

Bayesian spatio-temporal methods for
small-area estimation of HIV indicators

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Statement of Originality

This thesis, and the work presented in it, is work that I conducted myself. In all cases where I describe others' work, I provide appropriate references.

For someone, or something

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Abstract

Progress towards ending AIDS as a public health threat by 2030 is faltering. Effective public health response requires accurate, timely, high-resolution estimates of epidemic and demographic indicators. Limitations of available data make obtaining these estimates difficult. I develop and apply Bayesian spatio-temporal methods to meet this challenge. First, I examine models for area-level spatial structure. Second, I estimate district-level HIV risk group proportions, enabling behavioural prioritisation of prevention services, as put forward in the Global AIDS Strategy. Finally, I develop a novel Bayesian inference method, combining adaptive Gauss-Hermite quadrature with principal component analysis, motivated by the Naomi district-level model of HIV indicators. Together, the contributions in this thesis help to guide precision HIV policy in sub-Saharan Africa, as well as advancing Bayesian methods for spatio-temporal data.

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List of Abbreviations

HIV	Human Immunodeficiency Virus.
AIDS	Acquired ImmunoDeficiency Syndrome.
PEPFAR	President’s Emergency Plan for AIDS Relief.
HIV	Demographic and Health Surveys.
AIS	AIDS Indicator Survey.
PrEP	Pre-Exposure Prophylaxis.
PEP	Post-Exposure Prophylaxis.
FSW	Female Sex Worker(s).
MSM	Men who have Sex with Men.
PWID	People Who Inject Drugs.
ANC	Antenatal Clinic.
UNAIDS	United Nations Joint Programme on HIV/AIDS.
CDC	Centers for Disease Control and Prevention.
UAT	Unlinked Anonymous Testing.
PMTCT	Prevention of Mother-to-Child Transmission.
PLHIV	People Living with HIV.
MCMC	Markov Chain Monte Carlo.
VI	Variational Inference.
INLA	Integrated Nested Laplace Approximation.
GP	Gaussian Process.
CAR	Conditionally Auto-regressive.
ART	Antiretroviral Therapy.
SAE	Small Area Estimation.
GMRF	Gaussian Markov Random Field.
HMC	Hamiltonian Monte Carlo.

List of Abbreviations

GMRF	Gaussian Markov Random Field.
HMC	Hamiltonian Monte Carlo.
LGM	Latent Gaussian Model.
ELGM	Extended Latent Gaussian Model.

List of Notations

ρ	HIV prevalence.
λ	HIV incidence.
α	ART coverage.
\mathcal{S}	Spatial study region $\mathcal{S} \subseteq \mathbb{R}^2$.
$s \in \mathcal{S}$	Point location.
\mathcal{T}	Temporal study period $\mathcal{T} \subseteq \mathbb{R}$.
$t \in \mathcal{T}$	Time.
\mathbf{y}	Data, a n -vector.
ϕ	Parameters, a d -vector (ϕ_1, \dots, ϕ_d) .
\mathbf{x}	Latent field, a N -vector (x_1, \dots, x_N) .
θ	Hyperparameters, a m -vector $(\theta_1, \dots, \theta_m)$.
$x \sim p(x)$	x is distributed according to $p(x)$.
A_i	Areal unit.
$A_i \sim A_j$	Adjacency between areal units.
\mathbf{H}	Hessian matrix.
\mathbf{R}	Structure matrix.
\mathbf{Q}	Precision matrix.
Σ	Covariance matrix.
\mathcal{N}	Gaussian distribution.
$k : \mathcal{X} \times \mathcal{X} \rightarrow \mathbb{R}$	Kernel function on the space \mathcal{X} .
$A_i \sim A_j$	Adjacency between areal units.
\mathcal{Q}	A set of quadrature nodes.
$\omega : \mathcal{Q} \rightarrow \mathbb{R}$	A quadrature weighting function.
$\mathcal{Q}(m, k)$	Gauss-Hermite quadrature points in m dimensions with k nodes per dimension, constructed according to a product rule.

1

Introduction

This thesis is about applied and methodological Bayesian statistics. It is applied and methodological in that I am concerned first with real world questions and second with the means to answer them. It is Bayesian in that I arrive at conclusions based on data using probability models.

The real world questions relate to surveillance of the human immunodeficiency virus (HIV) epidemic in sub-Saharan Africa (SSA). The Global Burden of Disease Collaborative Network (2019) found HIV to be the largest cause of disability adjusted life years (DALYs) in SSA among those one year and older (Figure 1.1). Using statistics to quantify the epidemic is an important part of the public health response, and the path towards disease control and elimination. However, there are numerous challenges involved in obtaining suitable estimates of relevant indicators.

The estimates in this thesis are based on data recorded from national household surveys or routinely collected from healthcare facilities. An important feature of this data is its location and the time at which it was collected. While diverse, spatio-temporal data have distinctive commonalities. I draw on modelling techniques well suited for this kind of data.

Computation is an essential part of modern statistical practice. Each project in this thesis, as well as the thesis itself, is accompanied by R code, hosted on GitHub.

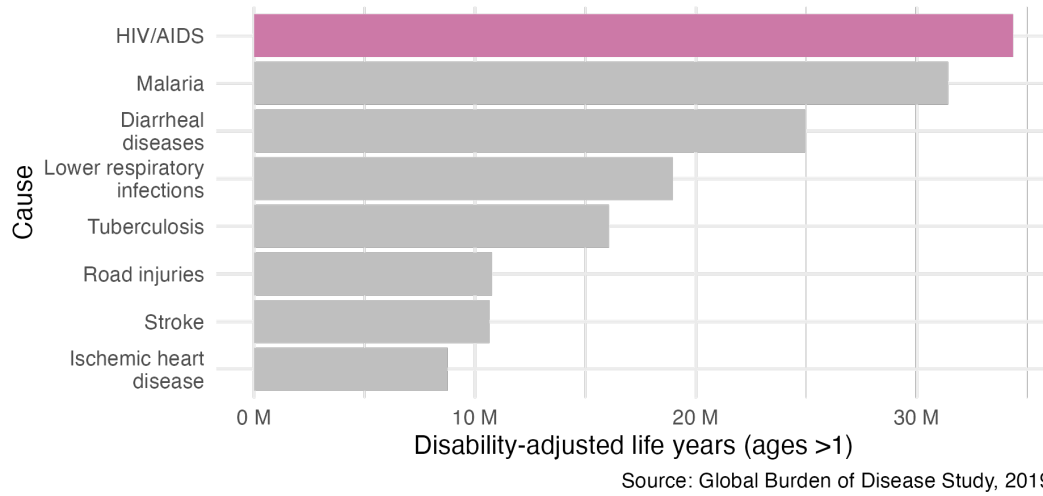


Figure 1.1: HIV/AIDS is the largest cause of DALYs for non-infects (>1 years) in SSA. One DALY represents the loss of the equivalent of one year of full health, and is calculated by the sum of years of life lost and years lost due to disability. The disability weights used in the 2019 Global Burden of Disease study vary depending on severity of the condition.

1.1 Chapter overview

The structure of this thesis (Figure 1.2) is as follows:

- Chapter 2: I begin by providing background on the HIV/AIDS epidemic, as well as describing the challenges faced by surveillance efforts.
- Chapter 3: I then introduce the statistical concepts and notation used throughout the thesis, focusing on Bayesian modelling and computation.
- Chapter 4: The prevailing model for spatial structure used in small-area estimation (Besag et al. 1991) was designed with analysis of a grid of pixels in mind. In disease mapping, we work using the districts of a country, which are not a grid. I evaluate the practical consequences of this concern (Howes, Eaton, et al. 2023+).
- Chapter 5: Adolescent girls and young women are a demographic group at disproportionate risk of HIV infection. The Global AIDS Strategy suggests prioritising interventions on the basis of behaviour to prevent the most new infections using available resources. I estimate the size of behavioural risk groups across priority countries to enable implementation of this strategy, and

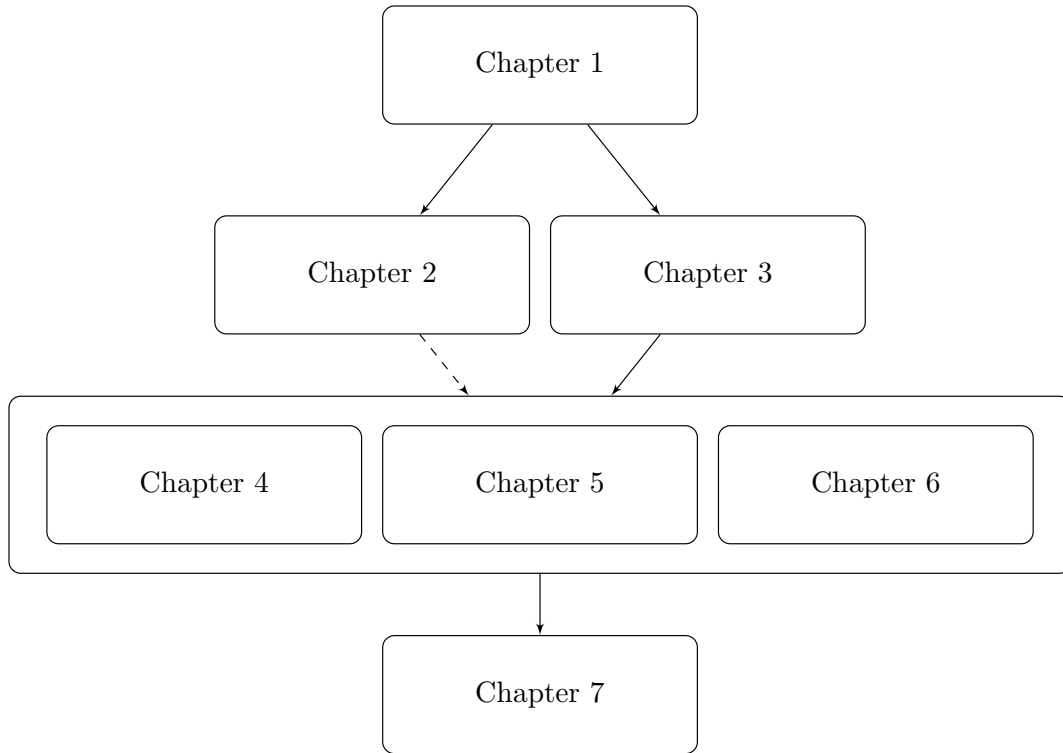


Figure 1.2: The chapters in this thesis are structured to depend on each other. The dashed line represents a recommended, but not required dependency. Though chronological order is recommended, Chapter 4, Chapter 5 and Chapter 6 may be read in any order as, for the most part, they correspond to separable research projects.

assess the potential benefits in terms of numbers of new infections prevented (Howes, Risher, et al. 2023).

- Chapter 6: The Naomi small-area estimation model (Eaton et al. 2021) is used by countries to estimate district-level HIV indicators. Motivated by this model, I develop an approximate Bayesian inference method combining adaptive Gauss-Hermite quadrature with principal components analysis (Howes, Stringer, et al. 2023+). I apply the method to data from Malawi, and analyse the consequences of inference method choice for policy relevant outcomes. Further, I open the door to a new class of fast, flexible, and accurate Bayesian inference algorithms.
- Chapter 7: Finally, I discuss avenues for future work, and my conclusions regarding the research, as well as its strengths and weaknesses.

2

The HIV/AIDS epidemic

2.1 Background

HIV is a retrovirus which infects humans. If untreated, HIV can develop into a more advanced stage known as acquired immunodeficiency syndrome (AIDS). HIV primarily attacks a type of white blood cell vital for the function of the immune system. As a result, AIDS is characterised by increased risk of developing opportunistic infections such as tuberculosis or *Pneumocystis* pneumonias.

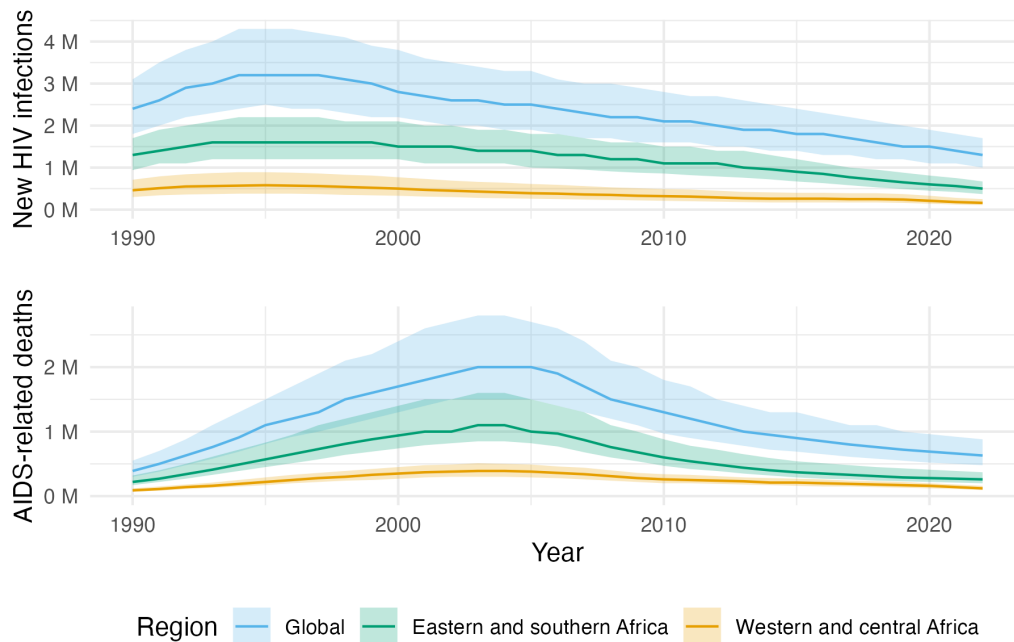
The first AIDS cases were reported in Los Angeles in the early 1980s. Since then, HIV has spread globally. Transmission occurs by exposure to specific bodily fluids of an infected person. The most common mode of transmission is via unprotected anal or vaginal sex, though transmission can also occur from a mother to her baby, or when drug injection equipment is shared. Approximately 86 million people have become infected with HIV, and of those 40 million have died of AIDS-related causes.

A major international effort has been made to address the epidemic, including by national governments, civil society organisations, pharmaceutical companies, research institutions, international agencies such as the Joint United Nations Programme on HIV/AIDS (UNAIDS), and global health initiatives such as the President's Emergency Plan for AIDS Relief (PEPFAR; \$85 billion invested) and the Global Fund to Fight AIDS, Tuberculosis, and Malaria (GFATM; \$29 billion

invested). A number of interventions have had a substantial impact on the course of the epidemic. These include:

- Antiretroviral therapy (ART) is a drug which stops the virus from replicating in the body. By taking ART daily, a person living with HIV can live a full and healthy life. Of the 39 million people living with HIV (PLHIV) in 2022, around 76% were accessing ART. ART is estimated to have averted 16.5 million AIDS-related deaths. Further, ART reduces the amount of virus in the blood and genital secretions. If the virus is undetectable then there is considerable evidence that it cannot be transmitted sexually (Cohen et al. 2011). For this reason, as well as saving lives, treatment also operates as prevention (TaSP).
- Condoms are an inexpensive and effective method for prevention of HIV and other sexually transmitted infections (STIs) such as *Chlamydia trachomatis*, *Neisseria gonorrhoeae*, syphilis, and *Trichomonas vaginalis*. There has been a significant scale-up of condom usage since 1990, which is estimated to have averted 117 million new HIV infections (Stover and Teng 2021). Even so, there remain significant and difficult to close gaps in the proportion of individuals reporting they used a condom during their last high-risk sexual encounter.
- Voluntary medical male circumcision (VMMC) has been found to provide partial protection against female-to-male HIV transmission. Complete surgical removal of the foreskin has been found to result in a reduction of HIV incidence in men by 50-60% (Auvert et al. 2005; Gray et al. 2007; Bailey et al. 2007). Based on this evidence, since 2007 VMMC has been recommended by WHO and UNAIDS as a key HIV intervention in high-prevalence settings. Scale up of VMMC across 15 priority countries between 2008 and 2019 was estimated to have already averted 340 thousand new HIV infections, though the future number of new HIV infections averted is likely to be much higher.
- Prevention of mother-to-child transmission

The HIV/AIDS epidemic



Source: UNAIDS epidemiological estimates, 2023

Figure 2.1: Globally, yearly new HIV infections peaked in 1995, and have since decreased by 59% and yearly AIDS-related deaths peaked in 2004, and have since decreased by 68%. Much of the disease burden is concentrated in eastern and southern Africa, as well as western and central Africa. These two regions combined comprise sub-Saharan Africa.

- Pre-exposure prophylaxis (PrEP) and post-exposure prophylaxis (PEP) are drugs which can be taken before and after exposure to prevent transmission. They are primarily useful in high risk settings.

The interventions above have had a significant impact on the course of the epidemic. Both the number of new HIV infections and number of AIDS-related deaths per year have fallen significantly since their peak (Figure 2.1). That said, there remains quite some distance to go in ending AIDS as a public health threat. In 2022 there were 1.3 million people newly infected with HIV and 630 thousand AIDS-related deaths (UNAIDS 2022).

There is substantial geographic variation in disease burden, across multiple spatial scales. In some countries, the epidemic is concentrated within small populations, and HIV prevalence is low. In others, transmission is sustained in the general population, and HIV prevalence is higher. Most countries severely affected

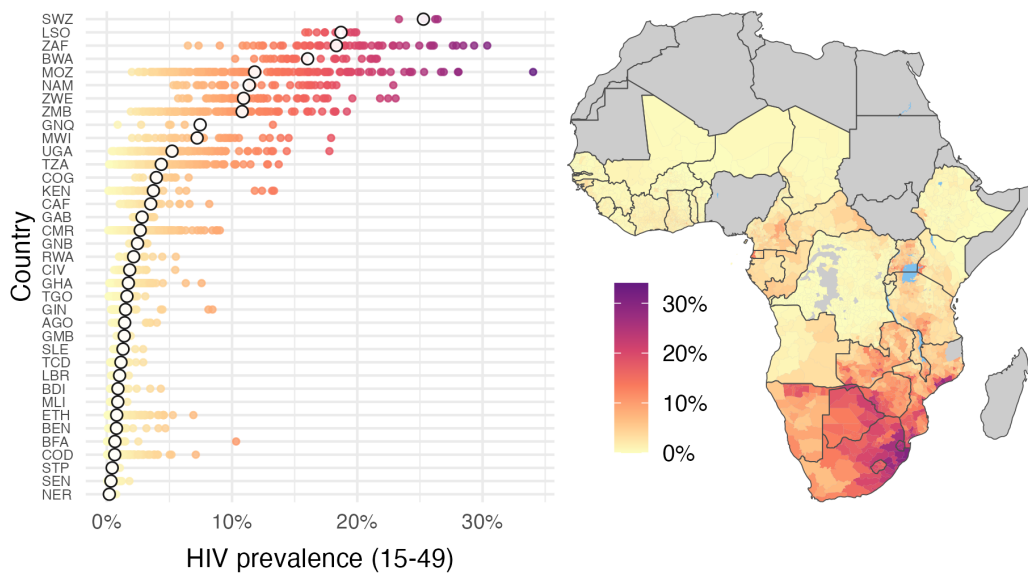


Figure 2.2: Adult (15-49) HIV prevalence varies substantially both within and between countries in SSA. These 2023 estimates were generated by country teams using the Naomi small-area estimation model in a process supported by UNAIDS and are available from aidsinfo.unaids.org. White filled points are country-level estimates, and coloured points are district-level estimates. Results from Nigeria are yet to be approved and have been redacted. The estimates process in the Cabo Delgado province of Mozambique was disrupted by conflict. Obtaining results Democratic Republic of the Congo required removing some districts from the model. Country names are given by three-letter codes as published by the International Organization for Standardization (ISO).

by HIV are in sub-Saharan Africa (SSA). It is estimated that 66% of the 39 million PLHIV worldwide live in SSA. For some countries in southern Africa, adult HIV prevalence (ages 15-49) is as high as 10% or more (Figure 2.2). As well as variation at between countries, there is significant variation within countries at the district-level. For example, adult HIV prevalence in South Africa ranges from 7% to 32%.

In all countries, some groups of people are at much higher risk than others. Groups of people at increased risk of HIV infection are known as key populations (KPs). Examples include men who have sex with men (MSM), female sex workers (FSW), people who inject drugs (PWID), and transgender people (TGP). KPs are often marginalised, and face legal and social issues. In concentrated settings, the majority of new HIV infections occur among KPs. In generalised settings like

SSA, risk is more diffuse accross the population. For example, adolescent girls and young women (AGYW) are a large demographic group at increased risk of HIV infection in SSA (Risher et al. 2021; Monod et al. 2023).

For available resources to have the greatest impact, it is important that HIV interventions are prioritised (Ord 2013). The precision public health paradigm (Khoury et al. 2016) looks to provide the right intervention, to the right population, in the right place, at the right time.

2.2 HIV surveillance

HIV surveillance refers to the collection, analysis, interpretation and dissemination of data relating to HIV/AIDS. Surveillance can used to track epidemic indicators, identify at-risk populations, find drivers of transmission, and evaluate the impact of prevention and treatment programs.

2.2.1 HIV indicators

Central indicators include:

- **HIV prevalence** is the proportion $\rho \in [0, 1]$ of the population who have HIV, typically written as a percentage. Interpreted alone, HIV prevalence can be misleading. HIV prevalence increasing could be as a result of new infections, or as a result of more PLHIV remaining alive by taking treatment. As such, HIV prevalence is primarily used in calculations of other indicators, rather than directly itself. In some circumstances those other indicators can be difficult to estimate, and HIV prevalence can be a useful proxy. The number of PLHIV is given by $N\rho$, where N is the population size.
- **HIV incidence** is the rate $\lambda \in \mathbb{R}$ of new HIV infections, typically written as number of new infections per 1000 person years. HIV incidence can be specified in terms of HIV prevalence by $\lambda = N\Delta\rho$, where $\Delta\rho$ is the change in HIV prevalence over some time period. Planning, delivery, and evaluation of prevention programming relies on estimates of HIV incidence.

- **ART coverage** is the proportion $\alpha \in [0, 1]$ of PLHIV who are on ART, typically written as a percentage. Estimates of ART coverage play a direct role in the provision of treatment services.

2.2.2 Challenges

There are significant difficulties associated with obtaining reliable, timely, estimates of these indicators at an appropriate resolution.

1. **Data sparsity** Collection of data is costly and time consuming. As a result, limited direct data might be available for the particular time, location, and sub-population of interest.
2. **Response biases** Under-reporting. Non-response. Measurement error.
3. **Hard to reach populations** Many of those at greatest risk, including KPs, are difficult to reach. May not appear on household survey sampling frames.
4. **Denominators and demography** Many indicators are rates or proportions, incorporating the population at risk in the denominator. However, accurately estimating these denominators is usually itself a challenging task, and taking a ratio of uncertain quantities only goes to amplify the uncertainty. For example, ART coverage has PLHIV as the denominator, which itself has the population size as its denominator.

2.2.3 Statistical methods

The above challenges make direct interpretation of data misleading, and foreground the importance of careful statistical modelling.

1. **Borrowing information** Although survey responses from women aged 20-24 are not directly informative about HIV prevalence in women 25-29, they are indirectly informative.
2. **Evidence synthesis** Multiple sources of evidence can be combined to overcome limitations of any one data source.

3. **Expert guidance** Expert epidemiological, demographic opinion can be used to improve estimates.

2.2.4 Future trends

Aims for HIV response going forward, and surveillance capabilities are needed to meet them.

1. **Greater reliance on routine health system data** Phasing out of nationally representative household surveys.
2. **Integration with other health programs**
3. **Use of mobile technologies**

3

Bayesian spatio-temporal statistics

3.1 Bayesian statistics

Bayesian statistics is a mathematical paradigm for learning from data. It is well suited for attending to the challenges in Section 2.2 because it allows principled, flexible incorporation of scientific knowledge. In this section I provide brief overview. For a more complete introduction, I recommend McElreath (2020) or Gelman, Carlin, et al. (2013).

3.1.1 Bayesian modelling

At its best, the Bayesian paradigm allows the analyst focus on how best to model the data. This is achieved by the construction of a generative model $p(\mathbf{y}, \boldsymbol{\phi})$ for the observed data $\mathbf{y} = (y_1, \dots, y_n)$ together with parameters $\boldsymbol{\phi} = (\phi_1, \dots, \phi_d)$, where n is the dimension of the data and d is the number of parameters. The model is generative in the sense that one can simulate from it to generate draws

$$(\mathbf{y}, \boldsymbol{\phi}) \sim p(\mathbf{y}, \boldsymbol{\phi}) \tag{3.1}$$

If these draws differ too greatly from what the analyst would expect, then the generative model does not capture their scientific understanding, and can be refined. In this way, models can be built iteratively, with complexity added gradually.

The model is usually constructed from two parts, known respectively as the likelihood $p(\mathbf{y} | \boldsymbol{\phi})$ and the prior distribution $p(\boldsymbol{\phi})$ whereby the joint distribution is obtained by the product $p(\mathbf{y}, \boldsymbol{\phi}) = p(\mathbf{y} | \boldsymbol{\phi})p(\boldsymbol{\phi})$. The likelihood, as a function of $\boldsymbol{\phi}$ with \mathbf{y} fixed, reflects the probability of observing the data when the true value of the parameters is $\boldsymbol{\phi}$. The prior distribution encapsulates beliefs about the parameters $\boldsymbol{\phi}$ before the data is observed.

Recommendations for specifying the prior distribution vary. A central issue is the extent to which subjective information should be incorporated into the prior distribution, and thereby influence the posterior distribution. Proponents of the objective Bayesian paradigm suggest that the prior distribution should be non-informative. That said, we shall see that the distinction between likelihood and prior distribution can be blurred (Section 3.3). As such, it may be argued that issues of subjectivity are not unique to the prior distribution, and ultimately the challenge of specifying the data generating process is better thought of more holistically (Gelman, Simpson, et al. 2017).

3.1.2 Bayesian computation

The posterior distribution $p(\boldsymbol{\phi} | \mathbf{y})$ encapsulates probabilistic beliefs about the parameters given the observed data. Using the eponymous Bayes' theorem, it is given by

$$p(\boldsymbol{\phi} | \mathbf{y}) = \frac{p(\mathbf{y} | \boldsymbol{\phi})p(\boldsymbol{\phi})}{p(\mathbf{y})}. \quad (3.2)$$

Unfortunately, it is usually intractable to calculate Equation 3.2 directly because of the integral $p(\mathbf{y}) = \int p(\mathbf{y}, \boldsymbol{\phi})d\boldsymbol{\phi}$ in the denominator, sometimes known as the marginal likelihood or evidence. As such, although the numerator is proportional to the posterior distribution $p(\boldsymbol{\phi} | \mathbf{y}) \propto p(\mathbf{y} | \boldsymbol{\phi})p(\boldsymbol{\phi})$ and easy to evaluate, it is typically difficult to evaluate the posterior distribution itself.

A great variety of computational methods have been developed to tackle this problem. Briefly, the main categories are:

1. **Sampling algorithms** Also known as Monte Carlo algorithms, these approaches look to generate samples from the posterior distribution. The most popular is Markov chain Monte Carlo (MCMC), which proceeds by simulating from a Markov chain with the posterior distribution as its stationary distribution. In this thesis, I make use of the No-U-Turn sampler (NUTS), a Hamiltonian Monte Carlo (HMC) algorithm, implemented in the Stan probabilistic programming language (PPL).
2. **Variational Bayes** In variational Bayes, the posterior distribution is assumed to belong to a particular class of functions and use optimisation to choose the best member of that class.

3.1.3 Interplay between modelling and computation

Bayesian computation aspires to abstract away calculation of the posterior distribution from the analyst. Modern computational techniques and software have made this aspiration a reality for many models. However, computation of the posterior distribution remains intractable for a majority of models. As such, the analyst need not only to be concerned with choosing a model suitable for the data, but also choosing a model for which the posterior distribution is tractable in reasonable time. As such, there is an important interplay between modelling and computation, wherein models are bound by the limits of computation. As computation improves, the space of models available to the analyst expands.

3.2 Spatio-temporal statistics

In spatio-temporal statistics (Cressie and Wikle 2015), we observe data indexed by spatial or temporal location. In this thesis we assume that the spatial study region $\mathcal{S} \subseteq \mathbb{R}^2$ has two dimensions, corresponding to latitude and longitude. Data may be associated to a point $s \in \mathcal{S}$ or area $A \subseteq \mathcal{S}$ in the study region. The temporal study period $\mathcal{T} \subseteq \mathbb{R}$ can more generally be assumed to be one dimensional. Similarly, data may be associated to a point $t \in \mathcal{T}$ or period of time $T \in \mathcal{T}$.

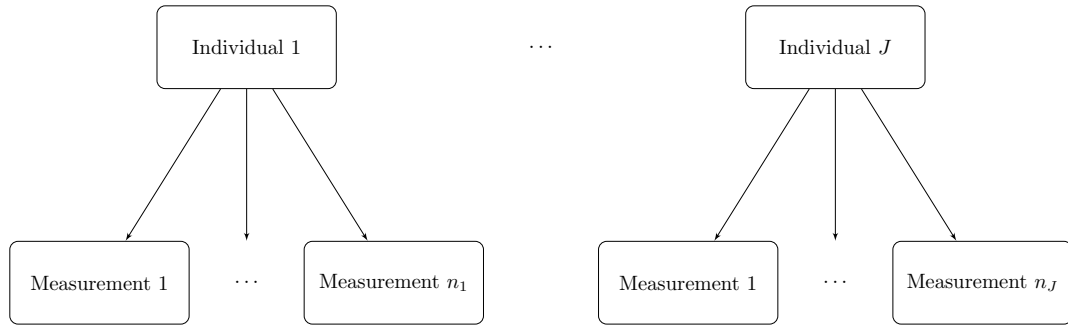


Figure 3.1: Data often has a hierarchical structure.

Spatio-temporal data has some important properties. Foremost among them is correlation structure. Tobler’s first law of geography, earlier expressed by Fisher (1936), is that “everything is related to everything else, but near things are more related than distant things” (Tobler 1970). This law extends not only to space but also time.

3.3 Model classes

3.3.1 Hierarchical models

Real world data usually has hierarchical structure. For example, we might have a study (Figure 3.1) where each individual $j = 1, \dots, J$ has a group of n_j observations. Then, it would be inappropriate to treat all $\sum_j n_j$ observations as being independent and identically distributed (IID). Observations belonging to the same group, that is of the same individual, are likely to be more similar than observations from different groups, which refer to different individuals.

Bayesian hierarchical or multilevel models, comprised of multiple stages, allow handling data of this sort. For example, in a three-stage hierarchical model, we partition the parameters so that $\boldsymbol{\phi} = (\mathbf{x}, \boldsymbol{\theta})$. I refer to $\mathbf{x} = (x_1, \dots, x_N)$ as the latent field, and $\boldsymbol{\theta} = (\theta_1, \dots, \theta_m)$ as the hyperparameters. The generative

model for data \mathbf{y} is then

$$\mathbf{y} \sim p(\mathbf{y} | \mathbf{x}, \boldsymbol{\theta}), \quad (3.3)$$

$$\mathbf{x} \sim p(\mathbf{x} | \boldsymbol{\theta}), \quad (3.4)$$

$$\boldsymbol{\theta} \sim p(\boldsymbol{\theta}), \quad (3.5)$$

with posterior distribution proportional to $p(\mathbf{x}, \boldsymbol{\theta} | \mathbf{y}) \propto p(\mathbf{y} | \mathbf{x}, \boldsymbol{\theta})p(\mathbf{x} | \boldsymbol{\theta})p(\boldsymbol{\theta})$. Referring back to the previous example, we may informally think about the latent field as modelling properties of individuals, and the hyperparameters as modelling shared properties of individuals or the observation process.

3.3.2 Mixed effects models

Fixed effects refer to parts of the latent field which are constant across groups. Random effects refer to parts of the latent field which vary across groups. Though these terms have notoriously many different definitions (Gelman 2005), I nonetheless find them useful to introduce here. No pooling, partial pooling, complete pooling. Structured random effects.

3.3.3 Latent Gaussian models

Latent Gaussian models [LGMs; Rue, Martino, et al. (2009)] are a class of three-stage Bayesian hierarchical models in which, loosely speaking, the middle layer is Gaussian. More specifically, in an LGM, the likelihood is given by

$$\begin{aligned} y_i &\sim p(y_i | \eta_i, \boldsymbol{\theta}_1), \quad i = 1, \dots, n, \\ \mu_i &= \mathbb{E}(y_i | \eta_i) = g(\eta_i), \\ \eta_i &= \beta_0 + \sum_{l=1}^p \beta_l z_{li} + \sum_{k=1}^r f_k(u_{ki}). \end{aligned}$$

The likelihood is given by a product $p(\mathbf{y} | \boldsymbol{\eta}, \boldsymbol{\theta}_1) = \prod_{i=1}^n p(y_i | \eta_i, \boldsymbol{\theta}_1)$, where $\boldsymbol{\eta} = (\eta_1, \dots, \eta_n)$. Each response has conditional mean μ_i with inverse link function $g : \mathbb{R} \rightarrow \mathbb{R}$ such that $\mu_i = g(\eta_i)$. The vector $\boldsymbol{\theta}_1 \in \mathbb{R}^{s_1}$, with s_1 assumed small, are additional parameters of the likelihood. The structured additive predictor η_i may

include an intercept β_0 , fixed effects β_j of the covariates z_{ji} , and random effects $f_k(\cdot)$ of the covariates u_{ki} . The parameters β_0 , $\{\beta_j\}$, $\{f_k(\cdot)\}$ are each assigned Gaussian prior distributions, and can be collected into a vector $\mathbf{x} \in \mathbb{R}^N$ such that $\mathbf{x} \sim \mathcal{N}(\mathbf{0}, \mathbf{Q}(\boldsymbol{\theta}_2)^{-1})$ where $\boldsymbol{\theta}_2 \in \mathbb{R}^{s_2}$ are further hyperparameters, again with s_2 assumed small. Let $\boldsymbol{\theta} = (\boldsymbol{\theta}_1, \boldsymbol{\theta}_2) \in \mathbb{R}^m$ with $m = s_1 + s_2$ be all hyperparameters, with prior distribution $p(\boldsymbol{\theta})$.

Spatio-temporal data are well suited to being modelled with LGMs.

3.3.4 Extended latent Gaussian models

Some models used in small-area estimation fall outside the LGM class. Many of these models do fit into the class of extended latent Gaussian models (ELGMs) as proposed by Stringer et al. (2021). By allowing many-to-one link functions, ELGMs facilitate modelling of non-linearities. In particular, the structured additive predictor is redefined as $\boldsymbol{\eta} = (\eta_1, \dots, \eta_{N_n})$, where $N_n \in \mathbb{N}$ is a function of n , and it is possible that $N_n \neq n$. Each mean response μ_i now depends on some subset $\mathcal{J}_i \subseteq [N_n]$ of indices of $\boldsymbol{\eta}$, with $\cup_{i=1}^n \mathcal{J}_i = [N_n]$ and $1 \leq |\mathcal{J}_i| \leq N_n$, where $[N_n] = \{1, \dots, N_n\}$. The inverse link function $g(\cdot)$ is redefined for each observation to be a possibly many-to-one mapping $g_i : \mathbb{R}^{|\mathcal{J}_i|} \rightarrow \mathbb{R}$, such that $\mu_i = g_i(\boldsymbol{\eta}_{\mathcal{J}_i})$. Put together, ELGMs are then of the form

$$\begin{aligned} y_i &\sim p(y_i | \boldsymbol{\eta}_{\mathcal{J}_i}, \boldsymbol{\theta}_1), \quad i = 1, \dots, n, \\ \mu_i &= \mathbb{E}(y_i | \boldsymbol{\eta}_{\mathcal{J}_i}) = g_i(\boldsymbol{\eta}_{\mathcal{J}_i}), \\ \eta_j &= \beta_0 + \sum_{l=1}^p \beta_l z_{lj} + \sum_{k=1}^r f_k(u_{kj}), \quad j = 1, \dots, N_n, \end{aligned}$$

with latent field and hyperparameter prior distributions as in the LGM case.

The ELGM class is well suited to small-area estimation of HIV indicators. Indeed, this class of models is used throughout the thesis. While it can be transformed to an LGM using the Poisson-multinomial transformation (Baker 1994), the multinomial logistic regression model used in Chapter 5 is naturally written as an ELGM in which each observation depends on the set of structured additive predictors corresponding to each multinomial observation. In Chapter 6, the Naomi small-area estimation

model used to produce estimates of HIV indicators across over 35 countries is shown to have the features of an ELGM.

3.4 Survey methods

Survey methods concern the design and analysis of survey data (Lohr 2009). In SSA, nationally representative household surveys, such as the Demographic and Health Surveys (DHS) (**dhs**) or Population-based HIV Impact Assessment (PHIA) (PHIA project n.d.) surveys, provide the highest quality information about HIV. Talk about the DDC (Meng 2018). With regard to the DDC, discuss how survey weighting is helpful. Discuss how we weight then model following Chen et al. (2014). Discuss how we ignore clustering structure. Discuss how all of this isn't great and that someone should figure this out.

4

Models for spatial structure

In this chapter, I present an investigation of spatial random effects specifications. This investigation was motivated by a particular instance of a fundamental question encountered during model construction. Namely, should the model be augmented to capture a feature we have prior belief exists? The results are presented in Howes, Eaton, et al. (2023+).

Code for the analysis in this chapter is available from [athowes/beyond-borders](https://github.com/athowes/beyond-borders).

4.1 Background

4.2 Models based on adjacency

Spatial structure can be encoded using a symmetric relation between areas. Let $i \sim j$ if the areas A_i and A_j are adjacent or neighbouring. Adjacency is often defined by a shared border, though other choices are possible (Paciorek et al. 2013). The Besag model (Besag et al. 1991) is an improper conditional auto-regressive (ICAR) model where the full conditional distribution of the i th spatial random effect is given by

$$u_i \mid \mathbf{u}_{-i} \sim \mathcal{N} \left(\frac{1}{n_{\delta i}} \sum_{j: j \sim i} u_j, \frac{1}{n_{\delta i} \tau_u} \right), \quad (4.1)$$

where δi is the set of neighbours of A_i with cardinality $n_{\delta i} = |\delta i|$ and \mathbf{u}_{-i} is the vector of spatial random effects with the i th entry removed. The conditional mean of the random effect u_i is the average of its neighbours $\{u_j\}_{j \sim i}$ and the precision $n_{\delta i}\tau_u$ is proportional to the number of neighbours $n_{\delta i}$. By Brook's lemma (Rue and Held 2005) the set of full conditionals of the Besag model are equivalent to the Gaussian Markov random field (GMRF) given by

$$\mathbf{u} \sim \mathcal{N}(\mathbf{0}, \tau_u^{-1} \mathbf{R}^-), \quad (4.2)$$

where \mathbf{R}^- is the generalised inverse of the rank-deficient structure matrix \mathbf{R} , so-called because it defines the structure of the precision matrix, with entries

$$R_{ij} = \begin{cases} n_{\delta i}, & i = j \\ -1, & i \sim j \\ 0, & \text{otherwise.} \end{cases} \quad (4.3)$$

The Markov property arises due to the conditional independence structure $p(u_i | \mathbf{u}_{-i}) = p(u_i | \mathbf{u}_{\delta i})$ whereby each area only depends on its neighbours. This is reflected in the sparsity of \mathbf{R} such that $u_i \perp u_j | \mathbf{u}_{-ij}$ if and only if $R_{ij} = 0$. The structure matrix \mathbf{R} may also be expressed as the Laplacian of the adjacency graph $\mathcal{G} = (\mathcal{V}, \mathcal{E})$ with vertices $v \in \mathcal{V}$ corresponding to each area and edges $e \in \mathcal{E}$ between vertices i and j when $i \sim j$.

Rewriting Equation @ref{eq:gmr}, the probability density function of \mathbf{u} is

$$p(\mathbf{u}) \propto \exp\left(-\frac{\tau_u}{2} \mathbf{u}^\top \mathbf{R} \mathbf{u}\right) \propto \exp\left(-\frac{\tau_u}{2} \sum_{i \sim j} (u_i - u_j)^2\right). \quad (4.4)$$

This density is a function of the pairwise differences $u_i - u_j$ and so is invariant to the addition of a constant c to each entry $p(\mathbf{u}) = p(\mathbf{u} + c\mathbf{1})$, leading to an improper uniform distribution on the average of the u_i . If \mathcal{G} is connected, in that by traversing the edges, any vertex can be reached from any other vertex, then there is only one impropriety in the model and $\text{rank}(\mathbf{R}) = n - 1$, while if \mathcal{G} is disconnected, and composed of $n_c \geq 2$ connected components with index sets I_1, \dots, I_{n_c} , then the corresponding structure matrix \mathbf{R} has rank $n - n_c$ and the density is invariant to the addition of a constant to each of the connected components $p(\mathbf{u}_I) = p(\mathbf{u}_I + c\mathbf{1})$ where $I = I_1, \dots, I_{n_c}$.

4.3 Models using kernels

4.4 Simulation study

4.5 HIV prevalence study

4.6 Discussion

5

A model for risk group proportions

In this chapter I describe an application of Bayesian spatio-temporal statistics to small-area estimation of HIV risk group proportions. This work was conducted in collaboration with colleagues from the MRC Centre for Global Infectious Disease Analysis and UNAIDS. My primary role was to develop the statistical model, building upon an earlier version of the analysis conducted by Kathryn Risher. The results are presented in Howes, Risher, et al. (2023). Kathryn has also created a spreadsheet tool using the estimates which is now being used by countries to guide policy. Code for the analysis in this chapter is available from [athowes/multi-agyw](https://github.com/athowes/multi-agyw).

5.1 Background

Adolescent girls and young women (AGYW, defined here as females aged 15-29) are a demographic group at increased risk of HIV infection. Though AGYW are only 28% of the population, they comprise 44% of new infections (UNAIDS 2021a). HIV incidence for AGYW is 2.4 times higher than for similarly aged males. Table 5.1 gives the social and biological reasons for this disparity.

Table 5.1: AYW are at higher risk of HIV infection as a result of interacting social and biological reasons.

Reason	Description
Structural vulnerability and power imbalances	Gender inequality, lack of agency, economic factors, limited healthcare access, stigma and discrimination
Age patterns of sexual mixing, including a younger age at first sex	Older male partners have higher prevalence of HIV
Increased susceptibility to HIV infection	Immature reproductive tract, including a thinner cervical lining and higher levels of immune cells

On this basis, AGYW have been identified as a priority population for HIV prevention services (Saul et al. 2018; The Global Fund 2018). The Global AIDS Strategy 2021-2026 (UNAIDS 2021b) proposed stratifying HIV prevention packages to AGYW based on 1) local population-level HIV incidence and 2) individual-level sexual risk behaviour. As risk depends on both factors, prioritisation of prevention services is more efficient if both are taken into account (Figure 5.1). The strategy encourages programmes to define targets for the proportion of AGYW to be reached with a range of interventions. To implement the strategy, estimates of the size of each risk group are required.

5.2 Data

A model for risk group proportions

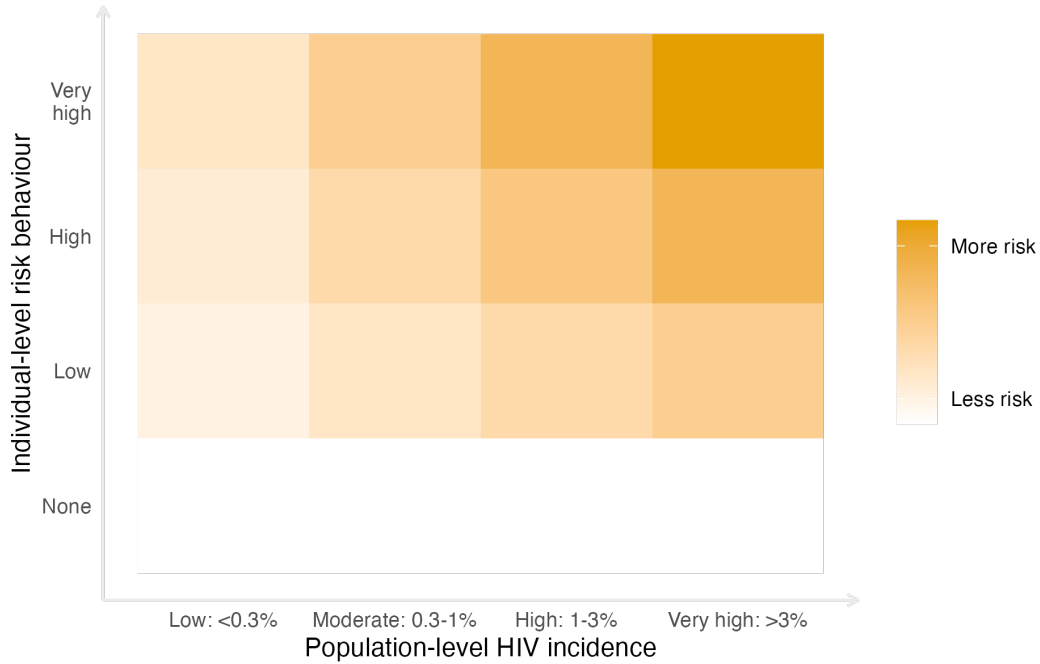


Figure 5.1: Risk depends on both individual-level risk behaviour and population-level HIV incidence.

Table 5.2: HIV risk groups and assumed HIV incidence rate ratio for each risk group relative to AGYW with one cohabiting sexual partner. Among FSW, the incidence rate ratio depended on the level of HIV incidence among the general population. The incidence rate ratio for women with non-regular or multiple sexual partner(s) was derived from analysis of ALPHA network data. Non-regular partners are defined to be non-cohabiting. The transactional sex risk group is adjusted during analysis to correspond to female sex worker, and incidence rate ratios among FSW were derived based on patterns of relative HIV prevalence among FSW compared to general population prevalence. When the local HIV incidence in the general population is higher, the incidence rate ratio for FSW is lower.

Risk group	Description	Local HIV incidence	Incidence ratio
None	Not sexually active	–	0.0
Low	One cohabiting partner	–	1.0 (Baseline)
High	Non-regular or multiple partner(s)	–	1.72
Very High	Transactional sex (adjusted to correspond to female sex workers)	0.1-0.3%, 0.3-1.0%, 1.0-3.0%, 1.0-3.0%	13.0, 9.0, 6.0, 3.0



Figure 5.2: Surveys.

I used household survey data from 13 designated AGYW priority countries: Botswana, Cameroon, Kenya, Lesotho, Malawi, Mozambique, Namibia, South Africa, Eswatini, Tanzania, Uganda, Zambia and Zimbabwe. I included surveys conducted in these countries between 1999 and 2018 in which women were interviewed about their sexual behaviour and sufficient geographic information was available to locate survey clusters to health districts. Demographic and Health Surveys (DHS), AIDS Indicator Surveys (AIS), Population-based HIV Impact Assessment (PHIA) surveys, and the Botswana AIDS Impact Survey 2013 (BAIS) were included.

For each survey, I classified respondents into one of four behavioural risk groups according to reported sexual risk behaviour in the past 12 months. These risk groups were (Table 5.2):

1. Not sexually active
2. One cohabiting sexual partner
3. Non-regular or multiple sexual partner(s), and
4. AGYW who report transactional sex.

In the case of inconsistent responses, women were categorised according to the highest risk group they fell into, ensuring that the categories were mutually

exclusive. Exact survey questions varied slightly across survey types and between survey phases. Questions captured information about whether the respondent had been sexually active in the past twelve months, and if so how with many partners. For their three most recent partners, respondents were also asked about the type of partnership. Possible partnership types included spouse, cohabiting partner, partner not cohabiting with respondent, friend, sex worker, sex work client, and other.

Some surveys included a specific question asking if the respondent had received or given money or gifts for sex in the past twelve months. In these surveys, 2.64% of women reported transactional sex. In surveys without such a question, women almost never (0.01%) answered that one of their three most recent partners was a sex work client. Due to this incomparability across surveys, I did not include surveys without a specific transactional sex question when estimating the proportion of the population who engaged in transactional sex. I instead focused on estimating the proportion of women who reported transactional sex at a district level, and subsequently adjusted these proportions to align to national estimates for the number of female sex workers.

I used estimates of population, PLHIV and new HIV infections stratified by district and age group from HIV estimates published by UNAIDS that were developed using the Naomi model (Eaton et al. 2021). The administrative area hierarchy and geographic boundaries I used correspond to those used for health service planning by countries. Exceptions are Cameroon and Kenya, where I conducted analysis one level higher at the department and county levels, respectively. I used the most recent 2022 estimates for all countries, apart from Mozambique where, due to data accuracy concerns, I used the 2021 estimates (in which the Cabo Delgado province is excluded due to disruption by conflict).

5.3 Model for risk group proportions

I took a two-stage modelling approach to estimate the four risk group proportions. Let the four risk groups be indexed by $k \in \{1, 2, 3, 4\}$, and denote being in either

the third or fourth risk group as $k = 3^+$. First, using all the surveys, I used a spatio-temporal multinomial logistic regression model (Section 5.3.1) to estimate the proportion of AGYW in the risk groups $k \in \{1, 2, 3^+\}$. Then, using only those surveys with a specific transactional sex question, I fit a spatial logistic regression model (Section 5.3.2) to estimate the proportion of those in the $k = 3^+$ risk group that were in the $k = 3$ and $k = 4$ risk groups respectively.

5.3.1 Spatio-temporal multinomial logistic regression

Let $i \in \{1, \dots, n\}$ denote districts partitioning the 13 studied AGYW priority countries $c[i] \in \{1, \dots, 13\}$. Consider the years 1999-2018 denoted as $t \in \{1, \dots, T\}$, and age groups $a \in \{15-19, 20-24, 25-29\}$. Let $p_{itak} > 0$ with $\sum_{k=1}^{3^+} p_{itak} = 1$, be the probabilities of membership of risk group k .

Multinomial logistic regression

A baseline category multinomial logistic regression model is specified by

$$\mathbf{y}_{ita} = (y_{ita1}, \dots, y_{ita3^+})^\top \sim \text{Multinomial}(m_{ita}; p_{ita1}, \dots, p_{ita3^+}), \quad (5.1)$$

$$\log\left(\frac{p_{itak}}{p_{ita1}}\right) = \eta_{itak}, \quad k = 2, 3^+, \quad (5.2)$$

where the number in risk group k is y_{itak} , the fixed sample size is $m_{ita} = \sum_{k=1}^{3^+} y_{itak}$, and $k = 1$ is chosen as the baseline category. This model is not an LGM, and is not possible to fit in R-INLA.

The multinomial-Poisson transformation

I use the multinomial-Poisson transformation to enable inference with R-INLA. The transformation reframes a given multinomial logistic regression model as an equivalent Poisson log-linear model of the form

$$y_{itak} \sim \text{Poisson}(\kappa_{itak}), \quad (5.3)$$

$$\log(\kappa_{itak}) = \eta_{itak}. \quad (5.4)$$

The basis of the transformation is that, conditional on their sum, Poisson counts are jointly multinomially distributed (McCullagh and Nelder 1989) as follows

$$\mathbf{y}_{ita} | m_{ita} \sim \text{Multinomial} \left(m_{ita}; \frac{\kappa_{ita1}}{\kappa_{ita}}, \dots, \frac{\kappa_{ita3^+}}{\kappa_{ita}} \right), \quad (5.5)$$

where $\kappa_{ita} = \sum_{k=1}^{3^+} \kappa_{itak}$. Category probabilities are then obtained by the softmax function

$$p_{itak} = \frac{\exp(\eta_{itak})}{\sum_{k=1}^{3^+} \exp(\eta_{itak})} = \frac{\kappa_{itak}}{\sum_{k=1}^{3^+} \kappa_{itak}} = \frac{\kappa_{itak}}{\kappa_{ita}}. \quad (5.6)$$

Under the equivalent model, the sample sizes $m_{ita} = \sum_k y_{itak}$ are treated as random $m_{ita} \sim \text{Poisson}(\kappa_{ita})$ rather than fixed. The joint distribution of $p(\mathbf{y}_{ita}, m_{ita}) = p(\mathbf{y}_{ita} | m_{ita})p(m_{ita})$ is then

$$p(\mathbf{y}_{ita}, m_{ita}) = \exp(-\kappa_{ita}) \frac{(\kappa_{ita})^{m_{ita}}}{m_{ita}!} \times \frac{m_{ita}!}{\prod_k y_{itak}!} \prod_k \left(\frac{\kappa_{itak}}{\kappa_{ita}} \right)^{y_{itak}} \quad (5.7)$$

$$= \prod_k \left(\frac{\exp(-\kappa_{itak}) (\kappa_{itak})^{y_{itak}}}{y_{itak}!} \right) \quad (5.8)$$

$$= \prod_k \text{Poisson}(y_{itak} | \kappa_{itak}). \quad (5.9)$$

corresponding to the product of independent Poisson likelihoods as in Equation (5.3).

This model, including random sample sizes, is equivalent to the multinomial logistic regression only when the normalisation constants m_{ita} are recovered exactly. To ensure that this is the case, one approach is to include observation-specific random effects θ_{ita} in the equation for the linear predictor. Multiplying each of $\{\kappa_{itak}\}_{k=1}^{3^+}$ by $\exp(\theta_{ita})$ has no effect on the category probabilities, but does provide the necessary flexibility for κ_{ita} to recover m_{ita} exactly. Although in theory an improper prior $\theta_{ita} \propto 1$ should be used, I found that in practise, by keeping η_{ita} otherwise small using appropriate constraints, so that arbitrarily large values of θ_{ita} are not required, it is sufficient (and practically preferable for inference) to instead use a vague prior.

Model specifications

I considered four models for η_{ita} in Equation (5.4) of the form

$$\eta_{ita} = \theta_{ita} + \beta_k + \zeta_{c[i]k} + \alpha_{ac[i]k} + \phi_{ik} + \gamma_{tk}. \quad (5.10)$$

A model for risk group proportions

Observation random effects $\theta_{ita} \sim \mathcal{N}(0, 1000^2)$ were included in all models I considered, and are required for the multinomial-Poisson transformation to be valid. To capture country-specific proportion estimates for each category, I included category random effects $\beta_k \sim \mathcal{N}(0, \tau_\beta^{-1})$ and country-category random effects $\zeta_{ck} \sim \mathcal{N}(0, \tau_\zeta^{-1})$. Heterogeneity in risk group proportions by age was allowed by including age-country-category random effects $\alpha_{ack} \sim \mathcal{N}(0, \tau_\alpha^{-1})$.

Spatial random effects For the space-category ϕ_{ik} random effects I considered two specifications:

1. Independent and identically distributed (IID) $\phi_{ik} \sim \mathcal{N}(0, \tau_\phi^{-1})$,
2. Besag (Besag et al. 1991) grouped by category

$$\boldsymbol{\phi} = (\phi_{11}, \dots, \phi_{n1}, \dots, \phi_{13+}, \dots, \phi_{n3+})^\top \sim \mathcal{N}(\mathbf{0}, (\tau_\phi \mathbf{R}_\phi^*)^-).$$

The scaled structure matrix $\mathbf{R}_\phi^* = \mathbf{R}_b^* \otimes \mathbf{I}$ is given by the Kronecker product of the scaled Besag structure matrix \mathbf{R}_b^* and the identity matrix \mathbf{I} , and $-$ denotes the generalised matrix inverse. Scaling of the structure matrix to have generalised variance one ensures interpretable priors may be placed on the precision parameter (Sørbye and Rue 2014). I followed the further recommendations of Freni-Sterrantino et al. (2018) with regard to disconnected adjacency graphs, singletons and constraints. The Besag structure matrix \mathbf{R}_b was obtained by the precision matrix of the random effects $\mathbf{b} = (b_1, \dots, b_n)^\top$ with full conditionals

$$b_i \mid \mathbf{b}_{-i} \sim \mathcal{N}\left(\frac{\sum_{j:j \sim i} b_j}{n_{\delta i}}, \frac{1}{n_{\delta i}}\right), \quad (5.11)$$

where $j \sim i$ if the districts A_i and A_j are adjacent, and $n_{\delta i}$ is the number of districts adjacent to A_i .

In preliminary testing, I excluded spatial random effects from the model, but found that this negatively effected performance. I also tested using the BYM2 model (Simpson et al. 2017) in place of the Besag, but found that the proportion

parameter posteriors tended to be highly peaked at the value one. For simplicity and to avoid numerical issues, by using Besag random effects I effectively decided to fix this proportion to one.

Temporal random effects For the year-category γ_{tk} random effects I considered two specifications:

1. IID $\phi_{tk} \sim \mathcal{N}(0, \tau_\phi^{-1})$,
2. First order autoregressive (AR1) grouped by category

$$\boldsymbol{\gamma} = (\gamma_{11}, \dots, \gamma_{13+}, \dots, \gamma_{T1}, \dots, \gamma_{T3+})^\top \sim \mathcal{N}(\mathbf{0}, (\tau_\phi \mathbf{R}_\gamma^*)^-).$$

The scaled structure matrix $\mathbf{R}_\gamma^* = \mathbf{R}_r^* \otimes \mathbf{I}$ is given by the Kronecker product of a scaled AR1 structure matrix \mathbf{R}_r^* and the identity matrix \mathbf{I} . The AR1 structure matrix \mathbf{R}_r is obtained by precision matrix of the random effects $\mathbf{r} = (r_1, \dots, r_T)^\top$ specified by

$$r_1 \sim \left(0, \frac{1}{1 - \rho^2}\right), \tag{5.12}$$

$$r_t = \rho r_{t-1} + \epsilon_t, \quad t = 2, \dots, T, \tag{5.13}$$

where $\epsilon_t \sim \mathcal{N}(0, 1)$ and $|\rho| < 1$.

Priors All random effect precision parameters $\tau \in \{\tau_\beta, \tau_\zeta, \tau_\alpha, \tau_\phi, \tau_\gamma\}$ were given independent penalised complexity (PC) priors (Simpson et al. 2017) with base model $\sigma = 0$ given by $p(\tau) = 0.5\nu\tau^{-3/2} \exp(-\nu\tau^{-1/2})$ where $\nu = -\ln(0.01)/2.5$ such that $\mathbb{P}(\sigma > 2.5) = 0.01$. For the lag-one correlation parameter ρ , I used the PC prior, as derived by Sørbye and Rue (2017), with base model $\rho = 1$ and condition $\mathbb{P}(\rho > 0 = 0.75)$. I chose the base model $\rho = 1$ corresponding to no change in behaviour over time, rather than the alternative $\rho = 0$ corresponding to no correlation in behaviour over time, as I judged the former to be more plausible a priori.

Constraints

To ensure interpretable posterior inferences relating to the random effects, I applied sum-to-zero constraints such that none of the category interaction random effects altered overall category probabilities. For the space-year-category random effects, I applied analogous sum-to-zero constraints to maintain roles of the space-category and year-category random effects. Together, these were:

1. Category $\sum_k \beta_k = 0$,
2. Country $\sum_c \zeta_{ck} = 0, \forall k$,
3. Age-country $\sum_a \alpha_{ack} = 0, \forall c, k$,
4. Spatial $\sum_i \phi_{ik} = 0, \forall k$,
5. Temporal $\sum_t \gamma_{tk} = 0, \forall k$.

Survey weighted likelihood

I included surveys which use a complex design, in which each individual has an unequal probability of being included in the sample. For example the DHS often employs a two-stage cluster design, first taking an urban rural stratified sample of enumeration areas, before selecting households from each enumeration area using systematic sampling (DHS 2012).

To account for this aspect of survey design, I use a weighted pseudo-likelihood where the observed counts y are replaced by effective counts y^\star calculated using the survey weights w_j of all individuals j in the corresponding strata. I multiplied direct estimates produced using the `survey` package (Lumley 2004) by the Kish effective sample size (Kish 1965)

$$m^\star = \frac{\left(\sum_j w_j\right)^2}{\sum_j w_j^2} \quad (5.14)$$

to obtain y^\star . These counts may not be integers, and as such the Poisson likelihood I used in Equation (5.3) is not appropriate. Instead, I used a generalised Poisson pseudo-likelihood $y^\star \sim \text{xPoisson}(\kappa)$, given by

$$p(y^\star) = \frac{\kappa^{y^\star}}{[y^\star!]} \exp(-\kappa), \quad (5.15)$$

as implemented by `family = "xPoisson"` in R-INLA, which accepts non-integer input.

5.3.2 Spatial logistic regression

To estimate the proportion of those in the $k = 3^+$ risk group that were in the $k = 3$ and $k = 4$ risk groups respectively, I fit logistic regression models of the form

$$y_{ia4} \sim \text{Binomial}(y_{ia3} + y_{ia4}, q_{ia}), \quad (5.16)$$

$$q_{ia} = \text{logit}^{-1}(\eta_{ia}), \quad (5.17)$$

where $q_{ia} = p_{ia4}/(p_{ia3} + p_{ia4}) = p_{ia4}/p_{ia3+}$. Taking this two-step approach allowed me to include all surveys in the multinomial regression model, but only those surveys with a specific transactional sex question in Equation (5.16). As all such surveys occurred in 2013-2018, in the logistic regression model I assumed q_{ia} to be constant over time.

Model specifications

I considered six logistic regression models. Each included a constant $\beta_0 \sim \mathcal{N}(-2, 1^2)$, country random effects $\zeta_c \sim \mathcal{N}(0, \tau_\zeta^{-1})$, and age-country random effects $\alpha_{ac} \sim \mathcal{N}(0, \tau_\alpha^{-1})$. The prior on β_0 placed 95% prior probability on the range 2-50% for the percentage of those with non-regular or multiple partners who report transactional sex. I considered two specifications (IID, Besag) for the spatial random effects ϕ_i . To aid estimation with sparse data, I also considered national-level covariates for the proportion of men who have paid for sex ever `cfswever` or in the last twelve months `cfswrecent`, available from Hodgins et al. (2022). For both random effect precision parameters $\tau \in \{\tau_\alpha, \tau_\zeta\}$ I used the PC prior with base model $\sigma = 0$ and $\mathbb{P}(\sigma > 2.5) = 0.01$. For the regression parameters $\beta \in \{\beta_{\text{cfswever}}, \beta_{\text{cfswrecent}}\}$ I used the prior $\beta \sim \mathcal{N}(0, 2.5^2)$.

Survey weighted likelihood

As with the multinomial regression model, I used survey weighted counts $\{y_{itak}^*\}$ and sample sizes $\{m_{itak}^*\}$. I used a generalised binomial pseudo-likelihood $y^* \sim \text{xBinomial}(m^*, q)$, as implemented by `family = "xBinomial"` in R-INLA, given by

$$p(y^* | m^*, q) = \binom{\lfloor m^* \rfloor}{\lfloor y^* \rfloor} q^{y^*} (1 - q)^{m^* - y^*}. \quad (5.18)$$

to extend the binomial distribution to non-integer weighted counts and sample sizes.

5.3.3 Female sex worker population size adjustment

Responding “yes” to the question “have you had sex in return for gifts, cash or anything else in the past 12 months” is not considered sufficient to constitute sex work. Recognising this, I adjusted the estimates obtained based on the survey to match FSW population size estimates obtained via alternative methods.

Stevens et al. (2022) used a Bayesian meta-analysis of key population specific data sources to estimate adult (15-49) FSW population size by country. I disaggregated these estimates by age as follows. First, I calculated the total sexually debuted population in each age group, in each country. To describe the distribution of age at first sex, I used skew logistic distributions (Nguyen and Eaton 2022) with cumulative distribution function given by

$$F(x) = (1 + \exp(\kappa_c(\mu_c - x)))^{-\gamma_c}, \quad (5.19)$$

where $\kappa_c, \mu_c, \gamma_c > 0$ are country-specific shape, shape and skewness parameters respectively. Next, I used the assumed $\text{Gamma}(\alpha = 10.4, \beta = 0.36)$ FSW age distribution in South Africa from the Thembisa model (Johnson and Dorrington 2020) to calculate the implied ratio between the number of FSW and the sexually debuted population in each age group. I assumed these ratios in South Africa were applicable to every country, allowing calculation of the number of FSW by age group in all 13 countries. The results obtained are shown in Figure 5.3.

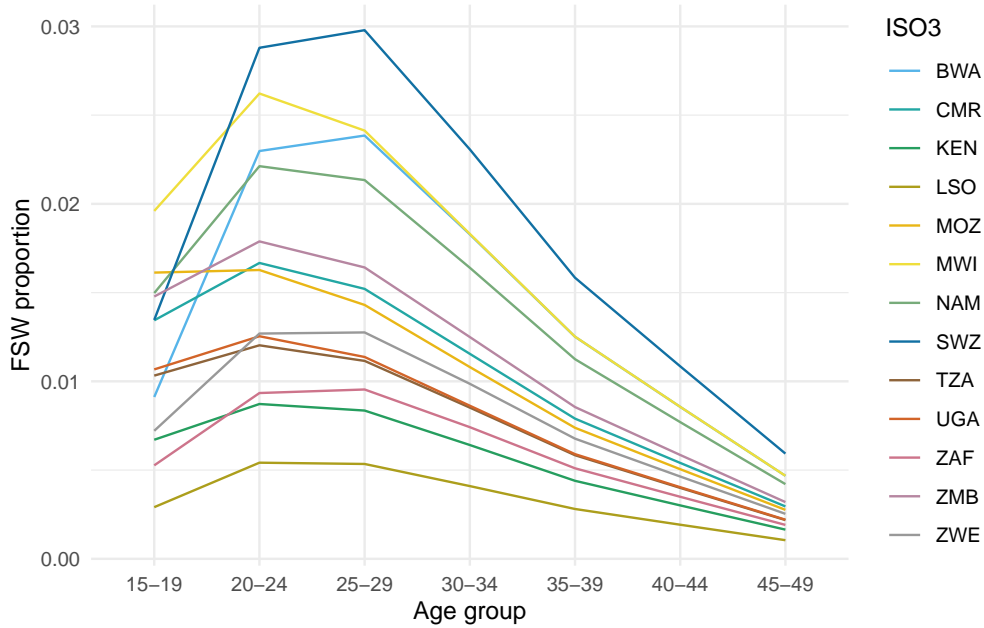


Figure 5.3: The disaggregation procedure I used produces a plausible age distribution of FSW by country.

5.3.4 Results

Coverage assessment

To assess the calibration of the fitted model, I calculated the quantile q of each observation within the posterior predictive distribution. For calibrated models, these quantiles, known as probability integral transform (PIT) values (Dawid 1984; Bosse et al. 2022), should follow a uniform distribution $q \sim \mathcal{U}[0, 1]$. To generate samples from the posterior predictive distribution, I applied the multinomial likelihood to samples from the latent field, setting the sample size to be the floor of the Kish effective sample size.

Using the PIT values, it is possible to calculate the empirical coverage of all $(1 - \alpha)100\%$ (equal-tailed) posterior predictive credible intervals. These empirical coverages can be compared to the nominal coverage $(1 - \alpha)$ for each value of $\alpha \in [0, 1]$ to give empirical cumulative distribution function (ECDF) difference values. This approach has the advantage of considering all possible confidence values at once. **sailynoja2021graphical** develop binomial distribution-based simultaneous confidence bands for ECDF difference values which test uniformity.

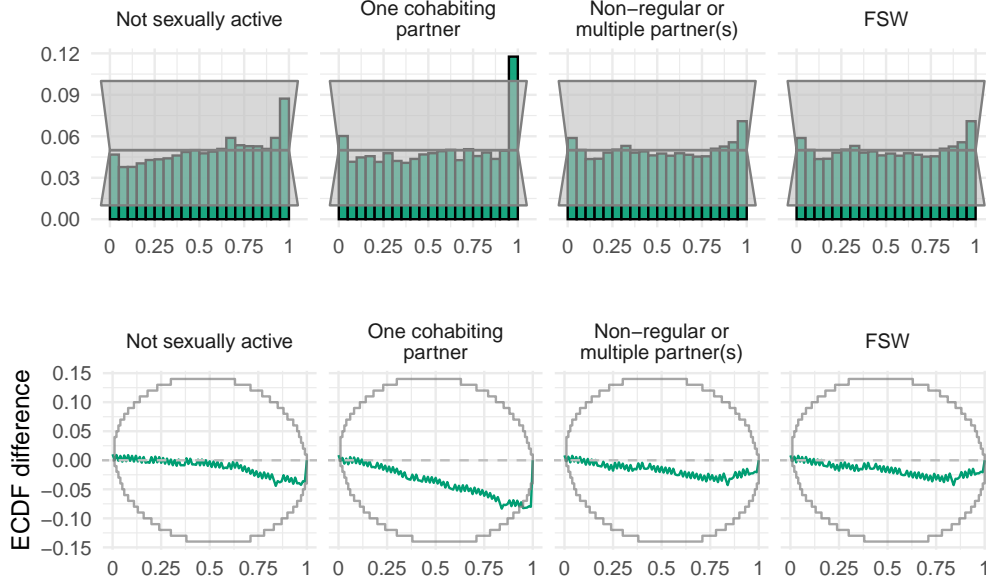


Figure 5.4: Coverage.

Variance decomposition

Estimates

5.4 Prevalence and incidence by risk group

5.4.1 Disaggregation of Naomi estimates

I calculated HIV incidence λ_{iak} and number of new HIV infections I_{iak} stratified according to district, age group and risk group by linear disaggregation

$$I_{ia} = \sum_k I_{iak} = \sum_k \lambda_{iak} N_{iak} \quad (5.20)$$

$$= 0 + \lambda_{ia2} N_{ia2} + \lambda_{ia3} N_{ia3} + \lambda_{ia4} N_{ia4} \quad (5.21)$$

$$= \lambda_{ia2} (N_{ia2} + \text{RR}_3 N_{ia3} + \text{RR}_4 (\lambda_{ia}) N_{ia4}) . \quad (5.22)$$

Risk group specific HIV incidence estimates are then given by

$$\lambda_{ia1} = 0, \quad (5.23)$$

$$\lambda_{ia2} = I_{ia} / (N_{ia2} + \text{RR}_3 N_{ia3} + \text{RR}_4 (\lambda_{ia}) N_{ia4}) , \quad (5.24)$$

$$\lambda_{ia3} = \text{RR}_3 \lambda_{ia2}, \quad (5.25)$$

$$\lambda_{ia4} = \text{RR}_4 (\lambda_{ia}) \lambda_{ia2}. \quad (5.26)$$

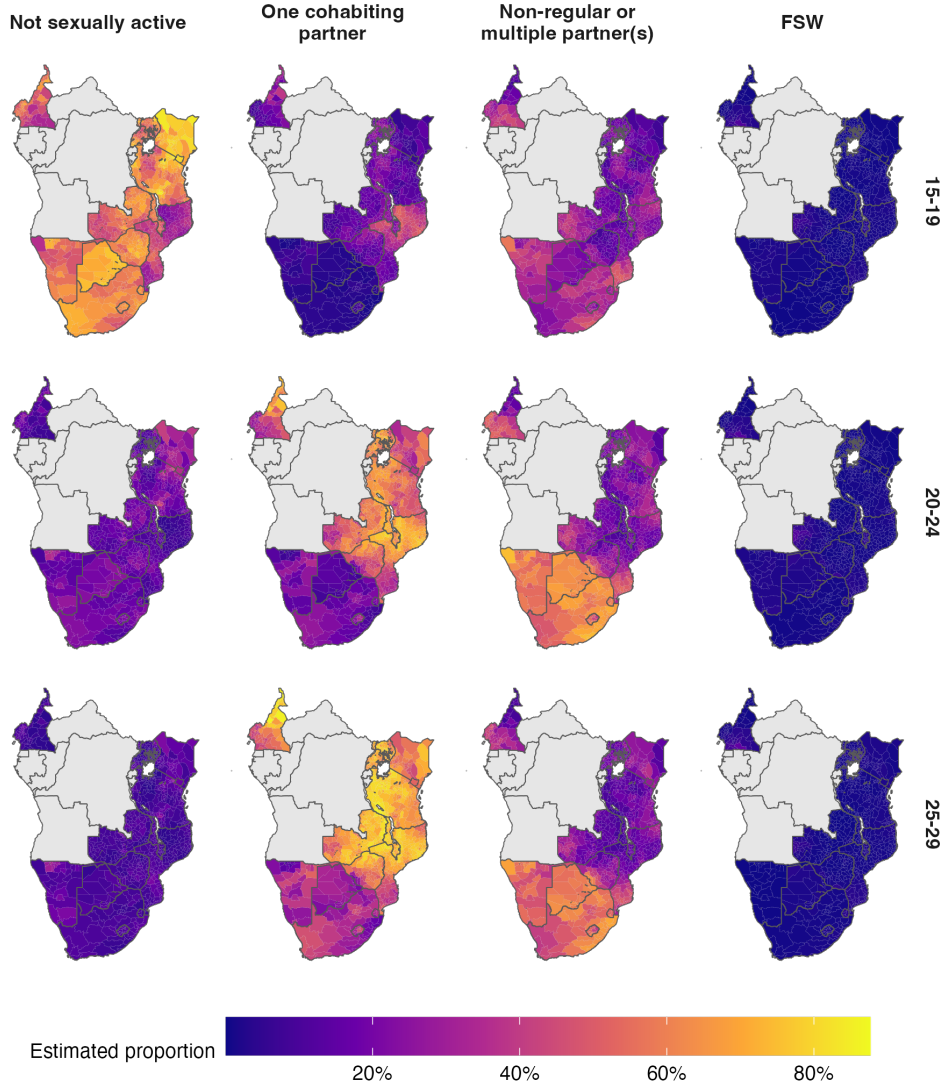


Figure 5.5: Map.

which I evaluated using Naomi model estimates of the number of new HIV infections $I_{ia} = \lambda_{ia}N_{ia}$, HIV infection risk ratios $\{RR_3, RR_4(\lambda_{ia})\}$, and risk group population sizes as above. The risk ratio $RR_4(\lambda_{ia})$ was defined as a function of general population incidence. The number of new HIV infections are then $I_{iak} = \lambda_{iak}N_{iak}$.

5.4.2 Expected new infections reached

I calculated the number of new infections that would be reached prioritising according to each possible stratification of the population—that is for all $2^3 = 8$ possible combinations of stratification by location, age, and risk group. As an illustration,

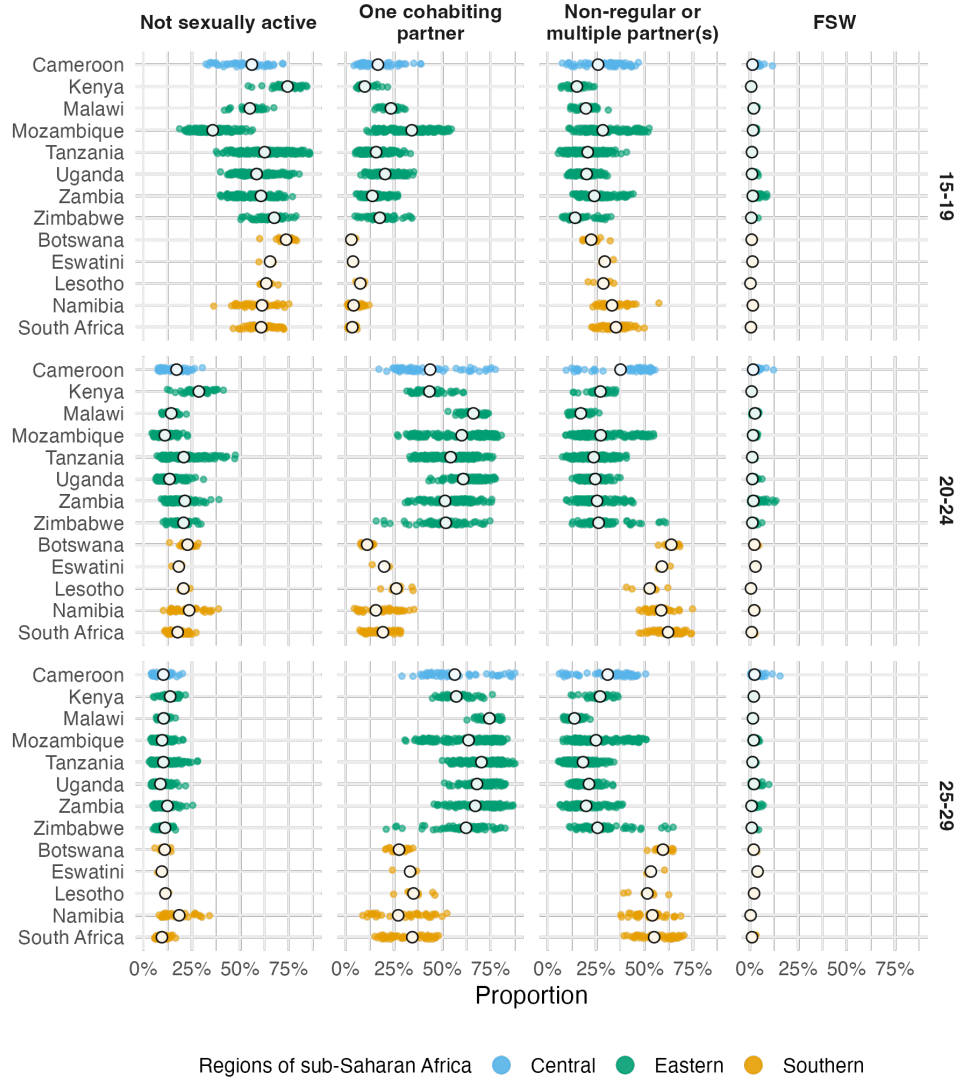


Figure 5.6: Dots.

for stratification just by age, I aggregated the number of new HIV infections and HIV incidence as such

$$I_a = \sum_{ik} I_{iak}, \quad (5.27)$$

$$\lambda_a = I_a / \sum_{ik} N_{iak}. \quad (5.28)$$

Under this stratification, individuals in each age group a are prioritised according to the highest HIV incidence λ_a . By cumulatively summing the expected infections, for each fraction of the total population reached I calculated the fraction of total expected new infections that would be reached.

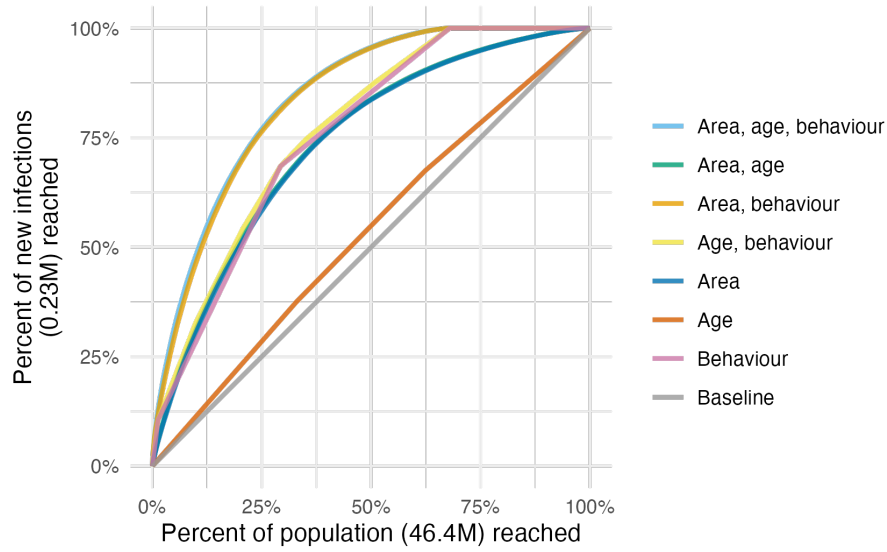


Figure 5.7: Surveys.

5.4.3 Results

5.5 Discussion

I estimated the proportion of AGYW who fall into different risk groups at a district level in 13 sub-Saharan African countries. These estimates support consideration of differentiated prevention programming according to geographic locations and risk behaviour, as outlined in the Global AIDS Strategy. Systematic differences in risk by age groups, and variation within and between countries, explained the large majority of variation in risk group proportions. Changes over time were negligible in the overall variation in risk group proportions. The proportion of 15-19 year olds who are sexually active, and among women aged 20-29 years, norms around cohabitation especially varied across districts and countries. This variation underscores the need for these granular data to implement HIV prevention options aligned to local norms and risk behaviours.

I considered four risk groups based on sexual behaviour, the most proximal determinant of risk. Other factors, such as condom usage or type of sexual act, may account for additional heterogeneity in risk from sexual behaviour. However, I did not include these factors in view of measurement difficulties, concerns

A model for risk group proportions

about consistency across contexts, and the operational benefits of describing risk parsimoniously. Sexual behaviour confers risk only when AGYW reside in geographic locations where there is unsuppressed viral load among their potential partners. I did not include more distal determinants, such as school attendance, orphanhood, or gender empowerment, as I expect their effects on risk to largely be mediated by more proximal determinants. However, to effectively implement programming, it is crucial to understand these factors, as well as the broader structural barriers and limits to personal agency faced by AGYW. Importantly, programs must ensure that intervention prioritisation occurs without stigmatising or blaming AGYW.

Brugh et al. (Brugh et al. 2021) previously geographically mapped AGYW HIV risk groups using biomarker and behavioural data from the most recent surveys in Eswatini, Haiti and Mozambique to define and subsequently map risk groups with a range of machine learning techniques. My work builds on Brugh et al. (Brugh et al. 2021) by including more countries, integrating a greater number of surveys, and connecting risk group proportions with HIV epidemic indicators to help inform programming.

By considering a range of possible risk stratification strategies, I showed that successful implementation of a risk-stratified approach would allow substantially more of those at risk for infections to be identified before infection occurs. A considerable proportion of estimated new infections were among FSW, supporting the case for HIV programming efforts focused on key population groups (Baral et al. 2012). There is substantial variation in the importance of prioritisation by age, location and behaviour within each country. This highlights the importance of understanding and tailoring HIV prevention efforts to country-specific contexts. By standardising our analysis across all 13 countries, I showed the additional efficiency benefits of resource allocation between countries.

I found a geographic delineation in the proportion of women cohabiting between southern and eastern Africa, calling attention to a divide attributable to many cultural, social, and economic factors. The delineation does not represent a boundary between predominately Christian and Muslim populations, which is further north.

I also note that the high numbers of adolescent girls aged 15-19 cohabiting in Mozambique is markedly different from the other countries (UNICEF n.d.).

My modelled estimates of risk group proportions improve upon direct survey results for three reasons. First, by taking a modular modelling approach, I integrated all relevant survey information from multiple years, allowing estimation of the FSW proportion for surveys without a specific transactional sex question. Second, whereas direct estimates exhibit large sampling variability at a district level, I alleviated this issue using spatio-temporal smoothing. Third, I provided estimates in all district-years, including those not directly sampled by surveys, allowing estimates to be consistently fed into further analysis and planning pipelines (such as our analysis of risk group specific prevalence and incidence).

The final surveys included in our risk model model were conducted in 2018. The analysis may be updated with more surveys as they become available. I do not anticipate that the risk group proportions will change substantially, as I found that they did not change significantly over time.

My analysis focused on females aged 15-29 years, and could be extended to consider optimisation of prevention more broadly, accounting for the 0% of new infections among adults 15-49 which occur in women 30-49 and men 15-49. Estimating sexual risk behaviour in adults 15-49 would be a crucial step toward greater understanding of the dynamics of the HIV epidemic in sub-Saharan Africa, and would allow incidence models to include stratification of individuals by sexual risk.

Phylogenetic results from BDI are about transmission rather than incidence. Only age-sex structured not age-sex-behaviour. Does not undermine my work.

5.5.1 Limitations

My analysis was subject to challenges shared by most approaches to monitoring sexual behaviour in the general population (Cleland et al. 2004). In particular, under-reporting of higher risk sexual behaviours among AGYW could affect the validity of my risk group proportion estimates. Due to social stigma or disapproval,

respondents may be reluctant to report non-marital partners (Nnko et al. 2004; Helleringer et al. 2011) or may bias their reporting of sexual debut (Zaba et al. 2004; Wringe et al. 2009; Nguyen and Eaton 2022). For guidance of resource allocation, differing rates of under-reporting by country, district, year or age group are particularly concerning to the applicability of my results; and, while it may be reasonable to assume a constant rate over space-time, the same cannot be said for age, where aspects of under-reporting have been shown to decline as respondents age (Glynn et al. 2011), suggesting that the elevated risks I found faced by younger women are likely a conservative estimate. If present, these reporting biases will also have distorted the estimates of infection risk ratios and prevalence ratios I used in my analysis, likely over-attributing risk to higher risk groups.

I have the least confidence in my estimates for the FSW risk group. As well as having the smallest sample sizes, my transactional sex estimates do not overcome the difficulties of sampling hard to reach groups. I inherit any limitations of the national FSW estimates (Stevens et al. 2022) which I adjust my estimates of transactional sex to match. Furthermore, I do not consider seasonal migration patterns, which may particularly affect FSW size. More generally, I did not consider covariates potentially predictive of risk group proportions (such as sociodemographic characteristics, education, local economic activity, cultural and religious norms and attitudes), which are typically difficult to measure spatially. Identifying measurable correlates of risk, or particular settings in which time-concentrated HIV risk occurs, is an important area for further research to improve risk prioritisation and precision HIV programme delivery.

The efficiency of each stratified prevention strategy depends on the ability of programmes to identify and effectively reach those in each strata. My analysis of new infections potentially averted assumed a “best-case” scenario where AGYW of every strata can be reached perfectly, and should therefore be interpreted as illustrating the potentially obtainable benefits rather than benefits which would be obtained from any specific intervention strategy. In practice, stratified prevention strategies are likely to be substantially less efficient than this best-case scenario.

Factors I did not consider include the greater administrative burden of more complex strategies, variation in difficulty or feasibility of reaching individuals in each strata, variation in the range or effectiveness of interventions by strata, and changes in strata membership that may occur during the course of a year. Identifying and reaching behavioural strata may be particularly challenging. Empirical evaluations of behavioural risk screening tools have found only moderate discriminatory ability (Jia et al. 2022), and risk behaviour may change rapidly among young populations, increasing the challenge to effectively deliver appropriately timed prevention packages. This consideration may motivate selecting risk groups based on easily observable attributes, such as attendance of a particular service or facility, rather than sexual behaviour.

I did not engage with country experts or civil society organisations. This led to problems, including in Malawi. Future work should be better.

5.5.2 Conclusion

I estimated the proportion of AGYW aged 15-19, 20-24 and 25-29 years in four sexual risk groups at a district-level in 13 priority countries and analyzed the number of infections that could be reached by prioritisation based upon location, age and behaviour. Though subject to limitations, these estimates provide data that national HIV programmes can use to set targets and implement differentiated HIV prevention strategies as outlined in the Global AIDS Strategy. Successfully implementing this approach would result in more efficiently reaching a greater number of those at risk of infection.

Among AGYW, there was systematic variation in sexual behaviour by age and location, but not over time. Age group variation was primarily attributable to age of sexual debut (ages 15-24). Spatial variation was particularly present between those who reported one cohabiting partner versus non-regular or multiple partners. Risk group proportions did not change substantially over time, indicating that norms relating to sexual behaviour are relatively static. These findings underscore

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the importance of providing effective HIV prevention options tailored to the needs of particular age groups, as well as local norms around sexual partnerships.

6

Fast approximate Bayesian inference

In this chapter I describe a novel Bayesian inference method I developed. The method is motivated the Naomi small-area estimation model. Over 35 countries have used the Naomi model software (<https://naomi.unaids.org>) to produce estimates of HIV indicators. Ensuring fast and accurate Bayesian inferences in this setting is challenging.

I began working on this project at the start of my PhD. However, it was only after I read Stringer et al. (2021) that I began making progress. Alex Stringer later supervised by visit to the University of Waterloo. The results of this work are presented in Howes, Stringer, et al. (2023+). Code for the analysis in this chapter is available from `athowes/naomi-aghq`.

6.1 Inference methods

In a Bayesian analysis, the primary goal is to perform inference. That is, to obtain the posterior distribution

$$p(\boldsymbol{\phi} | \mathbf{y}) = \frac{p(\boldsymbol{\phi}, \mathbf{y})}{p(\mathbf{y})}. \quad (6.1)$$

Inference is a reasonable goal because the posterior distribution is sufficient for use in decision making. Given a loss function, the posterior loss of a decision depends

on the data only via the posterior distribution.

It is usually intractable to directly obtain the posterior distribution. This is because the denominator contains an intractable integral called the posterior normalising constant

$$p(\mathbf{y}) = \int_{\mathbb{R}^d} p(\mathbf{y}, \phi) d\phi \quad (6.2)$$

For this reason, approximations to the posterior distribution $\tilde{p}(\phi | \mathbf{y})$ are typically used in place of the exact posterior distribution. Some approximate Bayesian inference methods avoid directly calculate the posterior normalising constant, instead working with the unnormalised posterior distribution

$$p(\phi | \mathbf{y}) \propto p(\phi, \mathbf{y}). \quad (6.3)$$

Other approximate Bayesian inference methods can more directly be thought of as ways to estimate the posterior normalising constant.

6.1.1 The Laplace approximation

Laplace's method (Laplace 1774) is a technique used to approximate integrals of the form

$$\int \exp(Ch(z)) dz, \quad (6.4)$$

where $C > 0$ is a large constant and h is a function which is twice-differentiable. The Laplace approximation (Tierney and Kadane 1986) is obtained by application of Laplace's method to calculate the posterior normalising constant. Let $h(\phi) = \log p(\phi, \mathbf{y})$ such that

$$p(\mathbf{y}) = \int_{\mathbb{R}^d} p(\mathbf{y}, \phi) d\phi = \int_{\mathbb{R}^d} \exp(h(\phi)) d\phi. \quad (6.5)$$

Laplace's method involves approximating the function h by its second order Taylor expansion evaluated at a maxima of h to eliminate the first order term. Let

$$\hat{\phi} = \arg \max_{\phi} h(\phi) \quad (6.6)$$

be the posterior mode, and

$$\hat{\mathbf{H}} = -\frac{\partial^2}{\partial \boldsymbol{\phi} \partial \boldsymbol{\phi}^\top} h(\boldsymbol{\phi})|_{\boldsymbol{\phi}=\hat{\boldsymbol{\phi}}} \quad (6.7)$$

be the Hessian matrix evaluated at the posterior mode. The Laplace approximation is then

$$\tilde{p}_{\text{LA}}(\mathbf{y}) = \int_{\mathbb{R}^d} \exp \left(h(\hat{\boldsymbol{\phi}}, \mathbf{y}) - \frac{1}{2}(\boldsymbol{\phi} - \hat{\boldsymbol{\phi}})^\top \hat{\mathbf{H}}(\boldsymbol{\phi} - \hat{\boldsymbol{\phi}}) \right) d\boldsymbol{\phi} \quad (6.8)$$

$$= p(\hat{\boldsymbol{\phi}}, \mathbf{y}) \cdot \frac{(2\pi)^{d/2}}{|\hat{\mathbf{H}}|^{1/2}}. \quad (6.9)$$

Equation (6.8) is calculated using the known normalising constant of the Gaussian distribution

$$p_{\text{G}}(\boldsymbol{\phi} | \mathbf{y}) = \mathcal{N}(\boldsymbol{\phi} | \hat{\boldsymbol{\phi}}, \hat{\mathbf{H}}^{-1}) = \frac{|\hat{\mathbf{H}}|^{1/2}}{(2\pi)^{d/2}} \exp \left(-\frac{1}{2}(\boldsymbol{\phi} - \hat{\boldsymbol{\phi}})^\top \hat{\mathbf{H}}(\boldsymbol{\phi} - \hat{\boldsymbol{\phi}}) \right). \quad (6.10)$$

It follows that the Laplace approximation may be thought of as approximating the posterior distribution by a Gaussian distribution $p(\boldsymbol{\phi} | \mathbf{y}) \approx p_{\text{G}}(\boldsymbol{\phi} | \mathbf{y})$ such that

$$\tilde{p}_{\text{LA}}(\mathbf{y}) = \frac{p(\boldsymbol{\phi}, \mathbf{y})}{p_{\text{G}}(\boldsymbol{\phi} | \mathbf{y})} \Big|_{\boldsymbol{\phi}=\hat{\boldsymbol{\phi}}}. \quad (6.11)$$

The marginal Laplace approximation

It may be inaccurate to approximate the full joint posterior distribution using a Gaussian distribution. An alternative is to instead approximate the marginal posterior distribution of some subset of the parameters. As before, let $\boldsymbol{\phi} = (\mathbf{x}, \boldsymbol{\theta})$ where \mathbf{x} is the latent field, and $\boldsymbol{\theta}$ are the hyperparameters. Applying an equivalent Laplace approximation to the latent field, we have $h(\mathbf{x}, \boldsymbol{\theta}) = \log p(\mathbf{x}, \boldsymbol{\theta}, \mathbf{y})$ with posterior mode

$$\hat{\mathbf{x}}(\boldsymbol{\theta}) = \arg \max_{\mathbf{x}} h(\mathbf{x}, \boldsymbol{\theta}) \quad (6.12)$$

and Hessian matrix evaluated at the posterior mode

$$\hat{\mathbf{H}}(\boldsymbol{\theta}) = -\frac{\partial^2}{\partial \mathbf{x} \partial \mathbf{x}^\top} h(\mathbf{x}, \boldsymbol{\theta})|_{\mathbf{x}=\hat{\mathbf{x}}(\boldsymbol{\theta})}. \quad (6.13)$$

For both quantities cases dependence on the hyperparameters $\boldsymbol{\theta}$ is made explicit. The resulting marginal Laplace approximation is then

$$\tilde{p}_{\text{LA}}(\boldsymbol{\theta}, \mathbf{y}) = \int_{\mathbb{R}^N} \exp \left(h(\hat{\mathbf{x}}(\boldsymbol{\theta}), \boldsymbol{\theta}, \mathbf{y}) - \frac{1}{2}(\mathbf{x} - \hat{\mathbf{x}}(\boldsymbol{\theta}))^\top \hat{\mathbf{H}}(\boldsymbol{\theta})(\mathbf{x} - \hat{\mathbf{x}}(\boldsymbol{\theta})) \right) d\mathbf{x} \quad (6.14)$$

$$= \exp(h(\hat{\mathbf{x}}(\boldsymbol{\theta}), \mathbf{y})) \cdot \frac{(2\pi)^{d/2}}{|\hat{\mathbf{H}}(\boldsymbol{\theta})|^{1/2}} \quad (6.15)$$

$$= \frac{p(\mathbf{x}, \boldsymbol{\theta}, \mathbf{y})}{\tilde{p}_{\text{G}}(\mathbf{x} | \boldsymbol{\theta}, \mathbf{y})} \Big|_{\mathbf{x}=\hat{\mathbf{x}}(\boldsymbol{\theta})}, \quad (6.16)$$

where $\tilde{p}_{\text{G}}(\mathbf{x} | \boldsymbol{\theta}, \mathbf{y}) = \mathcal{N}(\mathbf{x} | \hat{\mathbf{x}}(\boldsymbol{\theta}), \hat{\mathbf{H}}(\boldsymbol{\theta})^{-1})$ is a Gaussian approximation to the marginal posterior of the latent field.

6.1.2 Quadrature

Quadrature is an approach which can be used to approximate integrals like the posterior normalising constant. As with the Laplace approximation, it is deterministic in that the computational procedure is not intrinsically random.

Let \mathcal{Q} be a set of quadrature points $\mathbf{z} \in \mathcal{Q}$ and $\omega : \mathbb{R}^d \rightarrow \mathbb{R}$ be a weighting function. Then a quadrature approximation to the posterior normalising constant is given by

$$\tilde{p}_{\mathcal{Q}}(\mathbf{y}) = \sum_{\mathbf{z} \in \mathcal{Q}} p(\mathbf{y}, \mathbf{z}) \omega(\mathbf{z}). \quad (6.17)$$

Quadrature methods are most effective when integrating over small dimensions. The reason why is that exponentially more quadrature points are required to cover each additional dimension. For even moderate dimension, this quickly becomes intractable.

Gauss-Hermite quadrature

Gauss-Hermite quadrature [GHQ; Davis and Rabinowitz (1975)] is a quadrature rule designed to integrate a certain class of functions exactly. These functions are the form $f(z) = \phi(z)P_\alpha(z)$, where $\phi(\cdot)$ is a standard normal density, and $P_\alpha(\cdot)$ is a polynomial of degree α .

Following the notation for GHQ established by Bilodeau et al. (2022) for $z \in \mathbb{R}$, let $H_k(z)$ be the k th Hermite polynomial

$$H_k(z) = (-1)^k \exp(z^2/2) \frac{d}{dz^k} \exp(-z^2/2). \quad (6.18)$$

The univariate GHQ rule has nodes $z \in \mathcal{Q}(1, k)$ given by the k zeroes of the k th Hermite polynomial. The corresponding weighting function $\omega : \mathcal{Q}(1, k) \rightarrow \mathbb{R}$ is given by

$$\omega(z) = \frac{\phi(z) \cdot k!}{[H_{k+1}(z)]^2}. \quad (6.19)$$

Multivariate GHQ rules are usually constructed using the product rule over identical univariate GHQ rules in each dimension. In d dimensions, the multivariate GHQ nodes $\mathbf{z} \in \mathcal{Q}(d, k)$ are defined by $\mathcal{Q}(d, k) = \mathcal{Q}(1, k)^d = \mathcal{Q}(1, k) \times \cdots \times \mathcal{Q}(1, k)$. The corresponding weighting function $\omega : \mathcal{Q}(d, k) \rightarrow \mathbb{R}$ is given by $\omega(\mathbf{z}) = \prod_{j=1}^d \omega(z_j)$.

Adaptive quadrature

In adaptive quadrature, the quadrature nodes and weights depend on the specific integrand. Adaptive quadrature rules are particularly important for Bayesian inference problems because the posterior normalising function is a function of the data. No fixed quadrature rule can be expected to perform well in integrating any posterior distributions produced by observation of particular data.

In adaptive GHQ (AGHQ).

6.1.3 Integrated nested Laplace approximation

The integrated nested Laplace approximation (INLA) method (Rue, Martino, et al. 2009) combines marginal Laplace approximations with quadrature to enable approximation of posterior marginal distributions.

6.2 Software

6.2.1 TMB

Template Model Builder [TMB, or when referring to the software TMB; Kristensen et al. (2016)] is an R package which implements the Laplace approximation. In TMB

derivatives are obtained using automatic differentiation (Baydin et al. 2017).

6.2.2 R-INLA

The R-INLA software implements the INLA method. R-INLA uses a formula interface (e.g. $y \sim 1 + x$) to facilitate use of INLA for common models. This is a beneficial design choice for new users. For more advanced users, the formula interface can impose constraints on model choice.

6.3 A universal INLA implementation

In this section, I implement the INLA method from scratch, using the TMB package. The result is universal in that it is compatible with any model with a TMB C++ template. This opens the door for application of INLA to models like Naomi which are not compatible with R-INLA. Indeed, Martino and Riebler (2019) note that “implementing INLA from scratch is a complex task”, and as a result “applications of INLA are limited to the (large class of) models implemented [in R-INLA]”. The potential benefits of a more flexible INLA implementation based on automatic differentiation were noted by Skaug (2009) in discussion of Rue, Martino, et al. (2009).

6.3.1 Epilepsy example

I use the epilepsy generalised linear mixed model example from Spiegelhalter et al. (1996) to demonstrate the implementation. This model is based on that of Breslow and Clayton (1993), a modification of Thall and Vail (1990), and the data are from an epilepsy drug double-blind clinical trial (Leppik et al. 1985). Rue, Martino, et al. (2009) (Section 5.2) demonstrate the INLA method using this example, and find a significant difference in approximation error depending on use of either the Gaussian or Laplace approximation for some parameters.

In the trial, patients $i = 1, \dots, 59$ were each assigned either the new drug $\text{Trt}_i = 1$ or placebo $\text{Trt}_i = 0$. Each patient made four visits the clinic $j = 1, \dots, 4$, and the observations y_{ij} are the number of seizures of the i th person in the two

weeks preceding their j th visit. The covariates used in the model were age \mathbf{Age}_i , baseline seizure counts \mathbf{Base}_i and an indicator for the final clinic visit \mathbf{V}_4 , which were all centered. The observations were modelled using a Poisson distribution $y_{ij} \sim \text{Poisson}(e^{\eta_{ij}})$ with linear predictor

$$\begin{aligned} \eta_{ij} = & \beta_0 + \beta_{\mathbf{Base}} \log(\mathbf{Baseline}_j/4) + \beta_{\mathbf{Trt}} \mathbf{Trt}_i + \beta_{\mathbf{Trt} \times \mathbf{Base}} \mathbf{Trt}_i \times \log(\mathbf{Baseline}_j/4) \\ & + \beta_{\mathbf{Age}} \log(\mathbf{Age}_i) + \beta_{\mathbf{V}_4} \mathbf{V}_{4j} + \epsilon_i + \nu_{ij}, \quad i \in [59], \quad j \in [4], \end{aligned}$$

where the prior distribution on each of the regression parameters, including the intercept, was $\mathcal{N}(0, 100^2)$. The random effects are IID $\epsilon_i \sim \mathcal{N}(0, 1/\tau_\epsilon)$ and $\nu_{ij} \sim \mathcal{N}(0, 1/\tau_\nu)$ with precision prior distributions $\tau_\epsilon, \tau_\nu \sim \Gamma(0.001, 0.001)$.

6.4 The Naomi model

The Naomi small-area estimation model (Eaton et al. 2021) synthesises data from multiple sources to estimate HIV indicators at a district-level, by age and sex.

6.4.1 Model structure

I consider a simplified version of Naomi defined only at the time of the most recent household survey with HIV testing. This version omits nowcasting and temporal projection. These time points involve limited inferences.

Household survey component

Consider a country in sub-Saharan Africa where a household survey with complex survey design has taken place. Let $x \in \mathcal{X}$ index district, $a \in \mathcal{A}$ index five-year age group, and $s \in \mathcal{S}$ index sex. For ease of notation, let i index the finest district-age-sex division included in the model. Let $I \subseteq \mathcal{X} \times \mathcal{A} \times \mathcal{S}$ be a set of indices i for which an aggregate observation is reported, and \mathcal{I} be the set of all I such that $I \in \mathcal{I}$.

Let $N_i \in \mathbb{N}$ be the known, fixed population size. HIV prevalence $\rho_i \in [0, 1]$, antiretroviral therapy (ART) coverage $\alpha_i \in [0, 1]$, and annual HIV incidence rate $\lambda_i > 0$ are modelled using linked regression equations.

Independent logistic regression models are specified for HIV prevalence and ART coverage in the general population such that $\text{logit}(\rho_i) = \eta_i^\rho$ and $\text{logit}(\alpha_i) = \eta_i^\alpha$. HIV incidence rate is modelled on the log scale as $\log(\lambda_i) = \eta_i^\lambda$, and depends on adult HIV prevalence and adult ART coverage. Let κ_i be the proportion recently infected among HIV positive persons. This proportion is linked to HIV incidence via

$$\kappa_i = 1 - \exp\left(-\lambda_i \cdot \frac{1 - \rho_i}{\rho_i} \cdot (\Omega_T - \beta_T) - \beta_T\right), \quad (6.20)$$

where the mean duration of recent infection Ω_T and the proportion of long-term HIV infections misclassified as recent β_T are strongly informed by priors for the particular survey.

These processes are each informed by household survey data. Weighted aggregate survey observations are calculated as

$$\hat{\theta}_I = \frac{\sum_j w_j \cdot \theta_j}{\sum_j w_j},$$

with individual responses $\theta_j \in \{0, 1\}$ and design weights w_j for each of $\theta \in \{\rho, \alpha, \kappa\}$. The design weights are provided by the survey and aim to reduce bias by decreasing possible correlation between response and recording mechanism (Meng 2018). The index j runs across all individuals in strata $i \in I$ within the relevant denominator i.e. for ART coverage, only those individuals who are HIV positive. The weighted observed number of outcomes is $y_I^\theta = m_I^\theta \cdot \hat{\theta}_I$ where

$$m_I^\theta = \frac{(\sum_j w_j)^2}{\sum_j w_j^2},$$

is the Kish effective sample size (ESS) (Kish 1965). As the Kish ESS is maximised by constant design weights, in exchange for reducing bias the ESS is reduced and hence variance increased. The weighted observed number of outcomes are modelled using a binomial working likelihood (Chen et al. 2014) defined to operate on the reals

$$y_I^\theta \sim \text{xBin}(m_I^\theta, \theta_I),$$

where θ_I are the following weighted aggregates

$$\rho_I = \frac{\sum_{i \in I} N_i \rho_i}{\sum_{i \in I} N_i}, \quad \alpha_I = \frac{\sum_{i \in I} N_i \rho_i \alpha_i}{\sum_{i \in I} N_i \rho_i}, \quad \kappa_I = \frac{\sum_{i \in I} N_i \rho_i \kappa_i}{\sum_{i \in I} N_i \rho_i}.$$

6.4.2 Connection to ELGMs

6.5 Extension of AGHQ to moderate dimensions

The Naomi model has $m = 24$ hyperparameters.

6.5.1 AGHQ with variable levels

6.5.2 Principal components analysis

6.6 Malawi case-study

6.6.1 NUTS convergence

6.6.2 Use of PCA-AGHQ

6.6.3 Model assessment

6.6.4 Inference comparison

6.6.5 Exceedance probabilities

6.7 Discussion

7

Future work and conclusions

7.1 Strengths

7.1.1 Chapter 4

- I designed experiments to thoroughly compare models for spatial structure using tools for model assessment such as proper scoring rules and posterior predictive checks.

7.1.2 Chapter 5

- I estimated HIV risk group proportions for AGYW, enabling countries to prioritise their delivery of HIV prevention services.
- I analysed the number of new infections that might be reached under a variety of risk stratification strategies.
- I used R-INLA to specify multinomial spatio-temporal models via the Poisson-multinomial transformation. This includes complex two- and three-way Kronecker product interactions defined using the `group` and `replicate` options.

7.1.3 Chapter 6

- I developed a novel Bayesian inference method, motivated by a challenging and practically important problem in HIV inference.
- The method enables integrated nested Laplace approximations to be fit to and studied on a wider class of models than was previously possible.
- My implementation of the method was straightforward, building on the **TMB** and **aghq** packages, and described completely and accessibly in Howes, Stringer, et al. (2023+).

7.2 Future work

Avenues for future work include:

1. Extending the risk group model described in Chapter 5 to include all adults 15-49. This may involve modelling of age-stratified sexual partnerships (Wolock et al. 2021). Such a model would likely fall out of the scope of **R-INLA**, but would be possible to write with **TMB** and therefore amenable to the methods discussed in Chapter 6.
2. Speeding up the implementation of Laplace marginals using the matrix algebra approximations described in Wood (2020).
3. Evaluating the accuracy of deterministic Bayesian inference methods for a broader variety of extended latent Gaussian models.

7.3 Conclusions

- Modelling complex data, more often than not, pushes the boundaries of the statistical toolkit available.
- A challenge I encountered was the difficulty of implementing identical models across multiple frameworks with the aim of studying the inference method. Or, of a similarly fraught nature, comparing different models implemented in different frameworks with the aim of studying model differences. The

Conclusions

frequently asked questions section of the **R-INLA** website (Rue 2023) notes that “the devil is in the details”. I have resolved this challenge by using a given **TMB** model template to fit models using multiple inference methodologies. The benefits of such a ecosystem of packages are noted by Stringer (2021). I particularly highlight the benefit of enabling analysts to easily vary their choice of inference method based on the stage of model development that they are in.

- To the best of my abilities, I have written this thesis, and the work described within it, in keeping with the principles of open science. I hope that doing so allows my work to be scrutinised, and optimistically built upon. This would not have been possible without a range of tools from the R ecosystem such as **rmarkdown** and **rticles**, as well as those developed within the MRC Centre for Global Infectious Disease Analysis such as **orderly** and **didehpc**.

Appendices



Spatial structure supplement

B

A model for risk group proportions supplement

B.1 Household survey data

				Sample size			
	Type	Year	Transactional sex question	15-19	20-24	25-29	Total
Botswana							
	BAIS	2013	✓	557	588	649	1794
Total				557	588	649	1794
Cameroon							
	DHS	2004	✗	2675	2207	1732	6614
	DHS	2011	✗	3588	3115	2655	9358
	PHIA	2017	✗	2620	2339	2259	7218
	DHS	2018	✓	3349	2463	2345	8157
Total				12232	10124	8991	31347
Kenya							
	DHS	2003	✗	1819	1709	1391	4919
	DHS	2008	✗	1767	1743	1419	4929
	DHS	2014	✗	2861	2534	2858	8253
Total				6447	5986	5668	18101
Lesotho							
	DHS	2004	✗	1761	1455	1026	4242
	DHS	2009	✗	1833	1543	1194	4570

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	DHS	2014	X	1537	1292	1067	3896
	PHIA	2017	✓	1156	1202	1054	3412
Total				6287	5492	4341	16120
Mozambique							
	AIS	2009	X	1031	1106	987	3124
	DHS	2011	X	2932	2299	2206	7437
	AIS	2015	X	1552	1389	1080	4021
Total				5515	4794	4273	14582
Malawi							
	DHS	2000	X	2914	2998	2358	8270
	DHS	2004	X	2407	2823	2135	7365
	DHS	2010	X	5031	4387	4309	13727
	DHS	2015	✓	5273	5094	3976	14343
	PHIA	2016	✓	1646	1934	1511	5091
Total				17271	17236	14289	48796
Namibia							
	DHS	2000	X	1427	1313	1098	3838
	DHS	2006	X	2203	1869	1544	5616
	DHS	2013	X	1852	1709	1481	5042
	PHIA	2017	✓	1491	1525	1370	4386
Total				6973	6416	5493	18882
Eswatini							
	DHS	2006	X	1265	1027	731	3023
	PHIA	2017	X	1031	895	811	2737
Total				2296	1922	1542	5760
Tanzania							
	AIS	2003	X	1466	1377	1270	4113
	AIS	2007	X	2137	1676	1509	5322
	DHS	2010	X	2221	1860	1613	5694
	AIS	2012	X	2474	1923	1815	6212
	PHIA	2016	✓	2999	2845	2521	8365
Total				11297	9681	8728	29706
Uganda							
	DHS	2000	X	1687	1541	1326	4554
	DHS	2006	X	1948	1660	1404	5012
	AIS	2011	X	2451	2164	1921	6536
	DHS	2011	X	2025	1664	1614	5303
	DHS	2016	✓	4276	3782	3014	11072
	PHIA	2016	X	3289	3059	2574	8922

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Total				15676	13870	11853	41399
South Africa							
	DHS	2016	✓	1505	1408	1397	4310
Total				1505	1408	1397	4310
Zambia							
	DHS	2007	✗	1598	1405	1373	4376
	DHS	2013	✗	3685	3036	2789	9510
	PHIA	2016	✓	2120	2045	1619	5784
	DHS	2018	✓	3112	2687	2166	7965
Total				10515	9173	7947	27635
Zimbabwe							
	DHS	1999	✗	1467	1230	1011	3708
	DHS	2005	✗	2128	1943	1438	5509
	DHS	2010	✗	1963	1796	1679	5438
	DHS	2015	✓	2154	1777	1646	5577
	PHIA	2016	✓	2114	1817	1573	5504
Total				9826	8563	7347	25736
Total				106397	95253	82518	284168

Table B.1: All of the surveys that we used in our analysis and their sample sizes, disaggregated by respondent age.

Survey	Exclusion reason
MOZ2003DHS	No GPS coordinates available to place survey clusters within districts.
TZA2015DHS	Insufficient sexual behaviour questions.
UGA2004AIS	Unable to download region boundaries.
ZMB2002DHS	No GPS coordinates available to place survey clusters within districts.

Table B.2: All of that surveys that were excluded from our analysis.

B.2 Spatial analysis levels

B. A model for risk group proportions supplement

Country	Number of areas	Analysis level
Botswana	27	3
Cameroon	58	2
Kenya	47	2
Lesotho	10	1
Mozambique	161	3
Malawi	33	5
Namibia	38	2
Eswatini	4	1
Tanzania	195	4
Uganda	136	3
South Africa	52	2
Zambia	116	2
Zimbabwe	63	2

Table B.3: The number of areas and analysis levels for each country that were used in our analysis.

C

Fast approximate Bayesian inference supplement

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