

# Modelling the allocation of Hepatitis C treatment: relationship to methadone maintenance

Irmgard Zeiler, Trevor Langlands, John M Murray & Alison Ritter

Drug Policy Modelling Program, UNSW  
School of Mathematics and Statistics, UNSW  
Institute of Mathematical Methods in Economics, Vienna  
University of Technology

Funding: Colonial Foundation Trust (DPMP), UNSW & NHMRC

# Background

- HCV significant cause of morbidity and mortality
- 60% of IDU in Australia HCV
- Treatment is available (with 50% success rate)
- HCV treatment rates in Australia low (1%)
- Interest in improving access to Hep C treatment and availability of Hep C treatment

# Motivation

- Allocating HCV treatment?
- Query regarding treatment for current injectors
- Uncertainty regarding reinfection
- Focus on people in opioid pharmacotherapy maintenance:
  - ↑ referrals into HCV treatment
  - Provide ongoing support during treatment
  - Improve compliance
- But: no evidence to date of improved access and outcomes in context of OST

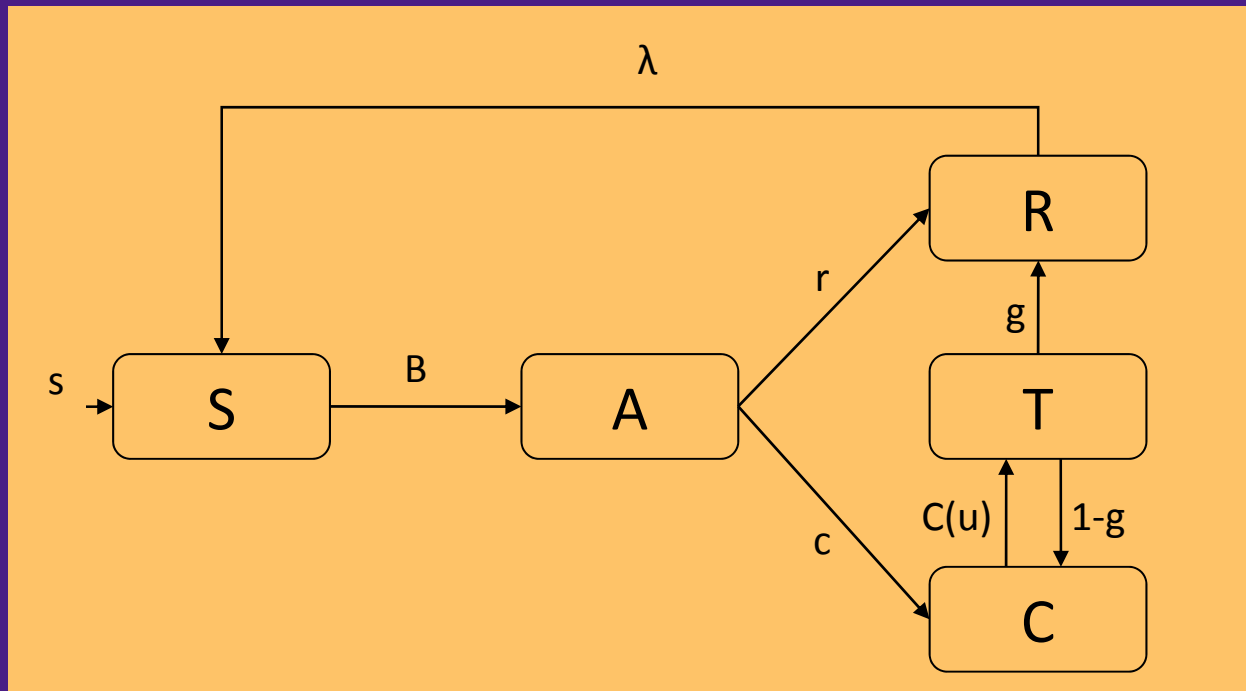


# Aim

- In context of:
  - a limited number of treatment spots; and
  - lack of current evidence re role of opioid pharmacotherapy maintenance in improving HCV treatment outcomes
- Explore the optimal allocation of HCV treatment between those people in pharmacotherapy maintenance and those outside pharmacotherapy maintenance treatment
  - How should treatment places be distributed?

# The model

- Simple infectious diseases model: SACTR



# Features

- Population model – all IDU in Australia
- Single state model = all IDU. Two state model splits the IDU group into two treatment grps
- Methadone maintenance only (MMT)
- Optimisation = minimum steady state HCV population prevalence (outcome = achieving the least no. of infected people within the model)

# Parameters

- Population estimates: # IDU, #'s in MMT etc.
- HCV infection rates: based on rate of injecting (in and out of MMT) and probability of becoming infected
- HCV treatment parameters: duration of treatment, success rate
- MMT treatment parameters: duration of treatment
- Reinfection rate

# Sensitivity analysis

- Model reproduces expected population prevalence of HCV (60%)
- Parameters that had the largest impact on steady state HCV prevalence were:
  - rate of infection and the exit rate
- changing these by +/-10% resulted in a 4-6% change in the predicted prevalence.
- Changing other parameters by +/- 10% resulted in changes in steady state HCV prevalence of less than 1%.

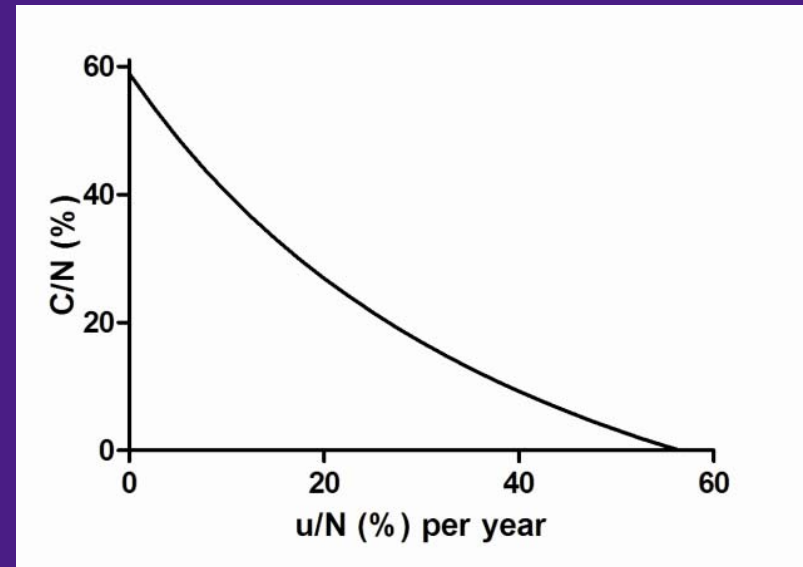
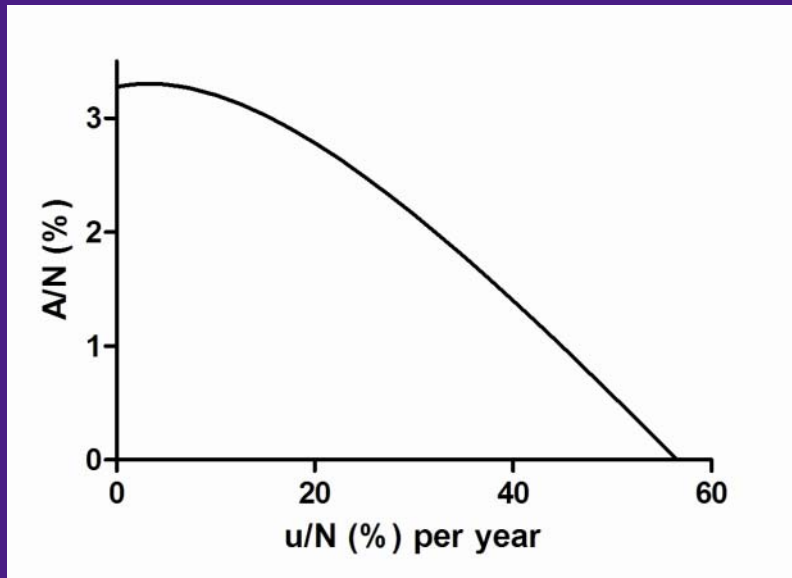


# Results

- Single state model without MMT split
- Examines:
  - Impact on HCV prevalence of increasing HCV treatment levels
  - Optimal HCV treatment rate to achieve minimum steady state prevalence

# Results

The effect of the treatment rate (percentage of the total population per year),  $u$ , on the percentage of acutely infected,  $A$ , (left) and chronically infected,  $C$ , (right) users in the endemic steady state.



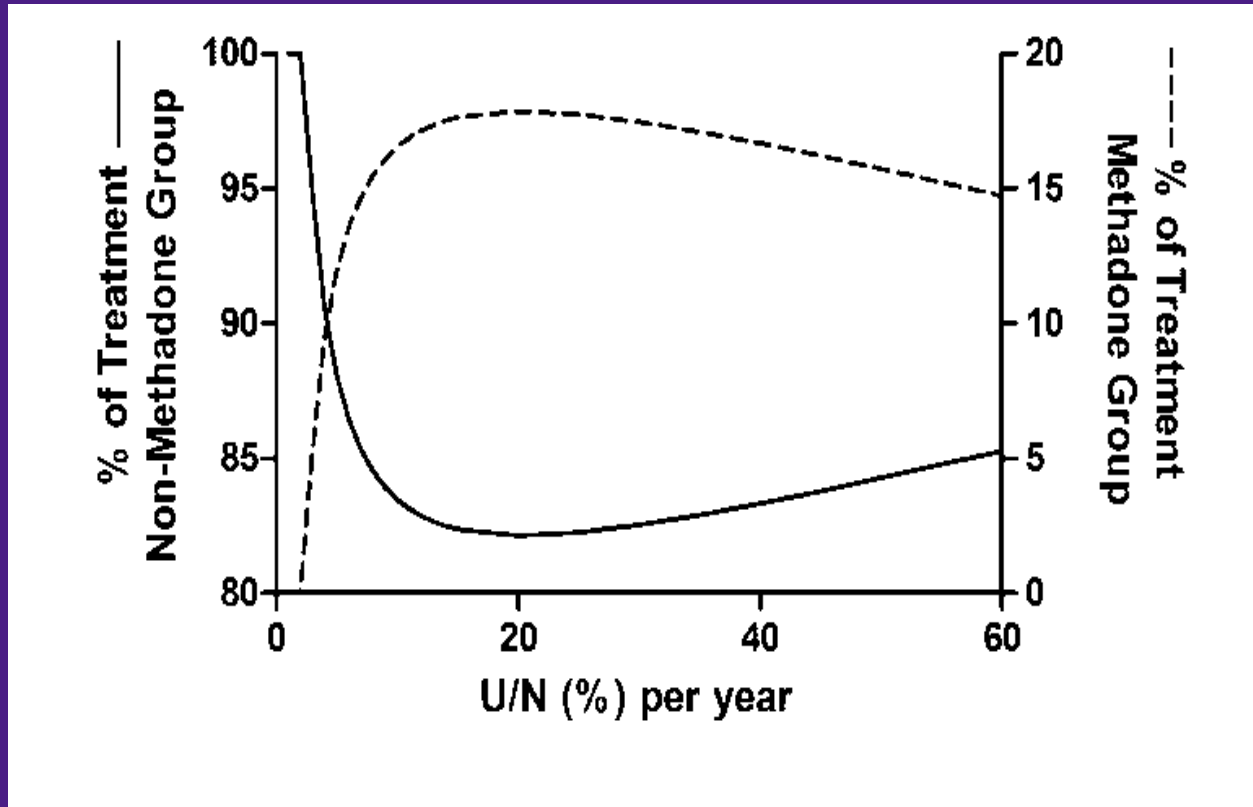
Never fully eliminate HCV (always new infections)

- As increasing numbers are treated, prevalence decreases:
  - But with an initial increase in acute
- Treatment level to minimise HCV is 56.5%
- Rate of infection continues (due to sharing) so never eliminated with treatment alone
- Slow: 3.3 yrs for chronic infections to decrease by half; 11.1 yrs for acute infections to decrease by half

# Optimal mix: MMT or non-MMT

- Assuming treatment rate of 50% (derived from single state model) optimal mix between MMT and non-MMT HCV treatment is:
  - 84% of HCV treatment places to non MMT
  - 16% of HCV treatment places to MMT
- This mix results in lowest steady state HCV in the population
- Varies as the HCV treatment rate changes (but not by much.....)

# Optimal allocation between MMT and non-MMT



The effect of varying the percentage of the total population treated per year on the optimal treatment allocation between non-methadone (solid line) and methadone (dashed line) users.

# Limitations

- As with all maths models, parameters are essential:
  - Reinfection rate too low?
  - MMT cycling behaviour too high?
  - Length of HCV treatment averaged across genotypes
- HCV treatment rate of 56% of total population unrealistic
- Does not account for potential advantages of MMT

# Conclusions

1. Programs that minimise sharing must continue (cannot eliminate HCV with treatment alone)
2. Optimal allocation is to treat those not in MMT (at significantly wide margins)
  - Focus efforts on treating current injectors
3. More research on whether MMT does improve HCV treatment compliance and outcomes
  - Then can re-run model with those new parameters

## Further information

Assoc Prof Alison Ritter  
Drug Policy Modelling Program, Director  
National Drug and Alcohol Research Centre  
UNSW, Sydney, NSW, 2052, Australia

E: [alison.ritter@unsw.edu.au](mailto:alison.ritter@unsw.edu.au)  
T: + 61 (2) 9385 0236

DPMP Website: <http://www.dpmp.unsw.edu.au>

Project funding sources: Colonial Foundation  
Trust (DPMP), UNSW, NHMRC