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# Spatial Data Analysis

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## Keywords

Bayesian hierarchical modeling, conditional autoregressive (CAR) models, cure rate models, disease mapping, multivariate CAR models, multivariate disease mapping, spatial survival analysis

## Abstract

With increasing accessibility to geographic information systems (GIS) software, statisticians and data analysts routinely encounter scientific data sets with geocoded locations. This has generated considerable interest in statistical modeling for location-referenced spatial data. In public health, spatial data routinely arise as aggregates over regions, such as counts or rates over counties, census tracts, or some other administrative delineation. Such data are often referred to as areal data. This review article provides a brief overview of statistical models that account for spatial dependence in areal data. It does so in the context of two applications: disease mapping and spatial survival analysis. Disease maps are used to highlight geographic areas with high and low prevalence, incidence, or mortality rates of a specific disease and the variability of such rates over a spatial domain. They can also be used to detect hot spots or spatial clusters that may arise owing to common environmental, demographic, or cultural effects shared by neighboring regions. Spatial survival analysis refers to the modeling and analysis for geographically referenced time-to-event data, where a subject is followed up to an event (e.g., death or onset of a disease) or is censored, whichever comes first. Spatial survival analysis is used to analyze clustered survival data when the clustering arises from geographical regions or strata. Illustrations are provided in these application domains.

## INTRODUCTION

The emergence of highly efficient geographical information systems (GIS) databases and associated computational resources has transformed the way spatial or geographical data are collected, stored, managed, and analyzed. Researchers in diverse disciplines within the physical, social, and environmental sciences and in public health are increasingly faced with the task of analyzing data that are geographically referenced and often presented as maps. Consequently, the past decade has seen significant development in statistical modeling of complex spatial data; for a variety of methods and applications, see the texts by Cressie (16), Webster & Oliver (41), Cromley & McLafferty (18), Møller (32), Schabenberger & Gotway (37), Waller & Gotway (40), Cressie & Wikle (17), and Banerjee et al. (5), among others.

Following convention, spatial data are often classified into one of three basic types: point-referenced data, point pattern data, and areal data. Point-referenced data sets consist of variables (e.g., outcomes and predictors) that are linked to a specific point location, customarily referenced by a coordinate system (e.g., longitude-latitude, easting-northing). Point-referenced data sets are not uncommon in environmental monitoring for public health, where pollutants are often measured at spatial fixed locations or monitoring stations. The spatial locations are considered fixed, and investigators are usually interested in the spatial distribution of the measurements and in predicting their levels at new spatial locations. Point pattern data refer to situations where the spatial locations themselves correspond to random events. Examples include locations being reported as sites of the occurrence of a particular disease. Areal data consist of variables that are aggregated over regions as counts or rates. Areal data are more common in public health applications, where geospatial referencing is not performed at very fine scales, such as GPS locations of households or small neighborhoods, to protect the privacy of human subjects.

The *Annual Review of Public Health* has published two excellent reviews on spatial analytic methods by Rushton (36) and Auchincloss et al. (1). This review differs from the previous *ARPH* articles because of its emphasis on the advances made in formal statistical modeling and inference for spatial data. It is beyond the scope of a single article to review all such methods. The aforementioned texts offer more comprehensive coverage. This review focuses primarily on areal data analysis because areal data are most conspicuous in public health. In fact, point patterns are often reported as areal aggregates, i.e., counts, rates of other summaries over well-delineated spatial regions such as counties or census tracts or zip codes, and subsequently modeled as areal data. Within this context, the review briefly discusses disease mapping for single diseases and for multiple diseases that may be associated with each other, as well as modeling of areally referenced survival data.

## SPATIAL MODELS FOR DISEASE MAPPING

In the fields of medicine and public health, researchers often seek a better understanding of regional patterns of disease. In the United States, publicly available data on precise locations of disease cases are fairly uncommon owing to strict confidentiality regulations. Summaries of disease at a regional level, however, are often relatively easy to obtain. Disease mapping is an epidemiological technique used to highlight geographic areas with high and low incidence or mortality rates of a specific disease and to map how such rates vary over the study region. Disease maps are often used to detect spatial clusters, which may generate hypotheses regarding common underlying environmental, demographical, or cultural factors shared by neighboring regions.

Although one could easily map the crude incidence and mortality rates, such maps can lead to spurious conclusions when the population sizes for some of the areal regions are small. Sparse populations usually result in large variability in the estimated rates and impair our ability to distinguish

chance variability from genuine differences. Statistical models that allow a more accurate depiction of true disease rates by borrowing information from neighboring regions will help mitigate the effects of sparsely populated regions and deliver better inference.

Perhaps the most conspicuous manner of modeling spatial dependence is to introduce spatially associated random effects within a Bayesian hierarchical setting [see, for example, Banerjee et al. (5)]. The Bayesian modeling and inferential framework is flexible and extremely rich in its capabilities to accommodate various scientific hypotheses and assumptions. In particular, it provides a cohesive framework for combining complex data models and external knowledge or expert opinion. This review discusses spatial modeling within a Bayesian context. The models and illustrations that follow are produced using Markov chain Monte Carlo (MCMC) simulation methods. Again, it is beyond the scope of this review to discuss MCMC algorithms. Details on established MCMC and other computational algorithms for spatial data can be found in the books by Møller (32), Gelman et al. (22), and Robert & Casella (35).

## Spatial Modeling of a Single Disease: A Brief Review

A popular class of models for areal data come from Markov random fields (MRF). These models are based on a Markov property, where the conditional distribution of the health outcome from a region, given the observations from all the other regions, depends only on the observations in the neighborhood. Here, we define the neighborhood by area adjacency, such that two regions are neighbors if they share a common boundary (or perhaps even meet at a point). Other definitions are sometimes used (e.g., regions with centroids within a given fixed distance).

Let  $Y_i$  be the observed number of cases of a certain disease in region  $i$ ,  $i = 1, \dots, n$ , and let  $E_i$  be the expected number of cases in this same region. A popular likelihood for mapping a single disease is

$$Y_i \stackrel{\text{ind}}{\sim} \text{Poisson}(E_i e^{\mu_i}), \quad i = 1, \dots, n, \quad 1.$$

where  $\mu_i = \mathbf{x}_i^\top \beta + \phi_i$  represents the log-relative risk expressed in terms of departures of the observed from expected counts, each  $\mathbf{x}_i$  is a vector of explanatory variables or covariates associated with region  $i$  having parameter coefficient  $\beta$ , and  $\phi_i$ s are spatially correlated random effects. We place a form of Gaussian MRF model, commonly referred to as the conditionally autoregressive (CAR) prior, on the random effects  $\phi = (\phi_1, \dots, \phi_n)^\top$ , i.e.,

$$\phi \sim N_n(0, [\tau(D - \alpha W)]^{-1}), \quad 2.$$

where  $N_n$  denotes the  $n$ -dimensional normal distribution,  $D$  is an  $n \times n$  diagonal matrix with diagonal elements  $m_i$  that denote the number of neighbors of region  $i$ , and  $W$  is the adjacency matrix of the map (i.e.,  $W_{ii} = 0$ , and  $W_{ii'} = 1$  if  $i'$  is adjacent to  $i$  and 0 otherwise). In Equation 2,  $\tau^{-1}$  is the spatial dispersion parameter, and  $\alpha$  is the spatial autocorrelation parameter. The CAR prior corresponds to the following conditional distribution of  $\phi_i$ :

$$\phi_i | \phi_j, j \neq i, \sim N \left( \frac{\alpha}{m_i} \sum_{i \sim j} \phi_j, \frac{1}{\tau m_i} \right), \quad i, j = 1, \dots, n, \quad 3.$$

where  $i \sim j$  denotes that region  $j$  is a neighbor of region  $i$ . The CAR structure (2) reduces to the well-known intrinsic conditionally autoregressive (ICAR) model [described in Besag et al. (10)] if  $\alpha = 1$  or an independence model if  $\alpha = 0$ . The ICAR model induces local smoothing by borrowing strength from the neighbors, whereas the independence model assumes independence of spatial rates and induces global smoothing. The smoothing parameter  $\alpha$  in the CAR prior (2) controls the strength of spatial dependence among regions, though it has long been appreciated

that a fairly large  $\alpha$  may be required to deliver significant spatial correlation [see Wall (39) for details on this]. Other variants of CAR models have been developed and applied to public health problems by Leroux et al. (30) and Dean et al. (19).

## Spatial Modeling of Multiple Diseases

Turning to multiple diseases, let  $Y_{ij}$  be the observed number of cases of disease  $j$  in region  $i$ ,  $i = 1, \dots, n$ ,  $j = 1, \dots, p$ , and let  $E_{ij}$  be the expected number of cases for the same disease in this same region. As in the previous section above, the  $Y_{ij}$ s are thought of as random variables, whereas the  $E_{ij}$ s are thought of as fixed and known. For the first level of the hierarchical model, conditional on the random effects  $\phi_{ij}$ , we assume the  $Y_{ij}$ s are independent of each other such that

$$Y_{ij} \stackrel{ind}{\sim} \text{Poisson}(E_{ij} e^{x_{ij}^T \beta_j + \phi_{ij}}), \quad i = 1, \dots, n, j = 1, \dots, p, \quad 4.$$

where each  $x_{ij}$  is a vector of region-specific explanatory variables for disease  $j$  having (possibly region-specific) parameter coefficients  $\beta_j$ . The key problem here is to specify rich and flexible spatial distributions for the  $\phi_{ij}$ s.

Carlin & Banerjee (11) and Gelfand & Vounatsou (21) generalized the univariate CAR (2) to a joint model for the random effects  $\phi_{ij}$ , which permits modeling of correlation among the  $p$  diseases while maintaining spatial dependence for each of the diseases. These models were subsequently subsumed by more general, and flexible, Bayesian hierarchical frameworks developed and implemented by Jin et al. (27, 28).

The idea in Jin et al. (28) is best expounded with  $p = 2$  diseases. Let  $\phi_1$  be the  $n \times 1$  vector of spatial random effects for the first disease, and let  $\phi_2$  be the same for the second disease. Jin et al. (28) specify a joint spatial model for  $\phi_1$  and  $\phi_2$  by specifying a conditional distribution of  $\phi_1$  given  $\phi_2$  and a marginal distribution for  $\phi_2$ . To achieve spatial smoothing, we assume that both these distributions are CARs. More precisely, we write the joint density as

$$p(\phi_1, \phi_2) = N(\phi_2 | 0, [\tau_2(D - \alpha_2 W)]^{-1}) \times N(\phi_1 | (\eta_0 I + \eta_1 W)\phi_2, [\tau_1(D - \alpha_1 W)]^{-1}), \quad 5.$$

where  $\eta_0$  and  $\eta_1$  are the bridging parameters associating the spatial effect for disease 1 in region  $i$  with disease 2 in region  $i$ . With disease 2 in a neighboring region,  $\rho_1$  and  $\rho_2$  are smoothing parameters associated with the conditional distribution of  $\phi_1 | \phi_2$  and the marginal distribution of  $\phi_2$  respectively, and  $\tau_1$  and  $\tau_2$  scale the precision of  $\phi_1 | \phi_2$  and  $\phi_2$ , respectively. The model in Equation 5 yields a legitimate probability density as long as the two CAR distributions on the right-hand side are valid, which means that the two dispersion matrices for  $\phi_1 | \phi_2$  and  $\phi_2$  must be positive definite. Jin et al. (28) provide conditions for these matrices to be positive definite.

Models where the spatial random effects are shown as in Equation 5 are known as generalized multivariate conditionally autoregressive (GMCAR) models. The specification in Equation 5 subsumes several special cases in the multivariate disease mapping literature. Setting  $\rho_1 = \rho_2 = \rho$  and  $\eta_1 = 0$  produces a model showing that the association between the two diseases remains the same across the regions. If we assume  $\rho_1 \neq \rho_2$  and  $\eta_0 = \eta_1 = 0$ , then we ignore dependence between the multivariate components, and the model turns out to be equivalent to fitting two separate univariate CAR models. Finally, if we assume  $\rho_1 = \rho_2 = 0$ ,  $\eta_0 \neq 0$ , and  $\eta_1 = 0$ , then the model becomes a simple bivariate normal model with no spatial association.

The above approach is appealing for two diseases, or perhaps at most for three diseases, but using it to model several diseases at once has its limitations. An inherent problem with these methods is that their conditional specification imposes a potentially arbitrary order on the variables being modeled, as they lead to different marginal distributions depending on the conditioning sequence [i.e., whether to model  $p(\phi_1 | \phi_2)$  and then  $p(\phi_2)$ , or  $p(\phi_2 | \phi_1)$  and then  $p(\phi_1)$ ]. This problem is

somewhat mitigated in certain (e.g., medical and environmental) contexts where a natural order is reasonable, but in many disease mapping contexts this is not the case.

To obviate the ordering issue, Jin et al. (27) developed an order-free, joint framework for multivariate areal modeling that allows versatile spatial structures, yet is computationally feasible for many outcomes. These are called coregionalized MCAR models, named after linear models of coregionalization in multivariate geostatistics [see, e.g., Wackernagel (38)]. The underlying idea here is to develop richer spatial association models using linear transformations of much simpler spatial distributions. The objective is to allow explicit smoothing of cross-covariances without being hampered by conditional ordering. In particular, suppose we assume a common proximity specification for each component of the random effects vector,  $\phi$ . Then, we could write  $\phi = A\psi$ , where  $\psi_j$ , the  $j$ th component of  $\psi$ , is a univariate intrinsic CAR with precision parameter  $\tau_j^2$  and each of the component CAR models are independent. The matrix  $A$  represents the linear transformation that maps independent CAR effects for each disease to correlated CAR (or multivariate CAR) effects for the diseases.

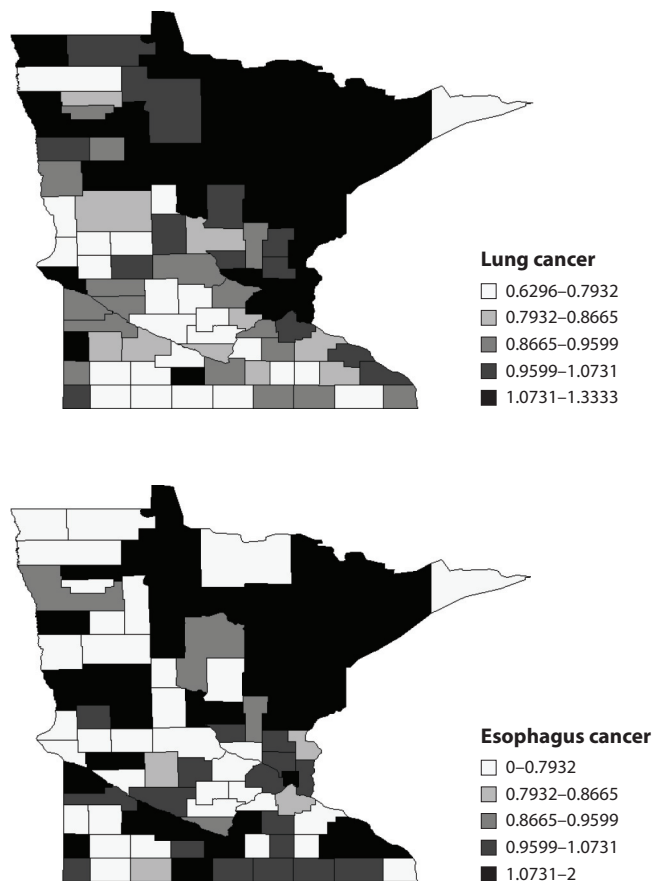
MCAR models are not the only available option for analyzing multivariate areal data. Zhang et al. (43) have developed an arguably simpler approach by adapting smoothed ANOVA (SANOVA) models (24) for areal data. The underlying idea is to extend SANOVA to cases in which one factor is a spatial map, which is smoothed using a CAR model, and a second factor is, for example, a type of disease. Data sets routinely lack enough information to identify the additional structure of MCAR. SANOVA offers a simpler and more intelligible structure than MCAR while performing equally well. Nevertheless, the MCAR and more general CAR-based models offer a rich inferential framework for capturing complex spatial associations. We focus on MCAR models and their variants within the disease mapping context in the remainder of this article.

## Illustration

We illustrate with a brief example from Jin et al. (28), who modeled the numbers of deaths due to cancers of the lung and esophagus between 1991 and 1998 across the 87 counties in Minnesota. The county-level maps of the raw standardized mortality ratios (i.e.,  $SMR_{ij} = Y_{ij}/E_{ij}$ ) shown in **Figure 1** exhibit evidence of correlation both across space and between cancers, motivating use of our proposed GMCAR models. The bottom row shows the smoothed maps obtained from the GMCAR model specified using a CAR prior for the conditional distribution [lung|esophagus] and another CAR for the marginal distribution [esophagus].

We fit the model Banerjee & Carlin (4) to this data set. To determine  $E_{ij}$ , we account for each county's age distribution by calculating the expected age-adjusted number of deaths due to cancer  $j$  in county  $i$  as  $E_{ij} = \sum_{k=1}^m \omega_{jk} N_{ik}$  for  $i = 1, \dots, 87$  and  $j = 1, 2$ , where  $\omega_{jk} = (\sum_{i=1}^{87} D_{ijk}) / (\sum_{i=1}^{87} N_{ik})$  is the age-specific death rate for cancer  $j$  and age group  $k$  over all Minnesota counties,  $D_{ijk}$  is the number of deaths in age group  $k$  for county  $i$  and cancer  $j$ , and  $N_{ik}$  is the total population at risk in age group  $k$  for county  $i$ . Jin et al. (28) conducted exploratory analysis on the basis of least-squares estimation as well as formal Bayesian model comparison methods to show that a GMCAR model specified using CAR distributions for [lung|esophagus] and [esophagus] was preferable to modeling [esophagus|lung]. The GMCAR models are easily implemented in the Bayesian modeling language BUGS (see <http://www.biostat.umn.edu/~brad/software.html> for the code and the data). **Figure 2** presents maps of the smoothed standardized mortality ratios (SMRs) for lung and esophagus cancer in Minnesota from the GMCAR.

Jin et al. (28) also reported that the estimate of the parameter  $\eta_1$  was statistically significant for the GMCAR with [lung|esophagus] and not significant in the reverse order. We also saw that the posterior distribution of the linking parameters  $\eta_0$  and  $\eta_1$  had mostly positive support,

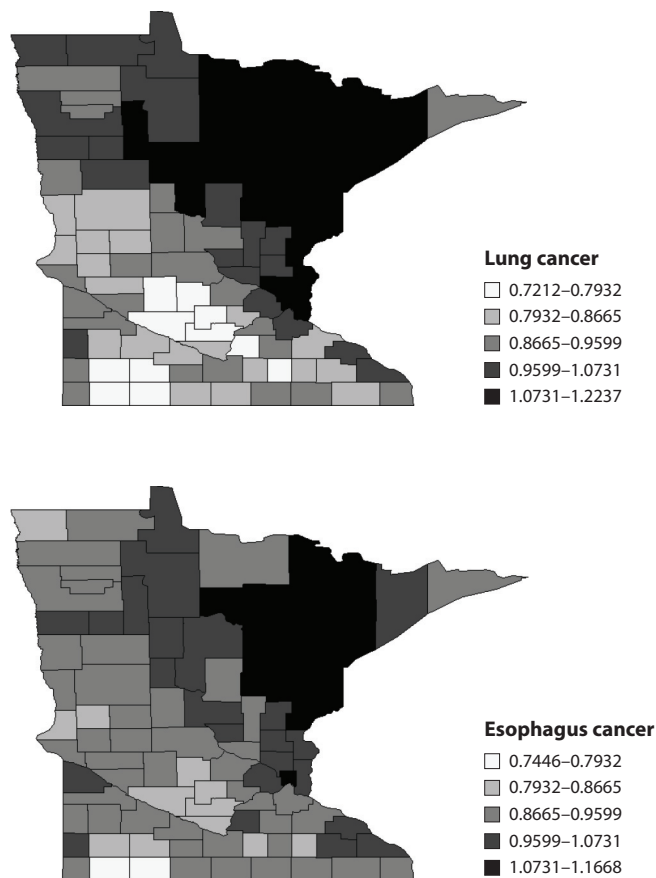


**Figure 1**

Maps of raw standard mortality ratios (SMRs) of lung and esophagus cancer in Minnesota between 1991 and 1998.

meaning that the two cancers had positive spatial correlation. This is also evident from the maps of the posterior means of the SMRs for the two cancers under the full model shown in **Figure 2**. Incidence of the two cancers is clearly strongly correlated, with higher fitted ratios extending from the Twin Cities metro area (eastern side, about one-third of the way up) to the mining- and tourism-oriented north and northeast, regions where conventional wisdom suggests that cigarette smoking may be more common.

The GMCAR delivered point and 95% equal-tail interval estimates of 0.602 and (0.0267, 0.979) for  $\rho_1$ , and 0.699 and (0.0802, 0.973) for  $\rho_2$ . These are spatial parameters, but while their values are between 0 and 1 they are not “correlations” in the usual sense; the moderate point estimates and wide confidence intervals suggest a relatively modest degree of spatial association in the random effects. Note also that in this setup,  $\rho_2$  measures spatial association in the esophagus random effects  $\phi_1$ , whereas  $\rho_1$  measures spatial association in the lung random effects  $\phi_1$  given the esophagus random effects  $\phi_2$ . Turning to  $\tau_1$  and  $\tau_2$ , under the GMCAR we obtained 32.65 (16.98, 66.71) and 13.73 (4.73, 38.05) as our point and interval estimates, respectively. Because these parameters measure spatial precision for each disease, they suggest slightly more variability in the



**Figure 2**

Maps of posterior means of standardized mortality ratios (SMRs) of lung and esophagus cancer in Minnesota between 1991 and 1998 from the generalized multivariate conditionally autoregressive (GMCAR) model with conditioning order [lung|esophagus].

esophagus random effects, although again comparison is difficult here because  $\tau_2$  is a marginal precision for  $\phi_2$  whereas  $\tau_1$  is a conditional precision for  $\phi_1$  given  $\phi_2$ .

## SPATIAL SURVIVAL ANALYSIS

Survival models, such as in Cox & Oakes (15), are widely used in biostatistics and epidemiology for analyzing time-to-event data, where a subject is followed up to an event (e.g., death or onset of a disease) or is “censored,” whichever comes first. Right censoring refers to situations where the event does not occur for a subject during the period of the study and the subject’s time to event is censored at the study end point. Certain study designs can produce left-censored or interval-censored data, defined analogously. As opposed to modeling disease incidence and mortality, survival models focus on how many are expected to survive after a certain period of time and the rate of failure, as well as to ascertain which underlying factors (e.g., gender, race, age, type of cancer, treatment obtained, and access to health care facilities) generate shortened or prolonged survival.



The past decade has seen much demand for the analysis of spatially referenced survival data. When each subject can be referenced with respect to a clinical site or geographical region, we might suspect that random effects corresponding to proximate regions will be similar in magnitude. Models for spatially arranged survival data customarily introduce spatial frailties, such as in Banerjee et al. (7). How these spatial frailties are introduced in survival models depends on the specific model. We briefly discuss a few alternate spatial survival models. Apart from the spatial distribution for the frailties, one needs to model a spatial hazard function with the understanding that expected survival times (or hazard rates) will be more similar in neighboring regions, owing to underlying factors (access to care, willingness of the population to seek care, etc.) that vary spatially. This expectation is in contrast to the similarity observed among survival times from subjects in proximate regions, which is not necessarily implied by spatially associated frailties.

## Survival Models with Spatial Frailties

Let  $T$  be the waiting time for a subject to experience an event (e.g., disease onset, relapse, death). The subject's survival function is defined as  $S(t) = P(T \geq t)$  and the hazard function as  $b(t) = f(t)/S(t)$ , where  $f(t)$  is the probability density function of  $T$ . Let  $(i, j)$  index the  $j$ -th subject in region  $i$  and let  $\{(t_{ij}, \delta_{ij}) : i = 1, 2, \dots, I; j = 1, 2, \dots, n_i\}$  be observations from  $n$  subjects in a study, where  $t_{ij}$  indicates the time at which either subject  $(i, j)$  experienced the event or the subject was censored. Associated with each  $t_{ij}$  is an event indicator,  $\delta_{ij}$ , where  $\delta_{ij} = 1$  if the event occurred before the termination of the study and  $\delta_{ij} = 0$  if the subject was censored. For right-censored data, we have the likelihood

$$\prod_{j=1}^{n_i} f(t_{ij})^{\delta_{ij}} S(t_{ij})^{1-\delta_{ij}} = \prod_{j=1}^{n_i} b(t_{ij})^{\delta_{ij}} S(t_{ij}). \quad 6.$$

If  $\delta_{ij} = 1$ , then subject  $j$  contributes  $f(t_{ij}) = b(t_{ij})S(t_{ij})$  to the likelihood, whereas if  $\delta_{ij} = 0$ , then it contributes  $S(t_{ij})$  to the likelihood. Cox & Oakes (15) provide the corresponding expressions for left-censored and interval-censored data.

Let  $\mathbf{x}_{ij}$  be a  $p \times 1$  vector of observed explanatory variables associated with subject  $(i, j)$ . To account for heterogeneity in the population, most survival models will introduce these explanatory variables in Equation 6 in the hazard function. For example, the proportional hazards model stipulates that

$$b(t_{ij}; \mathbf{x}_{ij}) = b_0(t_{ij}) \exp(\mathbf{x}_{ij}^\top \boldsymbol{\beta}), \quad 7.$$

where  $b_0(t)$  is a baseline hazard function affected only multiplicatively by the exponential term involving the explanatory variables. Another option is a "proportional odds" model (9), which requires the survival function for subject  $(i, j)$  to satisfy

$$\frac{S(t|\mathbf{x}_{ij})}{1 - S(t|\mathbf{x}_{ij})} = \frac{S_0(t)}{1 - S_0(t)} \exp(\mathbf{x}_{ij}^\top \boldsymbol{\beta}). \quad 8.$$

Yet another alternative is the accelerated failure time model. Here, the survival function for subject  $(i, j)$  is  $S(t) = S_0(t/\gamma_{ij})$ , where  $S_0(t)$  is any parametric survival function and  $\gamma_{ij} = \exp\{\mathbf{x}_{ij}^\top \boldsymbol{\beta}\}$ . The corresponding hazard function for subject  $(i, j)$  is  $b(t) = b_0(t/\gamma_{ij})/\gamma_{ij}$ , where  $b_0(t)$  is the hazard derived from  $S_0(t)$ . In each of the above situations, the hazard function can be modeled using parametric or nonparametric statistical methods. The data-analytic settings where the above specifications are appropriate, or not, have been comprehensively explored and documented in the survival analysis literature. For example, the proportional odds model posits that the hazard ratio approaches unity over time, i.e., the covariate effects on the hazards disappear over time, which is clearly distinct from the proportional hazards model. The interpretation of the regression



component significantly differs. The term  $\exp\{x^\top \beta\}$  in the proportional odds model reflects the change in the odds of survival (or failure, depending on the parameterization) given the observed covariates or risk factors.

Li & Ryan (31) provided the basis for legitimate likelihood-based inference from semiparametric spatial survival models. They proposed modeling the hazard function nonparametrically and the spatially correlated frailties using different spatial covariance functions. These models were applied to the East Boston Asthma Study to detect prognostic factors leading to childhood asthma. Henderson et al. (23) proposed using multivariate Gamma distributions to investigate spatial association and variation in the survival of acute myeloid leukemia patients in northern England. Banerjee et al. (7) proposed a Bayesian hierarchical framework to introduce spatially correlated frailties and compared performances between frailties modeled using Markov random field and geostatistical covariance functions. Data from a large infant mortality study in the state of Minnesota were analyzed. Subsequent papers explored Bayesian semiparametric modeling (2), spatiotemporal modeling (3, 8), semiparametric proportional odds models with spatial frailties (6), joint survival and longitudinal modeling with frailties (44), and parametric accelerated failure time models (42). Finally, we refer the reader to Lawson et al. (29) for spatial survival models that do not deploy spatial frailties.

## Spatial Cure Rate Models

In light of significant progress in medical and health sciences, scientists and health professionals increasingly encounter data sets in which patients are expected to be cured. Models accounting for cure are important for understanding prognosis in potentially terminal diseases. Traditional parametric survival models such as Weibull or Gamma [see, e.g., Cox & Oakes (15)] do not account for cure, assuming instead that individuals who do not experience the event are censored. The subtle distinction between censoring and cure is worth noting: A subject who does not fail within the time window of the experiment is considered censored, whereas a subject is cured if he will never relapse. The latter is clearly a more abstract concept because we are never able to observe a cure, yet there is interest in estimating the probability of such an outcome, especially in various disease-relapse settings.

Cure models, such as survival models, also enjoy a rich literature too vast to be comprehensively reviewed here. The reader should see Ibrahim et al. (26) for a methodological introduction, whereas Othus et al. (34) offer a more recent review and practical introduction. Cooner et al. (14) build on their previously proposed flexible framework [13; also see Hurtado Rúa & Dey (25)] to introduce spatial frailties in cure models for geographically referenced data. Banerjee & Carlin (4) propose a spatial extension of earlier work by Chen et al. (12), which assumes that some latent biological process is generating the observed data. Suppose that subject  $(i, j)$  has  $N_{ij}$  potential latent (unobserved) risk factors, the presence of any of which (i.e.,  $N_{ij} \geq 1$ ) will ultimately manifest the event. Chen et al. (12) consider the case of multiple latent factors, assuming that the  $N_{ij}$  are distributed as independent Poisson random variables with mean  $\theta_{ij}$ , i.e.,  $p(N_{ij}|\theta_{ij})$  is  $Poi(\theta_{ij})$ . For example, in cancer settings, these factors may correspond to metastasis-competent tumor cells within the individual. Subjects who do not experience the event during the observation period are considered censored. Thus, if  $U_{ijk}$ ,  $k = 1, 2, \dots, N_{ij}$  is the time to an event arising from the  $k$ -th latent factor for subject  $(i, j)$ , the observed time to event for an uncensored individual is generated by  $T_{ij} = \min\{U_{ijk}, k = 1, 2, \dots, N_{ij}\}$ .

Given  $N_{ij}$ , the  $U_{ijk}$ s are independent with survival function  $S(t|\Psi_{ij})$  and corresponding density function  $f(t|\Psi_{ij})$ . The parameter  $\Psi_{ij}$  is a collection of all the parameters (including possible regression parameters) that may be involved in a parametric specification for the survival function  $S$ .

In this section, we work with a two-parameter Weibull distribution specification for the density function  $f(t|\Psi_{ij})$ , where we allow the Weibull scale parameter  $\rho$  to vary across the regions, and  $\eta$ , which may serve as a link to covariates in a regression setup, to vary across individuals. Therefore,  $f(t|\rho_i, \eta_{ij}) = \rho_i t^{\rho_i-1} \exp(\eta_{ij} - t^{\rho_i} \exp(\eta_{ij}))$ .

Banerjee & Carlin (4) analyze smoking cessation data using interval-censored spatial cure rate models. The outcome of interest is the time for a subject to relapse into smoking. Here, we observe only a time *interval*  $(t_{ijL}, t_{ijU})$  within which the event (smoking relapse) is known to have occurred. For patients who did not resume smoking prior to the end of the study, we have  $t_{ijU} = \infty$ , yielding the case of right-censoring at time point  $t_{ijL}$ . Thus we now set  $v_{ij} = 1$  if subject  $ij$  is interval-censored (i.e., the subject has experienced the event) and  $v_{ij} = 0$  if the subject is right-censored.

Following Finkelstein (20), the general interval-censored cure rate likelihood is given by

$$\begin{aligned} & \prod_{i=1}^I \prod_{j=1}^{n_i} [S(t_{ijL}|\rho_i, \eta_{ij})]^{N_{ij}-v_{ij}} \{N_{ij}[S(t_{ijL}|\rho_i, \eta_{ij}) - S(t_{ijU}|\rho_i, \eta_{ij})]\}^{v_{ij}} \\ &= \prod_{i=1}^I \prod_{j=1}^{n_i} [S(t_{ijL}|\rho_i, \eta_{ij})]^{N_{ij}} \left\{ N_{ij} \left( 1 - \frac{S(t_{ijU}|\rho_i, \eta_{ij})}{S(t_{ijL}|\rho_i, \eta_{ij})} \right) \right\}^{v_{ij}}. \end{aligned}$$

If  $N_{ij} \stackrel{iid}{\sim} \text{Ber}(\theta_{ij})$ , then the marginal likelihood obtained by summing over the  $N_{ij}$ s is  $L(\{(t_{ijL}, t_{ijU})\}|\{\rho_i\}, \{\theta_{ij}\}, \{\eta_{ij}\}, \{v_{ij}\})$  and can be written as

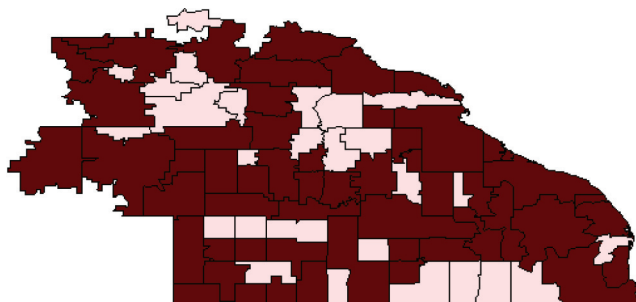
$$\prod_{i=1}^I \prod_{j=1}^{n_i} S^*(t_{ijL}|\theta_{ij}, \rho_i, \eta_{ij}) \left\{ 1 - \frac{S^*(t_{ijU}|\theta_{ij}, \rho_i, \eta_{ij})}{S^*(t_{ijL}|\theta_{ij}, \rho_i, \eta_{ij})} \right\}^{v_{ij}}. \quad 9.$$

As with the covariates, we introduce the frailties  $\phi_i$  through the Weibull link as intercept terms in the log-relative risk; that is, we set  $\eta_{ij} = \mathbf{x}_{ij}^\top \beta + \phi_i$ . Here we allow the  $\phi_i$  to be spatially correlated across the regions; similarly we would like to permit the Weibull baseline hazard parameters,  $\rho_i$ , to be spatially correlated. A natural approach in both cases is to use a univariate CAR prior. Although one may certainly employ separate, independent CAR priors on  $\phi \equiv \{\phi_i\}$  and  $\zeta \equiv \{\log \rho_i\}$ , another option is to use a bivariate CAR model for the  $\delta_i = \{\phi_i, \zeta_i\} = \{\phi_i, \log \rho_i\}$ . For further details, see Banerjee & Carlin (4).

## Illustration

We present part of a more elaborate data analysis as part of a smoking cessation study reported by Murray et al. (33), which is of particular relevance to studies of lung health and primary cancer control. For our illustration here, we restrict attention to 223 subjects from 54 zip codes in southeastern Minnesota. These subjects were all smokers at study entry and were randomized into either a smoking intervention (SI) group or a usual care (UC) group, which received no antismoking intervention. On the basis of a consecutive five-year monitoring period between 1994 and 1998, each of these subjects were known to have quit smoking at least once during these five years. The event of interest is whether they relapse into smoking (resume smoking). The raw data revealed that 29.7% resumed smoking, producing an empirical cure fraction of 0.703. Additional information available for each subject includes sex, years as a smoker, and the average number of cigarettes smoked per day prior to the quit attempt.

As is not unusual in spatial data sets, the 54 zip codes that contributed the data were not contiguous, which made it difficult to fit neighborhood-based models. Banerjee & Carlin (4) considered 81 contiguous zip codes shown in **Figure 3**, which included the 54 dark-shaded regions



**Figure 3**

Map showing a missingness pattern for the smoking cessation data between 1994 and 1998 from 54 zip codes in southeastern Minnesota: Lightly shaded regions are those having no responses.

that had patients in the data set; the 27 regions that did not contribute patients were treated as if the data were missing.

**Table 1** presents estimated posterior quantiles for the fixed effects  $\beta$ , cure fraction  $\theta$ , and hyperparameters. Smoking intervention, expectedly, produces a significant decrease in the log relative risk of relapse. Women seem to be more likely to relapse than men. This result is often attributed to the (real or perceived) risk of weight gain following smoking cessation. The number of cigarettes smoked per day seems to be less significant; however, what is perhaps somewhat counterintuitive is that shorter-term smokers relapse sooner, perhaps attributable to subjects being better able to quit smoking as they age.

## CONCLUDING REMARKS

This article has provided a glimpse of the different types of statistical spatial models available for analyzing regionally aggregated data (or areal data) and the type of statistical inference that is obtained from such models. Although the illustrations provided here aggregated the data over a number of years and did not attempt to model associations across time, such associations can also be modeled by allowing the spatial random effects to vary across time. Also, this review has restricted attention to the CAR models, which are especially congruous with Bayesian statistical

**Table 1** Posterior quantiles, full model, interval-censored case

| Parameter                                | Median | (2.5%, 97.5%)    |
|--|--------|------------------|
| Intercept                                | −2.720 | (−4.803, −0.648) |
| Sex (male = 0)                           | 0.291  | (−0.173, 0.754)  |
| Duration as smoker                       | −0.025 | (−0.059, 0.009)  |
| SI/UC (usual care = 0)                   | −0.355 | (−0.856, 0.146)  |
| Cigarettes smoked per day                | 0.010  | (−0.010, 0.030)  |
| $\theta$ (cure fraction)                 | 0.694  | (0.602, 0.782)   |
| $\rho_\phi$                              | 0.912  | (0.869, 0.988)   |
| $\rho_\zeta$                             | 0.927  | (0.906, 0.982)   |
| Spatial variance component for $\phi_i$  | 0.005  | (0.001, 0.029)   |
| Spatial variance component for $\zeta_i$ | 0.007  | (0.002, 0.043)   |

Abbreviations: SI, smoking intervention; UC, usual care.

inference. Other types of spatial dependence structures, such as simultaneous autoregressive (SAR) models, are very popular, and perhaps better suited, for maximum-likelihood-based inference. Comparisons between these models can be found in Wall (39). Several other variants of such models, including spatiotemporal extensions, can be found in Banerjee et al. (5) and references therein.

### SUMMARY POINTS

1. Statistical modeling and scientific inference using spatially referenced data sets are becoming increasingly common in public health research. Examples include disease mapping and spatial survival analysis.
2. Researchers are formulating more complex spatially oriented hypotheses that require formal model-based testing and inference.
3. Statistical models for spatial data introduce dependence on the basis of whether the data are point referenced or areally referenced. The latter, which are usually presented as aggregates or summaries over regions, are more common in public health research and practice because they protect patients' privacy.
4. Much of the statistical research over the past decade has focused on stochastic models for spatial dependence and how they can be introduced as random effects within Bayesian hierarchical models. These models are estimated using computationally intensive MCMC methods and have been applied to diverse data-analytic settings, including multiple disease mapping and spatial survival analysis.

### FUTURE ISSUES

1. As the accessibility to GIS and related computational resources continues to expand, spatial statisticians are encountering increasingly complex data sets with more demanding research questions. The scope for spatial modeling and analysis within public health will continue to expand, ushering in new domains of application.
2. A large part of methodological research will be devoted to the development of probability models, estimation methods, and computational algorithms for analyzing such data sets.
3. Statistical methods for analyzing spatially referenced data sets are computationally expensive and become unfeasible for large data sets. As spatial data sets become larger, statisticians start encountering the so-called "big data" problems in geostatistics. This area has started to garner much attention over the past five years or so and is seeing increasing research activity with regard to statistical models, methods, and algorithms for massive spatial data sets.

### DISCLOSURE STATEMENT

The author is not aware of any affiliations, memberships, funding, or financial holdings that might be perceived as affecting the objectivity of this review.

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# Contents

## Epidemiology and Biostatistics

|   |     |
|---|-----|
| Improved Designs for Cluster Randomized Trials<br><i>Catherine M. Crespi</i> .....  | 1   |
| Mediation Analysis: A Practitioner's Guide<br><i>Tyler J. VanderWeele</i> .....   | 17  |
| Nutritional Determinants of the Timing of Puberty<br><i>Eduardo Villamor and Erica C. Jansen</i> .....  | 33  |
| Spatial Data Analysis<br><i>Sudipto Banerjee</i> .....  | 47  |
| Using Electronic Health Records for Population Health Research:<br>A Review of Methods and Applications<br><i>Joan A. Casey, Brian S. Schwartz, Walter F. Stewart, and Nancy E. Adler</i> ..... | 61  |
| Metrics in Urban Health: Current Developments and Future Prospects<br><i>Amit Prasad, Chelsea Bettina Gray, Alex Ross, and Megumi Kano</i> .....  | 113 |
| A Transdisciplinary Approach to Public Health Law: The Emerging<br>Practice of Legal Epidemiology<br><i>Scott Burris, Marice Ashe, Donna Levin, Matthew Penn, and Michelle Larkin</i> .....     | 135 |

## Environmental and Occupational Health

|  |     |
|--|-----|
| Cumulative Environmental Impacts: Science and Policy to Protect<br>Communities<br><i>Gina M. Solomon, Rachel Morello-Frosch, Lauren Zeise, and John B. Faust</i> .....   | 83  |
| Heat, Human Performance, and Occupational Health: A Key Issue for<br>the Assessment of Global Climate Change Impacts<br><i>Tord Kjellstrom, David Briggs, Chris Freyberg, Bruno Lemke, Matthias Otto,<br/>and Olivia Hyatt</i> ..... | 97  |
| Metrics in Urban Health: Current Developments and Future Prospects<br><i>Amit Prasad, Chelsea Bettina Gray, Alex Ross, and Megumi Kano</i> .....   | 113 |
| One Hundred Years in the Making: The Global Tobacco Epidemic<br><i>Heather Wipfli and Jonathan M. Samet</i> .....  | 149 |



## Public Health Practice

|   |     |
|---|-----|
| A Transdisciplinary Approach to Public Health Law: The Emerging<br>Practice of Legal Epidemiology<br><i>Scott Burris, Marice Ashe, Donna Levin, Matthew Penn, and Michelle Larkin</i> .....     | 135 |
| One Hundred Years in the Making: The Global Tobacco Epidemic<br><i>Heather Wipfli and Jonathan M. Samet</i> .....   | 149 |
| The Double Disparity Facing Rural Local Health Departments<br><i>Jenine K. Harris, Kate Beatty, J.P. Leider, Alana Knudson, Britta L. Anderson,<br/>and Michael Meit</i> .....                  | 167 |
| Using Electronic Health Records for Population Health Research:<br>A Review of Methods and Applications<br><i>Joan A. Casey, Brian S. Schwartz, Walter F. Stewart, and Nancy E. Adler</i> ..... | 61  |
| Defining and Assessing Public Health Functions: A Global Analysis<br><i>Jose M. Martin-Moreno, Meggan Harris, Elke Jakubowski, and Hans Kluge</i> .....   | 335 |

## Social Environment and Behavior

|  |     |
|--|-----|
| Civil Rights Laws as Tools to Advance Health in the Twenty-First<br>Century<br><i>Angela K. McGowan, Mary M. Lee, Cristina M. Meneses, Jane Perkins,<br/>and Mara Youdelman</i> .....  | 185 |
| Documenting the Effects of Armed Conflict on Population Health<br><i>Barry S. Levy and Victor W. Sidel</i> .....   | 205 |
| Latino Immigrants, Acculturation, and Health: Promising New<br>Directions in Research<br><i>Ana F. Abraído-Lanza, Sandra E. Echeverría, and Karen R. Flórez</i> .....  | 219 |
| Making Healthy Choices Easier: Regulation versus Nudging<br><i>Pelle Guldberg Hansen, Laurits Rohden Skov, and Katrine Lund Skov</i> .....   | 237 |
| Preventing Obesity Across Generations: Evidence for Early Life<br>Intervention<br><i>Debra Haire-Joshu and Rachel Tabak</i> .....  | 253 |
| Sugar-Sweetened Beverages and Children's Health<br><i>Rebecca J. Scharf and Mark D. DeBoer</i> .....   | 273 |
| Visible and Invisible Trends in Black Men's Health: Pitfalls and<br>Promises for Addressing Racial, Ethnic, and Gender Inequities<br>in Health<br><i>Keon L. Gilbert, Rashawn Ray, Arjumand Siddiqi, Shivan Shetty,<br/>Elizabeth A. Baker, Keith Elder, and Derek M. Griffith</i> ..... | 295 |

|  |     |
|--|-----|
| One Hundred Years in the Making: The Global Tobacco Epidemic<br><i>Heather Wipfli and Jonathan M. Samet</i> .....          | 149 |
| The Health Effects of Income Inequality: Averages and Disparities<br><i>Beth C. Truesdale and Christopher Jencks</i> ..... | 413 |

## Health Services

|  |     |
|--|-----|
| A Review of Opportunities to Improve the Health of People Involved<br>in the Criminal Justice System in the United States<br><i>Nicholas Freudenberg and Daliah Heller</i> ..... | 313 |
| Defining and Assessing Public Health Functions: A Global Analysis<br><i>Jose M. Martin-Moreno, Meggan Harris, Elke Jakubowski, and Hans Kluge</i> .....                          | 335 |
| Opportunities for Palliative Care in Public Health<br><i>Liliana De Lima and Tania Pastrana</i> .....  | 357 |
| Racial and Ethnic Disparities in the Quality of Health Care<br><i>Kevin Fiscella and Mechelle R. Sanders</i> .....   | 375 |
| Rural Health Care Access and Policy in Developing Countries<br><i>Roger Strasser, Sophia M. Kam, and Sophie M. Regalado</i> .....  | 395 |
| The Health Effects of Income Inequality: Averages and Disparities<br><i>Beth C. Truesdale and Christopher Jencks</i> .....   | 413 |

## Indexes

|   |     |
|---|-----|
| Cumulative Index of Contributing Authors, Volumes 28–37 ..... | 431 |
| Cumulative Index of Article Titles, Volumes 28–37 .....       | 437 |

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