

Risk group turnover in STI/HIV epidemics

Jesse Knight¹, Linwei Wang¹, Huiting Ma¹, Sheree Schwartz², Stefan Baral², Sharmistha Mishra¹

¹MAP Centre for Urban Health Solutions, Unity Health Toronto

²Dept. Epidemiology, Johns Hopkins Bloomberg School of Public Health

2019 July 17

STI & HIV 2019 World Congress
Vancouver, BC, Canada

Disclosures

None.

Acknowledgements



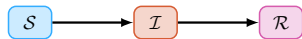
Background

Methods

Results

Conclusion

SIR Models & Heterogeneity in Risk



► SIR Model:

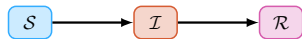
- \mathcal{S} = susceptible, \mathcal{I} = infectious, \mathcal{R} = recovered
- e.g. HIV

► 3 Risk Groups:

- e.g. female sex workers, multiple partners, monogamous
- fundamentally changes epidemic dynamics [1, 2]
- what about: “retirement” from sex work
“recruitment” into sex work } **Turnover**



SIR Models & Heterogeneity in Risk

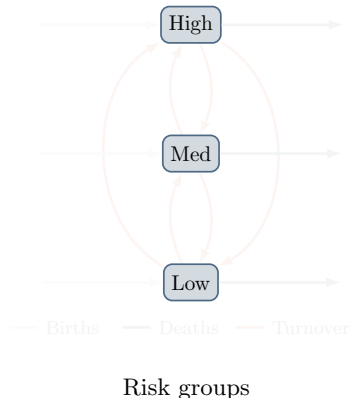


► SIR Model:

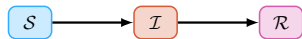
- \mathcal{S} = susceptible, \mathcal{I} = infectious, \mathcal{R} = recovered
- e.g. HIV

► 3 Risk Groups:

- e.g. female sex workers, multiple partners, monogamous
- fundamentally changes epidemic dynamics [1, 2]
- what about: “retirement” from sex work
“recruitment” into sex work } **Turnover**



SIR Models & Heterogeneity in Risk

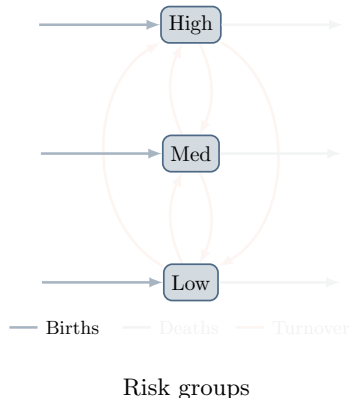


► SIR Model:

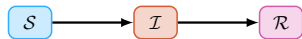
- \mathcal{S} = susceptible, \mathcal{I} = infectious, \mathcal{R} = recovered
- e.g. HIV

► 3 Risk Groups:

- e.g. female sex workers, multiple partners, monogamous
- fundamentally changes epidemic dynamics [1, 2]
- what about: “retirement” from sex work
“recruitment” into sex work } **Turnover**



SIR Models & Heterogeneity in Risk

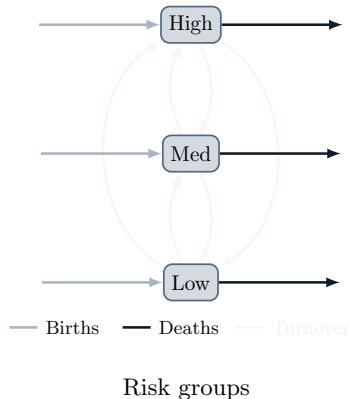


► SIR Model:

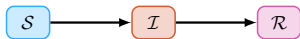
- \mathcal{S} = susceptible, \mathcal{I} = infectious, \mathcal{R} = recovered
- e.g. HIV

► 3 Risk Groups:

- e.g. female sex workers, multiple partners, monogamous
- fundamentally changes epidemic dynamics [1, 2]
- what about: “retirement” from sex work
“recruitment” into sex work } **Turnover**



SIR Models & Heterogeneity in Risk



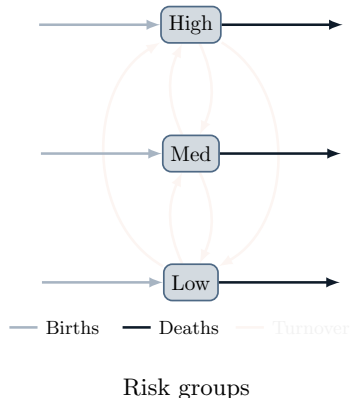
► SIR Model:

- \mathcal{S} = susceptible, \mathcal{I} = infectious, \mathcal{R} = recovered
- e.g. HIV

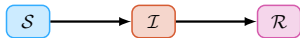
► 3 Risk Groups:

- e.g. female sex workers, multiple partners, monogamous
- fundamentally changes epidemic dynamics [1, 2]
- what about: “retirement” from sex work
“recruitment” into sex work

Turnover



SIR Models & Heterogeneity in Risk

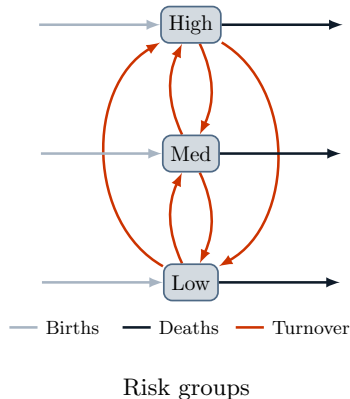


► SIR Model:

- \mathcal{S} = susceptible, \mathcal{I} = infectious, \mathcal{R} = recovered
- e.g. HIV

► 3 Risk Groups:

- e.g. female sex workers, multiple partners, monogamous
- fundamentally changes epidemic dynamics [1, 2]
- what about: “retirement” from sex work
“recruitment” into sex work } **Turnover**



Influence of Turnover in STI Epidemic Models

Research Questions:

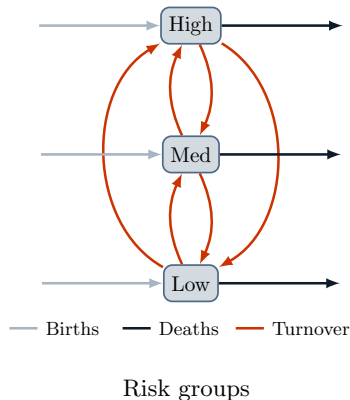
Influence of turnover on:

- ▶ Equilibrium incidence & prevalence by risk group
- ▶ TPAF* of high risk group

* TPAF: “Transmission Population Attributable Fraction” [3]

Proportion of cumulative new infections averted
if transmission to / from that group is stopped.

e.g. impact of perfect TasP in one group



Influence of Turnover in STI Epidemic Models

Research Questions:

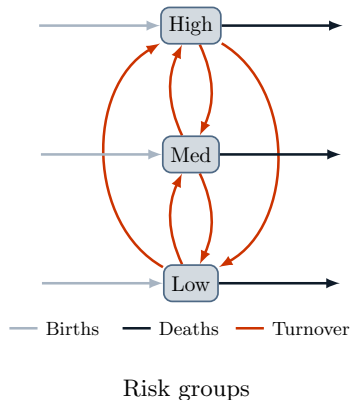
Influence of turnover on:

- ▶ Equilibrium incidence & prevalence by risk group
- ▶ TPAF* of high risk group

* TPAF: “Transmission Population Attributable Fraction” [3]

Proportion of cumulative new infections averted
if transmission to / from that group is stopped.

e.g. impact of perfect TasP in one group



Influence of Turnover in STI Epidemic Models

Research Questions:

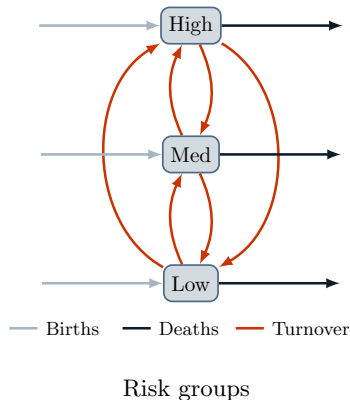
Influence of turnover on:

- Equilibrium incidence & prevalence by risk group
- TPAF* of high risk group

* TPAF: “Transmission Population Attributable Fraction” [3]

Proportion of cumulative new infections averted
if transmission to / from that group is stopped.

e.g. impact of perfect TasP in one group



Influence of Turnover in STI Epidemic Models

Research Questions:

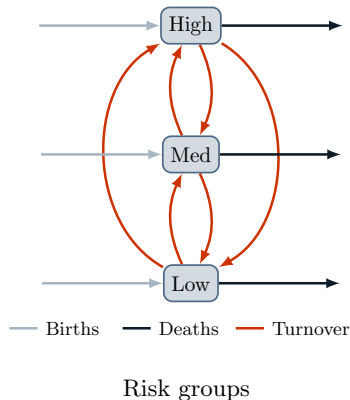
Influence of turnover on:

- Equilibrium incidence & prevalence by risk group
- TPAF* of high risk group

* TPAF: “Transmission Population Attributable Fraction” [3]

Proportion of cumulative new infections averted
if transmission to / from that group is stopped.

e.g. impact of perfect TasP in one group



Influence of Turnover in STI Epidemic Models

Research Questions:

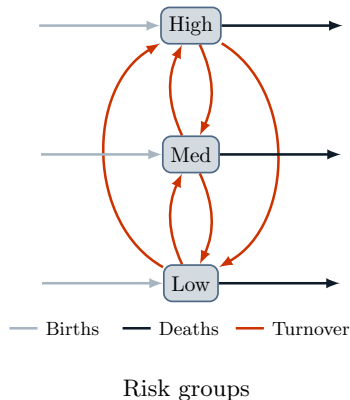
Influence of turnover on:

- Equilibrium incidence & prevalence by risk group
- TPAF* of high risk group

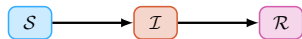
* TPAF: “Transmission Population Attributable Fraction” [3]

Proportion of cumulative new infections averted if transmission to / from that group is stopped.

e.g. impact of perfect TasP in one group



Illustrative Model of STI Transmission

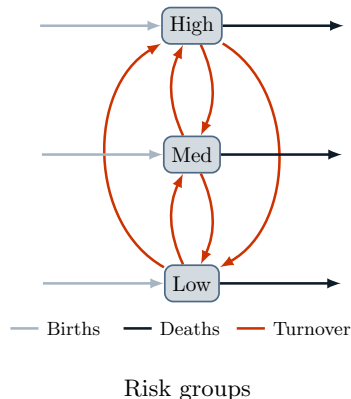


► SIR model:

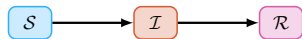
- 1-sex
- proportional mixing
- same mortality across risk groups

► Risk group turnover:

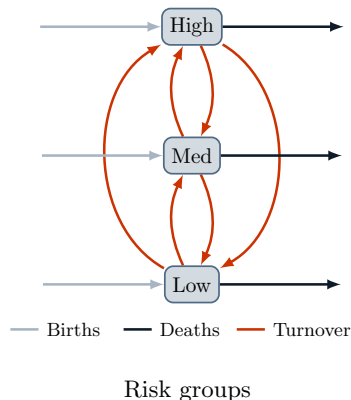
- Rates ensure group sizes don't change:
5% High Risk, 20% Medium Risk, 75% Low Risk
- All rates equal among: \mathcal{S} , \mathcal{I} , \mathcal{R}
- All rates scaled proportionally when varied



Illustrative Model of STI Transmission



- ▶ SIR model:
 - ▶ 1-sex
 - ▶ proportional mixing
 - ▶ same mortality across risk groups
- ▶ Risk group turnover:
 - ▶ Rates ensure group sizes don't change:
5% High Risk, 20% Medium Risk, 75% Low Risk
 - ▶ All rates equal among: \mathcal{S} , \mathcal{I} , \mathcal{R}
 - ▶ All rates scaled proportionally when varied



Experiments: Influence of Turnover on Model Outputs

1. Equilibrium outputs:

- ▶ **Vary:** Turnover magnitude
- ▶ **Compare:** a) prevalence, b) incidence (by risk group, at equilibrium)

2. TPAF after model fitting:

- ▶ **Fit:** Contact rates: High Risk; and Low Risk
- ▶ **Targets:** Prevalence: 25% in High Risk; and 5% in Low Risk
- ▶ **Vary:** No-turnover vs Turnover
- ▶ **Compare:** a) Fitted contact rates, b) TPAF of high risk group

Experiments: Influence of Turnover on Model Outputs

1. Equilibrium outputs:

- ▶ **Vary:** Turnover magnitude
- ▶ **Compare:** a) prevalence, b) incidence (by risk group, at equilibrium)

2. TPAF after model fitting:

- ▶ **Fit:** Contact rates: High Risk; and Low Risk
- ▶ **Targets:** Prevalence: 25% in High Risk; and 5% in Low Risk
- ▶ **Vary:** No-turnover vs Turnover
- ▶ **Compare:** a) Fitted contact rates, b) TPAF of high risk group

Experiments: Influence of Turnover on Model Outputs

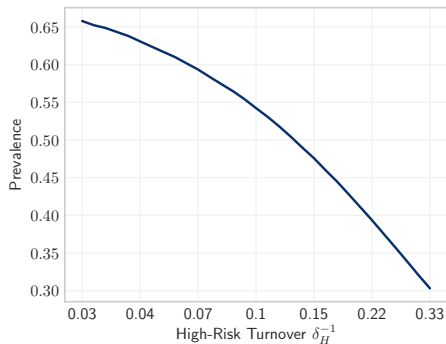
1. Equilibrium outputs:

- ▶ **Vary:** Turnover magnitude
- ▶ **Compare:** a) prevalence, b) incidence (by risk group, at equilibrium)

2. TPAF after model fitting:

- ▶ **Fit:** Contact rates: High Risk; and Low Risk
- ▶ **Targets:** Prevalence: 25% in High Risk; and 5% in Low Risk
- ▶ **Vary:** No-turnover vs Turnover
- ▶ **Compare:** a) Fitted contact rates, b) TPAF of high risk group

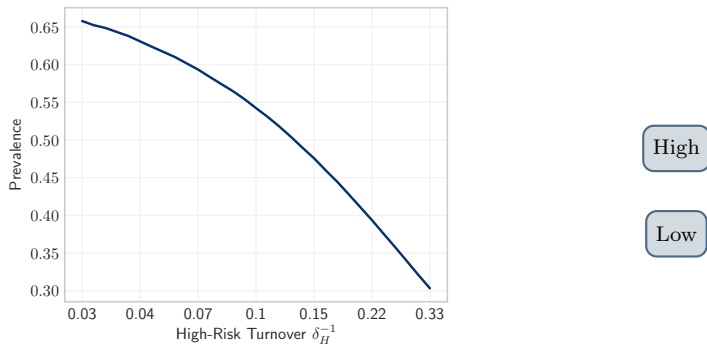
Turnover decreases high risk group equilibrium prevalence



High risk prevalence vs turnover

Turnover causes a net movement of infected: high \rightarrow low risk

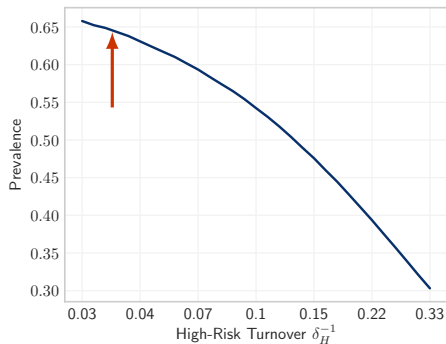
Turnover decreases high risk group equilibrium prevalence



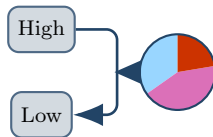
High risk prevalence vs turnover

Turnover causes a net movement of infected: high \rightarrow low risk

Turnover decreases high risk group equilibrium prevalence

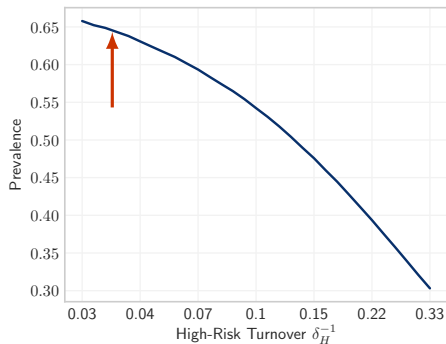


High risk prevalence vs turnover

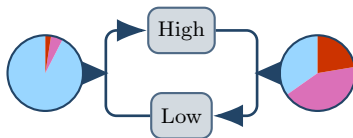


Turnover causes a net movement of infected: high \rightarrow low risk

Turnover decreases high risk group equilibrium prevalence

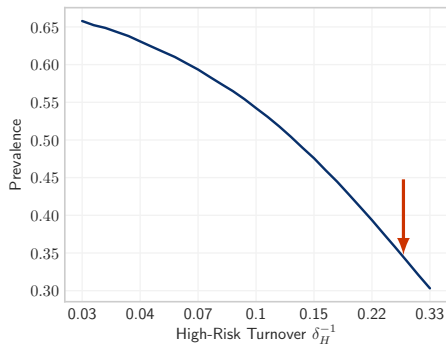


High risk prevalence vs turnover

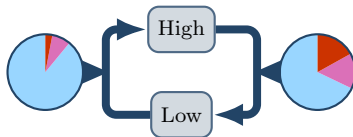


Turnover causes a net movement of infected: high \rightarrow low risk

Turnover decreases high risk group equilibrium prevalence

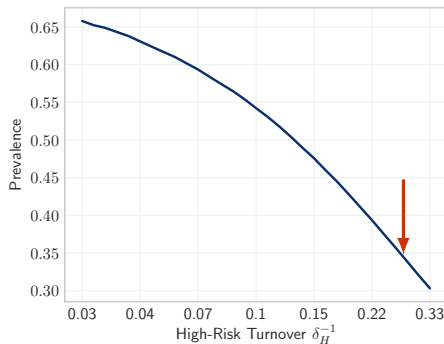


High risk prevalence vs turnover

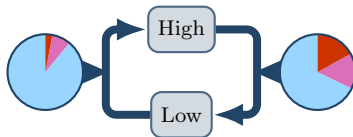


Turnover causes a net movement of infected: high \rightarrow low risk

Turnover decreases high risk group equilibrium prevalence

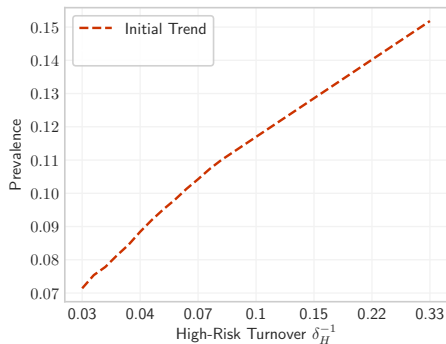


High risk prevalence vs turnover



Turnover causes a net movement of infected: high \rightarrow low risk

Turnover increases low-risk equilibrium prevalence ... at first



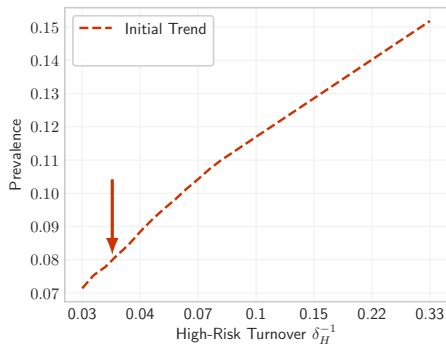
High

Low

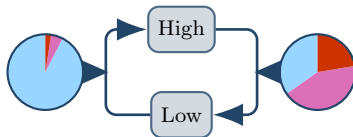
Low risk prevalence vs turnover

High turnover reduces prevalence in all groups ... but why?

Turnover increases low-risk equilibrium prevalence ... at first

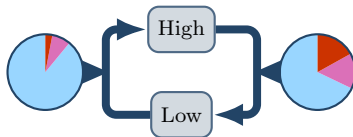
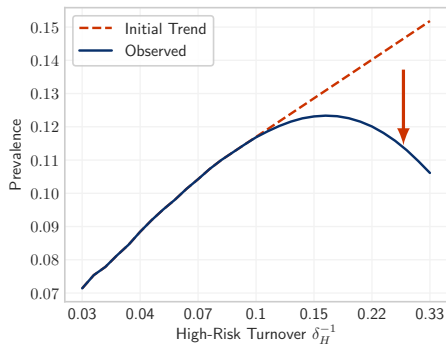


Low risk prevalence vs turnover



High turnover reduces prevalence in all groups ... but why?

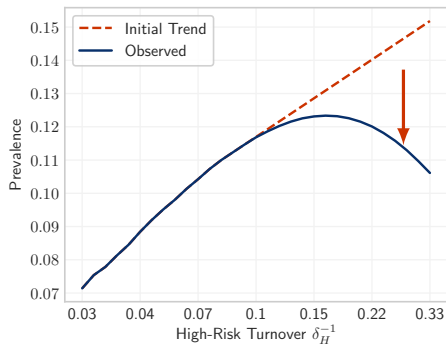
Turnover increases low-risk equilibrium prevalence ... at first



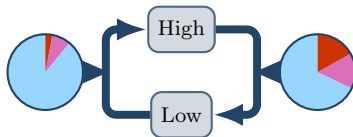
Low risk prevalence vs turnover

High turnover reduces prevalence in all groups ... but why?

Turnover increases low-risk equilibrium prevalence ... at first

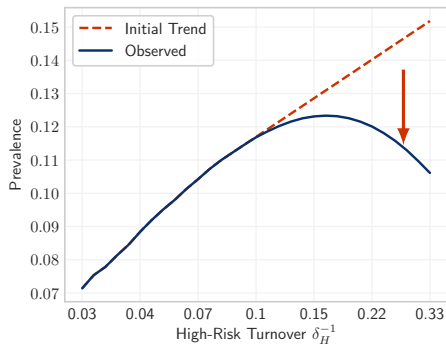


Low risk prevalence vs turnover

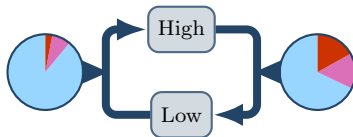


High turnover reduces prevalence in all groups ... but why?

Turnover increases low-risk equilibrium prevalence ... at first

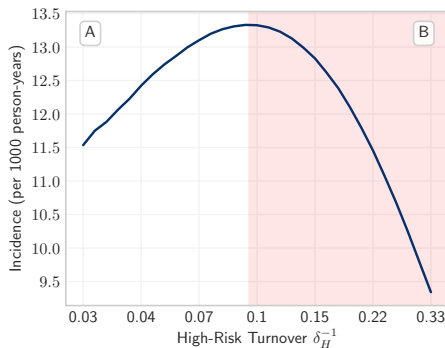


Low risk prevalence vs turnover



High turnover reduces prevalence in all groups ... but why?

Two competing effects of turnover on incidence



Overall incidence vs turnover

► Turnover \uparrow proportion who are infectious

Dominates at low turnover (A)

Incidence \uparrow

► Turnover \downarrow contact rate among infectious

Dominates at high turnover (B)

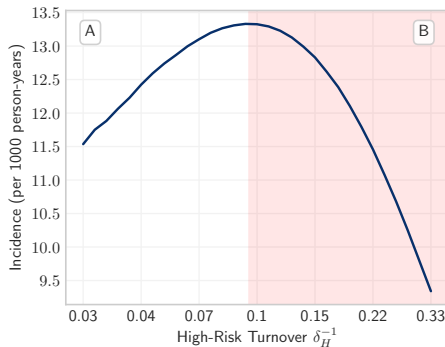
Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

► Turnover \uparrow proportion who are infectious

► Dominates at low turnover (A)

► Incidence \uparrow

► Turnover \downarrow contact rate among infectious

► Dominates at high turnover (B)

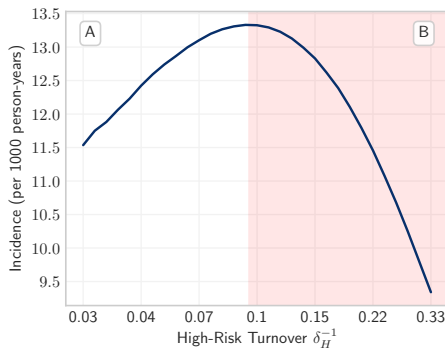
► Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

► Turnover \uparrow proportion who are infectious

► Dominates at low turnover (A)

► Incidence \uparrow

► Turnover \downarrow contact rate among infectious

► Dominates at high turnover (B)

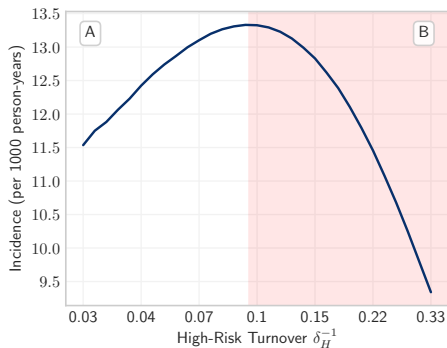
► Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

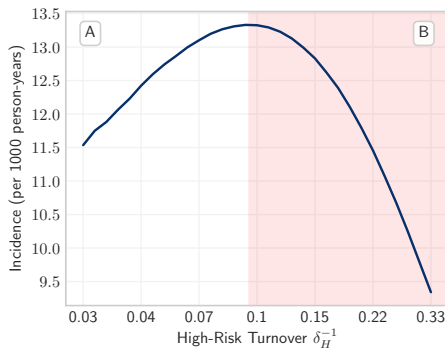
- ▶ Turnover \uparrow proportion who are infectious
 - ▶ Dominates at low turnover (A)
 - ▶ Incidence \uparrow
- ▶ Turnover \downarrow contact rate among infectious
 - ▶ Dominates at high turnover (B)
 - ▶ Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

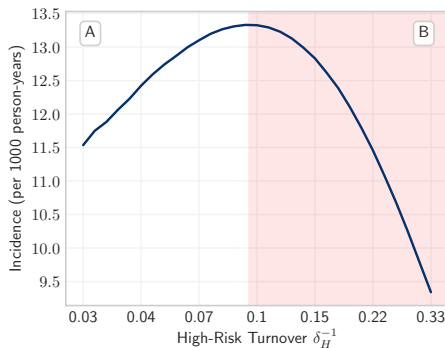
- ▶ Turnover \uparrow proportion who are infectious
 - ▶ Dominates at low turnover (A)
 - ▶ Incidence \uparrow
- ▶ Turnover \downarrow contact rate among infectious
 - ▶ Dominates at high turnover (B)
 - ▶ Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

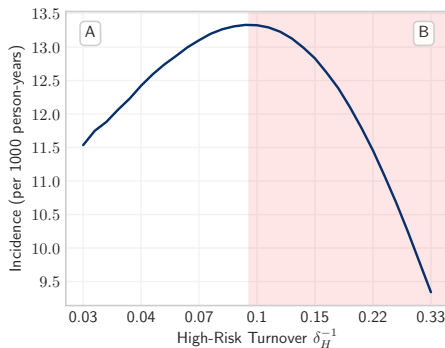
- ▶ Turnover \uparrow proportion who are infectious
 - ▶ Dominates at low turnover (A)
 - ▶ Incidence \uparrow
- ▶ Turnover \downarrow contact rate among infectious
 - ▶ Dominates at high turnover (B)
 - ▶ Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

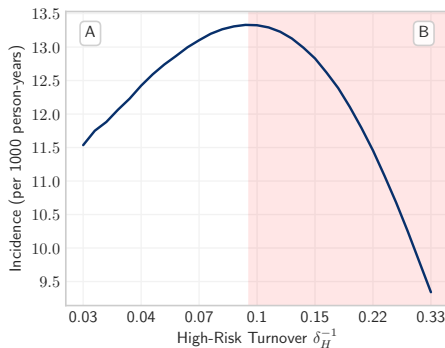
- ▶ Turnover \uparrow proportion who are infectious
 - ▶ Dominates at low turnover (A)
 - ▶ Incidence \uparrow
- ▶ Turnover \downarrow contact rate among infectious
 - ▶ Dominates at high turnover (B)
 - ▶ Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

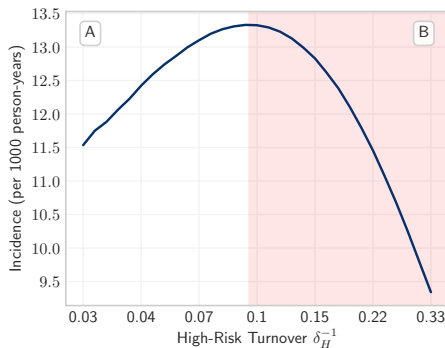
- ▶ Turnover \uparrow proportion who are infectious
 - ▶ Dominates at low turnover (A)
 - ▶ Incidence \uparrow
- ▶ Turnover \downarrow contact rate among infectious
 - ▶ Dominates at high turnover (B)
 - ▶ Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

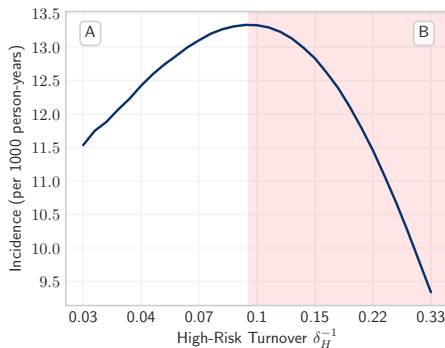
- ▶ Turnover \uparrow proportion who are infectious
 - ▶ Dominates at low turnover (A)
 - ▶ Incidence \uparrow
- ▶ Turnover \downarrow contact rate among infectious
 - ▶ Dominates at high turnover (B)
 - ▶ Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Two competing effects of turnover on incidence



Overall incidence vs turnover

- ▶ Turnover \uparrow proportion who are infectious
 - ▶ Dominates at low turnover (A)
 - ▶ Incidence \uparrow
- ▶ Turnover \downarrow contact rate among infectious
 - ▶ Dominates at high turnover (B)
 - ▶ Incidence \downarrow

(explains low risk prevalence decline with high turnover)

Low turnover: \uparrow incidence

High turnover: \downarrow incidence

Experiments: Influence of Turnover on Model Outputs

1. Equilibrium outputs:

- ▶ **Vary:** Turnover magnitude
- ▶ **Compare:** a) prevalence, b) incidence (by risk group, at equilibrium)

2. TPAF after model fitting:

- ▶ **Fit:** Contact rates: High Risk; and Low Risk
- ▶ **Targets:** Prevalence: 25% in High Risk; and 5% in Low Risk
- ▶ **Vary:** No-turnover vs Turnover
- ▶ **Compare:** a) Fitted contact rates, b) TPAF of high risk group

Turnover makes it “harder” to observe a high prevalence ratio

		No turnover	Turnover
Prevalence	High risk	25%	25%
	Low risk	5%	5%
	Ratio	5.0	5.0
Contact rate	High risk	15.8	16.9
	Low risk	2.49	0.28
	Ratio	6.3	60

To observe the same prevalence ratio:

Risk heterogeneity must be higher with turnover than without
(overcome “homogenizing” effect of turnover)

Turnover makes it “harder” to observe a high prevalence ratio

		No turnover	Turnover
Prevalence	High risk	25%	25%
	Low risk	5%	5%
	Ratio	5.0	5.0
Contact rate	High risk	15.8	16.9
	Low risk	2.49	0.28
	Ratio	6.3	60

To observe the same prevalence ratio:

Risk heterogeneity must be higher with turnover than without
(overcome “homogenizing” effect of turnover)

Turnover makes it “harder” to observe a high prevalence ratio

		No turnover	Turnover
Prevalence	High risk	25%	25%
	Low risk	5%	5%
	Ratio	5.0	5.0
Contact rate	High risk	15.8	16.9
	Low risk	2.49	0.28
	Ratio	6.3	60

To observe the same prevalence ratio:

Risk heterogeneity must be higher with turnover than without
(overcome “homogenizing” effect of turnover)

Turnover makes it “harder” to observe a high prevalence ratio

		No turnover	Turnover
Prevalence	High risk	25%	25%
	Low risk	5%	5%
	Ratio	5.0	5.0
Contact rate	High risk	15.8	16.9
	Low risk	2.49	0.28
	Ratio	6.3	60

To observe the same prevalence ratio:

Risk heterogeneity must be higher with turnover than without
(overcome “homogenizing” effect of turnover)

Turnover makes it “harder” to observe a high prevalence ratio

		No turnover	Turnover
Prevalence	High risk	25%	25%
	Low risk	5%	5%
	Ratio	5.0	5.0
Contact rate	High risk	15.8	16.9
	Low risk	2.49	0.28
	Ratio	6.3	60

To observe the same prevalence ratio:

Risk heterogeneity must be higher with turnover than without
(overcome “homogenizing” effect of turnover)

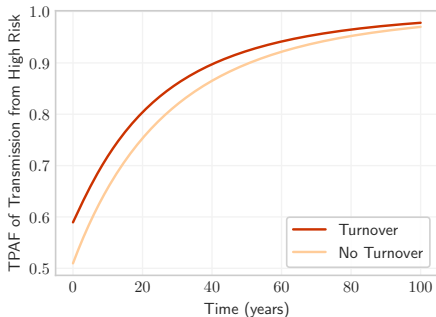
Turnover makes it “harder” to observe a high prevalence ratio

		No turnover	Turnover
Prevalence	High risk	25%	25%
	Low risk	5%	5%
	Ratio	5.0	5.0
Contact rate	High risk	15.8	16.9
	Low risk	2.49	0.28
	Ratio	6.3	60

To observe the same prevalence ratio:

Risk heterogeneity must be higher with turnover than without
(overcome “homogenizing” effect of turnover)

TPAF of high risk group is higher with turnover than without

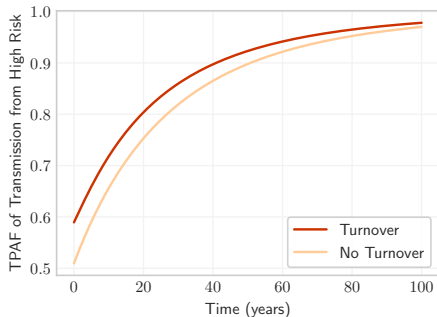


TPAF \approx impact of perfect TasP in one group

- ▶ Risk heterogeneity (contact rate ratio) is higher with turnover
- ▶ Impact of reaching high risk group is higher with turnover

TPAF of high risk group may be underestimated if turnover is not modelled

TPAF of high risk group is higher with turnover than without

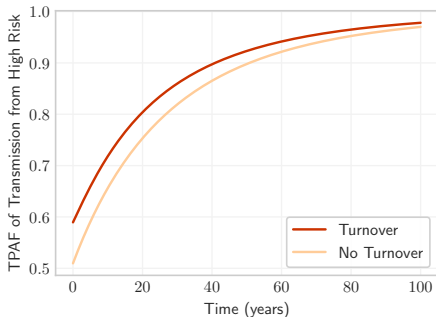


TPAF \approx impact of perfect TasP in one group

- Risk heterogeneity (contact rate ratio) is higher with turnover
- Impact of reaching high risk group is higher with turnover

TPAF of high risk group may be underestimated if turnover is not modelled

TPAF of high risk group is higher with turnover than without

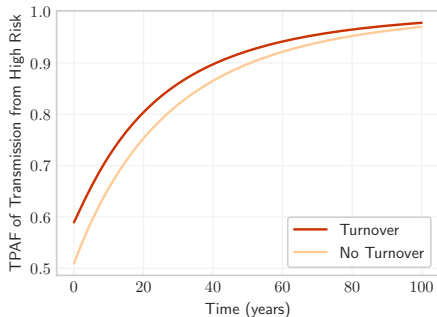


TPAF \approx impact of perfect TasP in one group

- ▶ Risk heterogeneity (contact rate ratio) is higher with turnover
- ▶ Impact of reaching high risk group is higher with turnover

TPAF of high risk group may be underestimated if turnover is not modelled

TPAF of high risk group is higher with turnover than without



TPAF \approx impact of perfect TasP in one group

- ▶ Risk heterogeneity (contact rate ratio) is higher with turnover
- ▶ Impact of reaching high risk group is higher with turnover

TPAF of high risk group may be underestimated if turnover is not modelled

Implications

Limitations:

- ▶ Results shown here conditional on model structure, assumptions, and parameters

① Turnover influences equilibrium prevalence & incidence

- ▶ Core prevalence always decreases (before fitting)
- ▶ Overall effect varies with context

② TPAF of high risk group may be underestimated if turnover is not modelled

- ▶ Prevalence ratios we observe are likely *in spite of* homogenizing effect of turnover

Implications

Limitations:

- ▶ Results shown here conditional on model structure, assumptions, and parameters

① Turnover influences equilibrium prevalence & incidence

- ▶ Core prevalence always decreases (before fitting)
- ▶ Overall effect varies with context

② TPAF of high risk group may be underestimated if turnover is not modelled

- ▶ Prevalence ratios we observe are likely *in spite of* homogenizing effect of turnover

Implications

Limitations:

- ▶ Results shown here conditional on model structure, assumptions, and parameters

① Turnover influences equilibrium prevalence & incidence

- ▶ Core prevalence always decreases (before fitting)
- ▶ Overall effect varies with context

② TPAF of high risk group may be underestimated if turnover is not modelled

- ▶ Prevalence ratios we observe are likely *in spite of* homogenizing effect of turnover

Implications

Limitations:

- ▶ Results shown here conditional on model structure, assumptions, and parameters

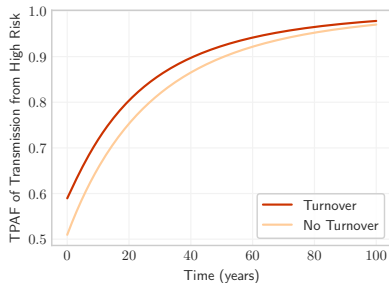
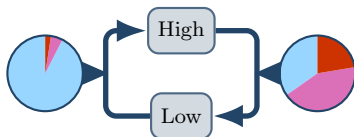
① Turnover influences equilibrium prevalence & incidence

- ▶ Core prevalence always decreases (before fitting)
- ▶ Overall effect varies with context

② TPAF of high risk group may be underestimated if turnover is not modelled

- ▶ Prevalence ratios we observe are likely *in spite of* homogenizing effect of turnover

Thank you



References

- [1] Hein Stigum, W. Falck, and P. Magnus. “The core group revisited: The effect of partner mixing and migration on the spread of gonorrhea, chlamydia, and HIV”. In: **Mathematical Biosciences** 120.1 (1994), pp. 1–23.
- [2] Marie Claude Boily and Benoît Mâsse. “Mathematical models of disease transmission: A precious tool for the study of sexually transmitted diseases”. In: **Canadian Journal of Public Health** 88.4 (1997), pp. 255–265.
- [3] Sharmistha Mishra et al. “Data and methods to characterize the role of sex work and to inform sex work programs in generalized HIV epidemics: evidence to challenge assumptions”. In: **Annals of Epidemiology** 26.8 (2016), pp. 557–569.
- [4] Xinyu Zhang et al. “Episodic HIV Risk Behavior Can Greatly Amplify HIV Prevalence and the Fraction of Transmissions from Acute HIV Infection”. In: **Statistical Communications in Infectious Diseases** 4.1 (2012).
- [5] Shah Jamal Alam et al. “Detectable signals of episodic risk effects on acute HIV transmission: Strategies for analyzing transmission systems using genetic data”. In: **Epidemics** 5.1 (2013), pp. 44–55.
- [6] Christopher J. Henry and James S. Koopman. “Strong influence of behavioral dynamics on the ability of testing and treating HIV to stop transmission”. In: **Scientific Reports** 5.1 (2015), p. 9467.