 5 6 7 8 9	3 0 1 2	3 2	2 1 0 2
5	0	2	
6	1	2	3
7	2	2	0
8	3	2	1
	0	1	1
10	3 0 1 2	1	3 0 1 1 0
11	2	1	3
12	3 0	1	2
13	0	2 2 1 1 1 1 0	3 2 0
14	1	0	1
15	2 3	0	2 3
16	3	0	3

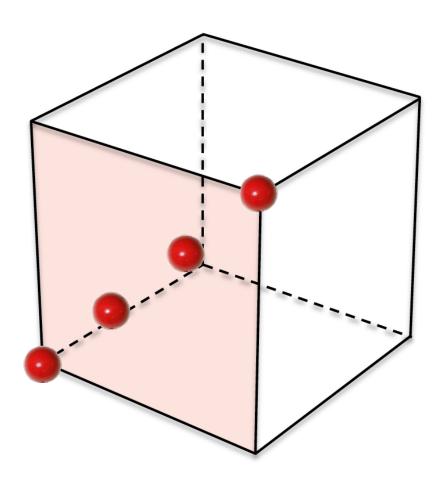


-			
1	0	3	3
2	1	3	2
3	2	3	1
4	3	3	0
1 2 3 4 5 6 7 8 9	1 2 3 0 1 3 0 1 2 3 3	3 2	3 2 1 0 2
6	1	2	3
7	2	2 2 1	
8	3	2	1
	0	1	0 1 1
10	1		0 3 2
11	2	1 1 1	3
12	3	1	2
13	0	0 0	0 1
14	1	0	1
15	0 1 2	0	2
16	3	0	3
I			

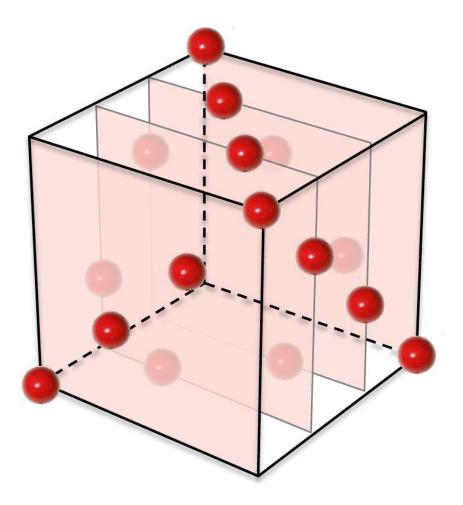
Orthogonal design (16 choice tasks)



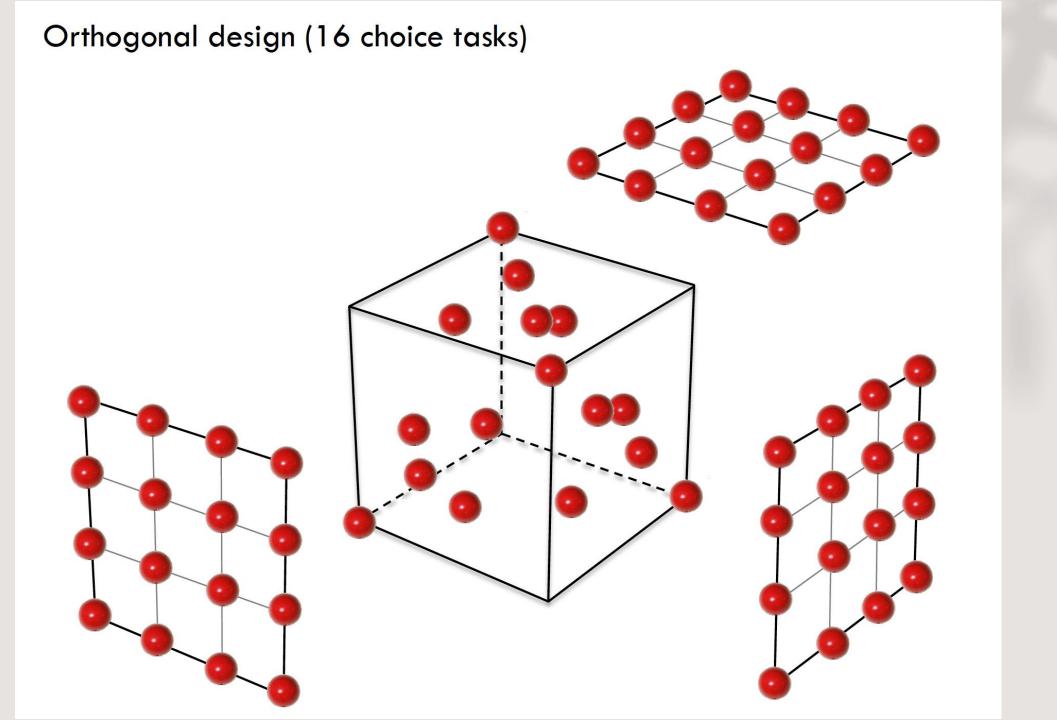
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2	1	3	3 2
3	1 2	3	1
4	3	3	0 2
1 3 4 5 6 7 8 9	3 0 1 2	2	
6	1	2	3 0
7	2	2	0
8	3	2	1
9	0	1	1
10	1	1	0
11	2	1	3
12	3	1	2
13	0	0	0
14	3 0 1 2 3 0 1	3 2 2 2 1 1 1 1 1 0 0	0 3 2 0 1
15	2	0	2
16	3	0	3
	2	0	3



0	3	3
1	3	2
2	3	1
3	3	0
0	2	2
1	2	3
2	2	0
3	2	1
0	1	1
	1	0
2	1	3
3	1	2
0	0	0
1	0	3 2 1 0 2 3 0 1 1 0 3 2 0 1 2 0 1 2
2	0	2
3	0	3
	0 1 2 3 () 1 2 3 () 1 2 3 () 1 2 3 () 1 2 3 () 1 2 3 () 1 2 3 () 1 2 3 () 1 2 3 () 1 3 () 1 () 1 () 1 () 1 () 1 () 1	1323330212223211112131001020



1	0	3	3
2		3	2
3	2	3	1
4	3	3	0
5	0	2	2
6	1	2	3 2 1 0 2 3
1 2 3 4 5 6 7 8 9 10	2	2	0
8	3	2	1
9	0	1	1
10	1	1	0
11	2	1	3
12	3	1	2
13	0	0	0
11 12 13 14	1 2 3 0 1 2 3 0 1 2 3 0 1 2 3 3	3 3 2 2 2 2 1 1 1 1 0 0 0 0 0 0	0 1 1 0 3 2 0 1 2 3
15	2	0	2
16	3	0	3
16	3	0	3



Types of experimental designs

- Fractional factorial design
 - Efficient designs
 - Captures more efficient per choice set answered => Less survey respondents needed
 - Smaller standard errors => More reliable estimates

asymptotic variance-
covariance (AVC) matrix
$$\Omega = \begin{pmatrix} se(\beta_1)^2 & \cdots \\ \vdots & \ddots \\ & se(\beta_K)^2 \end{pmatrix}, \text{ where } se(\beta_k) \text{ is the standard} \text{ error of parameter } \beta_k$$

Types of experimental designs

- Fractional factorial design
 - Efficient designs
 - Captures more efficient per choice set answered => Less survey respondents needed
 - Smaller standard errors => More reliable estimates
 - Bayesian D-efficient designs (gold standard)

How to generate designs?

- Generator developed designs
 - <u>https://onlinelibrary.wiley.com/doi/book/10.1002/9780470148563</u>
 - https://www.youtube.com/watch?v=xv3n2ZfRmYo

Experimental designs

- Applying "constraints"
 - Need to avoid unusual / impossible combinations

What do you think about:

- Type of PreP: "Injectable PrEP"
- Access: "Mailed to Home"

Experimental designs

- Applying "constraints"
 - Need to avoid unusual / impossible combinations

What do you think about:

- Type of PreP: "Injectable PrEP"
- Access: "Mailed to Home"
- (Depends on aim of study)
 - Have to carefully explain "unusual"/"new" combinations to participants

- iHEA Webinar June 1, 2020: An Introduction to the Construction of Discrete Choice Experiments
- <u>https://www.youtube.com/watch?v=xv3n2ZfRmYo</u>



2) Presentation of choice sets



Why do we care about *how*choice sets are presented?

Why care?

- Could influence choices
- Help reduce cognitive burden
- Reduce misunderstandings of what you want from them



What could influence someone's choices in a DCE survey? Presentation of choice sets *may* influence respondent choice

- Text
- Cognitive biases
- Mode of delivery
- Visual cues

Text

- "Cheap talk"
- Terminology
 - Terminology and concepts may be unfamiliar to respondents
 - Key assumption is that respondents understand all attributes and levels in the same way to make a rational choice
- How the choice scenario is asked

Cognitive biases

- Left-right bias
 - Testable / amenable
- Order bias
 - Amenable
- Anchoring effect
 - People swayed by statistics / numerical value presented
- Framing
 - Gain or loss
 - Lives saved v.s. lives lost
 - Probability of surviving vs. probability of dying

Mode of delivery

- Paper
 - static
- Interviewer assisted
- Online
 - Beware of small screens for Mobile phones



Visual cues

- Display format of choice set
 - The more realistic you can make it, the better
- Pictures, statistical values, graphs, videos
- Structure of choice sets
 - Number of alternatives, attributes, levels

Presentation of choice sets

- Table
- Pictures
- Videos / animation
 - Unexplored



What do you like or dislike about these examples?

Choosing a person to contact

In this survey, we are going to show you a number of hypothetical profiles of people from a dating website that you could potentially contact for a date.

We will show you the profiles of three potential contacts at a time.

Each time we show you the different profiles, we want you to choose the profile of the person that you would most likely contact in real life.

You will be shown nine scenarios which ask you to choose among the three potential contacts on offer.

Before we start, we will look at an example.

Next

An example

You will first be asked to choose between contacting Person A , B and C, or none.

We will then ask which you would prefer to contact if you had to choose among the three.

In making the choice, we want you to consider the following scenario:

If you were looking through a dating website and had a choice among the three people shown based on the descriptions listed, which person would you choose to contact?

	Ů Ř	Ů Ř	Ů Ř	
	Person A	Person B	Person C	None
Drinking Habit	Moderate drinker	Non drinker	Casual drinker	
Smoking Habit	Smoker	Non smoker	Ex smoker	
Children	None currently	Single parent	Single parent	
Job	White Collar	Blue Collar	Blue Collar	
Looks	Average	Above average	Below average	
Cost to contact	\$10	\$20	\$15	
l would choose to contact	0	۲	0	0
lf I had to choose, I would choose	0	۲	0	

You will be shown nine scenarios similar to the above one. Each scenario will show the profiles of different potential contacts.

We want you to select which person you would contact in each scenario based only on the profiles shown in that scenario.

That is, we want you to think only about the three people shown in the scenario and not other people who might have been shown in scenarios that you have seen previously.

Please make sure that you understand the task before proceeding. Once you go to the next screen, you will not be able to go back.

Back

Prostate cancer

	Active Surveillance	Radiotherapy	Surgery
Risk of permanent urinary incontinence due to treatment	Nobody (0%)	5 out of 100 (5%)	20 out of 100 (20%)
Risk of permanent erectile dysfunction due to treatment	Nobody (0%)	25 out of 100 (25%)	45 out of 100 (45%)
Risk of other permanent side-effects due to treatment	No	 Yes, substantial risk of Bowel problems Frequent urge to urinate 	Yes, small risk of mortality within 6 weeks
Main aim is cure	No, the tumour remains in the body	Yes, but the disease may return	Yes, but the disease may return
Frequency of PSA testing with a risk of new prostate biopsies	Four times in a year and at least one biopsy per year	Once in a year	Once in a year
Which alternative do you prefer?			

Meat choice scenario 4 of 4

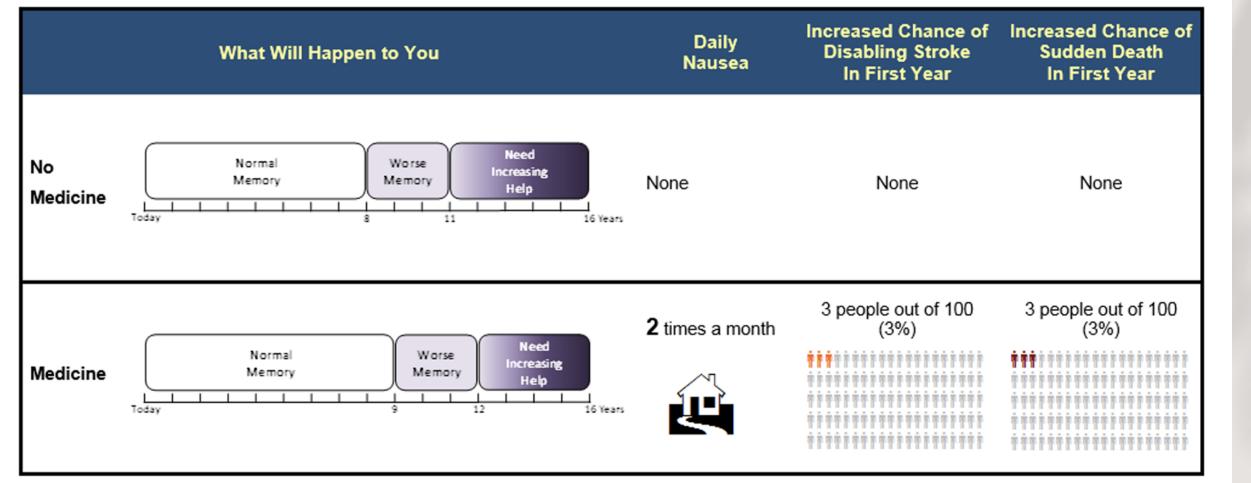
Looking at each of the meat products shown, we want you to tell us how much of each meat product you would purchase, assuming that you were shopping under similar circumstances to your last grocery shopping trip when you purchased meat (that is, if you were buying for a BBQ, then please pretend that you are buying for one now. If it were for a normal week, then pretend that it is a normal week now).

				Mince		Diced		
S	Y AN							
	Vacuum packed		Vacuum packed		Multi wrapped		Modified atmospher packed	
ountry of Origin	New Zealand	Country of Origin	China	Country of Origin	New Zealand	Country of Origin	United States	
00% Grass fed		100% Grass fed 100% Free of antibiotic 100% Free of hormones 100% Free of steroids 100% Free range		100% Grass fed 100% Free of hormon 100% Traceable 100% Free range	25	100% Free of antibiotik 100% Free of hormons 100% Traceable 100% Free of steroids		
	Average per 100g		Average per 100g		Average per 100g		Average per 100g	
Engergy	590 KJ	Engergy	520 KJ	Engergy	545 KJ	Engergy	610 KJ	
Protein	18 g	Protein	27 g	Protein	26 g	Protein	33 g	
Fat	9 g	Fat	18 g	Fat	11 g	Fat	13 g	
Omega 3	0.28 g	Omega 3	0.28 g	Omega 3	0.28 g	Omega 3	0.28 g	
Iron	5 mg	Iron	5 mg	Iron	3 mg	Iron	4 mg	
Zinc	9 mg	Zinc	11 mg	Zinc	10 mg	Zinc	14 mg	
est Before 3 days to KG \$18.00 let KG 0.25 kg otal Price \$4.50 iven the meat pr	日次日 35年3月 日後代	Best Before 1 week to \$KG \$19.00 Net KG 0.5 kg Total Price \$9.50 ed, how many of e		Best Before 1 week 1 \$KG \$18.00 Net KG 0.5 kg Total Price \$9.00 ne) would you mi	o expiry	Best Before 2 weeks SIKG \$14.00 Net KG 1 kg Total Price \$14.00 ssuming these we		

None	© None	© None	None
One	One	One	One
©тwo	©т⊮о	©т⊮о	©™
Othree	OThree	OThree	OThree
More than three	More than three	More than three	More than three



Iberia	IBERIA 🖡					Total	3 hr 40 min	€65	
Flight Ground Tran	sportation	From Amsterdam Girona Airport	To Girona Airport Barcelona City	Departure 6:00	Arrival 8:00	Stops 0	Travel Time 2 hr 00 min 1 hr 40 min	Price € 50 € 15	Choose this ticket
Transavia	transavia.com					Total	5 hr 00 min	€65	
Flight Ground Tran	sportation	From Amsterdam Girona Airport	To Girona Airport Barcelona City	Departure 12:00	Arrival 16:00	Stops 1 - (2 hr 00)	Travel Time 4 hr 00 min 1 hr 00 min	Price € 50 € 15	Choose this ticket
Vueling	vueling com					Total	4 hr 20 min	€84	
Flight Ground Tran	sportation	From Amsterdam Girona Airport	To Girona Airport Barcelona City	Departure 6:00	Arrival 9:00	Stops 1 - (1 hr 00)	Travel Time 3 hr 00 min 1 hr 20 min	Price €75 €9	Choose this ticket



Which would you choose if these were your only options?

L			
L			
_	_	_	

No medicine

Medicine

Johnson FR, et al, Value Health. 2019;22(9):1063-9.

Ways probabilities are presented matters

• Risk

• Probability or likelihood of an outcome

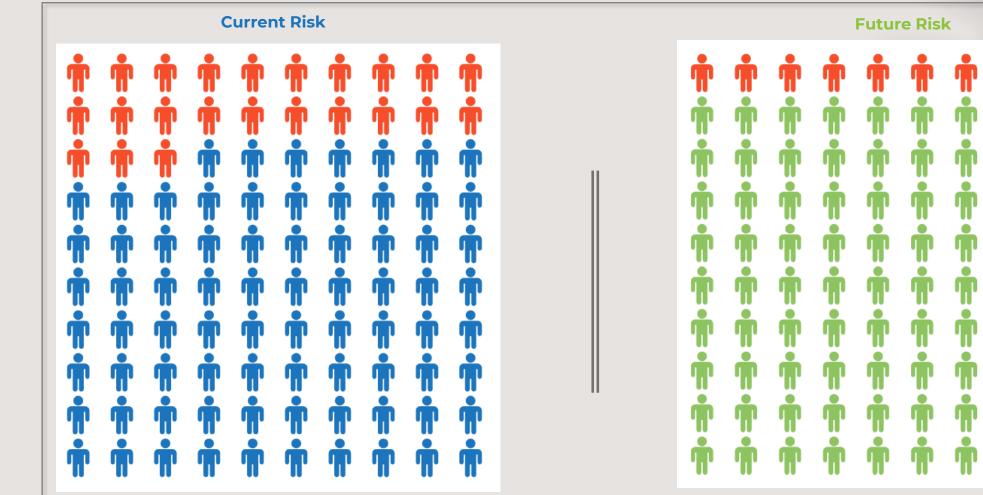
Phyu Mon Latt Monash University



MySTIRisk

Risk Display Options

Icon Array Option 1



Your highest risk of getting an STI is Chlamydia.

At present, in a group of 100 people who gave the same answers as you on this survey, 23 will have Chlamydia.

Over the next 12 months, in a group of 100 people who gave the same answers as you on this survey, 10 are likely to have Chlamydia.

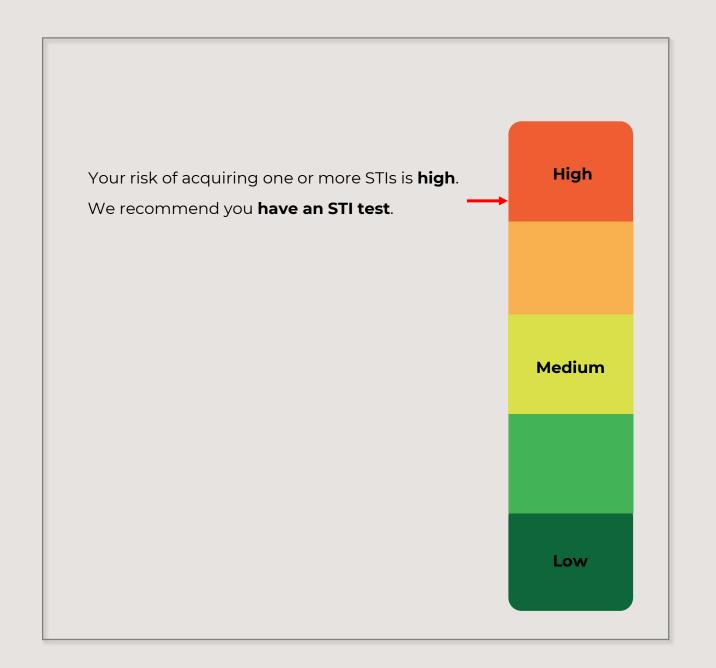
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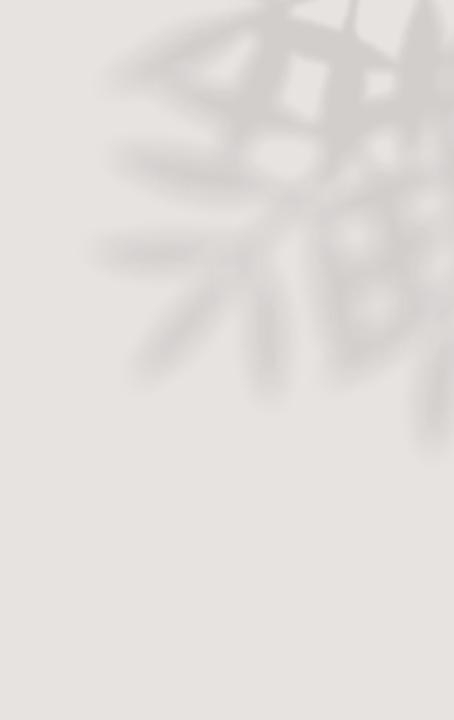
۲

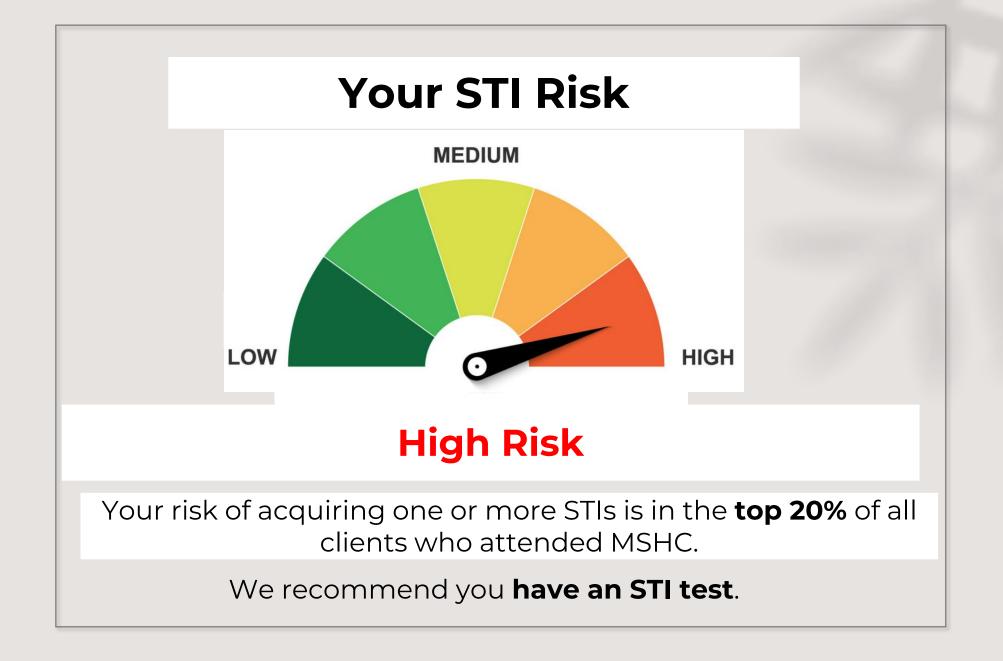


Your risk of acquiring one or more STIs is **high**.

Australian Guidelines recommend you have an STI testing.







Risk report

If a person had attended Melbourne Sexual Health Centre and provided the answers you did, their risk of infections are below.

Risk of HIV

- At present, in a group of **1000 people**, **4** will have HIV.
- Over the next 12 months, in a group of **1000 people**, **2** are likely to catch HIV.

Risk of Syphilis

- At present, in a group of 100 people, 9 will have syphilis.
- Over the next 12 months, in a group of **100 people**, **4** are likely to catch syphilis.

Risk of Gonorrhoea

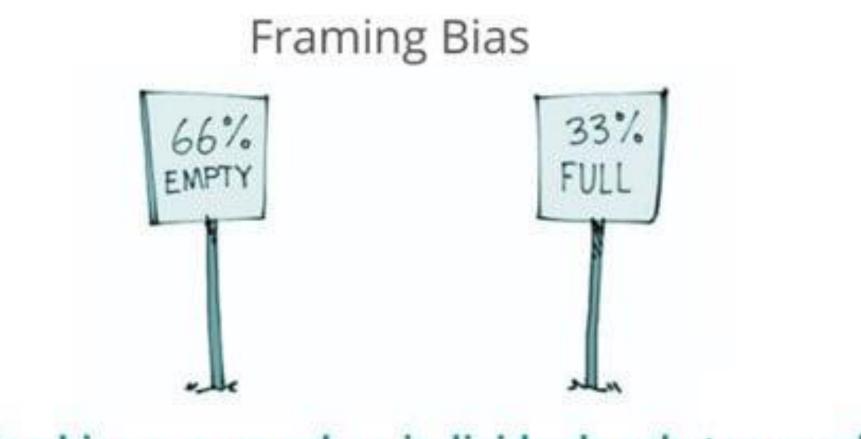
- At present, in a group of **100 people**, **14** will have gonorrhoea.
- Over the next 12 months, in a group of **100 people**, **9** are likely to catch gonorrhoea.

Risk of Chlamydia

- At present, in a group of 100 people, 23 will have chlamydia.
- Over the next 12 months, in a group of **100 people**, **10** are likely to catch chlamydia.

Which is the 'right' one?

- Test which is the most effective, preferred by your target population
- Ask Phyu Mon later...



Framing bias occurs, when individuals rely too much on "how information is presented."



Framing bias

- Individual's decisions influenced by the way information is presented (instead of facts and figures)
- "Positive" framing
 - 70% people saved
- "Negative" framing
 - 30% people died
- People more likely to "gamble" to avoid losses

Be careful

- Left-right bias
- Order bias
- Colours
- Imagery





How can you check if your choice set presentation is reasonable?

Test it!

Pre-test Think aloud interviews • Pilot Up to 20 people 10% of sample

Tips

- Think about your audience
 - How much background knowledge would they have?
 - Would they understand the terms consistently?
- Instruction page before DCE survey
- Account for or test for cognitive biases
 - Framing
 - Remove anchoring statements
 - Left-right / order bias

We will now show some choices to understand your HIV PrEP care preferences. You won't have experienced some of these things, but please try and decide what you think you would prefer based on the information shown. Below we detail the options you will select among:

PrEP modality:

- Daily oral pill: a daily tablet you take each day
- On-demand tablet: you take the pill depends on your sexual activity schedule 2 pills 2-24 hours before sex, 1 pill 24 hours after the first pill, and 1 pill 48 hours after the first pill.
- Long-acting injectable PrEP: injectable PrEP, after the first two injections in the first month, provided every 2 months.
- PrEP implant: small (2-4cm) tube inserted under the skin of upper arm, allows slow release to prevent HIV infection for 5-6 months. It can be removed if treatment needs to be stopped.

Getting your PrEP prescription:

- Same-day prescription: you don't need to wait the lab test result to get your prescription.
- 2-visit prescription: you wait till the lab result shows that you are eligible to take on PrEP (standardof-care).
- Telehealth prescription: you will still be required to have the same lab tests and upload results to the online platform.

In the example below, choosing Option 1 means you would prefer to use daily oral pill, 2-visit prescription, having text reminders about taking your pill and follow-up visit, and pay no out-of-pocket cost.

Choosing Option 2 means you would prefer to use on-demand oral pill, same-day prescription, having a smartphone application to remind you about taking pills and follow-up visit, and pay 700 RMB out-of-pocket.

If you do not prefer option 1 or option 2, you can choose "None of these options".

Option 1	Option 2
Daily oral pill	On-demand oral pill
2-visit prescription	Same-day prescription
Medication home delivery	Medication home delivery
Text reminder for medication adherence	Smartphone application
At no cost	30% off (700RMB)



Questions?

Session 4 – Live demonstration NGENE / Qualtrics

www.choice-metrics.com

- Experimental design generation software
- Can apply constraints
- Generate orthogonal and Bayesian efficient designs

gene

www.choice-metrics.com

• Syntax driven

🔝 Syntax - Singapore DCE Workshop 2023.ngs

```
Design
; alts = A, B, C
; rows = 60
; eff = (mnl,d)
; block = 10
; cond:
if(a.Mode = [3], a.Pickup=[0,1]),
if(b.Mode = [3], b.Pickup=[0,1])
; model:
U(A) = b1.effects[0|0|0] * Mode[1,2,3,0] +
       b2.effects[0|0]*Pres[1,2,0] +
       b3.effects[0|0|0|0]*Pickup[1,2,3,4,0] +
       b4.effects[0|0|0]*Supp[1,2,3,0] +
       b5.effects[-0.00001]-0.00001]-0.00001]*Cost[1,2,3,0]/
U(B) = b1.effects*Mode +
       b2.effects*Pres +
       b3.effects*Pickup +
       b4.effects*Supp +
       b5.effects*Cost/
U(C) = b6[0]
```



https://choice-metrics.com/NgeneManual120.pdf



Ngene 1.2 USER MANUAL & REFERENCE GUIDE

NGENE live demonstration

Qualtrics live demonstration



PRODUCTS V SOLUTIONS V CUSTOMERS V RESOURCES V COMPANY V



Platform > Core XM > Conjoint Analysis

CONJOINT ANALYSIS SOFTWARE TOOL

Optimize products and pricing with predictive insights

REQUEST DEMO

	· <u>·</u>			Option 1	Option 2
(1/5) Chor	se your pre	aferred		Garmin	Apple
	options fro		acker	Included	Not Included
			ker	Not Included	Included
frant	Option 1 Garrerin	Option 1 Apprin	ture	Weight Loss Coach	Immune System Tracking
Activity Tracker	Included	First Included		\$59	\$99
Seep Tracker	NativeLabel	Not included		0	0
Health Feature	Classify	System Track-Ing		0	0
Cent	\$50	- 810			Continue
	0	0			

Session 5. How to analyse choice data

Have you downloaded Nlogit and the 'PrEP DCE.lpj' file?

Warittha Tieosapjaroen (Nittha)

- PharmD, MBiotech
- PhD Candidate at Melbourne Sexual Health Centre, Alfred Health, Central Clinical School, Monash University
- HOPE Project Manager
- wtieosapjaroen@mshc.org.au

CLINICAL SCIENCE

Preferences for Weight Gain Compared With Other Antiretroviral Therapy Side Effects in People Living With HIV: A Discrete Choice Experiment

Warittha Tieosapjaroen, PharmD, MBiotech,^{a.b} Christopher K. Fairley, MBBS, PhD, FRACP, FAFPHM,^{a.b} Eric P.F. Chow, PhD, MPH, MBiostats, MApplSc,^{a.b.c} Ivette Aguirre, MClinPharm,^a Jennifer F. Hoy, AM, FRACP, FAHMS,^d and Jason J. Ong, PhD, MMed, MBBS, FAChSHM, FRACGP^{a,b,e}



Session 4: Abstract driven session Friday June 9, 2023 10.30-11.30 am

Objectives of the session

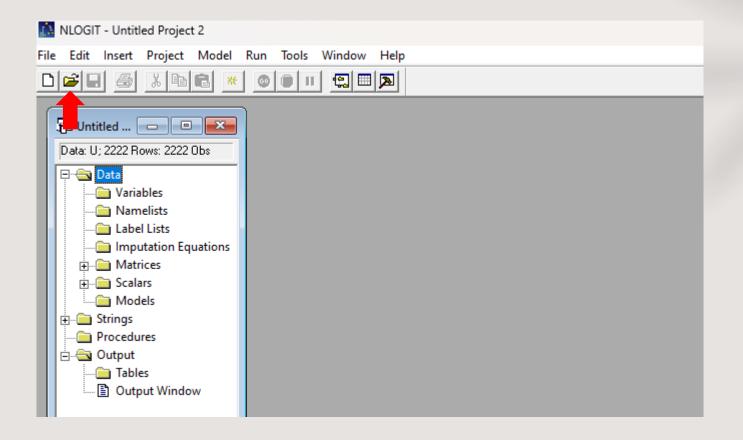
- Understand DCE data
- Demonstrate DCE syntax
- Demonstrate to Nlogit
- Understand DCE outputs

Common DCE models

Multinomial logit (MNL) - analyse preferences from each choice set independently

- Random parameter logit (RPL) analyse preferences by panelling all the choice sets from each respondent
- Latent class model (LCM) explore preferences shared between individuals and group them together.

Understand DCE data



Open the 'PrEP DCE.lpj' file

Understand DCE data

NLOGIT - 305 Asian DCE MSM - all completed the survey only three atttributes.lpj File Edit Insert Project Model Run Tools Window Help አ 🖻 💼 💥 🐑 🖽 🔊 🔂 305 Asia... 🗖 🔲 💌 Data: U; 5494 Rows: 5490 Obs 🖃 👈 🔁 Data 🙈 Variables Ē٠ PID TOTALROW CHOICEV QUESTION ALTIJ QUESTIO0 TYPE LOCATION COST SIDEEFFE FREQ EXTRA A_MERGE CSET TYPE1 TYPE2 Data

Understand DCE data

Choice indicator 1= chosen 0= not chosen Number of alternatives in each choice situation Alternative indicator 1= Alt. A 2= Alt. B 3= Alt. C or Opt out

	pid	totalrow	choicev	question	cset	altij	type	location	cost	sideeffects	freq	extra
1	168	18	0	1	3	1	2	6	4	4	2	3
2	168	18	1	1	3	2	1	1	1	3	3	1
з	168	18	0	1	3	3						
4	168	18	0	2	3	1	2	1	4	2	4	3
5	168	18	1	2	3	2	1	3	3	3	2	1
6	168	18	0	2	3	3			-		-	
7	168	18	1	3	3	1	5	2	1	4	4	1
8	168	18	0	3	3	2	3	4	3	1	1	3
9	168	18	0	3	3	3						
10	168	18	1	4	3	1	5	5	1	4	2	3

An example of a DCE question

Cset=3

```
Altij
1= Alt. A
2= Alt. B
3= Alt. C or Opt out
```

Question 1	Α	В	Opt out
Type of PrEP	Oral long-acting PrEP	Injectable PrEP	Х
Service location	Pharmacy	Hospital	Х
Cost	\$AU 25	Free	Х
Side effects	Rare chance of kidney problems	Mild	X
Visit frequency	Every 6 months	Every year	Х
Extra services	STI testing	None	Х
Which choice do you prefer?	0	0	0

Understand your DCE data

Number of alternatives in each choice situation Alternative indicator 1= Alt. A 2= Alt. B 3= Alt. C or Opt out

	pid	totalrow	choicev	question	cset	altij	type	location	cost	sideeffects	freq	extra
1	168	18	0	1	3	1	2	6	4	4	2	3
2	168	18	1	1	3	2	1	1	1	3	3	1
з	168	18	0	1	3	3						
4	168	18	0	2	3	1	2	1	4	2	4	3
5	168	18	1	2	3	2	1	3	3	3	2	1
6	168	18	0	2	3	3	-					
7	168	18	1	3	3	1	5	2	1	4	4	1
8	168	18	0	3	3	2	3	4	3	1	1	3
9	168	18	0	3	3	3						
10	168	18	1	4	3	1	5	5	1	4	2	3

An example of a DCE question

Question 1	Α	В	Opt out
Type of PrEP	Oral long-acting PrEP	Injectable PrEP	Х
Service location	Pharmacy	Hospital	Х
Cost	\$AU 25	Free	х
Side effects	Rare chance of kidney problems	Mild	Х
Visit frequency	Every 6 months	Every year	Х
Extra services	STI testing	None	Х
Which choice do you prefer?	0		0

Understand your DCE data

Choice indicator 1= chosen 0= not chosen Alternative indicator 1= Alt. A 2= Alt. B 3= Alt. C or Opt out

	pid	totalrow	choicev	question	cset	altij	type	location	cost	sideeffects	freq	extra
1	168	18	0	1	3	1	2	6	4	4	2	3
2	168	18	1	1	3	2	1	1	1	3	3	1
з	168	18	0	1	3	3						
4	168	18	0	2	3	1	2	1	4	2	4	3
5	168	18	1	2	3	2	1	3	3	3	2	1
6	168	18	0	2	3	3		-			-	
7	168	18	1	3	3	1	5	2	1	4	4	1
8	168	18	0	3	3	2	3	4	3	1	1	3
9	168	18	0	3	3	3						
10	168	18	1	4	3	1	5	5	1	4	2	3

Nlogit

- ;lhs=choicev,cset,altij
- ;choices=A,B,C

;model:

- U(A,B) = Type1*Type1+Type2*Type2+Type3*Type3+Type4*Type4+
 - Cost1*Cost1+Cost2*Cost2+Cost3*Cost3+
 - SE1*SE1+SE2*SE2+SE3*SE3+SE4*SE4/
- U(C)=neither

\$

Modify the code in red according to your DCE data

;=start the command / =separate utility functions \$= end command

Nlogit

- ;lhs=choicev,cset,altij
- ;choices=A,B,C

;model:

In the 'PrEP DCE' data, there are Alternatives A,B and opt out If you have four alternatives, ;choices=A,B,C,D

- U(A,B) = Type1*Type1+Type2*Type2+Type3*Type3+Type4*Type4+
- Cost1*Cost1+Cost2*Cost2+Cost3*Cost3+
- SE1*SE1+SE2*SE2+SE3*SE3+SE4*SE4/
- U(C)=neither

\$

Nlogit

- ;lhs=choicev,cset,altij
- ;choices=A,B,C

;model:

U (alternative name)= <parameter name>*<variable name>

;model:

U(A,B) = Type1*Type1+Type2*Type2+Type3*Type3+Type4*Type4+

Cost1*Cost1+Cost2*Cost2+Cost3*Cost3+

SE1*SE1+SE2*SE2+SE3*SE3+SE4*SE4/

U(C)=neither

\$

*baseline levels are not included

U (alternative name)= <parameter name>*<variable name>

Variables = attributes presented to respondent.

Parameters = estimated values that represent underlying preferences that individuals have for each level of variables

DCE syntax

Nlogit

;lhs=choicev,cset,altij

;choices=A,B,C

;model:

U(A,B) = Type1*Type1+Type2*Type2+Type3*Type3+Type4*Type4+

Cost1*Cost1+Cost2*Cost2+Cost3*Cost3+

SE1*SE1+SE2*SE2+SE3*SE3+SE4*SE4/

U(C)=neither

\$

*baseline levels are not included

Practice 5.1

(Question 5.1) Create a DCE syntax for the dataset below

pid	Totalrow	choicev	question	cset	altij	nausea	diarrhea	headache
111	4	1	1	2	1	1	2	3
111	4	0	1	2	2	2	3	1
111	4	1	2	2	1	3	2	1
111	4	0	2	2	2	2	1	3
112	4	0	1	2	1	1	3	2
112	4	1	1	2	2	3	1	2
112	4	0	2	2	1	2	1	3
112	4	1	2	2	2	3	2	1
113	4	1	1	2	1	2	1	3
113	4	0	1	2	2	1	2	3
113	4	1	2	2	1	1	3	2
113	4	0	2	2	2	2	3	1

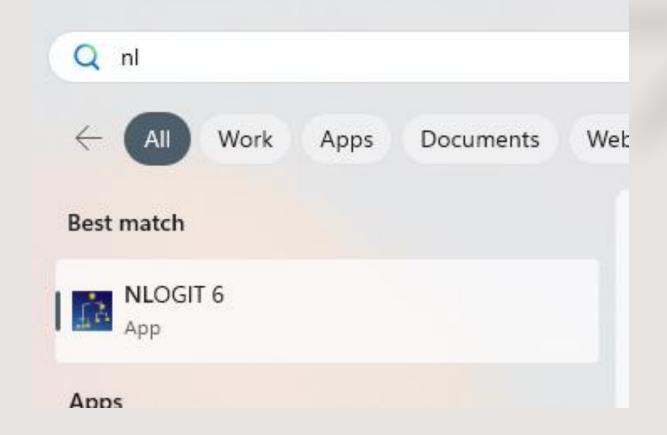
Please answer the questions 5.1 using data provided.

Antiretroviral A and B have side effects as mentioned below. Choose the antiretroviral that you prefer more.

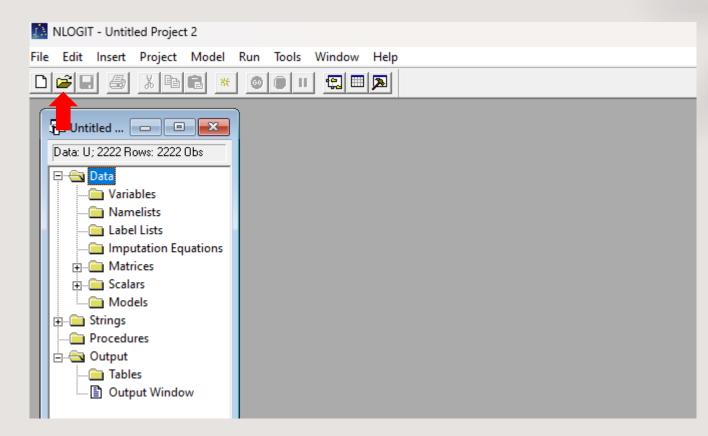
	Antiretroviral A	Antiretroviral B
Nausea	Not all all	Once a week
diarrhea	Three times a week	Not at all
headache	Not at all	Once a week
	0	\bigcirc

There are two DCE questions in this survey

Nc	ot at all	Attribute level 1
Or	nce a week	Attribute level 2
Th	ree times a week	Attribute level 3

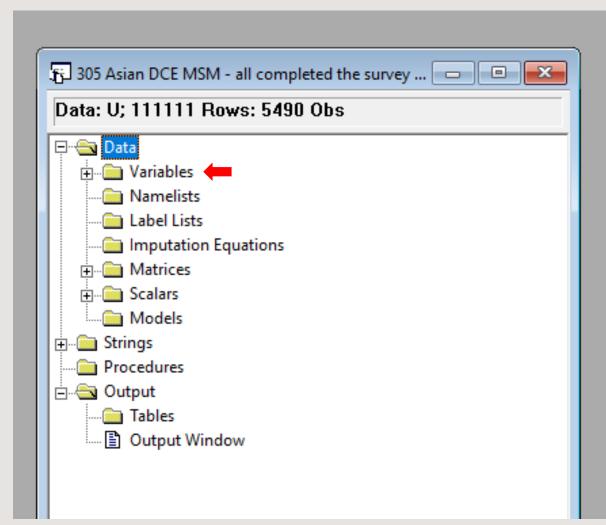


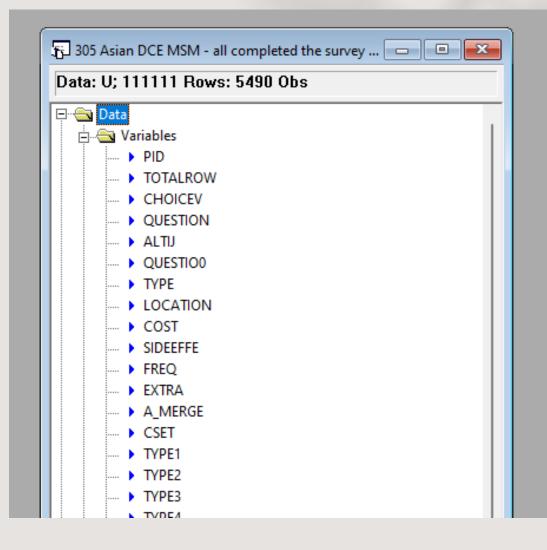
If the project window is already opened



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305 Asian DCE MSM - all completed the survey.lpj	Type: LPJ File Size: 1.04 MB Date modified
File name:	Open
Files of type: All Readable Files (*.lim; *.lpj; *.sav) 🖛 💌	Cancel





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305 Asia Data: U; 55555 Rows: 5490 Obs Data Dat	Untitled 3* Insert Name: Nlogit ; lhs=choicev,cset,altij ; choices=A,B,C ; check data ;model: U(A,B)=Type1*Type1+Type2*Type3*Type3+Type4*Type4+ Cost1*Cost1+Cost2*Cost2+Cost3*Cost3+ SE1*SE1+SE2*SE2+SE3*SE3+SE4*SE4/ U(C)=neither \$		

NLOGIT - Untitled 3 * — D				
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÷-	;choices=A,B,C ;check_data				
⊕ <mark>`</mark> Str	;model: U(A,B)=Type1*Type1+Type2*Type2+Type3*Type3+Type4*Type4+				
<u>(</u> Pr ⊡ (Ou	Cost1*Cost1+Cost2*Cost2+Cost3*Cost3+ SE1*SE1+SE2*SE2+SE3*SE3+SE4*SE4/				
	U(C)=neither \$				
····· (Ξ)	Inspecting the data set before estimation. These errors mark observations which will be skipped.				
	Rose Filots wark observations which will be skipped. Row Individual = 1st row then group number of data block				
	No bad observations were found in the sample				
	Iterative procedure has converged Normal exit: 5 iterations. Status=0, F= .1826378D+04				
	Discrete choice (multinomial logit) model Dependent variable Choice Log likelihood function -1826.37831				
	Estimation based on N = 1830, K = 12 Inf.Cr.AIC = 3676.8 AIC/N = 2.009				
	Log likelihood R-sqrd R2Adj				
Data	ASCs only model must be fit separately Use NLOGIT ;;RHS=ONE\$				
	Note: R-sqrd = 1 - logL/Logl(constants) Warning: Model does not contain a full set of ASCs. R-sqrd is problematic. Use				
	model setup with ;RHS=one to get LogL0.				
	Root Likelihood:Geom. Mean of P [*] .3686 Response data are given as ind. choices				
	Number of Obs. = 1830, skipped 0 obs				
	Standard Prob. 95% Confidence CHOICEV Coefficient Error z z >Z* Interval				
	TYPE1 .24388*** .06540 3.73 .0002 .11569 .37206 TYPE2 .08589 .07893 1.09 .2765 06881 .24058				
	TYPE304222 .0658464 .521417128 .08683 TYPE419615*** .06568 -2.99 .00283248706742				
	COST1 .04817 .05347 .90 .367705663 .15298 COST220063*** .05470 -3.67 .00023078409342				
	COST341327*** .05735 -7.21 .00005256730086 SE1 .02937 .06376 .46 .645009559 .15434				
	SE202724 .0672341 .685415901 .10454 SE304397 .0645968 .496117056 .08263				
	SE4 .09150 .12775 .72 .4738 15889 .34189 NEITHER 81453*** .06494 -12.54 .0000 94180 68725				
	****, ** ==> Significance at 1%, 5%, 10% level. Model was estimated on May 12 2023 at 03:03:47 PM				

Ln 157/157 Idle 15:06

Inspecting the data set before estimation. | These errors mark observations which will be skipped. | Row Individual = 1st row then group number of data block |

No bad observations were found in the sample

Iterative procedure has converged Normal exit: 5 iterations. Status=0, F= .1826378D+04 Check bad observations

Normal or abnormal convergence

Discrete choice (multinomial logit) model Dependent variable Choice Log likelihood function -1826.37831 Estimation based on N = 1830, K = 12 Inf.Cr.AIC = 3676.8 AIC/N = 2.009

Log likelihood R-sqrd R2Adj ASCs only model must be fit separately Use NLOGIT ;...;RHS=ONE\$ Note: R-sqrd = 1 - logL/Logl(constants) Warning: Model does not contain a full set of ASCs. R-sqrd is problematic. Use model setup with ;RHS=one to get LogL0. Root Likelihood:Geom. Mean of P[^].3686

Response data are given as ind. choices Number of obs.= 1830, skipped 0 obs Check model fit Lower LL and AIC/N = better model fit

Compare the number of observations loaded to Nlogit to your dataset

	Standard Prob. 95% Confidence CHOICEV Coefficient Error z z >Z* Interval
On-demand PrEP	TYPE1 .24388*** .06540 3.73 .0002 .11569 .37206
Injectable PrEP	TYPE2 .08589 .07893 1.09 .276506881 .24058
Long-acting oral PrEP	TYPE304222 .0658464 .521417128 .08683
Implant PrEP	TYPE419615*** .06568 -2.99 .00283248706742
Low cost	COST1 .04817 .05347 .90 .367705663 .15298
Fair cost	COST220063*** .05470 -3.67 .00023078409342
High cost	COST3 41327*** .05735 -7.21 .00005256730086
Interaction with other medications	SE1 .02937 .06376 .46 .645009559 .15434
Mild	SE2 02724 .0672341 .685415901 .10454
Rare chance of kidney problems	SE3 04397 .0645968 .496117056 .08263
Mild pain at injection	SE4 .09150 .12775 .72 .473815889 .34189
, ,	U(C)=NEI .81453*** .06494 12.54 .0000 .68725 .94180

***, **, * ==> Significance at 1%, 5%, 10% level. Model was estimated on Mar 21, 2023 at 06:41:17 PM

Level

Free PrEP

Low cost

Fair cost

High cost

	Standard Prob. 95% Confidence CHOICEV Coefficient Error z z >Z* Interval
	TYPE1 .24388*** .06540 3.73 .0002 .11569 .37206 TYPE2 .08589 .07893 1.09 .276506881 .24058
Coefficient	TYPE3 04222 .0658464 .521417128 .08683
= -sum(low + medium + high cost) =0.566	TYPE4I .19615*** .06568 -2.99 .0028 32487 06742 COST1 .04817 .05347 .90 .3677 05663 .15298
0.048	COST2 20063** .05470 -3.67 .00023078409342 COST3 41327** .05735 -7.21 .00005256730086
-0.201	SE11 .02937 .06376 .46 .645009559 .15434
-0.413	SE202724 .0672341 .685415901 .10454
	SE304397 .0645968 .496117056 .08263 SE4 .09150 .12775 .72 .473815889 .34189 U(C)=NEI .81453*** .06494 12.54 .0000 .68725 .94180

***, **, * ==> Significance at 1%, 5%, 10% level. Model was estimated on Mar 21, 2023 at 06:41:17 PM

Output interpretation

Like

Free	<pre>= - sum(low + medium + high cost) =0.566</pre>
Low cost	0.048
Fair cost	-0.201
High cost	-0.413

Dislike

Standard Prob. 95% Confidence CHOICEV Coefficient Error z z >Z* Interval
TYPE1 .24388*** .06540 3.73 .0002 .11569 .37206
TYPE2 .08589 .07893 1.09 .276506881 .24058
TYPE304222 .0658464 .521417128 .08683
TYPE419615*** .06568 -2.99 .00283248706742
COST1 .04817 .05347 .90 .367705663 .15298
COST2 20063*** .05470 -3.67 .00023078409342
COST3 41327*** .05735 -7.21 .00005256730086
SE1 .02937 .06376 .46 .645009559 .15434
SE2 02724 .0672341 .685415901 .10454
SE3 04397 .0645968 .496117056 .08263
SE4 .09150 .12775 .72 .473815889 .34189
U(C)=NEI .81453*** .06494 12.54 .0000 .68725 .94180

***, **, * ==> Significance at 1%, 5%, 10% level. Model was estimated on Mar 21, 2023 at 06:41:17 PM

Practice 5.2

No bad obs	ervations were				I	
	procedure has <u>c</u>					
Normal exi	t: 5 iteratio	ns. Status=0	, F=	.16716961)+05	
 Discrete c	hoice (multinom	ial logit) m	odel			
	variable hood function					
	based on N =					
mi.cr.Alc	33457.9 AI	C/N = 1.9				
	Log likelihoo	d R-sard R2A	.dj			
ASCs only	model must be					
Note: R-sa	Use NLOGIT rd = $1 - \log L/L$	_:;RHS=ON	2) 2)			
	Note: R-sgrd = 1 - logL/Logl(constants) Warning: Model does not contain a full					
set of ASCs. R-sgrd is problematic. Use						
set of ASC	s. R-sgrd is pr	oblematic. U	se			
set of ASC model setu	s. R-sgrd is pr p <u>with :RHS</u> =one	oblematic. U to get LogI	se 0.			
set of ASC model setu	s. R-sgrd is pr	oblematic. U to get LogI	se 0.			
set of ASC model setu Root Likel Response d	s. R- <u>sgrd</u> is pr p <u>with :RHS</u> =one ihood: <u>Geom.</u> Mea ata are given a	oblematic. U to get LogI n of P^ .37 	se 0. 48 			
set of ASC model setu Root Likel Response d	s. R-sgrd is pr p <u>with :RHS</u> =one ihood:Geom. Mea	oblematic. U to get LogI n of P^ .37 	se 0. 48 			
set of ASC model setu Root Likel Response d Number of	s. R-sgrd is pr p <u>with :RHS</u> =one <u>ihood:Geom.</u> Mea ata are given a obs.= 17034, sk	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard	se 0. 48 <u>es</u> 25	Prob.	95% Cor	nfidence
set of ASC model setu Root Likel Response d Number of	s. R-sgrd is pr p <u>with :RHS</u> =one <u>ihood:Geom.</u> Mea ata are given a obs.= 17034, sk	oblematic. U to get LogI n of P [^] .37 s ind. <u>choic</u> ipped 0 g Standard Error	se 0. 48 <u>es</u> 2	z >Z*	Inte	erval
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1	s. R- <u>sgrd</u> is pr p <u>with :RHS</u> =one ihood:Geom. Mea ata are given a obs.= 17034, sk <u>Coefficient</u> .18423***	oblematic. U to get LogI n of P [^] .37 s ind. <u>choic</u> ipped 0 g Standard Error	se 0. 48 <u>es</u> 2	z >Z*	Inte	erval
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1	s. R- <u>sgrd</u> is pr p <u>with :RHS</u> =one ihood:Geom. Mea ata are given a obs.= 17034, sk <u>Coefficient</u> .18423***	oblematic. U to get LogI n of P [^] .37 s ind. <u>choic</u> ipped 0 g Standard Error	se 0. 48 <u>es</u> 2	z >Z*	Inte	erval
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1	s. R- <u>sgrd</u> is pr p <u>with :RHS</u> =one ihood:Geom. Mea ata are given a obs.= 17034, sk <u>Coefficient</u> .18423***	oblematic. U to get LogI n of P [^] .37 s ind. <u>choic</u> ipped 0 g Standard Error	se 0. 48 <u>es</u> 2	z >Z*	Inte	erval
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1 TYPE1 TYPE3 TYPE3 TYPE4	s. R- <u>sgrd</u> is pr p <u>with :RHS</u> =one ihood:Geom. Mea ata are given a obs.= 17034, sk <u>Coefficient</u> .18423*** -10623*** .20243*** 33720***	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard Error .02209 .02681 .02177 .02297	se 0. 48 <u>z</u> <u>8.34</u> - <u>3.96</u> 9.30 -14.68	z >Z* 0000 0001 0000 0000	Inte .14092 15878 .15975 38222	22753 05367 .24510 29219
set of ASC model setu Root Likel Response d Vumber of CHOICEV CHOICEV TYPE1 TYPE2 TYPE3 TYPE3 TYPE4 COST1	<pre>s. R-sgrd is pr p with :RHS=one ihood:Geom_ Mea ata are given a obs.= 17034, sk Coefficient .18423*** .10623*** .20243*** .33720*** .15201***</pre>	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard Error .02209 .02681 .02177 .02297	se 0. 48 <u>z</u> <u>8.34</u> - <u>3.96</u> 9.30 -14.68	z >Z* 0000 0001 0000 0000	Inte .14092 15878 .15975 38222	22753 05367 .24510 29219
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1 TYPE1 TYPE2 TYPE3 TYPE4 COST1 COST1	s. R-sgrd is pr p with :RHS=one ihood:Geom. Mea ata are given a obs.= 17034, sk 	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard Error .02209 .02681 .02177 .02297 .01789 .01816	z <u>8.34</u> <u>8.34</u> <u>8.34</u> -3. <u>96</u> <u>9.30</u> -14. <u>68</u> <u>8.50</u> -5. <u>04</u> <u>24.055</u>	z >Z* .0000 .0001 .0000 .0000 .0000 .0000	14092 15878 .15975 38222 .11695 12710	.22753 05367 .24510 29219 .18707 05593
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1 TYPE1 TYPE2 TYPE3 TYPE4 COST1 COST1	s. R-sgrd is pr p with :RHS=one ihood:Geom. Mea ata are given a obs.= 17034, sk 	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard Error .02209 .02681 .02177 .02297 .01789 .01816	z <u>8.34</u> <u>8.34</u> <u>8.34</u> -3. <u>96</u> <u>9.30</u> -14. <u>68</u> <u>8.50</u> -5. <u>04</u> <u>24.055</u>	z >Z* .0000 .0001 .0000 .0000 .0000 .0000	14092 15878 .15975 38222 .11695 12710	.22753 05367 .24510 29219 .18707 05593
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1 TYPE1 TYPE2 TYPE3 TYPE4 COST1 COST1	s. R-sgrd is pr p with :RHS=one ihood:Geom. Mea ata are given a obs.= 17034, sk 	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard Error .02209 .02681 .02177 .02297 .01789 .01816	z <u>8.34</u> <u>8.34</u> <u>8.34</u> -3. <u>96</u> <u>9.30</u> -14. <u>68</u> <u>8.50</u> -5. <u>04</u> <u>24.055</u>	z >Z* .0000 .0001 .0000 .0000 .0000 .0000	14092 15878 .15975 38222 .11695 12710	.22753 05367 .24510 29219 .18707 05593
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1 TYPE1 TYPE2 TYPE3 TYPE4 COST1 COST1	s. R-sgrd is pr p with :RHS=one ihood:Geom. Mea ata are given a obs.= 17034, sk 	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard Error .02209 .02681 .02177 .02297 .01789 .01816	z <u>8.34</u> <u>8.34</u> <u>8.34</u> -3. <u>96</u> <u>9.30</u> -14. <u>68</u> <u>8.50</u> -5. <u>04</u> <u>24.055</u>	z >Z* .0000 .0001 .0000 .0000 .0000 .0000	14092 15878 .15975 38222 .11695 12710	.22753 05367 .24510 29219 .18707 05593
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1 TYPE1 TYPE2 TYPE3 TYPE4 COST1 COST1	s. R-sgrd is pr p with :RHS=one ihood:Geom. Mea ata are given a obs.= 17034, sk 	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard Error .02209 .02681 .02177 .02297 .01789 .01816	z <u>8.34</u> <u>8.34</u> <u>8.34</u> -3. <u>96</u> <u>9.30</u> -14. <u>68</u> <u>8.50</u> -5. <u>04</u> <u>24.055</u>	z >Z* .0000 .0001 .0000 .0000 .0000 .0000	14092 15878 .15975 38222 .11695 12710	.22753 05367 .24510 29219 .18707 05593
set of ASC model setu Root Likel Response d Number of CHOICEV TYPE1 TYPE1 TYPE2 TYPE3 TYPE4 COST1 COST1	s. R- <u>sgrd</u> is pr p <u>with :RHS</u> =one ihood:Geom. Mea ata are given a obs.= 17034, sk <u>Coefficient</u> .18423*** .10623*** .20243*** .33720*** .15201*** .09151***	oblematic. U to get LogI n of P^ .37 s ind. <u>choic</u> ipped 0 g Standard Error .02209 .02681 .02177 .02297 .01789 .01816	z <u>8.34</u> <u>8.34</u> <u>8.34</u> -3. <u>96</u> <u>9.30</u> -14. <u>68</u> <u>8.50</u> -5. <u>04</u> <u>24.055</u>	z >Z* .0000 .0001 .0000 .0000 .0000 .0000	14092 15878 .15975 38222 .11695 12710	.22753 05367 .24510 29219 .18707 05593

Attribute	Levels	Coefficient	Range of attribute	Relative Importance*100
Type of PrEP	Daily oral			
	On-demand			
	Injectable			
	Long-acting oral			
	Implant			
Cost	Free			
	Low			
	High			
	Very High			
Side effects	No			
	Interactions with			
	other medications			
	Mild			
	Rare change of			
	kidney problems			
	Mild pain at			
	injection			

The participants most preferred daily oral PrEP, followed by implant, long acting oral,.....

Relative importance

Attribute	Levels	Coefficient	Range of attribute	Relative Importance*100	
Type of PrEP	Daily oral On-demand Injectable Long-acting oral	-0.0914 0.24388 0.08589 -0.04222	0.24388-(-0.19615)= <mark>0.44003</mark>	0.44003/(0.44003+0.97900+0.1 4116)=28.2	
	Implant	-0.19615			
Cost	Free Low High Very High	0.56573 0.04817 -0.20063 -0.41327	<mark>0.56573</mark> -(<mark>-0.41327</mark>)= <mark>0.97900</mark>	0.97900/(<mark>0.44003+0.97900+0.1</mark> <mark>4116</mark>)=62.7	
Side effects	No Interactions with other medications Mild Rare change of kidney problems	-0.04966 0.02937 -0.02724 -0.04397	0.09150-(-0.04966)= <mark>0.14116</mark>	0.14116/(0.44003+0.97900+0.1 4116)=9.05	
	Mild pain at injection	0.09150			

Session 6. Application of choice data

Objectives of the session

- Demonstrate how DCE data can be applied
- RPL models
 - Heterogeneity overall
 - RPLX interactions with certain sociodemographic groups
- LCA models
 - Market shares
- Simulation
 - Uptake from best/worst combinations of attribute levels

Real world examples

Designing HIV testing and self-testing services for young people in Nigeria: A discrete choice experiment



The Patient - Patient-Centered Outcomes Research

An Official Journal of the International Academy of Health Preference Research

Motivation

• A third of new HIV infections occur among young people and the majority of young people living with HIV are in sub-Saharan Africa.

• Aim

• We examined the strength of Nigerian youth preferences related to HIV testing and HIV self-testing.

Method

- Participants completed one of two DCEs:
 - 1) preferred qualities of HIV testing (cost, location of test, type of test, person who conducts the test and availability of HIV medicine at the testing site)
 - 2) preferred qualities of HIVST kits (cost, test quality, type of test, extra items and support if tested positive).

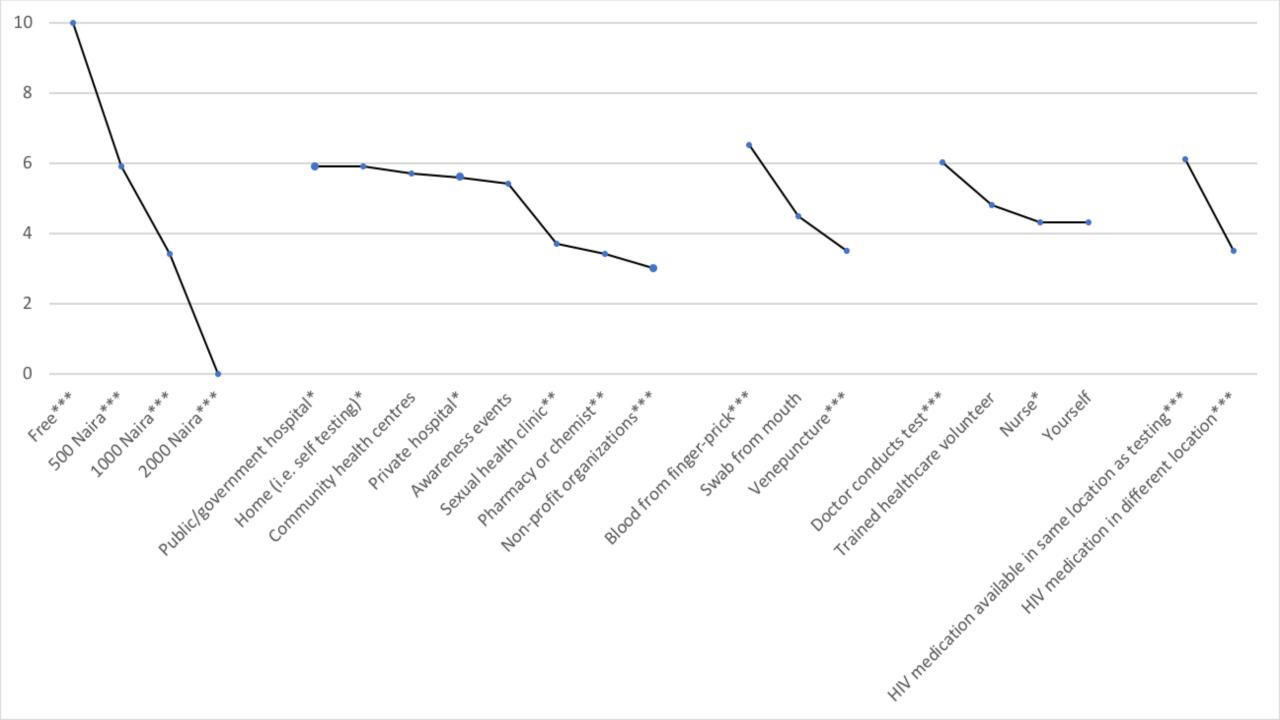
	Option 1
Cost	Free
Test quality	Approved by World Health Organization
Type of test	Swab from mouth
Extra items	Test for other sexually transmitted infections (syphilis, chlamydia, gonorrhoeae)
Support if test positive	Same-day in-person consultation with trained counsellor or youth health worker
Location to get Kit	Pharmacy or Chemist
Your choice	•

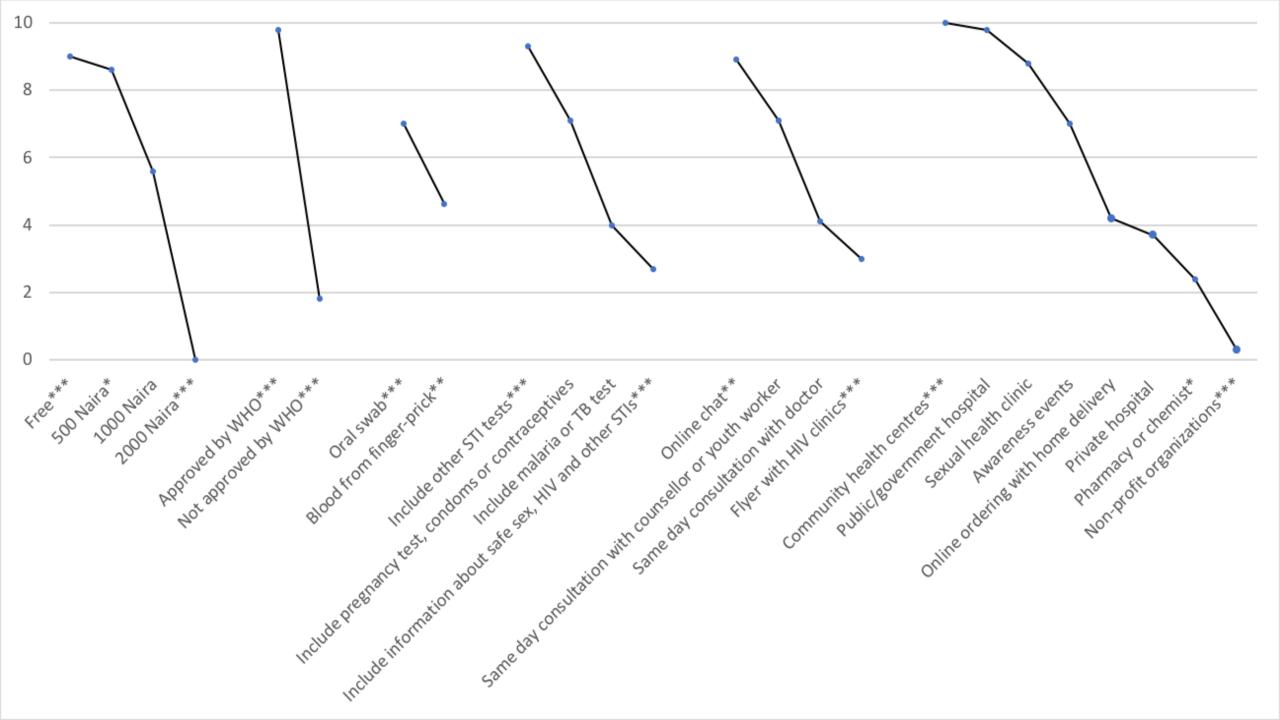
	Option 2	Option 3
	1000 naira	
N	ot approved by the World Health Organization	
	Blood from finger-prick	
	Information about safe sex, HIV and other	I would not test for HIV
	mormation about sale sex, my and other	
	sexually transmitted infections	using option 1 or 2
	Flyer with the list of HIV clinics	
	School	

•

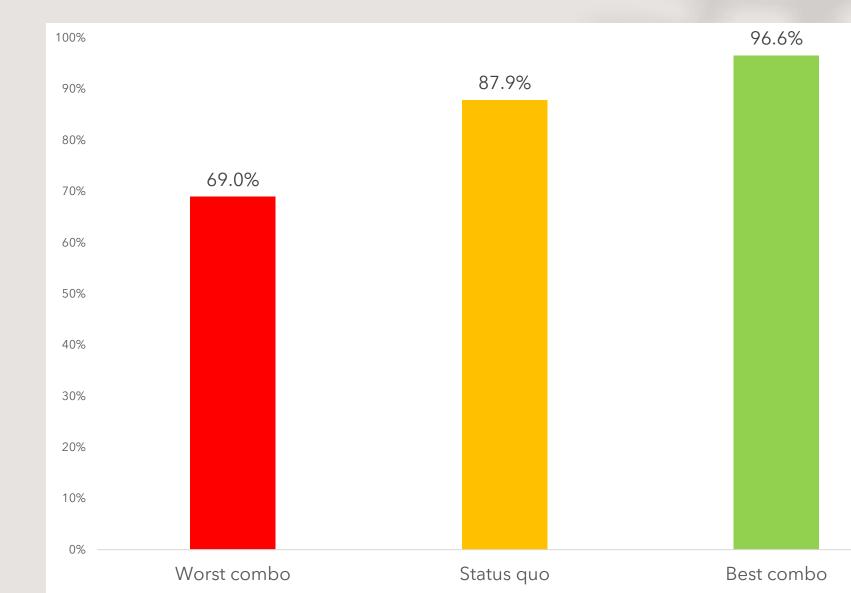
Results

- 504 youth participated
 - mean age 21 (SD 2) years
 - 38% men
 - 35% had higher than secondary school education

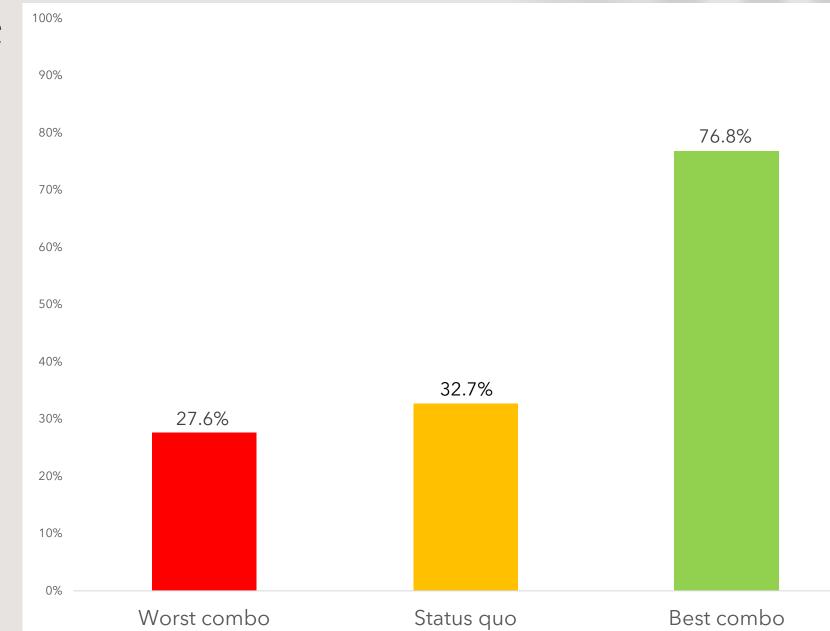




HIV testing uptake



HIVST uptake





THEMED SECTION: APPLICATIONS OF HEALTH PREFERENCES RESEARCH | VOLUME 23, ISSUE 7, P870-879, JULY 01, 2020

The Preferred Qualities of Human Immunodeficiency Virus Testing and Self-Testing Among Men Who Have Sex With Men: A Discrete Choice Experiment

Jason J. Ong, PhD 2 S I Richard De Abreu Lourenco, PhD Deborah Street, PhD ... Martin Holt, PhD John Kaldor, PhD Rebecca Guy, PhD Show all authors

Published: July 17, 2020 • DOI: https://doi.org/10.1016/j.jval.2020.04.1826 •



Check for updates

Aims

• Primary:

• To assess preferences of Australian GBM for HIVST relative to other testing methods, and for how to access HIVST.

• Secondary:

- Assess for **heterogeneous preferences** between subgroups
 - Young vs. older (>25 years)
 - Frequent vs. infrequent testers (>2 years ago or never)
 - Australian-born vs. recent migrants (arrived <5 years)
 - 1 vs. > 1 sexual partners in preceding 6 months

METHODS

Study population

- Australia-wide
- GBM
- 18 years and over
- HIV negative

Online Recruitment

• Weekly Grindr advertisements over 6 weeks with link to online survey

Sexual health clinic recruitment

- Two urban sexual health clinics
- Two community-based organizations



Which of these two options would you prefer?

	HIV self-testing Option 1	HIV self-testing Option 2
Information on how to use the kit	Written instruction leaflet (text and pictures)	Option of having an online chat with peer
Packaging	A small branded package (size of egg carton containing 2 eggs)	A small plain package (size of egg carton containing 2 eggs)
Access	Order online with kits mailed to your home	Kits available from staff of a medical clinic
Cost (out of pocket)	\$60	\$40
Which would you choose?		

DCE 1: HIV TESTING PREFERENCES

Imagine you have decided to have an HIV test. There are a number of different options for testing that may differ based on the following factors:

Attribute	Levels
Cost:	Free; \$20; \$40; \$60
Speed of results:	1 minute; 20 minutes; 1 day; 3 days
Window period (time it takes after a risky event before test shows if you've been infected):	4 weeks; 6 weeks; 12 weeks;
Mode of test:	Venepuncture; oral swab; finger prick
% of tests that are correct:	92%; 95%; 99%; 99.9%
Specimen collected by:	Healthcare worker; yourself; peer

DCE 2: SELF TEST KIT DISTRIBUTION

Imagine you have decided to have an HIV test and have been offered the option of an HIV self-test. Your choice may differ based on the following factors:

Attribute	Levels
Cost:	Free; \$20; \$40; \$60
	Order online; vending machine; pharmacy shelf;
Test Access:	pharmacy staff; medical clinic; community-based organization; sex-on-premises-venues
Packaging:	Large plain; large branded; small plain; small branded.
Information:	Written leaflet; online video; online chat.

Results

•

- 1,606 men participated
 - Mean age 36.1 (SD 11.6)
 - . 15% recently arrived in Australia (<5 years)
 - Median 4 (anal) sex partners in the last 6 months (IQR 2-9)
 - . 15% last test > 2 years ago or never tested

DCE 1: Preference for type of HIV test

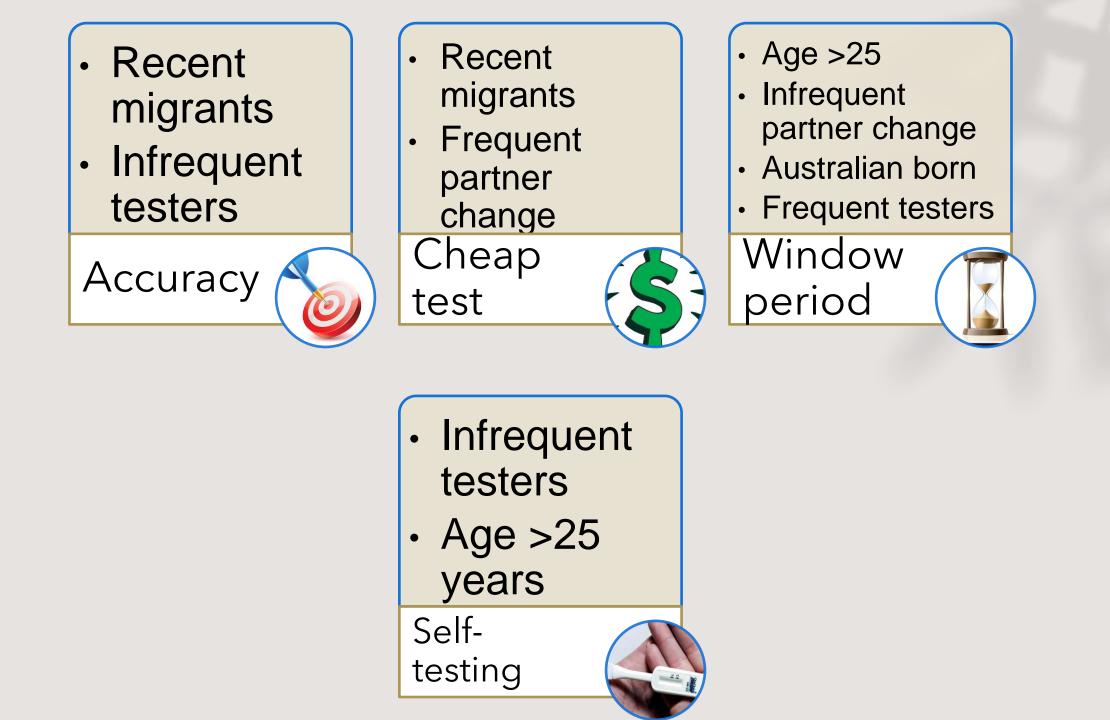
- Short window period (36%) • Self-testing (27%)
- High accuracy (22%)
- Cheap tests (15%)

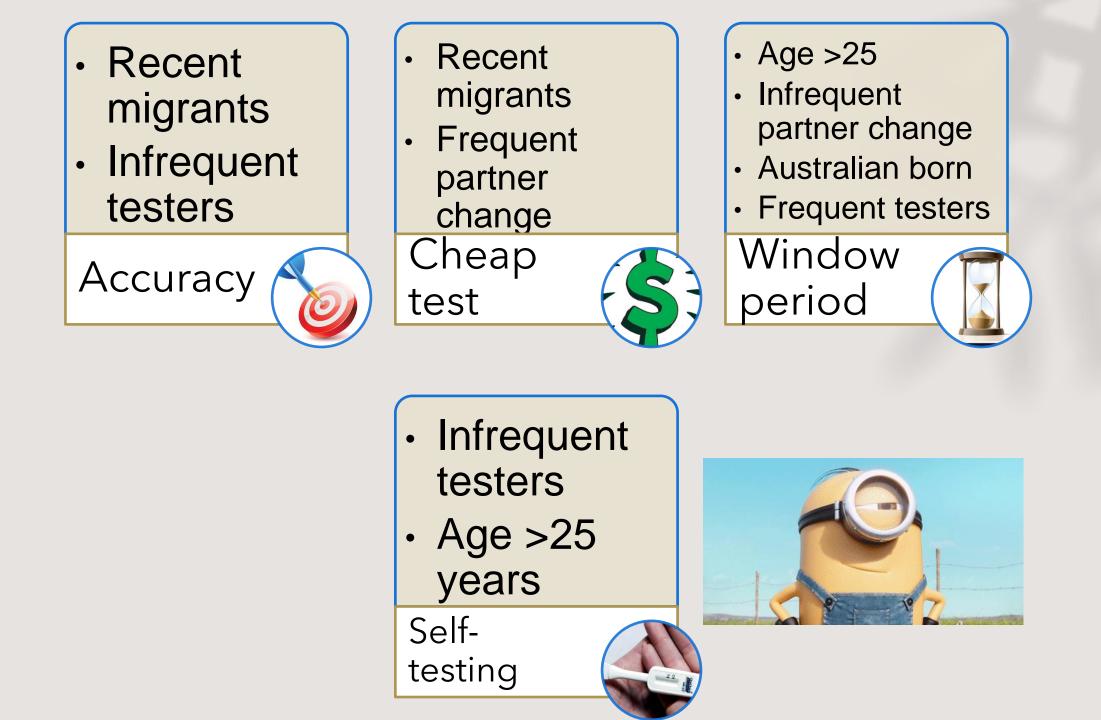


DCE 1: Preference for type of HIV test

Who are they?

- Short window period (36%)
- Self-testing (27%)
- High accuracy (22%)
- Cheap tests (15%)







DCE 2: Self-test kit distribution

- Price-sensitive (45%)
- Retail (Pharmacy or online) (29%)
- Sex-on-premises venues (14%)
- Buy from healthcare staff (12%)

DCE 2: Self-test kit distribution

Retail (29%)
 Online was 4x more
 preferred than pharmacy







Preference for PrEP among Men who Have Sex with Men and Transgender women in 16 Countries in the Asia-Pacific

A Discrete Choice Experiment

W. Tieosapjaroen^{1,2}, B.R. Bavinton³, H.-M. Schmidt^{4,5}, K.E. Green⁶, N. Phanuphak⁷, M. Poonkasetwattana⁸, N.S. Suwandi⁸, D. Fraser³, C. Chan³, M. Cassell⁹, L. Zhang^{1,2}, W. Tang¹⁰, J.J. Ong^{1,2,11}

1 Alfred Health, Melbourne Sexual Health Centre, Carlton, Australia, 2 Monash University, Central Clinical School, Melbourne, Australia, 3 University of New South Wales, Kirby Institute, Sydney, Australia, 4 UNAIDS Regional Office for Asia and the Pacific, Bangkok, Thailand, Global HIV, Hepatitis and STIs Programme, 5 World Health Organization, Geneva, Switzerland, 6 The Program for Appropriate Technology in Health, Hanoi, Vietnam, 7 Institute of HIV Research and Innovation, Bangkok, Thailand, 8 APCOM, Bangkok, Thailand, 9 Family Health International 360, Hanoi, Vietnam,

10 The University of North Carolina at Chapel Hill Project-China, Guangzhou, China,

11 London School of Hygiene and Tropical Medicine, Faculty of Infectious





STUDY SETTING

A cross-sectional survey was created and delivered to MSM in 16 countries in the Asia-Pacific region between May-November 2022.



Gay dating apps

Social media platforms Social media influencers Local MSM/TGW community's mailing lists

STUDY SETTING



- Cambodia
- China
- India
- Indonesia
- Laos
- Malaysia
- Myanmar
- Nepal
- The
- Philippines
- Thailand
- Vietnam

- China SAR Hong Kong
- China SAR Taiwan
- Japan
- Singapore
- Australia

INCLUSION CRITERIA

- MSM and TGW aged over 18 years
- no prior HIV diagnosis
- who lived in the included 16 countries in Asia-Pacific
- who self-identified as gay, bisexual or other men who have sex with men or transgender woman.

FINAL ATTRIBUTES

Type of PrEP

- Daily oral
- On-demand oral
- Injectable
- Long-acting oral
- implant

Service location

- Hospital
- STI clinic
- GP
- Peer-led community clinic
- pharmacy

Cost

- Free
- Low
- Moderate
- High

Side effects

- None
- Interaction with other medications
- Mild
- Rare chance of kidney problems
- Mild pain form injection

Extra services

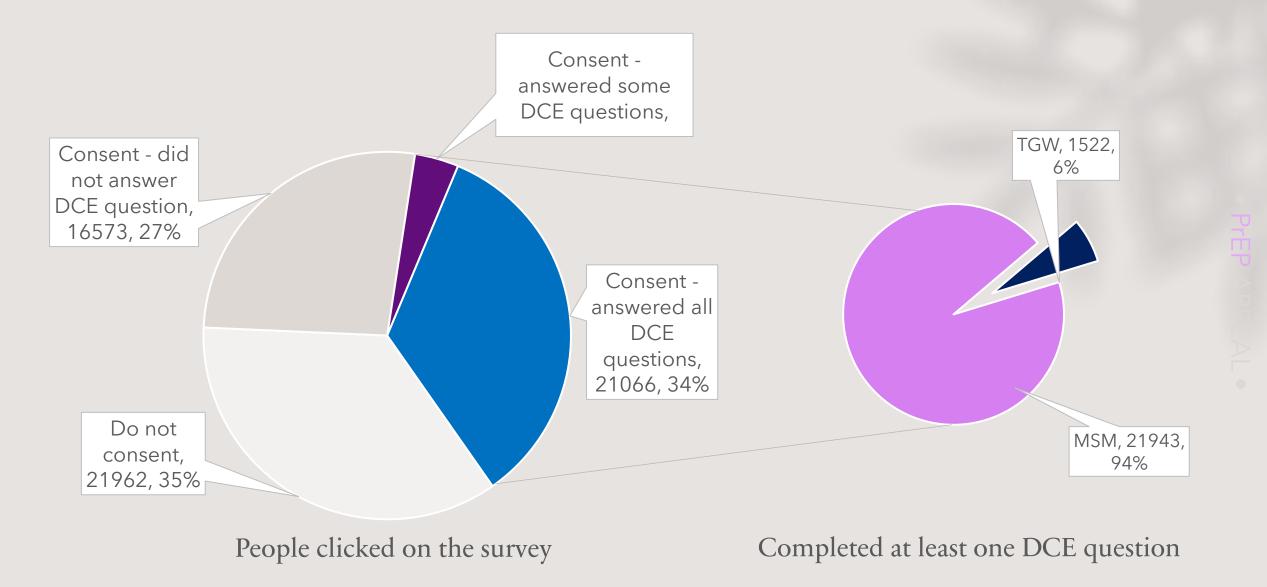
- None
- STI testing
- Mental health counselling

Suppose you want to take PrEP, knowing that it reduces your risk for HIV by 99%, which of the following products do you prefer:

	Α	В	Opt out
Type of PrEP	Oral long-acting PrEP	Injectable PrEP	X
Service location	Pharmacy	Hospital	Х
Cost	\$AU 25	Free	Х
Side effects	Rare chance of kidney problems	Mild	Х
Visit frequency	Every 6 months	Every year	Х
Extra services	STI testing	None	Х
Which choice do you prefer?	0	0	0

PREP APPEAL RESULT

RESPONDENTS INCLUDED



Demographic characteristics

	MSM	% or SD	TGW	% or SD
Ν	21,943		1,522	
Age	32	9.5	28	7.0
Country				
Thailand	1,552	7.1	256	16.8
Vietnam	1,451	6.6	253	16.6
Indonesia	1,428	6.5	80	5.3
The Philippines	2,289	10.4	98	6.4
China	1,853	8.4	28	1.8
Malaysia	1,035	4.7	13	0.9
Myanmar	561	2.6	145	9.5
India	2,768	12.6	145	9.5
Cambodia	821	3.7	127	8.3
Laos	312	1.4	34	2.2
Nepal	459	2.1	343	22.5
Australia	1,894	8.6	0	0
China, Taiwan	2,564	11.7	0	0
Singapore	770	3.5	0	0
China, Hong Kong	646	2.9	0	0
Japan	1,540	7.0	0	0

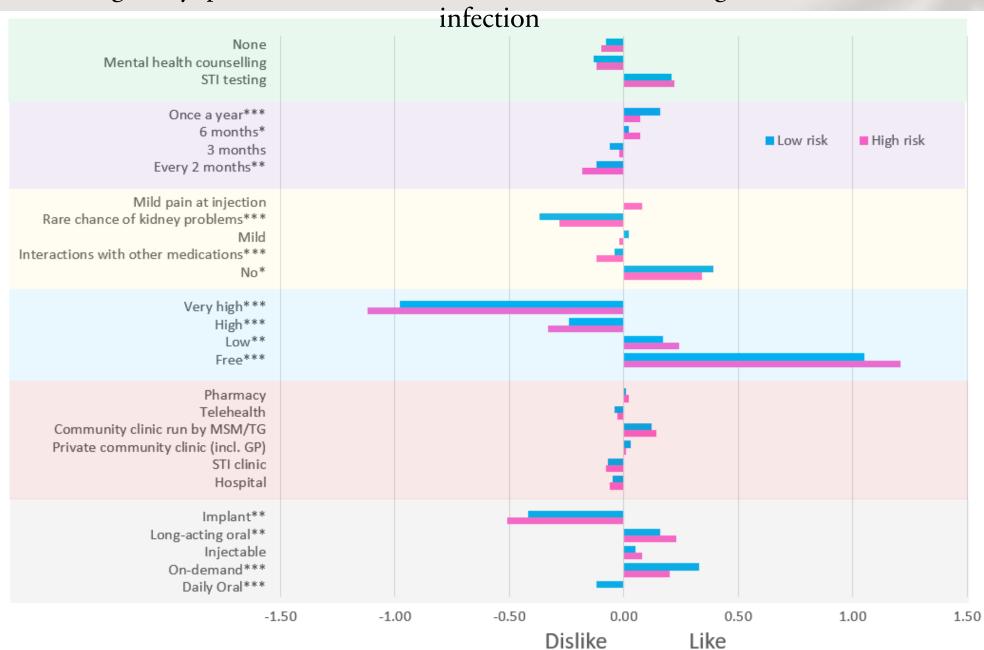
Demographic characteristics

	MSM	% or SD	TGW	% or SD
N	21,943		1,522	
Education level				
No schooling	120	0.6	57	3.8
Up to high school or equivalence	7,515	34.3	984	64.7
At least undergraduate degree	13,372	60.9	460	30.2
Others	936	4.3	3	0.2
Ever taken PrEP				
No	10,923	49.8	450	29.6
Yes	5,892	26.9	697	45.8
missing	5,128	23.4	7	0.5
Currently taking PrEP				
Yes	4,098	18.7	531	34.9
No	1,768	8.1	151	9.9
Missing	16,077	73.3	840	55.2
Sexual partner in the last 6 months				
None	2 1 9 9	10.0	180	11.8
	2,188			
One partner	3,751	17.1	237	15.6
Multiple partners	13,028	59.4	826	54.3
Missing	2,976	13.6	149	9.8

Drivers of choice (relative importance) of PrEP for Men who Have Sex with Men in Asian Countries (N=21,722)

VIETNAM	53.5				25.3	5.3	4.3 8.0	3.8	
THAILAND	49.5		1	4.8	14.4	9	.4 5.0	6.8	
CHINA, TAIWAN	53.9			17.	.3	14.3	6.6 2	.9 5.0	
SINGAPORE	42.7		20.8		14.1	6.8	8.9	6.8	
PHILIPPINES	57.2				15.6	11.5	6.9 4	.6 4.3	
NEPAL	50.1			14.7	9.5	3.4	17.2	5.0	
MYANMAR	28.9	25.0	4	5.8	14.0	10.2	16.2	2	
MALAYSIA	43.7		14.5		18.8	9.2	6.3	7.5	
LAOS	48.7			20.8	5.3	10.0	9.6	5.7	
JAPAN	45.7		14.1		21.9		6.6 6.0	5.7	
INDONESIA	51.6			12.3	18	.1	7.1 7.	7 3.2	
INDIA	41.4		18.9		16.9	8.8	5.9	8.1	
NA, HONG KONG	41.3		21.8		15.6	8.	2 7.3	5.8	
CHINA	57.2				18.3	12.	7 5.1	3.03.7	
CAMBODIA	30.1	23.9		19	9.5	14.5	6.7	4.4	
AUSTRALIA	42.3		17.2		13.2	9.4	9.4	8.5	
ASIA	51.6			16.1		15.0	7.5 4.8	8 5.0	
MIC	52.3			18.3		13.2	7.6 4	.4 4.2	
HIC	50.1			16.3	1	6.8	6.9 4.7	7 5.2	
		RE	LATIVE	IMPO	RTANCE	(%)			
	Cost Type of PrEP	Side e	effects	Extra	services	Locati	on 🗖 Vi	sit frequen	су

Heterogeneity preference for PrEP between individuals at high-vs low-risk of HIV



The uptake of the worst and best PrEP service package among Asian MSM

Country	Scenario	Uptak e (%)	Type of PrEP	Service location	Cost	Side effects	Visit frequency	Extra service
Asian MIC	Worst	71	Implant	Hospital	Very high	Rare chance of kidney problems	Every 2 months	Mental health counselling
	Status quo	95			0			6
	Best	98	On-demand	Community clinic run by MSM/TG	Free	No side effect	Once a year	STI testing
HIC S	Worst	50	Implant	STI clinic	Very high	Rare chance of kidney problems	Every 2 months	Mental health counselling
	Status quo	96			0			U
	Best	99	On-demand	Community clinic run by MSM/TG	Free	No side effect	Once a year	STI testing
Australia	Worst	48	Implant	Hospital	Very high	Rare chance of kidney problems	Every 2 months	None
	Status quo	97			0			
	Best	100	Long-acting oral	Pharmacy	Free	No side effect	Once a year	STI testing

The uptake of the worst and best PrEP service package among Asian MSM

Country	Scenario	-	Type of PrEP	Service location	Cost	Side effects	Visit	Extra service
		e (%)					frequency	
Asian MIC	Worst	71	Implant	Hospital	Very high	Rare chance of kidney problems	Every 2 months	Mental health counselling
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Asian HIC	Worst	50	Implant	STI clinic	Very high	Rare chance of kidney problems	Every 2 months	Mental health counselling
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	Status	97			U	7 1		
	quo Best	100	Long-acting	Pharmacy	Free	No side effect	Once a year	STI testing
			oral					

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	Status	97						
	quo Best	100	Long-acting oral	Pharmacy	Free	No side effect	Once a year	STI testing

Conclusions

- Cost is a major driver of choice for both MSM in Asia-Pacific
- Preference-sensitive PrEP services can improve uptake
 - Community-run clinics
 - Less frequent visits
 - Options beyond daily oral PrEP
- Key differences between countries



Questions?

Session 6 – Small group discussion

Session 6 – discussion

- Share with your group members a possible DCE you could construct within the next 12 months in your area of interest.
- What is your next step?

Next year

- Economic evaluations alongside trials
- Resources already on website
 - Crowdsourcing
 - Designathons







Post-workshop survey





Thank you

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