

Assessing MMR vaccination coverage gaps in US children with digital participatory surveillance

Received: 16 May 2025

Accepted: 24 November 2025

Published online: 15 January 2026

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Recent measles outbreaks in the USA have emerged despite the availability of the highly effective measles–mumps–rubella (MMR) vaccine. Current surveillance systems rely primarily on telephone surveys with provider verification or school-entry data, methods prone to incompleteness and systematic exclusion of vulnerable populations. Here, to address these limitations, we used a validated digital participatory surveillance platform to collect parental reports of ≥ 1 -dose MMR vaccination among children under 5 years of age. Applying Small Area Estimation methods to generate granular, county-level coverage estimates nationwide, we found substantial geographic variation, including areas with MMR coverage $< 60\%$. Analysis of spatial clustering revealed hotspots of undervaccination overlapping closely with recent measles outbreaks, particularly in Texas and New Mexico—where our model estimates substantially lower vaccine coverage than official data. These findings underscore the urgent need for surveillance systems to include more granular and timely data that accurately identify undervaccinated communities, enabling targeted, timely public health interventions.

The USA is experiencing a resurgence of measles¹, despite the widespread availability of the safe and effective measles–mumps–rubella (MMR) vaccine. Multiple states reported cases in 2025, notably concentrated in western Texas and New Mexico. Declining MMR coverage, fuelled by multifaceted vaccine hesitancy² and pandemic-related disruption³, has left national coverage below thresholds required to prevent sustained transmission^{4,5}. Differences in vaccination coverage by geographic, socioeconomic and demographic factors have further contributed to pockets of vulnerability^{6–8}, particularly in communities with lower MMR vaccine rates.


Effective public health interventions require timely, spatially granular surveillance data. However, existing US vaccination surveillance systems face notable limitations, including reporting delays,

coarse geographic resolution (often reported only at the state level⁴) and reliance on milestones assessment at 24 and 36 months or kindergarten entry. These estimates typically depend on healthcare provider-verified, school or health department records^{4,5,9,10}, which systematically underrepresent children who are homeschooled, uninsured or foreign-born, or face structural barriers to care—groups that historically have shown undervaccination^{11–13}. Consequently, the existing system provides an incomplete picture of the true vaccine coverage, omitting key subpopulations and underestimating local vulnerability¹⁴.

Most official vaccination estimates focus on kindergarten-entry requirements, subject to state-exemption policies⁴, leaving younger children—who are more vulnerable to severe measles complications—poorly presented. In Texas, for example, the statewide kindergarten

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MMR uptake for the 2024–2025 school year was reported at 93.2%, near the herd-immunity threshold¹⁵, yet West Texas is currently experiencing a measles outbreak. Such aggregated figures may obscure local immunity gaps, especially among children too young for school entry or those facing barriers to care. National case reports from the Centers for Disease Control and Prevention (CDC) echo this concern: in 2025, nearly 30% of US measles cases occurred among children under 5 years of age, who also had the highest hospitalization rate (21%), while over 90% of all cases were in unvaccinated individuals¹. The reliance on school-based reporting and state-specific data systems makes it difficult to construct a timely, unified national picture of measles immunity.

To address these gaps, we used a validated digital participatory surveillance platform OutbreaksNearMe (ONM)^{16,17} and Small Area Estimation (SAE) framework^{18,19} methods to generate publicly available county-level estimates of MMR vaccination coverage (≥ 1 dose) for children under 5 across the contiguous USA. We then leverage a geographic artificial intelligence (AI) foundation model to super-resolve these findings to a finer spatial scale.

Our approach complements existing surveillance systems, including recently published county-level reports of two-dose MMR coverage²⁰, by better capturing populations who might otherwise be absent from official reporting, including homeschooled and uninsured children. The analysis identifies clusters of undervaccination that aligned closely with recent measles outbreaks, offering actionable insights for targeted immunization strategies and outbreak preparedness.

Results

In a nationally representative sample of 22,062 US adults with children under 5 collected via the ONM participatory surveillance platform (fielded between July 2023 and April 2024), the survey-weighted estimate of MMR vaccine uptake (≥ 1 dose) was 64.0% (95% confidence interval (CI) 63.2–64.9%, representing approximately 71.1% (70.2–72.1%) of the MMR eligible population (children >6 months). As reported previously, uptake differed substantially by parental characteristics, including age, race/ethnicity and coronavirus disease 2019 (COVID-19) vaccination status¹⁶.

County-level vaccine uptake and geographic clustering

We applied a multilevel SAE framework to predict county-level MMR (≥ 1 dose) coverage across the contiguous USA. The estimates revealed substantial geographic variation in MMR uptake, with distinct patterns of spatial clustering, as shown in Fig. 1. For interpretability, counties were grouped into five risk categories based on predicted coverage: very high risk (<60%), high risk (60–69%), medium risk (70–79%), low risk (80–84%) and lowest risk ($\geq 85\%$). Higher coverage was observed across the Northeast, Midwest and Northwest and along the Pacific coast. Spatial autocorrelation was strong (global Moran's $I = 0.53$, $P < 0.0001$), indicating statistically significant geographical clustering of counties with similar vaccination rates. Local Moran's I analysis identified statistically significant clusters of low coverage—hot spots—in West Texas, in southern New Mexico, in parts of Mississippi and across the rural Southeast. By contrast, cold spots—clusters of high coverage—were concentrated in the Northeast and Upper Midwest. Notably, several high-risk hot spot counties were located in states experiencing active measles outbreaks.

At the state level, county-aggregated estimate ranges from 61.6% (95% CI 58.9–64.5%) in New Mexico to 79.1% (95% CI 76.5–81.6%) in Massachusetts, with a median of 71.3% (95% CI 69.4–73.4%). County-level estimates showed even greater variation, with a median MMR uptake of 71.4%, ranging from 35.8% (95% CI 35.8–42.0%) to 86.8% (95% CI 85.1–88.4%). Counties with the lowest modelled coverage were primarily in Georgia, Texas and Mississippi, while the highest coverage appeared in parts of New York, Indiana and Oregon (Fig. 2).

Model validation and local discrepancies

Multiple validation analyses supported our model-based coverage estimates. Shown in Table 1, the distribution of model-based estimates

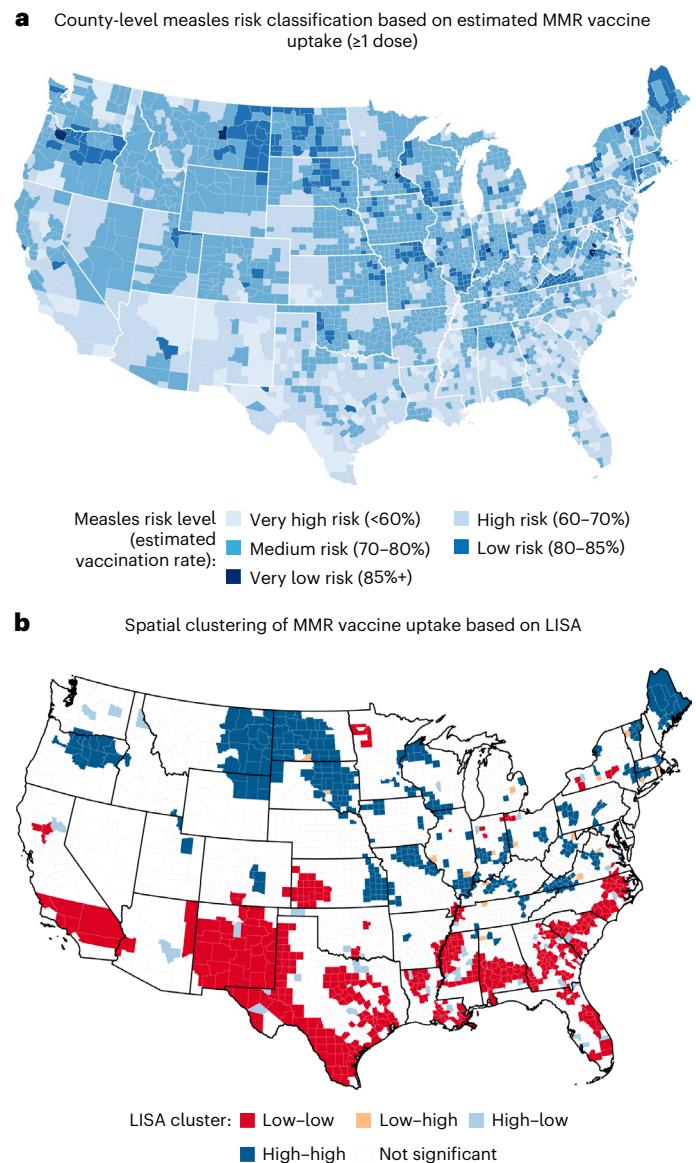


Fig. 1 | County-level estimates of MMR vaccine uptake and spatial clustering among US children under age 5. County-level estimates of ≥ 1 -dose MMR vaccine coverage among US children under age 5 were generated using a multilevel regression with poststratification (MRP) framework, based on digital surveillance data (ONM) collected between July 2023 and April 2024 ($n = 3,109$ counties; one modelled estimate per county). **a**, Modelled vaccine uptake categorized into five risk levels based on estimated vaccination rate: very high risk (<60%), high risk (60–69%), medium risk (70–79%), low risk (80–84%) and lowest risk ($\geq 85\%$), relative to the herd-immunity threshold for measles. Because these estimates include children under 6 months who are not yet vaccine-eligible, the upper threshold appears lower than the 92–95% benchmark typically cited for herd immunity. **b**, Results from a spatial clustering analysis using local indicators of spatial association (LISA), which identifies counties with vaccination rates statistically significantly higher or lower than their geographic neighbours (two-sided permutation test, 499 permutations, $P < 0.05$ after Benjamini–Hochberg correction). LISA cluster labels denote: high–high (counties with high uptake surrounded by high-uptake neighbours), low–low (low uptake surrounded by low-uptake neighbours), high–low (high uptake surrounded by low-uptake neighbours) and low–high (low uptake surrounded by high-uptake neighbours). Statistically significant clusters are highlighted; counties shown in white did not exhibit statistically significant spatial clustering. Figure adapted from TIGER/Line Shapefiles, US Census Bureau (2022).

closely aligned with that of direct survey data at both the county and state levels. SAE-predicted MMR uptake showed strong agreement with direct survey estimates at both the county level ($n = 932$; Pearson $r = 0.84$, Spearman $\rho = 0.86$, $R^2 = 0.71$; all $P < 2.2 \times 10^{-16}$) and

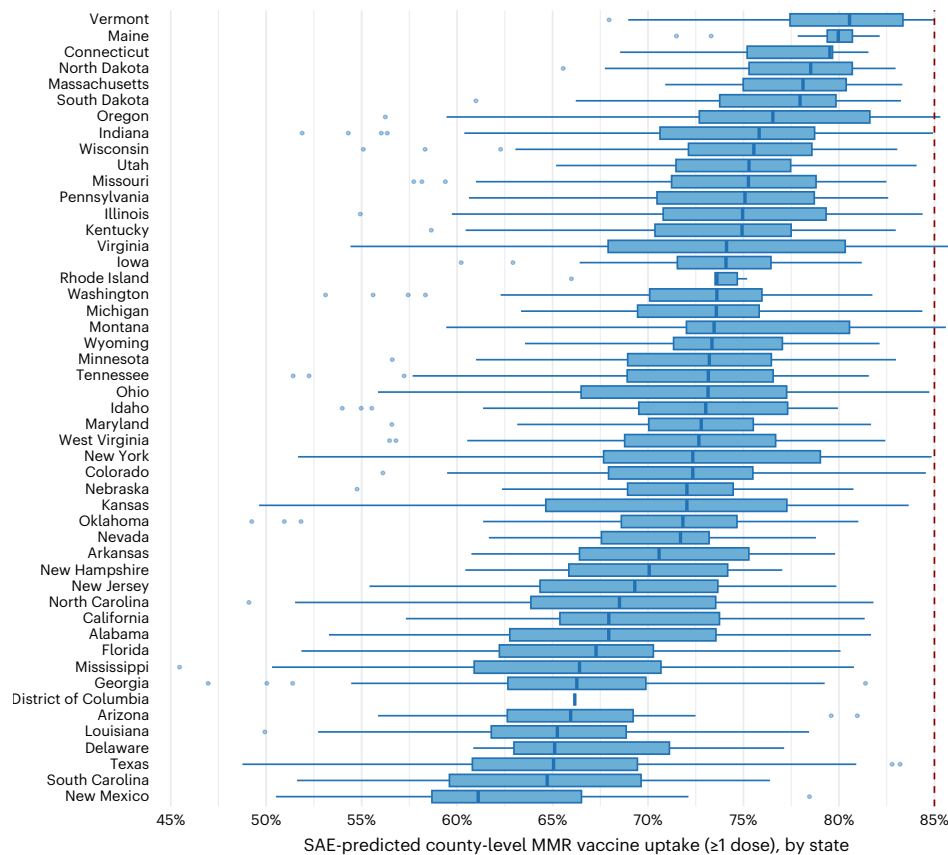


Fig. 2 | County-level estimates of MMR vaccine uptake (≥1 dose) among US children under age 5 by state. The box plot presents SAE-predicted county-level MMR vaccine uptake (≥1 dose) among children under age 5, grouped by the 48 contiguous states and the District of Columbia ($n = 3,109$ counties). Each box represents the interquartile range of county-level estimates, with the central line indicating the median. The horizontal lines extend to the minimum and maximum values within a typical range; counties with values far outside

this range are plotted individually. The dashed vertical line at 85% denotes the threshold for the lowest measles risk category used in Fig. 1. Because these estimates include all children under 5, including those under 6 months who are not yet vaccine-eligible, the upper threshold appears lower than the 92–95% benchmark typically cited for herd immunity. No formal hypothesis testing was performed; all values represent model-derived county-level estimates from a single fitted model.

state level ($n = 49$; Pearson $r = 0.88$, Spearman $\rho = 0.91$, $R^2 = 0.77$; $P = 7.0 \times 10^{-16}$), with substantial R^2 values in corresponding bivariate regressions. The estimated coverage and spatial clustering analysis are not sensitive to various alternative model specifications or spatial aggregation procedures.

Lastly, we compared state-level model estimates with the CDC's provider-verified 36-month one-dose MMR coverage data to assess overall coherence. Modelled and CDC estimates clustered closely along the 45° line, with modest differences (5–10 percentage points) and no signs of systematic differences (Fig. 3). However, only two states—Texas and New Mexico—were positioned well above the diagonal line, with model-based estimates substantially lower than the CDC-reported figures (New Mexico: 61.6% versus 90.3%; Texas: 62.9% versus 93.7%). These two states were also the only states experiencing substantial initial measles outbreaks during the study period. To better understand this discrepancy, we examined county-level patterns in Texas (Supplementary Information section 2.3.3) and found that measles cases were more than twice as likely to occur in 'low–low' counties (areas with both low estimated vaccination coverage and low-coverage neighbours), suggesting that spatial vulnerability and suboptimal vaccination rates among young children may help to explain the elevated risk of outbreaks.

AI super-resolution

Although public health surveillance typically aggregates data at the county level, measles outbreaks in the USA often emerge from tight-knit local communities. To extend our findings to a finer geographic scale, we leveraged Google's Population Dynamic Foundational Model

(PDFM)²¹ to produce MMR vaccination estimates at the ZIP Code Tabulation Area (ZCTA) level (see details in Supplementary Information section 2.4). PDFM is a multimodal AI system trained on privacy-preserving data sources (for example, search trends, map interactions, mobility and environmental signals) that captures neighbourhood-level context beyond standard census variables. Our subcounty estimates revealed similar regional patterns to the county-level model but identified more compact clusters of vaccine behaviour (Extended Data Figs. 1 and 2). The distribution of Local Indicators of Spatial Association (LISA) categories shifted slightly, preserving most low–low clusters in Texas, New Mexico and the southern and southeastern USA, as well as high–high clusters in parts of the Midwest, while diluting other mixed patterns.

Discussion

Our study provides the nationwide county-level MMR vaccine coverage among US children under age 5, leveraging a digital surveillance tool and advanced spatial modelling methods. These granular estimates reveal substantial gaps in coverage, highlighting the critical role of local variation in vaccine-induced immunity in shaping measles vulnerability. Importantly, by drawing on a digital participatory surveillance platform rather than administrative records, our approach captures children who are often absent from official reporting systems, including those who are homeschooled, uninsured or otherwise outside traditional health-care and school-based surveillance. Despite long-standing national recommendations for routine childhood vaccination, our findings show that MMR uptake remains low in many counties—particularly those

Table 1 | Summary statistics and agreement between SAE-predicted and survey-based MMR vaccine uptake estimates

Descriptive statistics	Min	First quartile	Median	Mean	Third quartile	Maximum
County-level ^a estimates (n=932)						
Survey estimate (unweighted)	0	0.6	0.7	0.69	0.8	1
Survey estimate (weighted)	0	0.54	0.67	0.66	0.8	1
SAE estimate	0.44	0.66	0.71	0.71	0.76	0.87
State-level ^b estimates (n=49)						
Survey estimate (unweighted)	0.59	0.67	0.7	0.7	0.74	0.85
Survey estimate (weighted)	0.54	0.62	0.66	0.66	0.7	0.82
SAE estimate	0.62	0.69	0.71	0.71	0.74	0.79

^aCounty refers to county-like geographic units used in estimation. ^bState (n=49) includes the 48 contiguous US states and the District of Columbia. Survey estimates are based on self-reported MMR vaccination status among parents of children under age 5, with and without survey weights applied. SAE estimates are derived from a SAE model using multilevel regression with poststratification (MRP). All values represent descriptive statistics only and were not derived from formal hypothesis testing.

with disadvantaged socioeconomic profiles—highlighting overlooked vulnerabilities within this at-risk population.

Notably, our model identified substantial clusters of undervaccination in locations across the US South and Southwest, including areas currently experiencing active measles outbreaks. These areas show sizable discrepancies between our model-based coverage estimates and official state-level data⁹, which probably reflect both age-composition differences (children under 5 years versus 36 months) and the exclusion of select populations in traditional surveillance methods (parent-reported coverage versus provider-verified records)^{4,22}. For example, Texas and New Mexico fall well below the national average in our county-level estimates, despite high reported state-level coverage at kindergarten entry. Both states subsequently reported early measles activity in early 2025, consistent with our model’s identification of lower effective coverage and suggesting that official figures (at kindergarten entry) might have not fully represented the community-level MMR vaccine uptake at that time. Recent analyses of electronic health records from Truveta²³ similarly document pandemic-related declines in MMR vaccination, although their data, derived solely from children with consistent healthcare access, suggested somewhat higher coverage among children engaged in routine care. These findings suggest that some children may be delaying rather than forgoing vaccination. Together, these differences underscore the critical need for surveillance systems capable of capturing delayed vaccinations and more comprehensively monitoring younger, harder-to-reach children.

Our spatial clustering analyses further identified considerable concentrations of low MMR vaccine uptake, counties with persistently low coverage surrounded by similarly undervaccinated areas, in regions such as West Texas, southern New Mexico, Mississippi and the rural Southeast. These clusters signal areas vulnerable to future outbreaks, even in the absence of current transmission. Given this elevated risk, public health officials may wish to revisit vaccination guidance for children aged 6–12 months⁹ residing in these regions²⁴. Although counties vary substantially in size and population density, they remain a central unit for public health planning and monitoring. Here, we were able to leverage an AI foundation model to produce MMR vaccination estimates at the ZCTA level. The finer-scale modelling revealed more compact local clusters and subtle shifts in LISA category distributions,

reflecting both improved precision from AI-enhanced local contextual factors and the inherent sensitivity of LISA statistics to spatial aggregation. However, the overall consistency of results across scales (county versus ZCTA; Supplementary Information section 2.4) suggests that county-level clustering captures broader underlying patterns of vaccine behaviour, while ZCTA-level maps provide complementary insight into smaller, community-based clusters that may be more relevant for localized behavioural dynamics. These maps do not define transmission boundaries but serve as practical tools to inform resource allocation and identify vulnerable regions. More systematic and inclusive data collection efforts at subcounty levels would greatly strengthen the ability to monitor undervaccination and design targeted interventions.

Beyond identifying spatial patterns of vaccination, an important contribution of this study lies in combining digital participatory surveillance with advanced statistical methods to improve the measurement of vaccination coverage, particularly among vulnerable populations. This methodology offers a timely and scalable complement to conventional immunization monitoring systems, such as vaccine registries and national surveys, by enhancing the detection of localized immunity gaps and increasing geographic granularity and inclusiveness to reach populations often missed by the existing systems. While digital surveillance data cannot replace traditional monitoring approaches, they can significantly improve timeliness, geographic resolution and representativeness via leveraging digital technology to enhance convenience, anonymity and outreach, features that are increasingly vital as social and economic interdependence accelerates and infectious-disease risks transcend local and national boundaries^{25,26}.

Several limitations should be acknowledged. First, parent-reported MMR uptake may be subject to recall bias; however, our estimates align closely with direct survey results, correlate strongly with official benchmarks and are consistent with findings from independent studies. Second, survey response volume was low in some counties; however, our modelling strategy addresses spatial sparsity through aggregation and hierarchical smoothing, with resulting uncertainty reflected in the CIs and robustness of results demonstrated in multiple analyses. Third, our estimates are based on ≥1-dose coverage and do not reflect full completion of the two-dose MMR schedule as usually reported in other data sources. Moreover, because our study population includes children under age 5, many had not yet reached the age of routine MMR eligibility (12 months generally; but 6 months for exceptions such as travel). As such, our coverage estimates are expectedly lower than administrative data, in part because we are capturing the

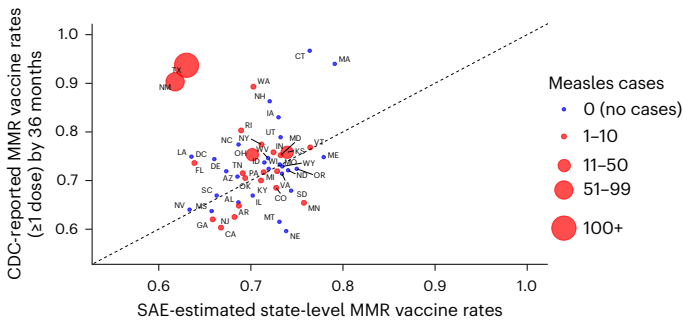


Fig. 3 | Discrepancies between model-predicted and official MMR coverage at the state level, with measles outbreak case counts overlay. Points represent the 48 contiguous states and the District of Columbia (n=49 states), with values corresponding to state-level SAE-predicted MMR vaccine uptake among children under age 5 (x axis) and CDC-reported 36-month MMR coverage for ≥1 dose (y axis). Point size and colour indicate the number of confirmed measles cases in each state as of 11 April 2025. The dashed diagonal line indicates perfect agreement between model-based estimates and reported coverage. States above the line have higher reported coverage than predicted by the model, while those below the line have lower reported coverage. Statistical comparisons were descriptive; no formal hypothesis testing was applied.

age group least likely to be fully vaccinated. These differences demonstrate how digital participatory surveillance and traditional monitoring approaches capture complementary populations and timeframes, and together could form a more robust, integrated monitoring system adapted to evolving population and disease dynamics. They also reinforce the urgency of adapting surveillance and outreach efforts to better include high-risk, undermonitored populations.

In summary, our work provides an innovative resource for improving immunization strategies and mitigating measles outbreaks through geographically targeted interventions. We developed an interactive website (<https://healthmap.org/measles/>) that enables users to explore county-level MMR vaccination estimates across the USA. Model-based surveillance can complement traditional systems by identifying at-risk communities earlier, guiding geographically targeted interventions and strengthening local preparedness, ultimately advancing national vaccine equity and disease prevention goals.

Methods

In a retrospective cohort study, we leveraged ONM, a previously validated digital health surveillance platform that collects anonymous, self-reported health information from a national sample of US adults. In brief, ONM utilized non-probability river sampling techniques to randomly deliver a survey to individuals in SurveyMonkey's diverse, multi-million-person user pool. From July 2023 to April 2024, participants provided demographics, residential ZIP code and information on children under 5, including parental report of MMR vaccine status (≥ 1 dose versus none). For households with multiple eligible children, one child was randomly selected at the time of survey completion to avoid intrahousehold clustering. We constructed a nationally representative analytic sample of 22,062 parents with children under 5 using a weighting procedure calibrated to US Census benchmarks (including age, gender, race/ethnicity, education and geography). This study is a quantitative analysis only. Additional methodological details on the ONM platform and survey design have been published previously¹⁶ and validated for various public health applications^{17,27}.

SAE with poststratification

To generate county-level MMR vaccine uptake (≥ 1 dose), we implemented a spatial multilevel logistic regression with poststratification (MRP) framework^{18,19}. The regression model included both individual- and county-level variables, with random intercepts for county-like areas and states to account for unobserved contextual variation in vaccination decisions. To address sparsity in counties with limited direct survey data, we implemented an iterative spatial aggregation algorithm that merged counties into larger county-like areas until each area contained a minimum of five valid observations. The distribution of direct survey responses and sensitivity analyses of the aggregation procedure are shown in the Supplementary Information section 2.2.

The outcome was parent-reported receipt of at least one dose of the MMR vaccine among children under 5. Individual-level controls included parent age group (18–29, 30–39, 40–49 and 50–59), gender (male or female) and race/ethnicity (Asian, Black, Hispanic, white or other). County-level covariates were selected on the basis of prior literature²⁸ and LASSO regression, and included: median household income, percentage of white residents, percentage of single-parent households, percentage enrolled in Medicaid from the American Community Survey (5-year estimates 2019–2023)²⁹, percentage of adults completing the primary COVID-19 vaccine series³⁰, and Democratic vote share in the 2020 US presidential election³¹.

Model predictions were then poststratified using US Census microdata³² to produce population-weighted county-level estimates of ≥ 1 -dose MMR coverage among children under 5. The fitted model was applied to all combinations of demographic strata (age group \times race/ethnicity \times gender) within each county. Poststratification ensured that estimates were aligned with the demographic and geographic

distribution of the US population, thereby generating population-representative county-level estimates, including for counties without direct survey responses. To calculate the 95% CIs, we used Monte Carlo simulation to generate 1,000 replicates of MMR uptake estimates for each county, state and demographic subgroup. State-level estimates were computed as population-weighted averages of county-level estimates based on US Census counts.

Benchmarking against traditional surveillance and outbreak data

To further evaluate alignment with established data sources, we compared our state-level estimates with CDC-published one-dose MMR coverage at 36 months³³. In parallel, we incorporated publicly reported measles case counts as of April 2025, corresponding to the initial phase of the current outbreak, to assess whether states with lower predicted coverage and larger gaps relative to official statistics overlapped with regions experiencing elevated outbreak activity. This benchmarking step enabled us to assess concordance between small area estimates and existing surveillance systems while examining the potential added value of our approach for identifying areas of public health concern.

Spatial clustering analysis and visualization

Given the central role of social and geographic clustering in measles transmission, we conducted spatial clustering analysis using LISA to detect statistically significant patterns of MMR uptake across counties. Although counties vary substantially in size and population density, this scale remains the most relevant administrative unit for many public health agencies. We used LISA to examine whether modelled undervaccination was geographically isolated or clustered across contiguous counties, providing descriptive insight into potential pockets of outbreak risk. This method identifies geographic clusters—such as counties with high or low uptake surrounded by neighbours with similar values—potentially indicative of localized outbreak risk. To support public health interpretation, we categorized counties into five risk groups based on predicted MMR vaccine coverage thresholds, informed by the herd immunity threshold for measles (typically estimated at 92–95%). Counties with estimated uptake below 60% were classified as very high risk, followed by high risk (60–69%), medium risk (70–79%), low risk (80–84%) and lowest risk ($\geq 85\%$). These thresholds reflect increasing proximity to the herd immunity benchmark and help prioritize areas for intervention. While the highest group cut-off ($\geq 85\%$) may appear low relative to the 92–95% herd immunity benchmark, this reflects the inclusion of infants under 6 months who are not yet eligible for MMR vaccination in the denominator of our estimates. Map boundaries are based on 2022 US Census Bureau TIGER/Line shapefiles, accessed via the R package *tigris* (version 2.1).

AI-based super-resolution with PDFM embeddings

Building on the county-level analysis, we extended our framework to the ZCTA scale to examine patterns of undervaccination across local communities, where measles transmission is shaped by social interactions and health behaviour. This extension was enabled by embeddings from PDFM²¹, an AI system pretrained on large-scale, multimodal data sources—including Google search and map activity, trends of mobility and busyness, geospatial data and environmental conditions—that capture fine-grained neighbourhood context while preserving privacy (Supplementary Information section 2.4). By incorporating these high-dimensional embeddings into our SAE framework, we applied an AI-based super-resolution approach to assess the consistency between county- and ZCTA-level estimates and to demonstrate how subcounty resolution can support more targeted public health strategies.

To further assess the robustness of our results, model performance was evaluated by comparing model-based estimates with direct survey estimates at both the county and state levels. In addition, we conducted extensive analyses evaluating model fit, the sensitivity

of results to alternative specifications and the stability of estimates across subgroups. We also validated predictions using independent external data sources and additional statistical tests. These steps were designed to show that our conclusions are not dependent on any single alternative analytical choice. Full methodological details and supplementary results are provided in the Supplementary Information. All analyses were conducted in R version 4.4.3 (RStudio). The study adheres to the STROBE reporting guideline, was approved by the institutional review board (IRB-P00023700) and received a waiver of informed consent. Use of the data in this study complied fully with the terms of use of the SurveyMonkey platform and associated data-use agreements.

Reporting summary

Further information on research design is available in the Nature Portfolio Reporting Summary linked to this article.

Data availability

For privacy, individual-level survey data cannot be made publicly available. Researchers affiliated with academic or public health institutions may request access to these data for non-commercial research purposes. Requests should be submitted to the corresponding author. Requests will typically be processed within 4–6 weeks. All derived data products from this study, including county- and ZCTA-level predicted MMR vaccination coverage, will be deposited in a public data repository and are also accessible through the interactive dashboard at <https://healthmap.org/measles/>

Code availability

All analyses were conducted in R (version 4.3.2). The code used to generate the estimates and figures is publicly available at Github (https://github.com/eric-gengzhou/MMR_vaccine_estimates)³⁴.

References

- Measles cases and outbreaks. *Centers for Disease Control and Prevention* <https://www.cdc.gov/measles/data-research/index.html> (2025).
- Grills, L. A. & Wagner, A. L. The impact of the COVID-19 pandemic on parental vaccine hesitancy: a cross-sectional survey. *Vaccine* **41**, 6127–6133 (2023).
- Desilva, M. B. et al. COVID-19 and completion of select routine childhood vaccinations. *Pediatrics* <https://doi.org/10.1542/peds.2024-068244> (2025).
- Seither, R. et al. Coverage with selected vaccines and exemption rates among children in kindergarten–United States, 2023–24 school year. *Morb. Mort. Wkly Rep.* **73**, 925–932 (2024).
- Seither, R. et al. Coverage with selected vaccines and exemption from school vaccine requirements among children in kindergarten–United States, 2022–23 school year. *Morb. Mort. Wkly Rep.* **72**, 1217–1224 (2023).
- Truelove, S. A. et al. Characterizing the impact of spatial clustering of susceptibility for measles elimination. *Vaccine* **37**, 732–741 (2019).
- Alvarez-Zuzek, L. G., Zipfel, C. M. & Bansal, S. Spatial clustering in vaccination hesitancy: the role of social influence and social selection. *PLoS Comput. Biol.* **18**, e1010437 (2022).
- Gromis, A. & Liu, K.-Y. Spatial clustering of vaccine exemptions on the risk of a measles outbreak. *Pediatrics* **149**, e2021050971 (2022).
- Measles vaccination. *Centers for Disease Control and Prevention* <https://www.cdc.gov/measles/vaccines/index.html> (2025).
- Hill, H. A., Yankey, D., Elam-Evans, L. D., Chen, M. & Singleton, J. A. Vaccination coverage by age 24 months among children born in 2019 and 2020–National Immunization Survey–Child, United States, 2020–2022. *Morb. Mort. Wkly Rep.* **72**, 1190–1196 (2023).
- Cordner, A. The health care access and utilization of homeschooled children in the United States. *Soc. Sci. Med.* **75**, 269–273 (2012).
- Ghildayal, N. et al. Public health surveillance in electronic health records: lessons from PCORnet. *Prevent. Chron. Dis.* **21**, E51 (2024).
- Mohanty, S. et al. Homeschooling parents in California: attitudes, beliefs and behaviors associated with child’s vaccination status. *Vaccine* **38**, 1899–1905 (2020).
- Masters, N. B. et al. Fine-scale spatial clustering of measles nonvaccination that increases outbreak potential is obscured by aggregated reporting data. *Proc. Natl Acad. Sci. USA* **117**, 28506–28514 (2020).
- Vaccination Coverage Levels in Texas Schools — Kindergarten* (Texas Department of State Health Services, 2025); <https://www.dshs.texas.gov/immunizations/data/school/coverage> (2025).
- Zhou, E. G. et al. Parental factors associated with measles–mumps–rubella vaccination in US children younger than 5 years. *Am. J. Public Health* **115**, 369–373 (2025).
- Rader, B. et al. Use of at-home COVID-19 tests—United States, August 23, 2021–March 12, 2022. *Morb. Mort. Wkly Rep.* **71**, 489 (2022).
- Mills, C. W., Johnson, G., Huang, T. T., Balk, D. & Wyka, K. Use of small-area estimates to describe county-level geographic variation in prevalence of extreme obesity among US adults. *JAMA Netw. Open* **3**, e204289 (2020).
- Zhang, X. et al. Multilevel regression and poststratification for small-area estimation of population health outcomes: a case study of chronic obstructive pulmonary disease prevalence using the behavioral risk factor surveillance system. *Am. J. Epidemiol.* **179**, 1025–1033 (2014).
- Dong, E., Saiyed, S., Nearchou, A., Okura, Y. & Gardner, L. M. Trends in county-level MMR vaccination coverage in children in the United States. *JAMA* <https://doi.org/10.1001/jama.2025.8952> (2025).
- Agarwal, M. et al. General geospatial inference with a population dynamics foundation model. Preprint at <https://arxiv.org/abs/2411.07207> (2024).
- Seeskin, Z. H. et al. Estimating county-level vaccination coverage using Small Area Estimation with the National Immunization Survey–Child. *Vaccine* **42**, 418–425 (2024).
- Latest US First-Time Measles Vaccination Trends for Children under 24 Months*, (Truveta, 2025); <https://www.truveta.com/blog/research/research-insights/first-time-measles-vaccination-trends-us-children-under-24-months/>
- Rader, B., Walensky, R. P., Rogers, W. S. & Brownstein, J. S. Revising US MMR vaccine recommendations amid changing domestic risks. *JAMA* **333**, 1201–1202 (2025).
- Khabbaz, R. F., Moseley, R. R., Steiner, R. J., Levitt, A. M. & Bell, B. P. Challenges of infectious diseases in the USA. *Lancet* **384**, 53–63 (2014).
- Baker, R. E. et al. Infectious disease in an era of global change. *Nat. Rev. Microbiol.* **20**, 193–205 (2022).
- Rader, B. et al. Mask-wearing and control of SARS-CoV-2 transmission in the USA: a cross-sectional study. *Lancet Digit. Health* **3**, e148–e157 (2021).
- Kempe, A. et al. Parental hesitancy about routine childhood and influenza vaccinations: a national survey. *Pediatrics* **146**, e20193852 (2020).
- American Community Survey 5-Year Estimates, 2019–2023* (US Census Bureau, 2023); <https://www.census.gov/programs-surveys/acs/data.html>
- COVID-19 Vaccinations in the United States (County)* (Centers for Disease Control and Prevention, 2023); https://data.cdc.gov/Vaccinations/COVID-19-Vaccinations-in-the-United-States-County/8xkx-amqh/about_data
- Baltz, S. et al. American election results at the precinct level. *Sci. Data* **9**, 651 (2022).

32. Schroeder, J. V. R. et al. IPUMS National Historical Geographic Information System: Version 20.0 (IPUMS, 2025).
33. *Vaccination Coverage Among Children Age 35 Months* (Kaiser Family Foundation, 2023); <https://www.kff.org/other-health/state-indicator/percent-of-children-aged-0-35-months-who-are-immunized/>
34. Zhou, E. G. MMR vaccine estimates. *Github* https://github.com/eric-gengzhou/MMR_vaccine_estimates (2025).

Acknowledgements

We thank A. Gertz, B. R. Anderson and C. Rimmel for their support and assistance.

Author contributions

E.G.Z. performed the statistical analyses. E.G.Z., J.S.B. and B.R. acquired, analysed and interpreted the data. E.G.Z. and B.R. drafted the article; they had full access to the study data and take responsibility for the data integrity and the accuracy of the data analysis. J.S.B. and B.R. supervised the study. All authors conceptualized and designed the study and critically revised the article for important intellectual content.

Competing interests

B.R. reports research funding from the Thrasher Research Fund (Boston Children's Hospital grant FP00000397). E.G.Z. receives salary support from the National Institutes of Health (R01 HL150044) outside the submitted work. All authors declare no other competing interests related to this study.

Additional information

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s44360-025-00031-8>.

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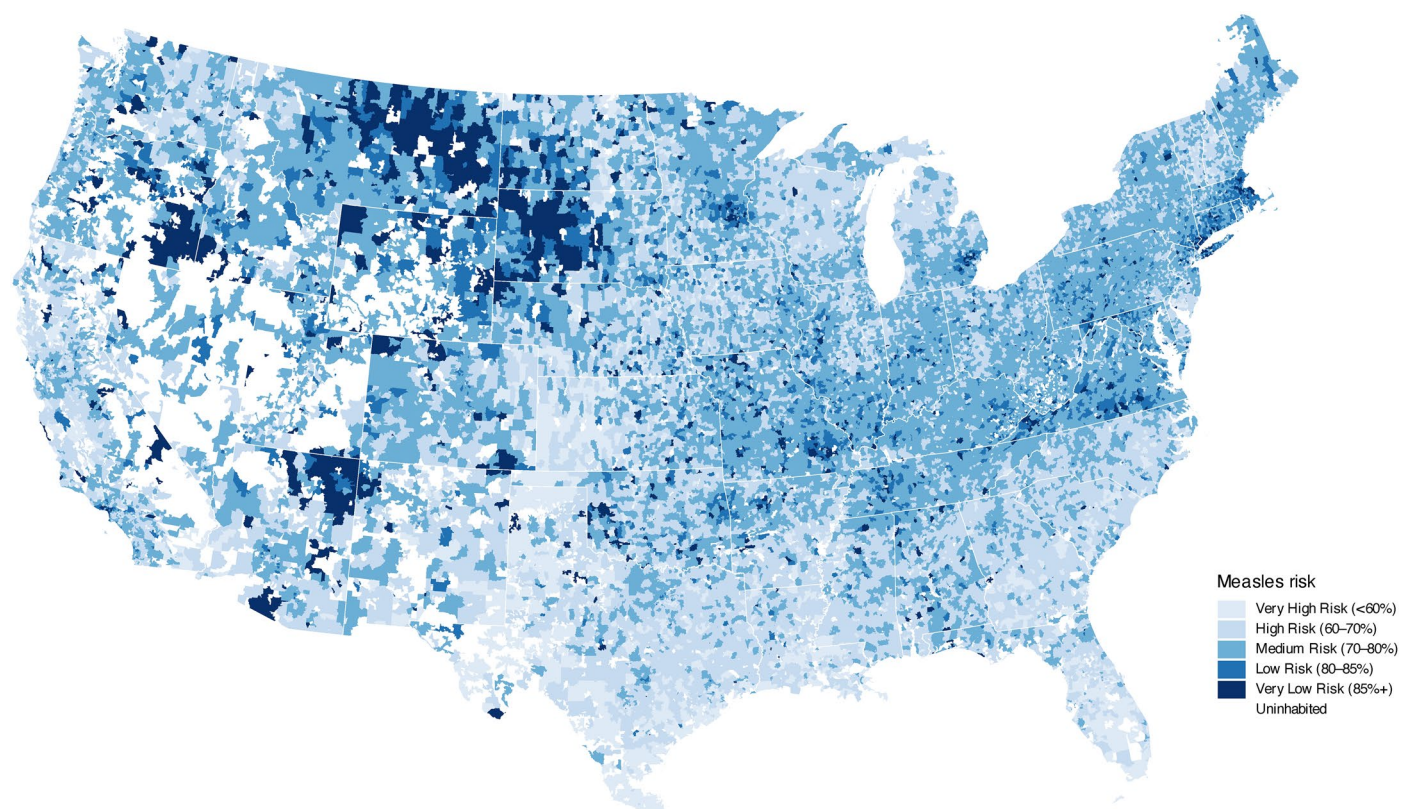
Peer review information *Nature Health* thanks Jon Zelner and the other, anonymous, reviewer(s) for their contribution to the peer review of this work. Primary Handling Editor: Lorenzo Righetto, in collaboration with the *Nature Health* team.

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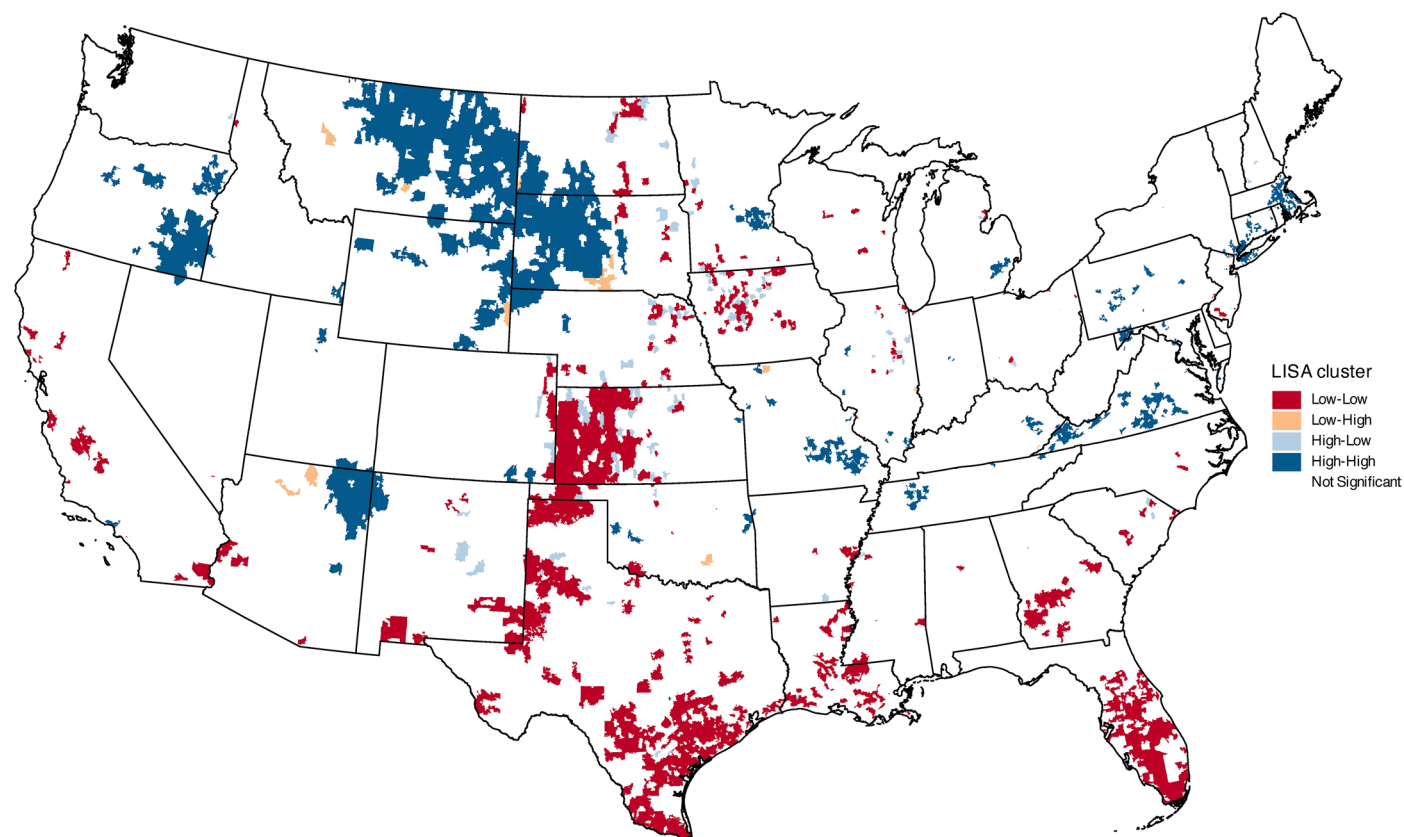
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Extended Data Fig. 1 | ZIP Code Tabulation Area (ZCTA)–Level Estimates of MMR Vaccine Uptake Among U.S. Children Under Age 5. Note: ZCTA-level estimates of ≥ 1 -dose measles–mumps–rubella (MMR) vaccine coverage among U.S. children under age 5 were generated using the same multilevel regression with post-stratification (MRP) framework as in the county-level analysis, extended with contextual embeddings from Google’s Population Dynamics Foundation Model (PDFM). Embedding components derived from principal component analysis were included as auxiliary covariates. Modeled vaccine

uptake is categorized into five risk levels—Very High Risk (<60%), High Risk (60–69%), Medium Risk (70–79%), Low Risk (80–84%), and Lowest Risk ($\geq 85\%$)—relative to the herd-immunity threshold for measles. Because estimates include children under 6 months who are not yet vaccine-eligible, the upper threshold appears lower than the 92–95% benchmark typically cited for herd immunity. ($n = 29,971$ ZCTAs; one modeled estimate per ZCTA.). Figure adapted from TIGER/Line Shapefiles, US Census Bureau (2022).



Extended Data Fig. 2 | Spatial Clustering of Predicted MMR Vaccine Uptake at the ZCTA Level. Note: Results from a Local Indicators of Spatial Association (LISA) analysis applied to modeled MMR vaccination rates at the ZCTA level ($n = 29,971$). LISA identifies ZCTAs with statistically significantly higher or lower uptake than their geographic neighbors using queen contiguity weights (two-sided permutation test, 499 permutations, $p < 0.05$ after Benjamini–Hochberg correction). Cluster categories include High–High (ZCTAs with high uptake

surrounded by high-uptake neighbors), Low–Low (low uptake surrounded by low-uptake neighbors), High–Low, and Low–High. Statistically significant clusters are highlighted; areas shown in white did not exhibit statistically significant spatial clustering. Patterns largely mirror those at the county level, though with finer spatial resolution and more localized cluster boundaries. Figure adapted from TIGER/Line Shapefiles, US Census Bureau (2022).

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Deidentified data used in this analysis were derived from a national digital health survey conducted in partnership with SurveyMonkey. Due to data use agreements,

the original individual-level survey data cannot be shared publicly. However, the analytical code and derived data outputs used in this study are available from the corresponding author upon reasonable request, subject to approval by the data providers and institutional review. Predicted county-level estimates are publicly accessible through an interactive dashboard [<https://healthmap.org/measles/>] and can be downloaded as a data file.

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The study uses the term “gender” throughout, consistent with the survey instrument, which asked respondents to self-identify their gender. The design and analysis acknowledge the distinction between gender and sex, and the variable reflects how participants chose to describe their own identity.

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The survey asked: “Which race or ethnicity best describes you? (Select all that apply)”, with the following options: White, Hispanic or Latino/a, Black or African-American, Asian or Asian-American, Middle Eastern or North African, Native Hawaiian or Pacific Islander, and Native American or American Indian. Respondents could select multiple categories. For analysis, we grouped race/ethnicity into five categories: White, Asian, Black, Hispanic, and Other. Individuals selecting multiple categories were assigned to “Other” unless one of the selected categories was Hispanic, which was prioritized to reflect common practice in demographic coding.

Although the primary goal of the study is prediction rather than causal inference, we incorporated a broad set of individual- and county-level covariates selected through a transparent and both data and theory-driven process. This included both demographic and contextual variables (e.g., income, insurance, political context, urbanicity) to account for observed sources of variation and reduce the influence of omitted variable bias in model estimation, which were included based on recent literature and then selected by the LASSO procedure.

Population characteristics

The analytic sample consisted of 22,062 U.S. parents or guardians of children under age 5, recruited through a national online survey. About 58% identified as female and 42% as male. The largest age group was 30–39 (45%), followed by 18–29 (35%), 40–49 (15%), and 50+ (5%). Racial and ethnic composition included 50% white, 26% Black, 14% Hispanic, 6% Asian, and 3% other or multiracial. Educational attainment ranged from 52% with a high school education or less to 21% with a college or postgraduate degree. 43% were covered by Medicaid or Medicare, 47% had private or employer-based insurance, and 10% were uninsured. A majority (56%) reported household incomes below \$50,000, and 72% were employed. Respondents identified as Democrats (27%), Republicans (29%), or Independents/Other (44%), with geographic representation from all U.S. Census regions.

Recruitment

Survey participants were randomly invited to participate anonymously on the multi-million user platform SurveyMonkey. Invitations were extended after users completed unrelated activities on the platform. Participation was voluntary, and no incentives were provided. Respondents were not required to have a SurveyMonkey account, and no personally identifiable information was collected.

Ethics oversight

This study was approved by the Boston Children’s Hospital institutional review board (IRB-P00023700) and received a waiver of informed consent.

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Study description

This is a cross-sectional, retrospective cohort study that combines individual-level survey data with county-level contextual data from the U.S. Census and other sources. Using a small area estimation approach, the study generates county-level estimates of MMR (measles, mumps, and rubella) vaccine uptake among children under age 5 across the United States.

Research sample

This cross-sectional, retrospective cohort study combines individual-level survey data with contextual county-level data to generate small area estimates of MMR (measles, mumps, and rubella) vaccine uptake among children under age 5 across the United States. The analytic sample consisted of 22,062 U.S. parents or guardians of children under 5, drawn from the OutbreaksNearMe digital health surveillance platform hosted by SurveyMonkey. Participants were recruited anonymously without incentives, following unrelated platform activity, and were not required to have an account to participate. The survey was administered between July 2023 and April 2024.

The survey instrument asked respondents to self-report their demographics and their child's vaccination status. All available respondents with a child under 5 years old were included in the analysis. The data was weighted to reflect national population distributions using standard demographic poststratification procedures. This weighting approach has been validated by the CD for

use in real-time surveillance and public health modeling.

In the final sample, approximately 58% identified as female and 42% as male. The largest age group was 30–39 years (45%), followed by 18–29 (35%), 40–49 (15%), and 50 or older (5%). Racial and ethnic composition included 50% white, 26% Black, 14% Hispanic, 6% Asian, and 3% other or multiracial. Educational attainment varied, with 52% reporting a high school education or less and 21% holding a college or postgraduate degree. Insurance coverage included Medicaid or Medicare (43%), private or employer-based insurance (47%), and uninsured (10%). A majority of respondents (56%) reported household incomes below \$50,000, and 72% were employed. Participants represented all four U.S. Census regions and identified politically as Democrats (27%), Republicans (29%), or Independents/Other (44%).

Sampling strategy

This study uses data from the OutbreaksNearMe–SurveyMonkey platform, a national digital health surveillance initiative developed by Boston Children’s Hospital and administered through SurveyMonkey. The survey employs a non-probability river sampling strategy, in which participants are recruited after completing unrelated online activity. This approach yields a large, demographically diverse respondent pool from across the U.S. Although river sampling is not random sampling, inverse probability weights based on U.S. Census 2019 data (age, sex, race/ethnicity, education, and geography) are applied to improve representativeness. This weighting approach has been validated in prior studies, including work supported by the CDC. The analytic sample includes all available respondents during the study period (2023–2024) who reported having a child under age 5. No statistical methods were used to pre-determine sample size, as the study was observational and used all eligible observations to maximize geographic coverage. Data saturation was not a relevant consideration due to the structured survey design and the use of predefined close-ended variables rather than qualitative or inductive data collection.

Data collection

Data were collected via the OutbreaksNearMe–SurveyMonkey online platform. The survey instrument consisted of a structured, self-administered questionnaire developed by researchers at Boston Children’s Hospital and delivered through the SurveyMonkey interface. All responses were entered directly by participants on their own devices (e.g., computers, tablets, smartphones). No interviewer was present during data collection, and participation was anonymous. Respondents completed the survey voluntarily after engaging in unrelated activities on the platform. No identifiable information was collected, and no experimental manipulation or random assignment was involved. Researchers were not present during data collection and were not blinded, as the study was observational and hypothesis-driven rather than experimental.

Timing

The survey data was collected between July 2023 and April 2024 with no gaps in between.

Data exclusions

After restricting the sample to respondents who reported having a child under age 5, we excluded 579 responses (approximately 2% of the eligible sample) due to missing data on the child’s MMR vaccination status. These exclusions were necessary to ensure a complete outcome variable for modeling.

Non-participation

Participation rate was approximately 11% among those randomly invited.

Randomization

No randomization was involved. Although the primary goal of the study is prediction rather than causal inference, we incorporated a broad set of individual- and county-level covariates selected through a transparent and theory-driven process. This included both demographic and contextual variables (e.g., income, insurance, political context, urbanicity) to account for observed sources of variation and reduce the influence of omitted variable bias in model estimation.

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