

Using BRFSS Data to Estimate County-level Colorectal Cancer Screening Prevalence in Missouri

METHODS: NOTATIONS Respondents were classified into 12 groups based on age (50–64, 65–74, 75+), gender and race (white, non-white). Let • n_{ijkl} : sample size (number of respondents) in county $i \in \{1, ..., 115\}$, age group $j \in \{1, 2, 3\}$, gender $k \in \{1, 2\}$ and race $l \in \{1, 2\}$; age. • *N_{ijkl}*: true population size for category *ijkl*, which is obtained from Census data; of people in Missouri aged 50 and older have had the screening. However, county-level CRCS prevalence cannot be directly obtained from BRFSS due to • y_{ijkl} : number of respondents who have had CRCS for category ijkl; small or even zero sample sizes. • *Y*_{*ijkl*}: true population total for people who have had CRCS for category *ijkl*; county-level estimates. Questions asked in CLS were similar to those in BRFSS. • p_{ijkl} : proportion of people who have had CRCS for category ijkl. Since much larger sample sizes were obtained for counties in Missouri, CLS could obtain direct estimates for CRCS; however, CLS is not regularly conducted. **METHODS: MODELS** for all US counties. The methods were used by CDC (https://www.cdc.gov/ A Bayesian binomial regression framework to estimate Y_{ijkl} : diabetes/atlas/obesityrisk/County_Methods.html). $y_{ijkl} \sim \text{Binomial}(n_{ijkl}, p_{ijkl})$ $logit(p_{ijkl}) = \alpha_{r(i)} + \beta_j + \gamma_k + \theta_l$ $+X_i\psi + u_i + \epsilon_{ijkl}$ where $\epsilon_{ijkl} \sim \text{Normal}(0, \delta_0)$ is the error term and • $\alpha_{r(i)}$ is the intercept for region r(i) where county *i* belongs to; • $\beta = (\beta_1, \beta_2, \beta_3)$ are the age effects; • $\gamma = (\gamma_1, \gamma_2)$ are the gender effects; • $\boldsymbol{\theta} = (\theta_1, \theta_2)$ are the race effects; • ψ contains some county attribute effects like medium income, percentage of people below high school, *etc*; and X_i is i^{th} row of the corresponding design matrix conclusive estimates by county. Therefore, those areas were clustered into seven The fixed effects above all follow Normal(0,100) priors. BRFSS regions (Figure 1). Numbers in parenthesis are the sample sizes for re-The county effects $u = (u_1, ..., u_{115})$ were modeled with a proper CAR prior spondents aged 50 or older which are suitable for our CRCS study after removing $\boldsymbol{u} \sim \operatorname{Normal}(\boldsymbol{0}, \delta_1 \boldsymbol{B}^{-1})$ respondents with unknown county, unknown response, etc. $B = I - \rho C$ – Kansas City Metro (767) where *C* is the adjacency matrix to describe the neighborhood structure for counties – St. Louis Metro (1030) in MO, ρ measures the spatial correlation strength and $\delta_0 = \delta_1/\eta_1$ measures the BRFSS Regions – Central (493) spatial variance. Their prior distributions (or densities) are: Kansas City Metro St. Louis Metro – Southwest (485) $\rho \sim \text{Unif}(0, \lambda_I^{-1}),$ $[\eta_1] = \frac{1}{(1+\eta_1)^2}, \quad \eta_1 > 0,$ Central – Southeast (433) Southwest Southeast – Northwest (355) Northwest Northeast – Northeast (244) where λ_I is the largest eigenvalue of **B**. **METHODS: ESTIMATION Figure 1:** BRFSS regions in MO A Markov Chain Monte Carlo algorithm was used to obtain posterior samples for p_{ijkl} . For category *ijkl*, there are $N_{ijkl}^- = N_{ijkl} - n_{ijkl}$ people outside MO–BRFSS data and $Y^{-}_{ijkl} \sim \text{Binomial}(N^{-}_{ijkl}, p_{ijkl})$ Figure 2 shows the sample size for each county. people who have had CRCS. Thirty-seven counties in For each posterior samples of p_{ijkl} we obtained posterior predictive samples for MO had zero sample $Y_{ijkl} = Y_{ijkl}^- + y_{ijkl}$ and (shown in white); only $p_{i} = \frac{\sum_{j=1}^{3} \sum_{k=1}^{2} \sum_{l=1}^{2} Y_{ijkl}}{\sum_{j=1}^{3} \sum_{k=1}^{2} \sum_{l=1}^{2} N_{ijkl}}.$ 15 counties had a sample more than 50.

BACKGROUND In the US, colorectal cancer (CRC) is the 3rd most common cancer in both men and women. • Colorectal cancer screening (CRCS) is recommended for people over 50 years of • Behavioral Risk Factor Surveillance System (BRFSS) data in 2012 show that 66.5% • Missouri conducted a County-level Study (CLS) in 2011 aimed for accurate • Cadwell, et al. (2010)* used Bayesian methods to estimate diabetes prevalence **OBJECTIVE** Use small area estimation techniques to estimate county-level CRCS prevalence in Missouri for people age 50+ with 2012 BRFSS data and compare with results from 2011 CLS. **DATA OVERVIEW** • Missouri is comprised of 114 counties and the City of St. Louis. • In 2012 MO–BRFSS, the sample size from each county was too small to make





predictive samples of p_i .

Figure 2: Sample sizes for counties in MO.

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Therefore, the estimated CRCS prevalence for county *i* is the mean of the posterior

The county-level CRCS prevalence estimates from BRFSS generally agree with those from CLS, with an average 5.05%-point difference across all counties in MO. Counties with large sample sizes tend to have more similar estimates to CLS.



Figure 4: Maps for CRCS prevalence estimates from CLS vs. BRFSS. The white color is the state average CRCS prevalence in Missouri. Our estimates from BRFSS generally have the same spatial variation as those from CLS, but the northern counties are mostly overestimated.

CONCLUSIONS

- lence are still noticeably large.
- several or tens of people.
- can still provide reasonable estimates at county level.
- uncertainty compared to our model.
- may help improve the results.

The Missouri Cancer Registry and Research Center (MCR-ARC) is supported in part by a cooperative agreement between the Centers for Disease Control and Prevention (CDC) and the Missouri Department of Health and Senior Services (DHSS) (5NU58DP003924-05) and a Surveillance Contract between DHSS and the University of Missouri (MU). We thank DHSS for access to CLS data





• The differences between BRFSS and CLS for counties with high/low CRCS preva-

• In BRFSS, small or zero sample sizes for counties in Missouri potentially produce biased estimates. It is hard to estimate a whole county's prevalence based only on

• When BRFSS is the only source to estimate county-level prevalence, our model

• We also used models in Cadwell, et al. (2010) to obtain CRCS prevalence estimates. However, due to small sample sizes in Missouri compared to all samples in US, covariances among classes of people were hard to estimate, which added more

• We classified people into 12 groups in our analysis. However, when detailed population sizes are available, finer clarification with more demographic variables

• In our evaluation of the results, we treat CLS (2011) as the true prevalence for comparison. However, the uncertainty from CLS itself was not considered.

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