





Getting better with evidence: Experiences of putting evidence into practice

10 December 2019 Auckland Medical Research Foundation Auditorium Grafton Campus, University of Auckland

PROGRAMME

Time	
8:00am	Registration opens
9:00am	Mihi whakatau Rawiri Wharemate (Ngāti Wai, Ngāpuhi, Te Kawerau a Maki), kaumātua
9:10am	Introduction and welcome Dr Ashley Bloomfield, director-general and chief executive, Ministry of Health
9:20am	Welcome from the hosts Professor Alan Merry, deputy dean, Faculty of Medical and Health Sciences, University of Auckland

9:30am	Keynote: Gathering and disseminating better evidence 20 minute presentations, followed by a 30-minute panel discussion			
	The path to independence: creating more trustworthy evidence	A critical lens on knowledge production	Harnessing the power of large national administrative datasets to create	
	 Fiona Godlee, editor-in-chief, British Medical Journal, London, UK A universal human aspiration is to use the most trustworthy evidence to inform our decisions about health care. Yet as the landmark Institute of Medicine report on conflicts of interest in medical research, education and practice highlighted, extensive industry influence may be jeopardising the integrity of scientific investigations, the objectivity of medical education, the quality of patient care and the public's trust in medicine. As the World Health Organization has observed, there is an 'inherent conflict of interest between the legitimate business goals of manufacturers and the social, medical and economic needs of providers and the public' It is time to ensure that scientific evaluation of tests and treatments, and communication and use of the resulting evidence, are conducted as independently as possible from industries that profit from their use. This talk will explore routes to greater independence from industry, calling on existing examples and ideas for how we can create a better, more trustworthy evidence base for health decisions. 	 production Prof Papaarangi Reid, tumuaki, deputy dean Māori, Faculty for Te Kupenga Hauora Māori (TKHM), University of Auckland Tēnā koutou katoa. Knowledge production systems are age-old and firmly bound in culture and contexts. Today we use words like science, knowledge and evidence uncritically and without acknowledging their theoretical foundations, and those of the institutions that generate them. In a conference that focuses on 'getting better with evidence' where many speakers will focus on methods, analytics, integration, utility and engagement, there must also be a space to question and debate theoretical issues. Do we dismiss knowledge from other knowledge production systems and what are the risks? Are there skeletons in the closet of westernised science and how might they be impacting on our work today? Do we produce evidence but wilful inaction? As we embrace 'big data' and 'artificial intelligence', do we understand that algorithms produced in a racist and biased society will be racist and biased? Once we know these things, what are our ethical responsibilities? Let's talk about these issues. Mauri Ora! 	 administrative datasets to create evidence and develop policy Prof Matthew Parsons, clinical chair in gerontology, Te Huataki Waiora Faculty of Health, University of Waikato Data science is rapidly emerging as an effective and powerful method of understanding complex problems and identifying solutions in health care. This presentation uses three case studies spanning 10 years where large data sets have been used to develop innovative health care models. The presentation will also focus on the use of classification arising from big data analytics to develop and implement alternative funding methodologies, namely case-mix across the health care continuum. Other benefits of big data analytics will also be explored, including the potential replacement of the randomised controlled trial. 	
11:00am	Morning tea			

 Getting to the heart of the evidence Catherine Marshall, independent health and guideline adviser, and consumer advocate People involved in the production of systematic reviews, guidelines and other evidence-based tools are usually passionate about their work and believe their evidence-based tools are usually passionate about their work and believe their evidence-based tools are usually passionate about their work and believe their evidence-based tools are usually passionate about their work and believe their evidence value about their work and believe their evidence based tools are usually passionate about their work and believe their evidence based tools are usually passionate about their work and believe their evidence based tools are usually passionate about their work and believe their evidence based tools are usually passionate about their work and believe their evidence based tools are usually passionate about their work and believe their evidence based tools are usually passionate about their work and believe their evidence based tools are usually passionate about their work and believe their evidence based is the systematic review. So whet factors other than 'scientific' evidence need to be taken into account? what factors other than 'scientific' evidence need to be taken into account? what is the special x-factor that can make evidence relevant to the everyday lives of consumers? Catherine will discuss both personal and policy approaches using examples from Catherine will discuss both personal and policy approaches using examples from Catherine will discuss both personal and policy approaches using examples from Catherine will abourds are using examples from Catherine will abourds and policy approaches for a being approaches using examples from Catherine will abourds approaches using examples from Catherine will abourds and provende that has attempted to estimate its
 and guideline adviser, and consumer advocate People involved in the production of systematic reviews, guidelines and other evidence-based tools are usually passionate about their work and believe their evidence will improve the quality of care of patients and consumers. Evidence based advice that is easy to understand and follow? what factors other than 'scientific' evidence need to be taken into account? what factors other than 'scientific' evidence need to be taken into account? what factors other than 'scientific' evidence relevant to the everyday lives of consumers? Catherine will discuss both personal and Catherine will
 policy approaches using examples from her own health care journey and her quest for evidence-based advice, along with evidence implementation projects designed to resonate with consumers. The presentation will also explore what might be driving this problem, such as broadening disease definitions and changes in diagnostic technology – often used in screening programmes – which can identify eversmaller abnormalities, many of which will highly likely never to go on to cause

1:00pm	Lunch			
1:45pm	Concurrent seminars			
	Room 505-007	Room 501-010		
	Implementation in a difficult environment	Public understanding of evidence		
	Medicine, University of Otago I Dr Rees Tapsell, director of clinical services, Mental Health & I Addictions Service Executive clinical director, PUAWAI: Midland I Regional Forensic Psychiatric Service I There is substantial information demonstrating that the provision of health care contributes to Māori health inequities through the underuse of evidence-based best practice and the overuse of inferior treatments. I Using examples from primary care and mental health we will I	Catherine Marshall		
		Deon York, programme manager – consumer engagement, Health Quality & Safety Commission		
		Louise Malone, committee member, Breast Cancer Aotearoa Coalition		
		Progressing towards a public understanding of evidence is more likely if true partnership and engagement with the development, production, and use of evidence takes place.		
		This session will focus on these elements, comparing and contrasting practical national and international examples of consumer engagement with developing, generating, presenting and using evidence to ultimately improve the quality of care.		
	difficult to implement evidence in these environments, and discuss potential solutions to these challenges.	This interactive session will explore questions including:		
		• Why is it important for consumers and communities to engage with health data?		
		 How can consumers contribute to the development of evidence? 		
		 What does the system need to provide to consumers for them to effectively use evidence for themselves and their whānau? 		
		 How can experience-based evidence be better curated and made available to consumers? 		
		How can consumers influence the research agenda with what matters to them?		
		 How can consumer engagement with evidence influence national policy and the delivery of quality health care? 		

	Room 503-028	Room 503-020	
Real-world appraisal		Shared decisions about medicines: the intersection between	
	Carl Heneghan, general practitioner and director of the Oxford Centre for Evidence-Based Medicine	people's preferences and evidence Jeff Harrison, head of school, pharmacy, University of Auckland	
	The growing harms caused by drugs and devices, the escalating costs of health care and the pace new technologies emerge means we need to be more effective in how we use evidence to inform health care.	This session will explore the identification, communication and use of best evidence in reaching shared decisions about the use of medicines. Health consumers (people) have a right to be fully informed, make informed choices about their treatment, enshrined in New Zealand in	
	How should we inform services; how should we improve post- marketing surveillance; do we need mandatory registries for all implantable devices? Some of the evidence gathering questions we face.	the Health & Disability Commissioner Code of Rights 6 and 7. The Code sets out that, among other things, people have the right to an explanation of the options available, including an assessment of the expected risks, side effects, benefits, and costs of each option.	
	Randomised trials have been the gold standard for evidence of effectiveness, but there is often a mismatch between the questions, the populations and the outcomes in the real world.	As health care practitioners and health consumers, how do we gather evidence on the benefits and harms of treatment? How do we incorporate people's beliefs about medicines and their preferences for	
	In this seminar, Professor Heneghan will discuss the types of evidence we require to aid decision making, what is working and what is not, and how can we contribute to better informing patients in the real-world?	treatments, including their cultural beliefs? How do we help patients make the trade-off between harms and benefits where the impacts may be unequal? How do we manage our own implicit biases, and respond when we are asked or feel the need to promote the 'right' choice?	
		For all this and a toolkit to address some of these problems in practice, you'll need to come along.	
3:15pm	Afternoon tea		

3:30pm	Keynote: Building solutions – the roles of big data, randomised controlled trials and real-world evidence 20-minute presentations, followed by a 30-minute panel discussion			
	Building solutions: avoiding harm and ensuring equity benefits for Māori	If not the randomised controlled trial, why not and then what?	Reliable evidence: The roles of real- world data, patient evidence and	
	Assoc Prof Sue Crengle The use of evidence obtained from clinical trials and real-world research has the potential to improve Māori health and reduce inequities in health, but this potential has yet to be fully realised. The use of big data in health and other sectors has the potential to do harm. This presentation reflects on both of these issues, and considers strategies to ensure that potential benefits are realised and harm avoided.	 Assoc Prof Rachael Parke, School of Nursing, University of Auckland; nurse senior research fellow, cardiothoracic and vascular intensive care unit, Auckland DHB Randomised controlled trials have long been considered the gold standard for clinical research. Why do promising new strategies leading to improved patient outcomes in single-centre randomised controlled trials fail to be replicated in large, definitive, multi-centre trials? Pivotal large-scale multi-centre trials in fields such as fluid resuscitation, sepsis and renal failure, have been conducted by experienced investigator-led clinical trials groups. However, few such trials undertaken in the setting of intensive care demonstrate improved patient outcomes. This has led to debate over the so-called 'negative trial' and the role of large-scale randomised controlled trials, with calls for them to be abandoned. This presentation will discuss the significant impact of neutral trial results on clinical practice and the health care system; debate the concept that it is time to move away from the idea of the 'negative trial'; and suggest some novel and innovative design 	randomised trials in judging treatment effectiveness Carl Heneghan New treatments are only slightly superior to established treatments when tested in randomised controlled trials. Also, results have remained stable over time and the success rate of new treatments has not changed over the last half century of clinical trials. Why does so little research translate into practice?	
5:00pm	Summary and close	methodologies that may improve success of future trials.		
5:15–	Professor Alan Merry Networking event			
7:00pm	Faculty of Medical and Health Sciences At	rium, University of Auckland		