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# Long term outcomes with Depot Cabotegravir/Rilpivirine in Australian General Practices

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# Disclosure

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David Baker received research funding from  
Gilead Sciences, ViiV, GSK

No funding was received for this current study

# Introduction

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Many people living with HIV in Australia are treated in General Practice (GP) settings

Long-acting injectable medication is widely used for many chronic health conditions

Long-acting antiretroviral (ARV) treatment in the form of depot treatment with Cabotegravir/Rilpivirine (LA CAB/RPV) is a treatment that can replace the need for daily ARV medication

LA CAB/RPV has been widely available in Australia since 1/4/2022 when it was listed on the Pharmaceutical Benefits Schedule (PBS)

# Method

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Retrospective cohort of patients treated with LA CAB/RPV IM through phase 3 trials and post PBS listing in 4 GP clinics

Collected as part of a GP treatment audit

# Results: Demographics

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192 patients were treated with LA CAB/RPV

96% were male

Age range of 23 – 87

403 patient years of treatment

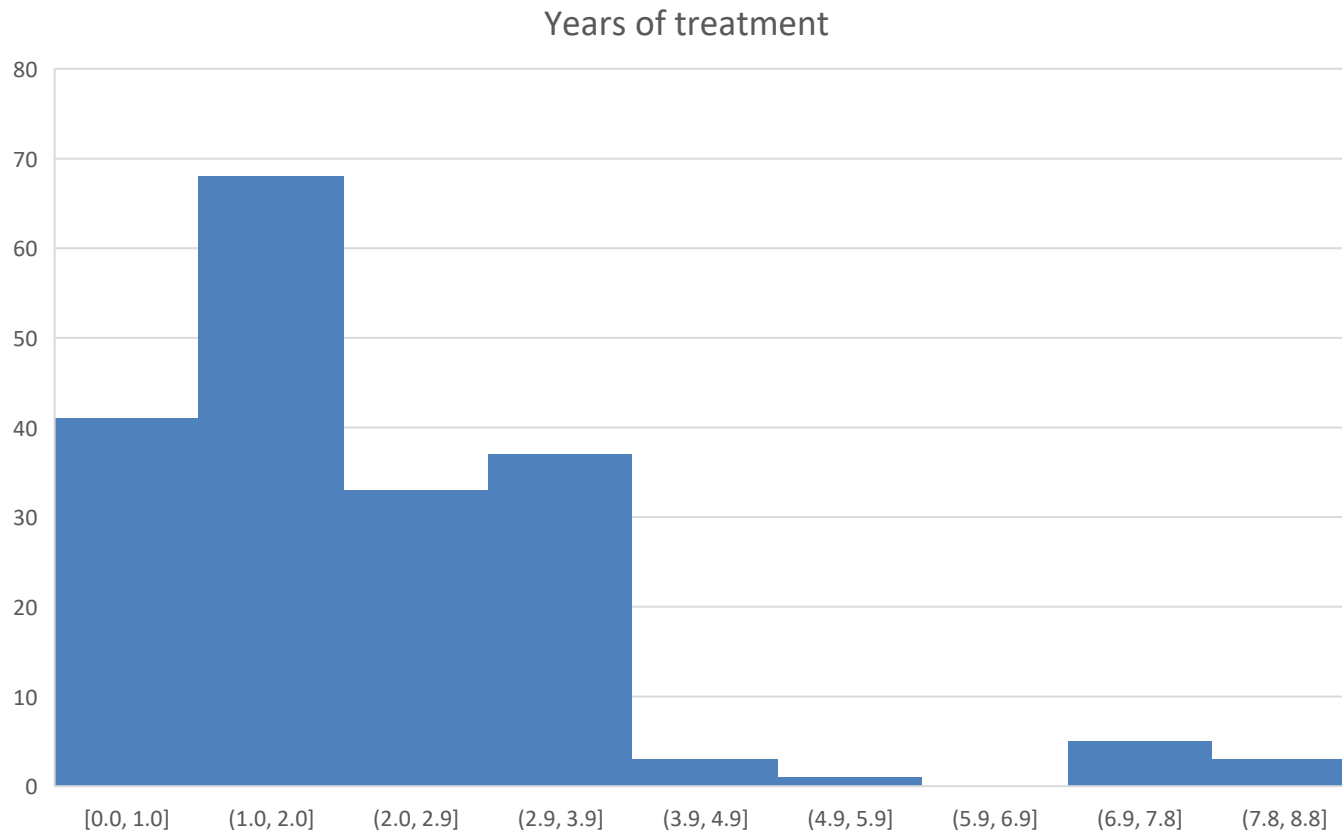
Median 22 months

Mean 25 months

# Results: length of treatment

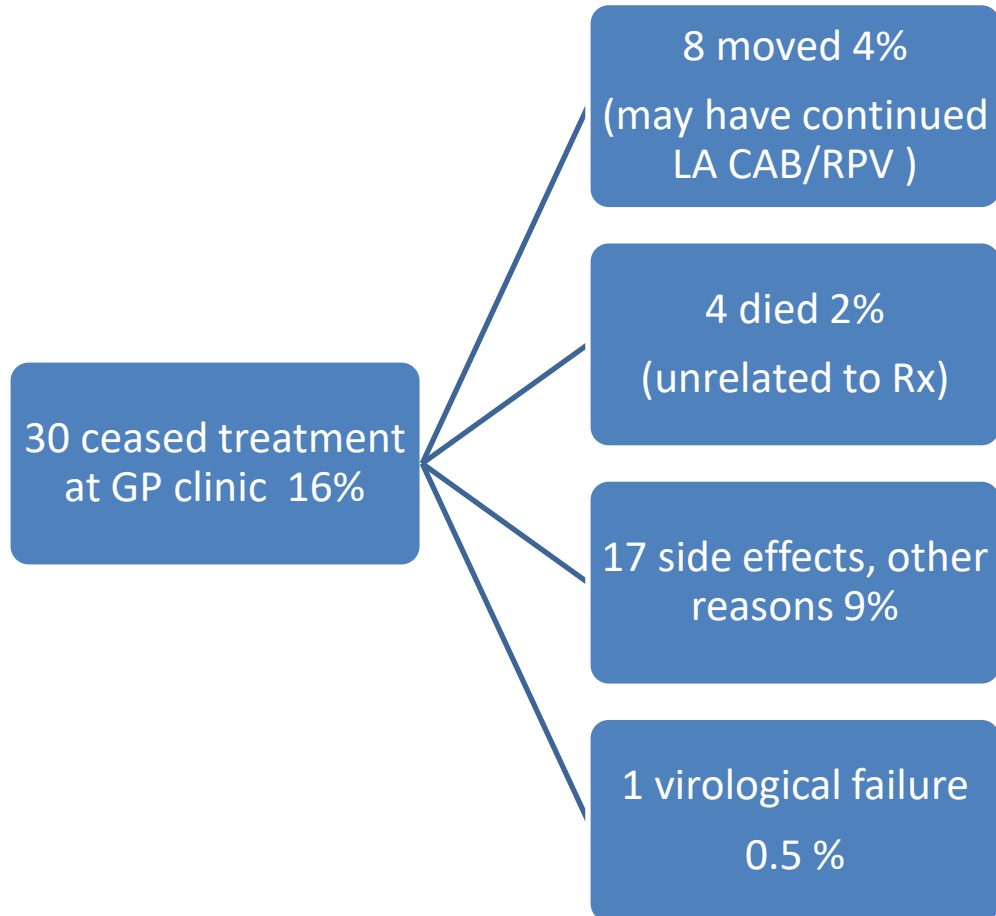
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Treatment length range was 0 – 9 years



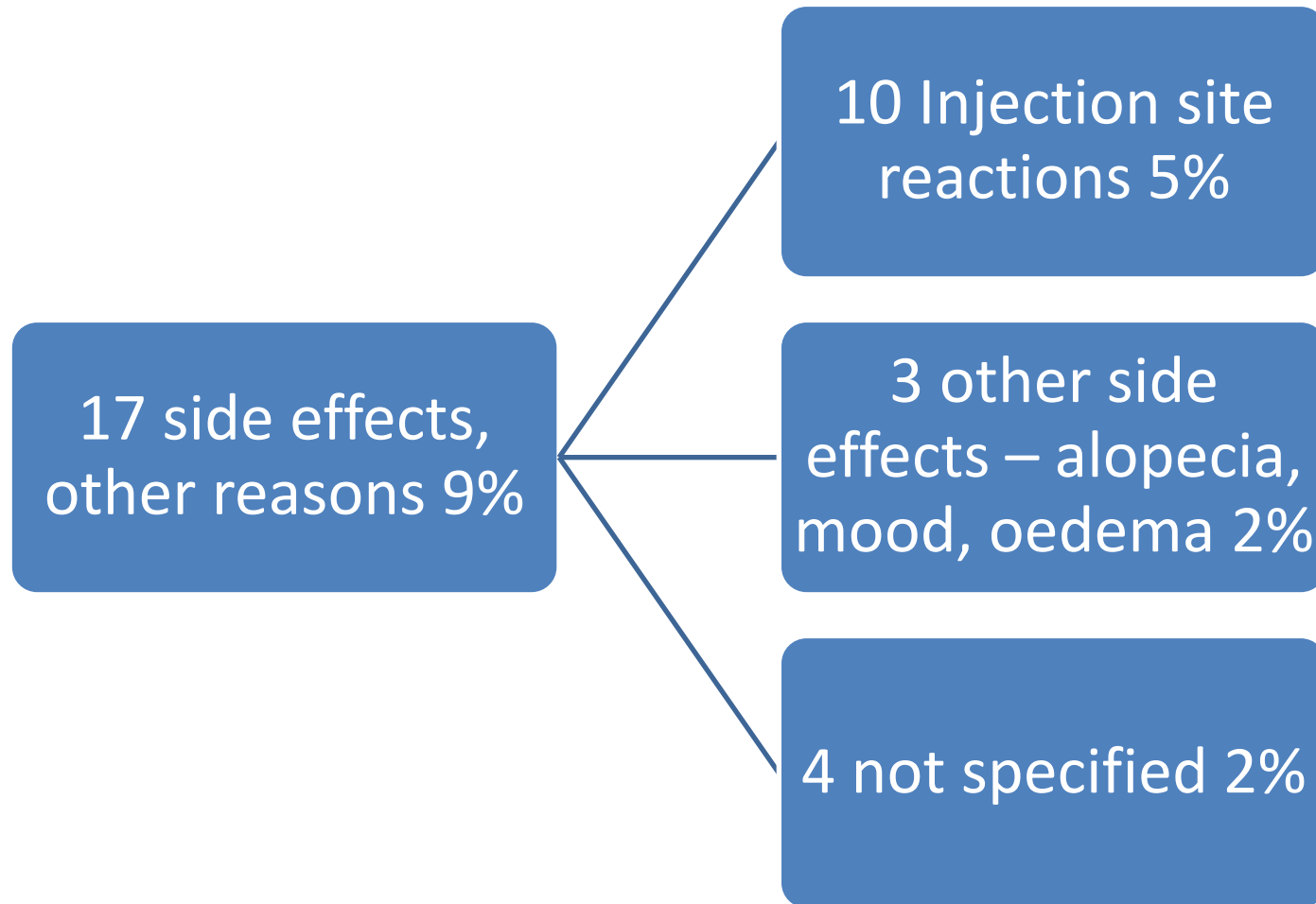
# Results: Patient outcomes

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# Results: reasons for stopping

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# Virological failure\*

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56-year-old male, HIV +ve 1993

Multiple regimens, stopped and started Rx

26/3/2008 resistance to EFV,NVP, 3TC (K103N, M184V)

28/6/2013 resistance to EFV,NVP not to RPV

21/2/2019 nil resistance

19/12/22 taking EVG/COB/FTC/TAF) (Genvoya) + darunavir  
admitted with atrial fibrillation

Switched to dolutegravir/rilpivirine (Juluca), given apixaban

6/1/23 VL bdl

\*patient provided consent for his data to be used in this presentation

# Virological failure

20/2/23 – LA CAB/RPV

12/5/23 VL 555

15/6/23 VL 5880

INSTI Major Mutations:

**E138EK • G140GS • Q148R**

INSTI Accessory Mutations:

None

IN Other Mutations:

V31I • K111T • T112V • V201I

## Integrase Strand Transfer Inhibitors

<b>bictegravir (BIC)</b>	High-Level Resistance
<b>cabotegravir (CAB)</b>	High-Level Resistance
<b>dolutegravir (DTG)</b>	High-Level Resistance
<b>elvitegravir (EVG)</b>	High-Level Resistance
<b>raltegravir (RAL)</b>	High-Level Resistance

NRTI Mutations:

None

NNRTI Mutations:

**K103N • E138K • P225H**

RT Other Mutations:

E6D • I142IV • S162C • T200I

## Nucleoside Reverse Transcriptase Inhibitors

<b>abacavir (ABC)</b>	Susceptible
<b>zidovudine (AZT)</b>	Susceptible
<b>stavudine (D4T)</b>	Susceptible
<b>didanosine (DDI)</b>	Susceptible
<b>emtricitabine (FTC)</b>	Susceptible
<b>lamivudine (3TC)</b>	Susceptible
<b>tenofovir (TDF)</b>	Susceptible

## Non-nucleoside Reverse Transcriptase Inhibitors

<b>doravirine (DOR)</b>	Intermediate Resistance
<b>efavirenz (EFV)</b>	High-Level Resistance
<b>etravirine (ETR)</b>	Potential Low-Level Resistance
<b>nevirapine (NVP)</b>	High-Level Resistance
<b>rilpivirine (RPV)</b>	Intermediate Resistance

Darunavir/r, TAF/FTC, last viral load = 54, CD4 = 482

# Conclusion

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LA CAB/RPV is an effective treatment for HIV in a General Practice setting with a low rate of treatment failure

LA CAB/RPV is generally well tolerated with the main reason for treatment discontinuation being injection site reaction

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