

# Prevalence of HIV indicator conditions in people with late diagnosis of HIV 25 years on:

## Are we still missing opportunities for earlier care?

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# Disclosures

- *Nil personal disclosures*
- *JSYL receives honoraria for participation in Advisory Boards and Consultancy roles for ViiV Healthcare and Gilead Sciences, and an investigator-initiated research grant from Merck, Sharp and Dohme which is unrelated to this project.*
- *IW has worked as an investigator on commercial and investigator-initiated studies with funding to institutions from Gilead, ViiV, MAS, Moderna and CSL. IW has worked on advisory boards for ViiV and Gilead. IW has received educational support from Gilead, ViiV, MSD and Pfizer.*



# Background – Late Diagnosis HIV

- Defined as CD4 count <350 cells/ $\mu$ L at first test
- 37% new HIV diagnosed late in Australia in 2023

Year of first ever HIV diagnosis	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023
Characteristic										
<b>Total cases<sup>a</sup></b>	1,079	1,029	1,006	962	840	895	626	541	553	722
<b>Diagnosed late</b>	277	263	260	285	269	276	225	225	215	247
<b>Late HIV diagnosis, %<sup>c</sup></b>	27.3%	27.7%	29.4%	32.5%	35.8%	35.8%	41.5%	47.5%	43.7%	37.0%

<sup>3</sup> Kirby Institute, 2024



# Background – Indicator Based Testing

WHO recommends reflexive, provider-initiated testing in conditions that may indicate presence of HIV<sup>4</sup>:

- AIDS-defining
- Linked to an undiagnosed HIV prevalence  $> 0.1\%$
- Where undetected HIV could result in serious negative outcomes e.g. pregnancy, immunosuppression

<sup>4</sup> World Health Organisation, 2007



**Neurology:** primary cerebral lymphoma, toxoplasmosis, cryptococcal meningitis, progressive multifocal leukoencephalopathy (PML), aseptic meningitis/encephalitis, Guillain-Barre syndrome, chronic inflammatory demyelinating polyneuropathy (CIDP), cerebral abscess, transverse myelitis, peripheral neuropathy, dementia, leucoencephalopathy

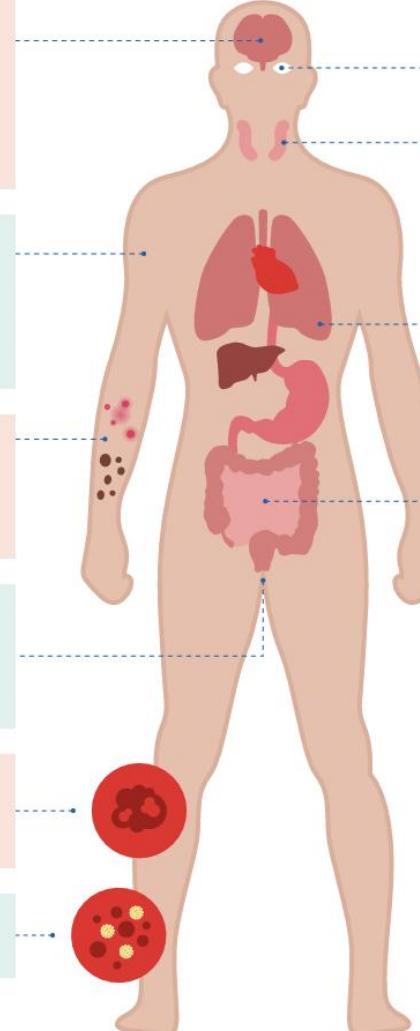
**Any AIDS-defining Illness:** oesophageal candidiasis, Kaposi's Sarcoma, pneumocystitis jiroveci pneumonia, histoplasmosis, cryptosporidiosis, toxoplasmosis, cryptococcal meningitis, cytomegalovirus (CMV), mycobacterium avium complex, non-Hodgkin lymphoma

**Skin:** Kaposi's Sarcoma, severe or recalcitrant seborrhoeic dermatitis or psoriasis, extensive warts or molluscum contagiosum, multidermatomal or recurrent varicella zoster, severe folliculitis

**Infections:** any STI (including syphilis, hepatitis B & C, chlamydia, gonorrhoea), refractory fungal infections, oral or oesophageal candidiasis, herpes varicella if multidermatomal or recurrent, toxoplasmosis

**Malignancies:** Hodgkin lymphoma, cervical, vaginal or anal intraepithelial neoplasia, Castleman disease and head & neck cancers

**Blood disorders:** unexplained thrombocytopenia, lymphopenia or neutropenia >4 weeks



**Eyes:** CMV retinitis, unexplained retinopathy or infective retinal disease (HSV & toxoplasma), syphilitic eye conditions

**Glandular fever type illness (could be HIV seroconversion):** pharyngitis, malaise, fever, lymphadenopathy, headache, maculopapular rash

**Persistent generalized lymphadenopathy**

**Respiratory conditions:** tuberculosis, recurrent bacterial pneumonia, aspergillosis, pneumocystitis jiroveci pneumonia

**Constitutional symptoms without an obvious cause:** pyrexia of unknown origin, unexplained weight loss, diarrhoea, myalgia, mononucleosis-like syndrome

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> [QJM. 2019 Jan 1;112\(1\):17-21. doi: 10.1093/qjmed/hcy223.](#)

## Prevalence of HIV indicator conditions in late presenting patients with HIV: a missed opportunity for diagnosis?

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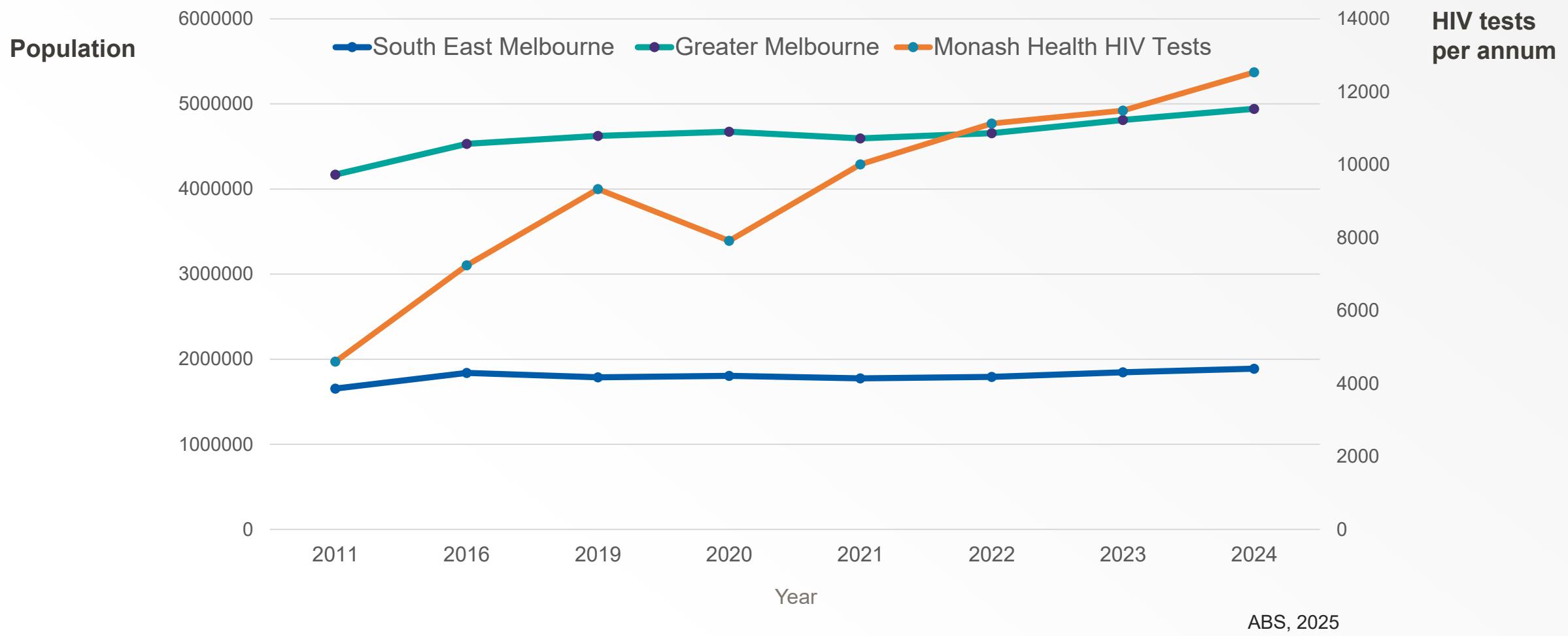
Affiliations + expand

PMID: 30295832 DOI: [10.1093/qjmed/hcy223](#)

<sup>6</sup>Lin et al, QJM, 2019



## Monash HIV testing 2011-2024, compared to ABS population data



# Study Aims – Are we *still* missing opportunities for earlier care?

- This study aimed to evaluate progress in provider-initiated, indicator-based testing over the last 10 years at Monash Health
- Analyse prevalence of indicator conditions in people living with HIV
  - In those with CD4 count <350 cells/ $\mu$ L at diagnosis, was there an opportunity for earlier diagnosis?
- Compare data from 2000-2014 study to recent results 2015-2025



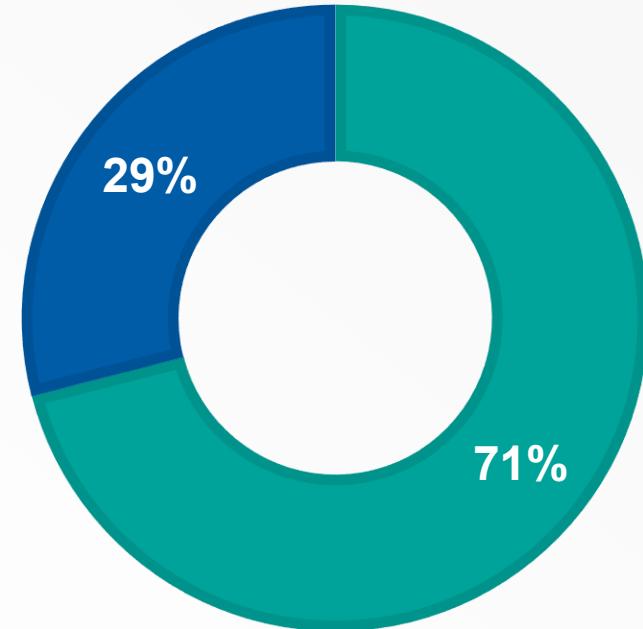
# Method

- Retrospective cohort study: 1<sup>st</sup> January 2015 – 1<sup>st</sup> January 2025
  - Study group: Individuals with late diagnosis HIV (defined by CD4<350cells/ $\mu$ L)
  - Control: Individuals with CD4 $\geq$ 350 at first presentation
- European AIDS Clinical Society guidelines used to identify indicator conditions in electronic health records
  - “Missed opportunity” defined as  $\geq$ 3 months
- Outcomes: Demographics, diagnostic setting and presence of indicator conditions.
- Statistical analysis: Chi-squared testing and two-proportion z-tests.



# Results

## HIV Diagnoses Monash Health 2000-2025



■ Total new, n = 676      ■ New dx CD4 <350cells/ $\mu$ L, n = 196

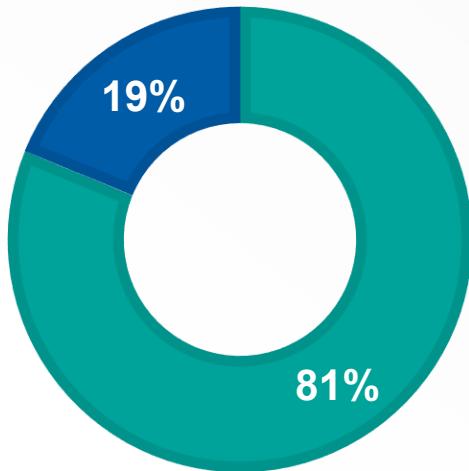
New cases per annum (mean) = 27



# Results

## HIV Diagnoses Monash Health 2000-2025

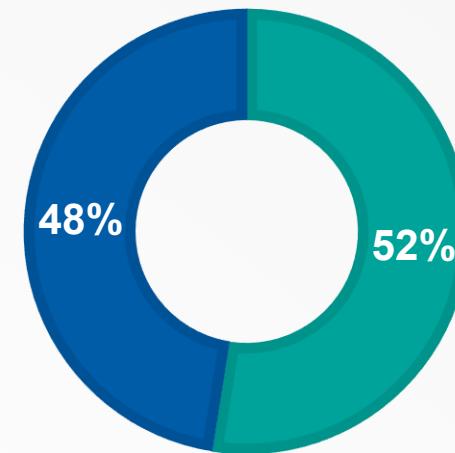
2000-2015



■ total, n = 436 ■ CD4 <350cells/µL, n = 82

Mean per annum = 29

2015-2025



■ total, n = 240 ■ CD4 <350cells/µL, n = 114

Mean per annum = 24

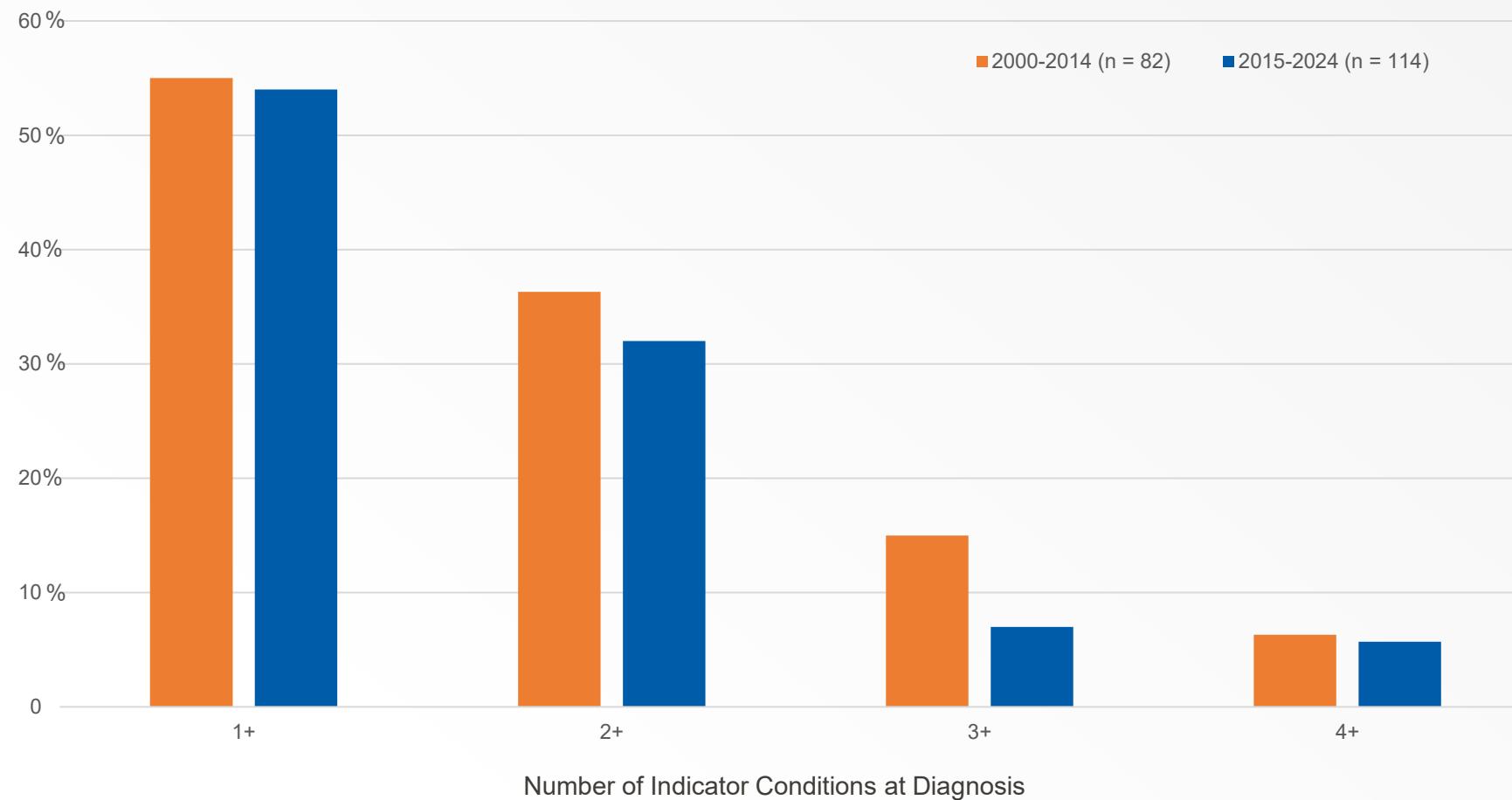


# Results: Demographics 2000-2025

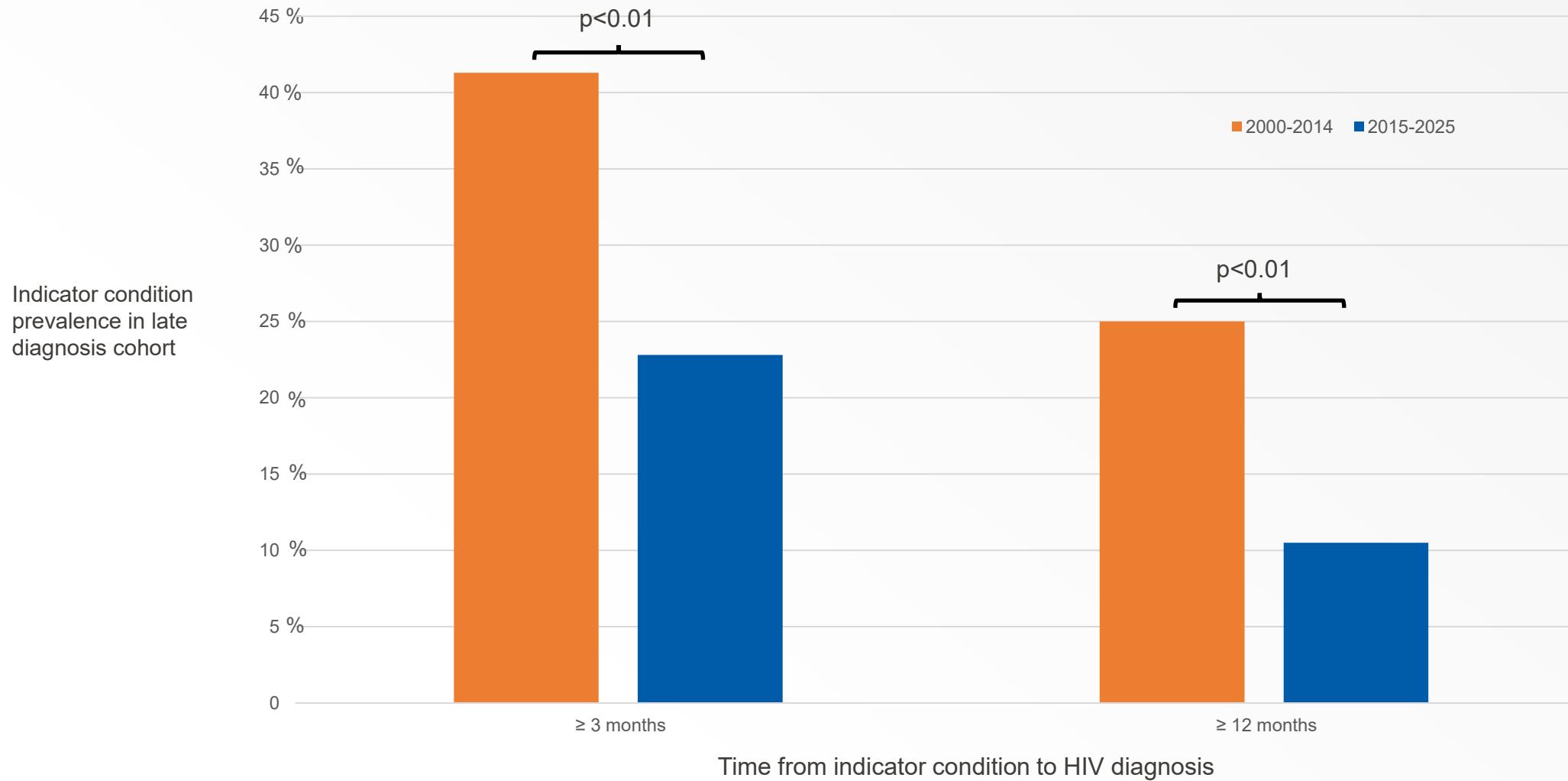
Demographics; Late Diagnosis			
	2000-2014	2015-2025	p value
Age (mean, years)	45	41.9	
Sex			0.096
Male	68 (83%)	83 (73%)	
Female	14 (17%)	31 (27%)	
Born Overseas	50 (61%)	83 (73%)	0.08
Total	82 (19%)	114 (48%)	p<0.005



## Prevalence of indicator conditions in late diagnosis cohort 2000 - 2025



## Proportion of individuals diagnosed late, who had an opportunity for earlier diagnosis



# Results: 2015-2025

<b>HIVIC</b>	<b>Individuals with indicator conditions diagnosed prior to HIV diagnosis</b>		
	<b>0-3 months</b>	<b>3-12 months</b>	<b>12 months or longer</b>
Unexplained weight loss	25	15	6
Hep B	16	0	0
STI	15	3	0
Candidiasis	15	7	3
HSV	9	2	2
Community Acquired Pneumonia	7	4	2
Unexplained lymphadenopathy	6	3	1
Unexplained leukocytopenia/ thrombocytopenia >4 weeks	4	1	1
Unexplained diarrhoea >4 weeks	3	2	1
Pregnancy	2	0	0
Cervical dysplasia	2	1	1
<b>Total</b>	<b>104</b>	<b>38</b>	<b>17</b>



# Discussion: Limitations

- Retrospective, single-centre design
- Causality cannot be inferred
- Analysis limited to cohort CD4 <350 cells/ $\mu$ L
- Reliance on medical records that may be incomplete / inaccurate
- Unmeasured confounders



# Key Messages

- Proportion of late diagnosis remains unacceptably high
- Within this cohort, clinician testing behaviour appears to have improved
- Next steps:
  - Continue to normalise HIV testing amongst colleagues, until it is routine
  - Integration of testing into guidelines is effective (pregnancy, hepatitis)
  - Scope for improvement: weight loss



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