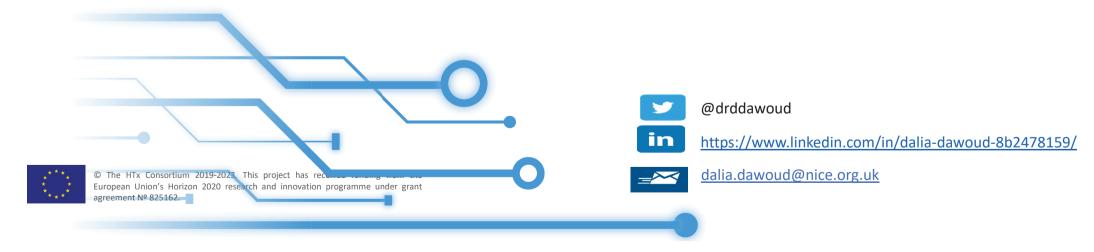


Future of Health Technology Assessment

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Disclaimers

- No Conflicts of Interest to declare
- Views expressed are my own and not those of NICE



HT Seneration Health Technology Assessment

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Outline

- Health Technology Assessment (HTA)
 - What, Why and How?
 - Current status and need for change
- Next Generation Health Technology Assessment (HTx) project
 - Project outline
 - Focus areas
 - Key outputs



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Outline



What is Health Technology Assessment (HTA)?



Health technology assessment

HTA Definitions

There are many definitions of HTA. All emphasize its role as a tool supporting decision making at different level of the healthcare system, its multidisciplinary nature and its strong reliance on transparent scientific rigorous methods.

Scientific

rigorous

Decision-making





Definition

 HTA is a multidisciplinary process that uses explicit methods to determine the value of a health technology at different points in its lifecycle. The purpose is to inform decision-making in order to promote an equitable, efficient, and high-quality health system.

International Journal of Technology Assessment in Health Care

cambridge.org/thc

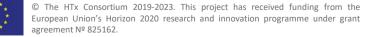
The new definition of health technology assessment: A milestone in international collaboration

Brian O'Rourke¹, Wija Oortwijn² ⁽ⁱ⁾, Tara Schuller³ ⁽ⁱ⁾ and the International Joint Task Group

https://pubmed.ncbi.nlm.nih.gov/32398176/







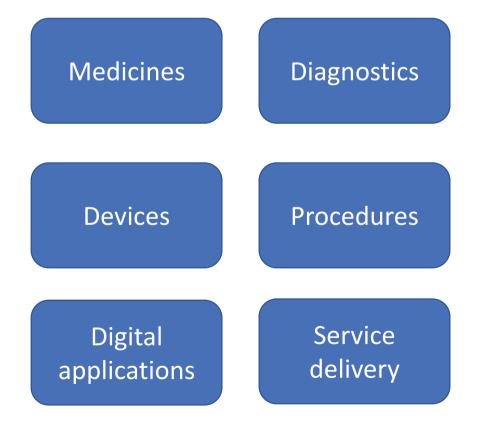
Health Technology

A health technology is an **intervention** developed to **prevent**, **diagnose** or **treat** medical conditions; **promote** health; provide **rehabilitation**; or **organize healthcare delivery**.

The intervention can be a **test**, **device**, **medicine**, **vaccine**, **procedure**, **program** Or **system**. (Definition from the HTA Glossary)







HTA process

The process is **formal**, **systematic** and **transparent**, and uses **state-of-the-art methods** to consider the **best available evidence**.







Developing technology appraisal guidance

An overview of the development process.

1. F	Provisional appraisal topics chosen	*
2.0	Consultees and commentators identified	~
3.5	Scope prepared	~
4./	Appraisal topics referred	*
5. E	Evidence submitted	*
6. E	Evidence Review Group (ERG) or assessment report prepared	~
7.0	Committee papers prepared	*
8.7	Appraisal committee	*
9.1	Appraisal consultation document (ACD) produced	*
10	Final appraisal determination (FAD) produced	*
11.	. Guidance issued	~



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NICE National Institute for Health and Care Excellence	NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE			
	Guide to the processes of technology appraisal			
	April 2018 Acknowledgements NICE is very grateful to everyone who contributed to the development of this guide (see section 7). Foreword The National Institute for Health and Care Excellence (NICE) provides guidance to the NHS in England on the clinical and cost effectiveness of selected new and established technologies. NICE carries out appraisals of health technologies at the request of the Department of Health and Social Care. Guidance produced by NICE on health technologies is also applied selectively in Northern Ireland and Wales. This document is one of a series describing the processes and methods that NICE uses to carry out technology appraisals. It focuses on the technology appraisal processes (and provides an overview for organisations invited to contribute to an appraisal).			
Guide to the methods of technology appraisal 2013				
Process and methods	The documents in the series are: Guide to the processes of technology appraisal (this document). <u>Guide to the methods of technology appraisal.</u> <u>Cancer Drugs Fund technology appraisal process and methods (addendum).</u>			
Published: 4 April 2013 nice.org.uk/process/pmg9				
Home Contact v Training v Appraisal Specific v Methods Devel				
Technical Support Documents	Search 9			



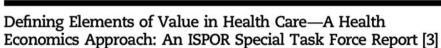
Value

- The dimensions of value for a health technology may be assessed by examining the intended and unintended consequences of using a health technology compared to existing alternatives.
- These dimensions often include clinical effectiveness, safety, costs and economic implications, ethical, social, cultural and legal issues, organisational and environmental aspects, as well as wider implications for the patient, relatives, caregivers, and the population.
- The overall value may vary depending on the perspective taken, the stakeholders involved, and the decision context.









CrossMark

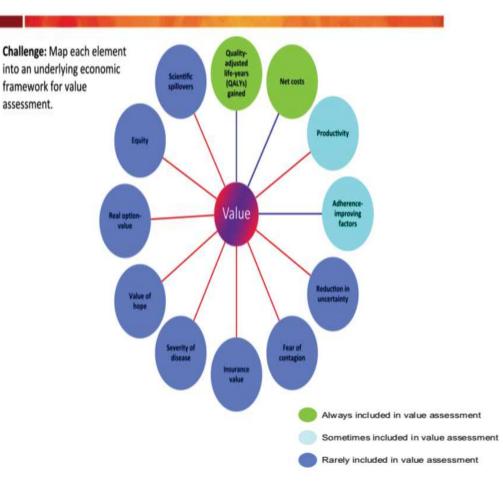
Darius N. Lakdawalla, PhD^{1,*}, Jalpa A. Doshi, PhD², Louis P. Garrison Jr, PhD³, Charles E. Phelps, PhD, MBA⁴, Anirban Basu, PhD³, Patricia M. Danzon, PhD⁵

¹Schaeffer Center for Health Policy and Economics, University of Southern California, Los Angeles, CA, USA; ²Division of General Internal Medicine, University of Pennsylvania, Philadelphia, PA, USA; ³The Comparative Health Outcomes, Policy, and Economics (CHOICE) Institute, University of Washington, Seattle, WA, USA; ⁴Economics, Public Health Sciences, Political Science, University of Rochester, Gualala, CA, USA; ⁵The Wharton School, University of Pennsylvania, Philadelphia, PA, USA



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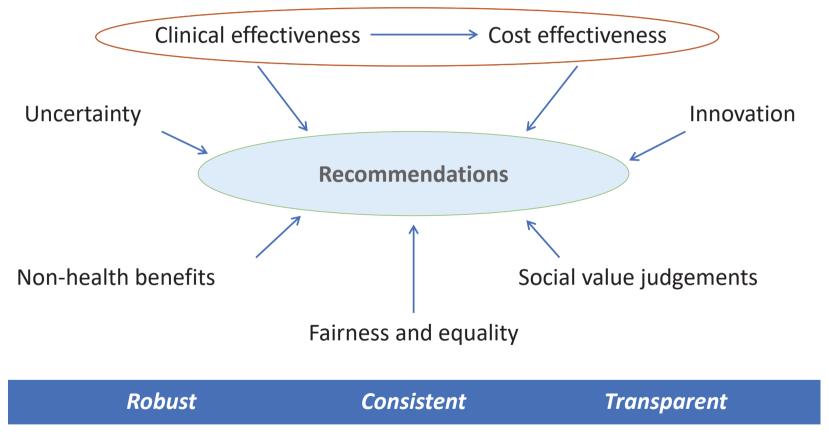
Elements of Value



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Timing

 HTA can be applied at different points in the lifecycle of a health technology, i.e. premarket, during market approval, post-market, through to the disinvestment of a health technology.

	Preclinical research	Clinical research			Market authorization	
		20–100 volunteers	100–500 volunteers Phase II		Market acces	
		Phase I			Phas IV	9
Basic research	Discovery research	Dev	Development research			eting)
	< 3–6 years	Î				0.5–1.5 years
5,000-10,0	000 compounds 250	5		1		

Figure I Schematic representation of the drug development process with timeline, attrition rate, and sample sizes of clinical studies. Notes: Timing of different stages and sample sizes vary according to different countries, manufacturers, and indications. Reprinted from *Drug Discov Today*, 17. van Nooten F, Holmstrom S, Green J, Wiklund I, Odeyemi IA, Wilcox TK. Health economics and outcomes research within drug development: challenges and opportunities for reimbursement and market access within biopharma research. 615–622. © 2012, with permission from Elsevier.¹⁰



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https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4234281/

Why?

To ensure the best use of the scarce healthcare system resources

- No health care system in the world can provide every effective intervention. Resources are limited and wants are limitless (Scarcity)
- If you provide more of one service, you have to provide less of another. (**Opportunity cost**)
- Choices and trade-offs have to be made.







HTA Recommendations

- Single intervention
- Target population level



Clinical effectiveness based on RCT evidence

Methods development







HTx: Next Generation Health Technology Assessment

Horizon 2020 project funded by the European Union (2019-2023)





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HTx 2nd General Assembly Bern, 6-7 February 2020

Next Generation HTA (HTx)

- To facilitate the development of methodologies to deliver more customized information on the effectiveness and cost-effectiveness of complex and personalised combinations of health technologies.
- To provide methods to support personalised treatment advice that will be shared with patients and their physicians



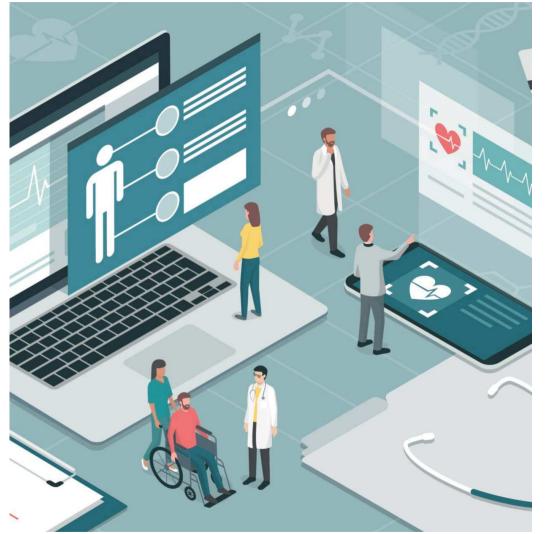
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Next generation

Health Technology Assessment

Patient-centred, societally oriented, real-time decision-making on access to and reimbursement for health technologies throughout Europe.



The HTx vision: a new generation of healthcare decision-making



Imagine an individual patient who visits the doctor for a medical problem. The her use of different health technologies, such as medical devices, e-health outcomes.

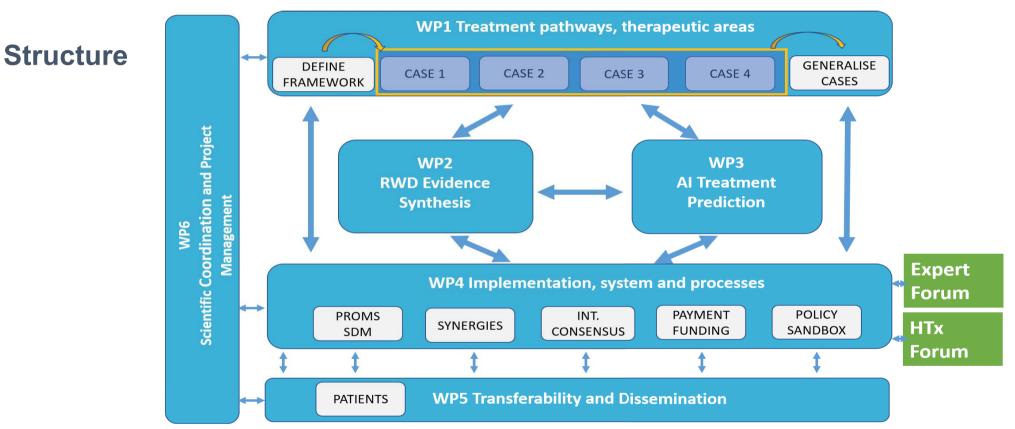


Adequate clinical studies and real-world data analysis have resulted in a **realpatient can use to obtain person-centered information** (in a user-friendly **associated with a range of possible strategies to manage the patient's ailment.**



The same information is made available to HTA agencies whose decisions are i level of individuals and summarised at the subgroup and population level for t is what we envision as HTx.







Focus areas

- Prediction modelling on the basis of data using different study designs (RCT, RWD etc) (WP2)
- Health-econometric tools to take into account effects and costs (WP2)
- Develop AI/ML methods to forecast individual patient treatment outcomes (WP3)

Focus on combinations (and/or sequences) of health technologies not evaluated in RCT as such





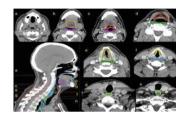


Focus areas

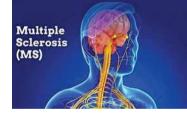
- Proton Therapy for head and neck cancer
- Monitoring and treatment pathways in **diabetes** (T1DM and T2DM)
- **Pharmacological treatments** for relapsing remitting **multiple sclerosis** (MS)
- Different treatment modalities in patients with myelodysplastic syndrome (MDS)

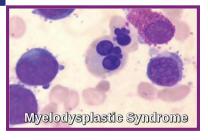














Outputs

- Clear methods developed for these disease areas:
 - Are **practically used** in healthcare practice
 - **By HTA organisations** to facilitate HTA for personalised treatments (including support appropriate use);
 - By healthcare providers as part of new guidelines For individual patients and their clinicians
 - Provides a general framework that can help other groups to develop methods for specific disease areas
 - Has a clear link to national reimbursement and pricing processes





Concluding remarks

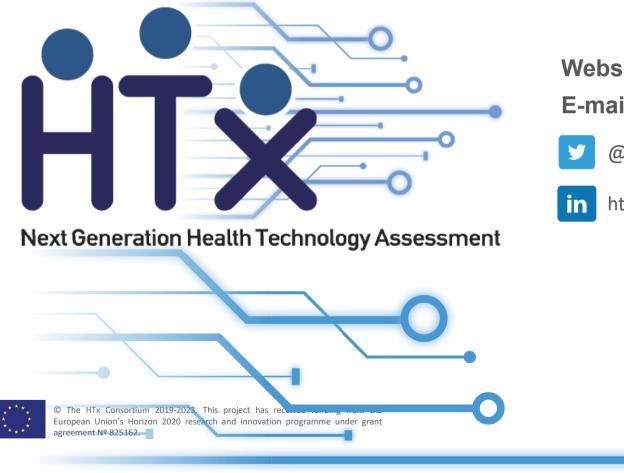
- Next generation HTA will increasingly consider:
 - Combinations and sequences of treatments
 - Individualised treatments using AI-based prediction models and genetic testing
- Methods' development is key to future-proofing HTA



 $[\]ensuremath{\mathbb{C}}$ The HTx Consortium 2019-2023. This project has received funding from the European Union's Horizon 2020 research and innovation programme under grant agreement Nº 825162.

Thank You!





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