

The future of health economic evaluation

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In the past seminar events, we learned....

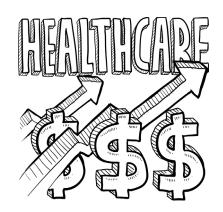
- Introduction to health economic evaluation (June)
- The greatest happiness of the greatest number? Experience use of economic evaluation evidence to inform policy decisions (July)
- Can we be more systematic? The role of MCDA in informing policy decisions (August)
- Can we be more consistent? The issues of the cost-effectiveness threshold (September)
- Ethical issues in health resource allocation (October)



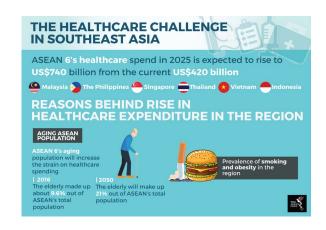
In this seminar, we will learn....















Sustainable Development Goals (SDGs) adopted by UN (Sept 2015): "Ensure healthy lives and promote well-being for all, at all ages" & "end poverty in all its forms everywhere"

▶ Goal 3 is for health

Includes target to: "achieve UHC, including financial risk protection, access to quality essential health care services and access to safe, effective quality and affordable essential medicines and vaccines for all."







health economic evaluation



Return of investment

a ratio
 between the
 net profit and
 cost of
 investment
 resulting from
 an investment
 of some
 resources.





Appraisal of ranibizumab (Lucentis) for diabetic macular oedema 2011

- Additional costs = £3,506 per patient with 23,000 eligible patients each year
- Opportunity cost threshold in the UK is £13,000 per QALY
- Incremental cost-effectiveness = £25,000 per QALY

Attributes	Investment
	Lucentis for diabetic macular oedema (£80m pa)
QALYs	3,225
Wider social benefits (net production)	£88.4m

Benefit-cost ratio = 88.4/80

= 1.105

But this is not the best option when considering opportunity costs

This slide was modified from the presentation of Prof. Karl Claxton, University of York



Appraisal of ranibizumab (Lucentis) for diabetic macular oedema 2011

- Additional costs = £3,506 per patient with 23,000 eligible patients each year
- Opportunity cost threshold in the UK is £12,936 per QALY
- Incremental cost-effectiveness = £25,000 per QALY
- Per capita GDP = £ 42,000

Attributes	Investment	Opportunity costs	
	Lucentis for diabetic macular oedema (£80m pa)	Expected effects of £80m pa	
QALYs	3,225	- 6,184	
Wider social benefits (net production)	£135 m	- £260m	

Benefit-cost ratio = 88.4/80

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Appraisal of ranibizumab (Lucentis) for diabetic macular oedema 2011

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Attributes	Investment	Opportunity costs	Opportunity cost of using Avastin	Net effect of using lucentis
	Lucentis for diabetic macular oedema (£80m pa)	Expected effects of £80m pa	Expected effects of £2m pa for avastin	
QALYs	3,225	- 6,184	-155	3,070
Wider social benefits (net production)	£135m	- £260m	£6m	£129m

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Methodological development

- Incorporating equity dimension
- Using real world data
- Automatisation of health economic evaluation

Applications

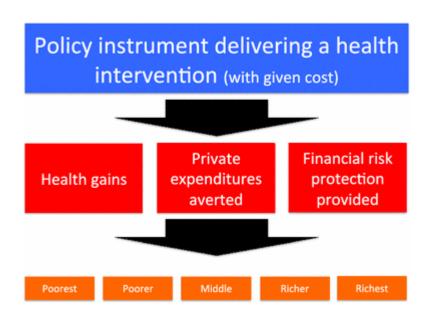
- HTA for developing health target/ goal setting
- Early HTA

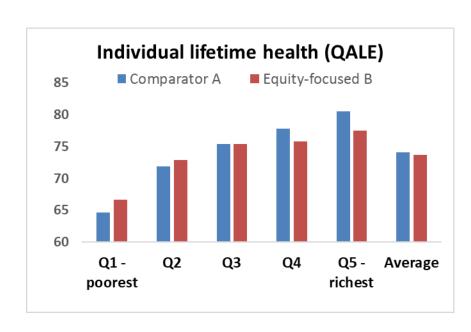
Policy

- Using HTA for price interventions
- Joint assessment or uniform HTA process



Efficiency-equity trade-off





Verguet et al, Cookson et al



What do we already know about RWD use?

- Many advantages of using RWD e.g. getting effectiveness (not efficacy),
 overcoming ethical barriers in conducting RCTs, reflecting real-life situation etc.
- Increasing potential for using RWD due to better establishment of digital health care systems
- The need for establishing **good process**, including <u>transparency</u> (the rationale for using RWD and the RWD used must be publicly accessible), <u>relevance</u> (there must be a reasonable explanation for a decision's rationale) and <u>fairness</u> (the RWD is used in similar fashion across technologies). Processes should also allow opportunity for <u>stakeholder participation</u>.

Garrison Jr LP, Neumann PJ, Erickson P, Marshall D, Mullins CD. Using real-world data for coverage and payment decisions: The ISPOR real-world data task force report. Value in health. 2007 Sep;10(5):326-35.

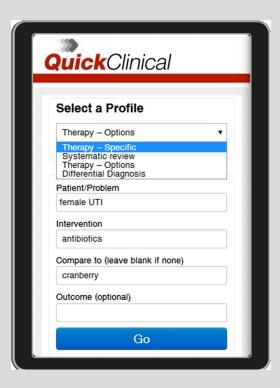
Berger ML, Lipset C, Gutteridge A, Axelsen K, Subedi P, Madigan D. Optimizing the leveraging of real-world data to improve the development and use of medicines. Value in Health. 2015 Jan 1;18(1):127-30.



Myth and facts of using RWD

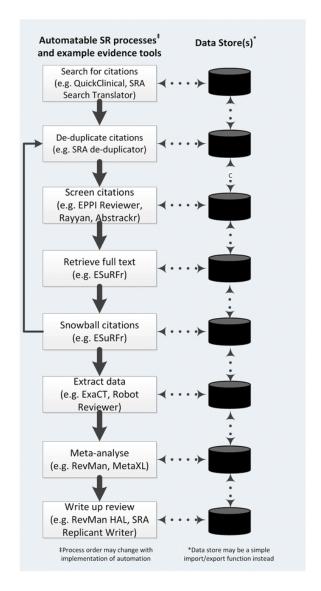
The myth	The facts
RWD is free	RWD needs serious investment
RWD is an alternative to traditional RCTs or experiment studies	RWD should be used to complement data from experimental studies
Only RWD can answer real world questions	Many careful designed experiments can answer real world questions Some studies using RWD cannot answer real world questions
Bias can be technically corrected	It's not possible to correct all biases

Automatisation of systematic review and meta-analysis



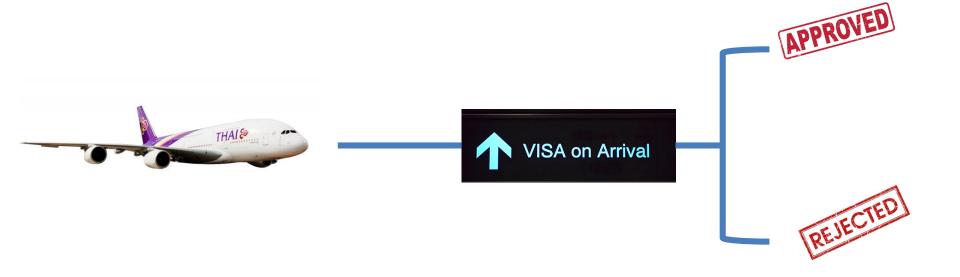
- Beller et al 2018. Making progress with the automation of systematic reviews: principles of the International Collaboration for the Automation of Systematic Reviews (ICASR)
- Tsafnat et al 2014 Systematic review automation technologies



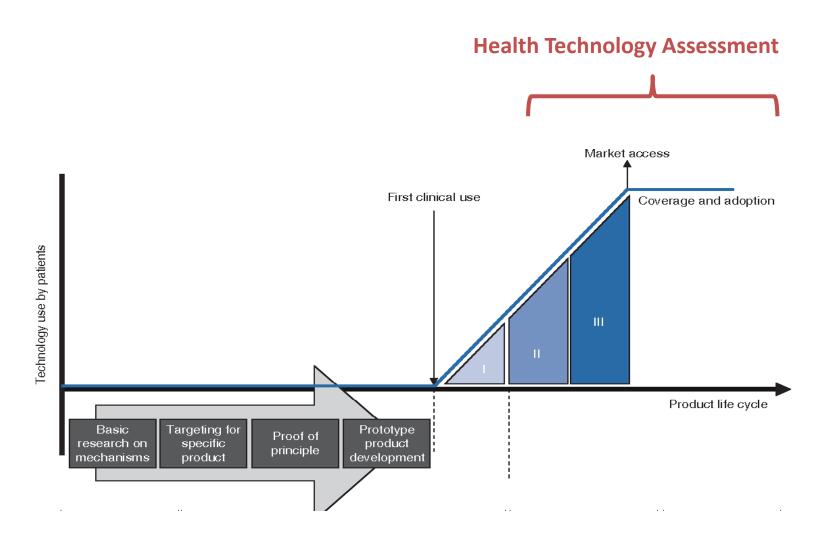




Early HTA or HTA for R&D









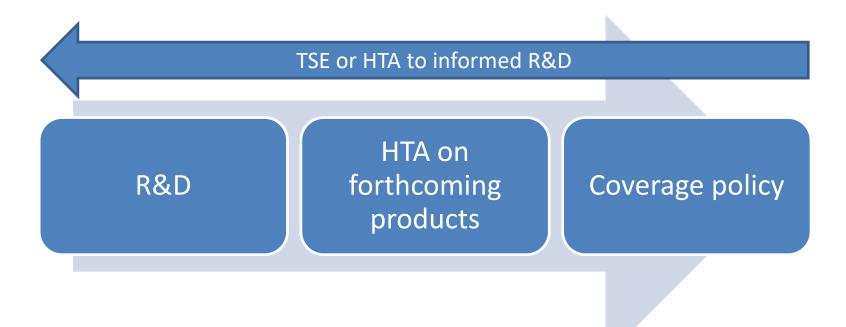
Traditional HTA

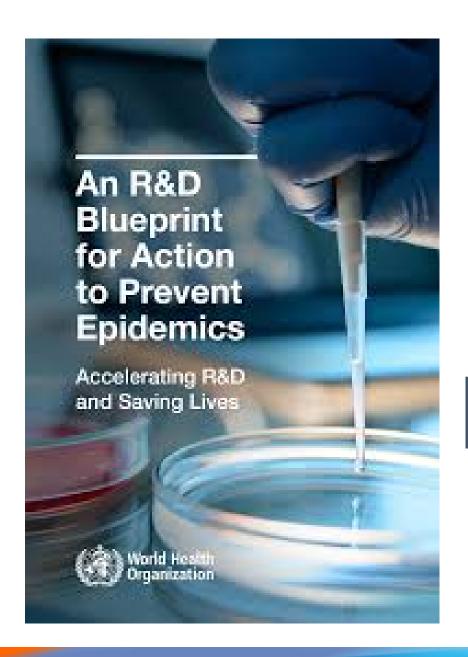
R&D

HTA on available products

Coverage policy



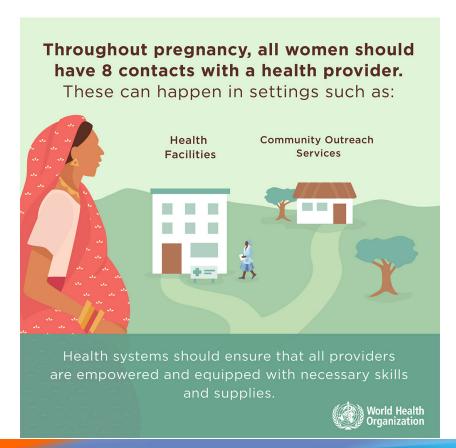


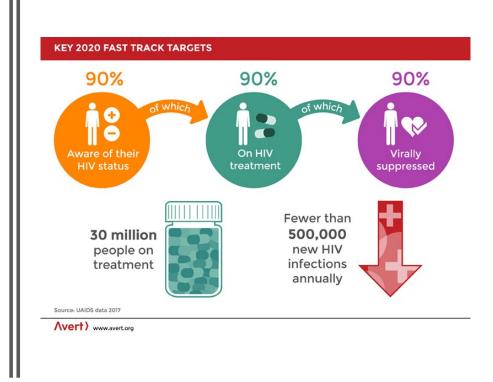




Thailand country pilotMarch – September 2018
DRAFT as of 08/03/18

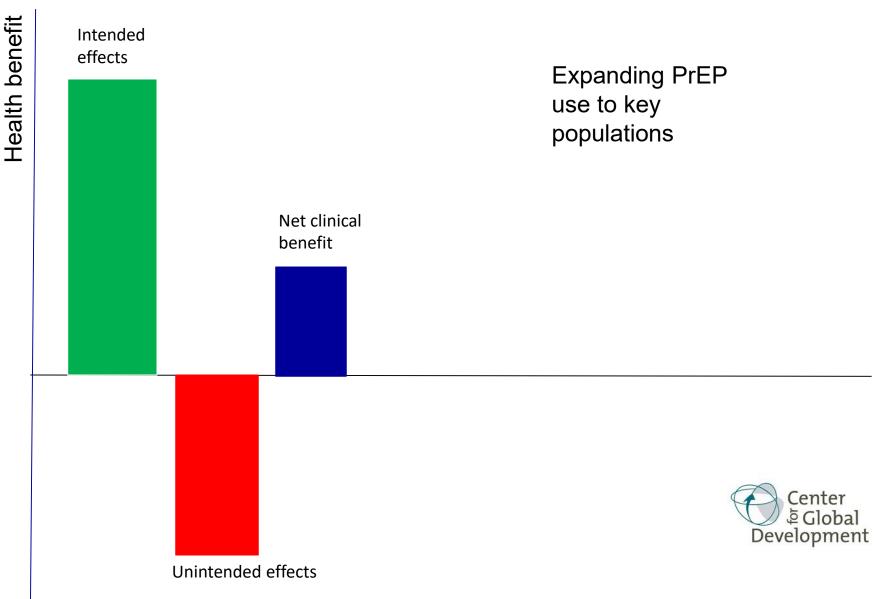
Setting the guidelines or global target without considering 'opportunity cost' can make more harm than good





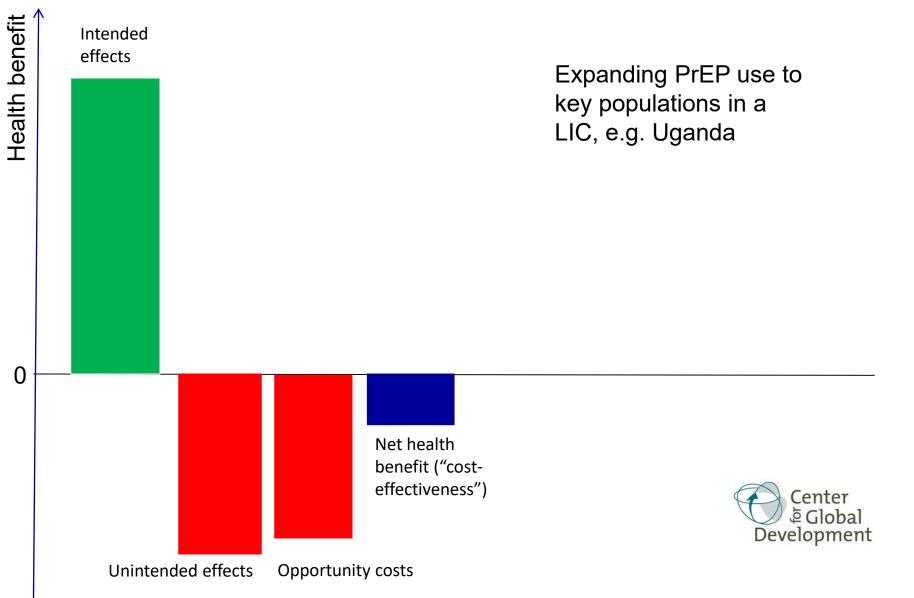
II. Understanding value in health care





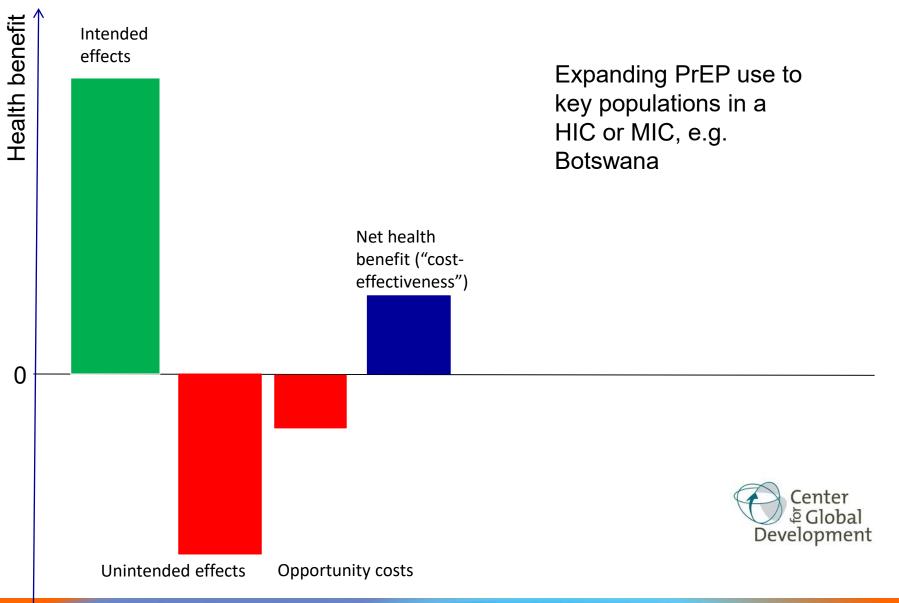
II. Understanding value in health care





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ngthening EU cooperation beyond

the European Commission started work on strengthening EU cooper tries, the European Parliament, and interested parties to ensure in Commission announced that this would extend to improving the fu

islative proposal

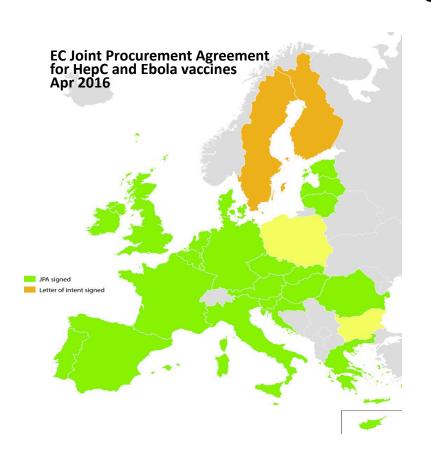
tive proposal was adopted by the European Commission on 31 Janu 3 the results of the impact assessment outlined below. It has been see 1 by 2019. The proposal and related information can be found here:

strengthening EU cooperation on Health Technology Assessment in response to calls from EU countries

- On 4 October 2018, the European Parliament adopted its Report on the Commission Proposal for a Regulation on Health Technology Assessment (HTA)
- The proposal seeks to ensure that when HTA is performed, the methodologies and procedures applied are more predictable across the EU and that joint clinical assessments are not repeated at national level, thereby avoiding duplication and discrepancies. The Report recognises that current approaches lead to "higher costs for industry, delays in access to technologies and a negative effect on innovation
- Concerns have been raised by some EU Member States that joint clinical assessments might lead to the loss of control of a Member States' ability to decide on prices. Deciding on reimbursement and pricing of medicinal products is a national competence of the Member States
- The Director General considers, however, that the HTA Regulation is foremost a matter of defining on an EU level which evidence to expect from innovator companies: "Clinical assessments you can do across Europe, cost effectiveness has to be done country by country."



As poorer countries graduate from purchasing clubs... rich countries come together to form new ones!











See you in Seoul!

