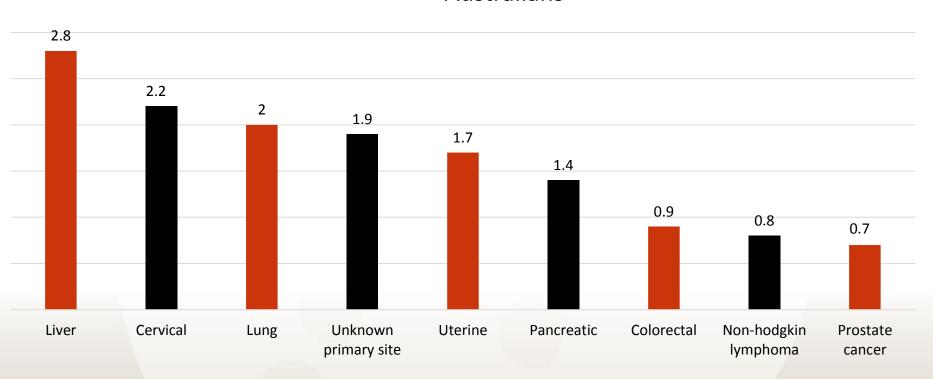
# Cancer and Aboriginal and Torres Strait Islander people - Gail Garvey

# Overall cancer incidence is higher amongst Indigenous Australians

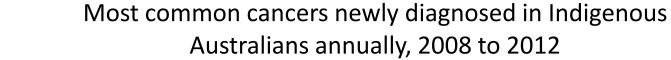
Cancer groups: Indigenous Australians vs non-Indigenous
Australians

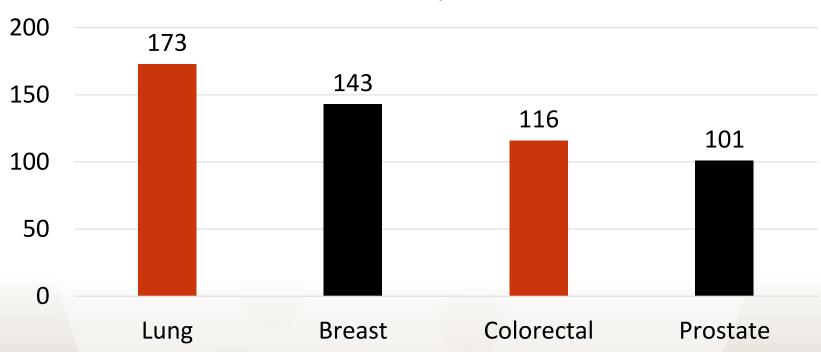


Australian Institute of Health and Welfare. (2017). *Cancer in Australia 2017.*Cancer series no. 101. Cat. no. CAN 100. Canberra: AIHW.

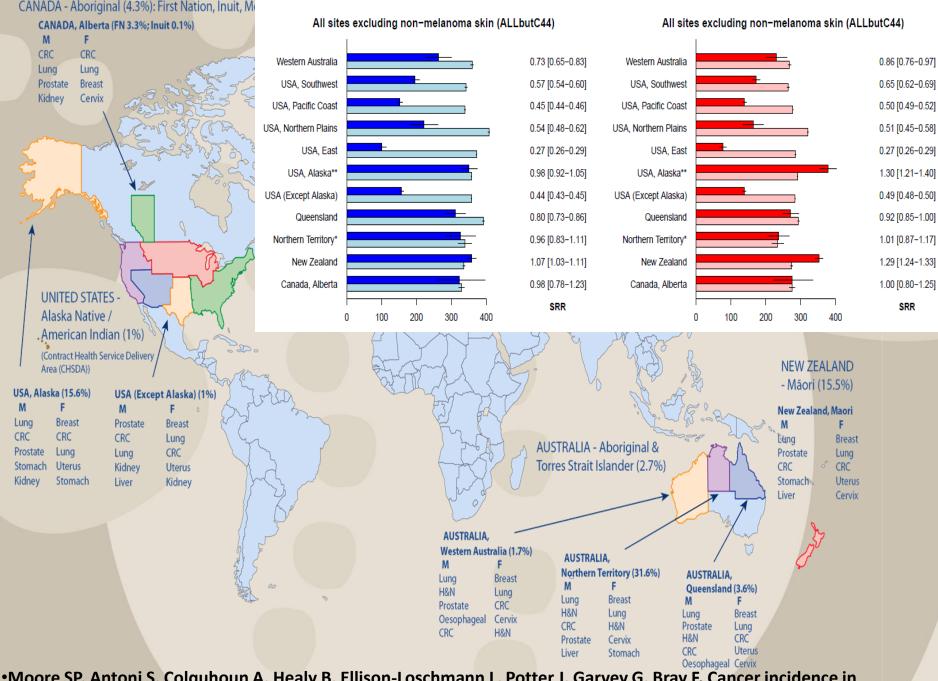
#### Most common cancers







Australian Institute of Health and Welfare. (2017). Cancer in Australia 2017.

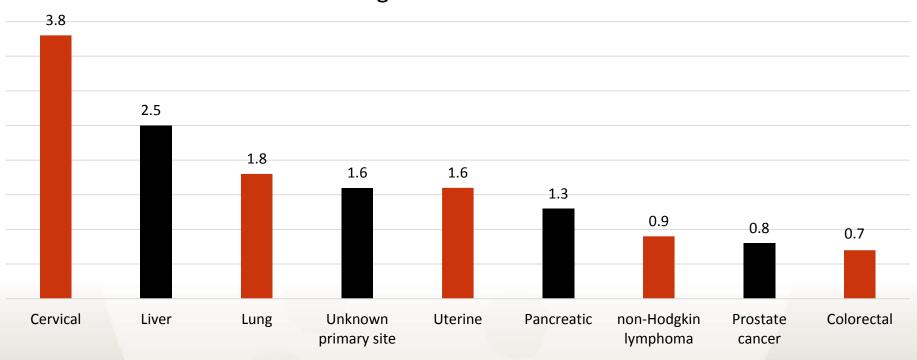


•Moore SP, Antoni S, Colquhoun A, Healy B, Ellison-Loschmann L, Potter J, Garvey G, Bray F. Cancer incidence in indigenous people in Aust, NZ, Canada, and the USA: a comparative population-based study. Lancet Oncology. 2015





Mortality by Cancer Group - Indigenous Australians vs non-Indigenous Australians

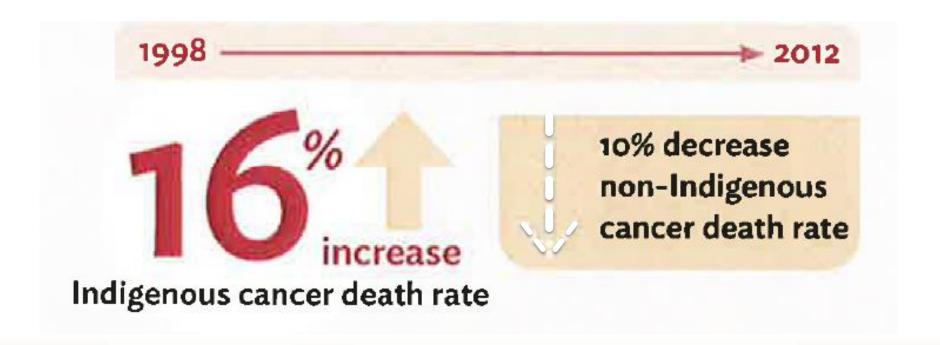


Australian Institute of Health and Welfare. (2017). Cancer in Australia 2017.

Cancer series no. 101. Cat. no. CAN 100. Canberra: AIHW.

## Cancer death rate increasing





## Reasons for poorer outcomes



#### **Indigenous Australians...**

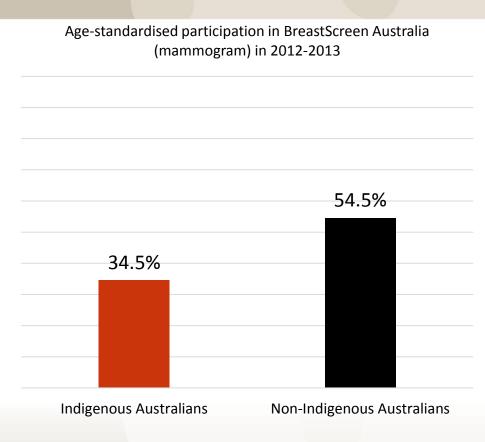
- •Higher incidence of cancers with poorer prognoses<sup>1</sup>
- •Diagnosed with cancers at later stage<sup>2-6</sup>
- •Higher rates of co-morbidities<sup>4,5</sup>
- •Lower cancer treatment uptake<sup>1,2</sup>
- •Receive less cancer treatment<sup>1</sup>

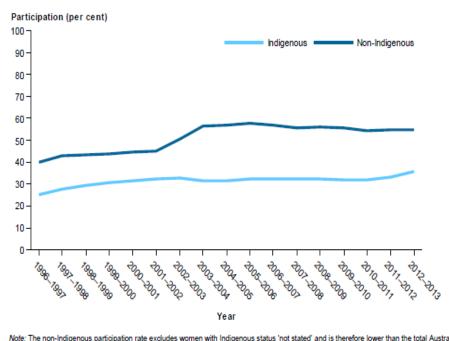


•Higher odds of receiving less cancer care concordant with clinical guidelines<sup>7</sup>

## Screening disparities







Note: The non-Indigenous participation rate excludes women with Indigenous status 'not stated' and is therefore lower than the total Australian participation rate.

Source: AIHW analysis of BreastScreen Australia data. Data for this figure are available in Table A.1.7.

Figure 3.6: Participation of women aged 50–69 in BreastScreen Australia, by Indigenous status, 1996-1997 to 2012-2013

AIHW 2015. BreastScreen Australia monitoring report 2012–2013. Cancer series no. 95. Cat. no. CAN 93. Canberra: AIHW.

## **Cancer risk factors**



- higher rates of lung cancer could be explained by the higher prevalence of smoking (38%, 18%)
- higher rates of cervical cancer could be explained in part by the higher prevalence of chronic infections (HPV), and lower participation in screening
- Liver cancer: higher incidence may be explained by higher rates of risky alcohol consumption and higher prevalence of hepatitis B infection

#### **Factors Impacting on Cancer Outcomes of Indigenous Cancer Patients**



disempower lang (interprimplicar colle implications lack of respect collective society mistrust and spiritual beliefs traditional healers experience of others support person (the right person)

patients' view and understanding of cancer and cancer treatment financial problems transportation attitudes towards service providers Out of town / country previous experiences access to supportive car eservices shame and fear of disease housing remoteness **literacy** patient disability co-morbidities family and community commitments

# Health Cares,

access to supportive care services, appropriateness of service / service

Droviders lack of confidence in system, staff and treatments late diagnosis (stage at diagnosis) communication (misunderstanding) medical practioner (gender, technical and communication skills) reduced uptake of treatment attitudes of staff

(language, body language, tone of voice, how questions are asked) Short consultation time no holistic treatment location of health facilities un-or under-insured (high co-pays) one system to fit all (no valuing diversity) waiting times for appointments when there

# Research strategies to investigate disparities in cancer outcomes

### **National Research Program Grants**



- NHMRC Centre of Research Excellence Discovering Indigenous Strategies to improve Cancer Outcomes Via Engagement, Research Translation and Training "DISCOVER-TT" – End date Oct 2018 Funding NHMRC
- 2. Strategic Research Partnership to improve Cancer Control for Indigenous Australians Funding Cancer Council NSW & WA End date Sept 2018

#### Western Australia collaborators

- Professor Sandra Thompson, Western Australia Centre for Rural Health, UWA
- Doctor Mick Adams, Australian Indigenous HealthInfoNet
- Professor Samar Aoun, Curtin University
- Doctor Shaouli Shahid, Curtin University
- Support an Aboriginal Cadet (Mikayla Hollows), Masters student (Leanne Pilkington) and PhD candidate (Belinda Davis)



## National Cervical Screening Program

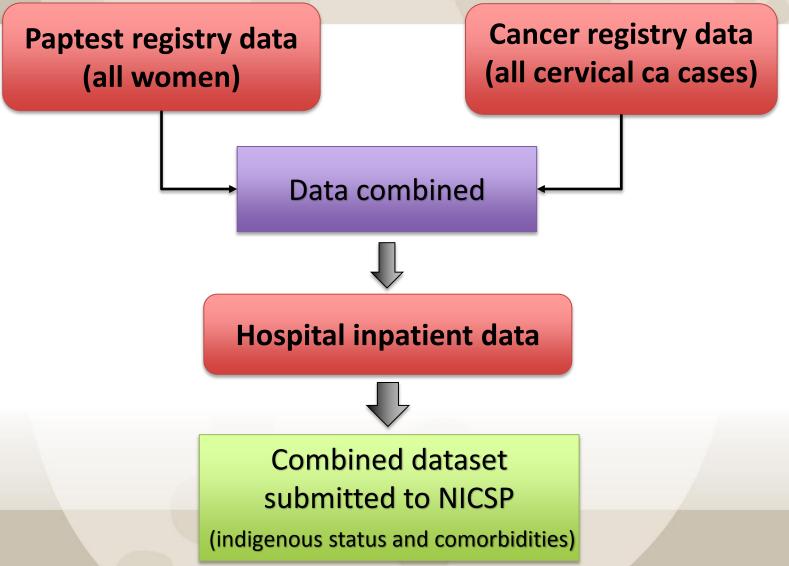
A joint Australian, State and Territory Government initiative

Cant report on Aboriginal and Torres Strait Islander women's participation

National data linkage Research project

Assess Indigenous women's participation, abnormalities after a pap test and timeliness to follow up

# Linkage within each state and territory (1997 to most recent)

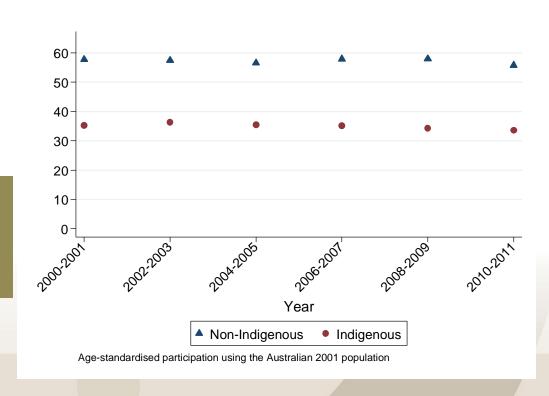


#### Indigenous women's cervical screening in Queensland

#### **Main Findings for Queensland**

- Approx. 1.4 million women & ~29000 linked to a record in the ever-Indigenous extract from hospital admissions
- 5 million Pap records available
- Only using 2000-2011 data

- Indigenous women participate less (20 percentage point diff)
- No change over time.



# Reviewing national population based screening programs





Developing education and training material for health professions and piloting the delivery of this program via primary health care services

Menzies contracted by DoH to pilot an Alternative Pathway to screening

# Development and validation of an Indigenous Quality of Life and Wellbeing Index for health decision making

The **Indigenous Quality of life and Wellbeing Index** will measure and value quality of life and wellbeing dimensions that are **important to Indigenous people** to improve the relevance and transparency of health decision making.

Specifically, this project will:

- 1. <u>Identify the dimensions of wellbeing and quality of life relevant for Indigenous people</u>
- 2. <u>Develop and validate a descriptive system</u> for a QOL/wellbeing index that appropriately captures these dimensions;
- 3. <u>Develop a scoring system for the new tool</u>

**NHMRC Funded: 2017-2021** 

Investigators: Howard, Garvey, Radcliffe, Tong, Cunningham, Cass, Whop

#### **Unmet Needs Assessment**



- Needs assessment is the <u>patient perspective</u>
   on concerns and degree to which they require help
- Solution-focused
- Measures gap between experience and expectations

## Most frequent unmet needs of Aboriginal and Torres Strait Islander Cancer Patients



Prevalence of unmet needs vary by jurisdiction

#### Queensland study

- 248 Indigenous Ca pts
- > 71% reported at least one unmet need.
- Domains
  - Mostly Physical/psychological

#### Items

- Money worries (22%)
- Concerns about the worries of those close to you (15%)
- Worrying about the illness spreading or getting worse (14%)
- Feeling down or sad (13%)
- Anxiety (12%)
- Worry about results of treatment (12%)
- Work around the home (11%)

#### Semi-National Study

- 148 Indigenous Ca pts
- ➤ 65% reported at least one unmet need.
- Domains
  - Physical/psychological
- NT and WA more unmet needs in the practical/cultural domain
- 'money worries'
- 'costs of accommodation'
- 'having an Indigenous person to talk to and support you, someone who understands your culture"

## Conclusions



- Continuing to build on this work
  - Under screened groups
  - Health literacy
  - Strategies to increase uptake and completion of treatment
  - Coordination of care
  - Building partnerships and networks
  - Hot North?????