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Research Letter

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Geographic access to United States SARS-CoV-2 testing sites highlights healthcare disparities and may bias transmission estimates

Benjamin Rader, MPH^{1,2,†}, Christina M. Astley, MD, ScD^{1,3,4,5,†}, Karla Therese L. Sy, MS^{2,6}, Kara Sewalk, MPH¹, Yulin Hswen, ScD, MPH^{1,7,8}, John S. Brownstein, PhD^{1,5} and Moritz U.G. Kraemer, DPhil^{1,5,8,9,*}

¹Computational Epidemiology Lab, Boston Children's Hospital, Boston, MA, USA, ²Department of Epidemiology, Boston University School of Public Health, Boston, MA, USA, ³Division of Endocrinology, Boston Children's Hospital, Boston, MA, USA, ⁴ Broad Institute of Harvard and MIT, Cambridge, MA, USA, ⁵Harvard Medical School, Harvard University, Boston, MA, USA, ⁶Department of Global Health, Boston University School of Public Health, Boston, MA, USA, ⁷Department of Epidemiology & Biostatistics, University of California San Francisco, CA, USA, ⁸Bakar Computational Health Sciences Institute, University of California San Francisco, CA, USA and ⁹Department of Zoology, University of Oxford, Oxford, UK

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Uniform access to SARS-CoV-2 testing is crucial for controlling the COVID-19 epidemic.¹ Lack of testing can result in the epidemic spreading undetected² and increase the risk of extensive local transmission. The USA has been slow to develop reliable diagnostic tests and, while there has been recent improvement in testing capabilities,³ large-scale testing remains a serious concern.

Inequalities in geographic accessibility to healthcare in the USA have been documented to cause negative health outcomes for seasonal influenza transmission and other diseases.⁴ Further, travel time negatively impacts healthcare-seeking behaviour.⁵ The deployment of SARS-CoV-2 testing within existing medical infrastructure, while logistically efficient, may exacerbate this disparity in health outcomes⁶ and underestimate disease burden in disadvantaged populations.

Geographic accessibility to SARS-CoV-2 testing sites, to our knowledge, has not been systematically quantified. Therefore, we evaluated whether testing sites were equally accessible to populations across the USA, leveraging two public SARS-CoV-2 testing site datasets and a high-resolution map of travel times.

American Community Survey (2014–2018) data for contiguous US states were used to tabulate county-level covari-

ates including population, population density ($\ln \frac{\text{Mean population}}{\text{Census block}}$), median income, percent uninsured and percent minority (1 – %non-Hispanic white).

A national database of SARS-CoV-2 testing sites was curated using the Carbon Health (N = 5376) and CodersAgainstCovid (N = 1547) datasets (accessed 7th April 2020). Carbon Health (carbonhealth.com/covid-19-testing-centers) prospectively called urgent care centers and hospitals on publicly listed telephone numbers starting 17th March 2020 to ask whether SARS-CoV-19 testing was being offered. Additionally, a verified, non-exhaustive collection of publicly documented and user-entered testing sites were included. CodersAgainstCovid identified urgent care centers, hospitals, drive-throughs, health departments and other facility types prospectively starting 15 March 2020, through volunteer-verified 'webscraping' and crowdsourcing (https://codersagainstcovid.org/).

We identified and geocoded (R v.3.6.2 ggmap v3.0.0) 6236 unique sites (687 excluded following manual de-duplication and cleaning). Related site ontologies were collapsed into metaontologies (e.g. Urgent with Immediate Care). To date, this is

^{*}To whom correspondence should be addressed: Email: moritz.kraemer@zoo.ox.ac.uk

[†]These authors contributed equally to this study.

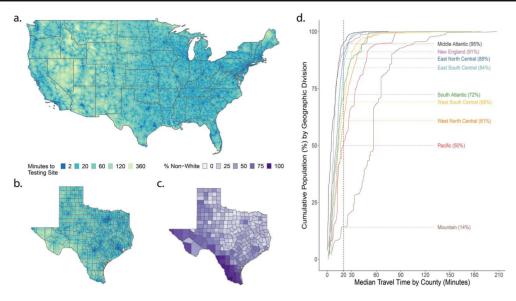


Figure 1. Distribution of SARS-CoV-2 testing sites. (A) Travel time to the nearest testing site per 1 km² area (shorter travel time in darker blue) in the 48 contiguous US states plus DC. (B) Travel time as in Panel A enlarged to show detail in the state of Texas. (C) Percent minority (1 – %non-Hispanic white) by county in Texas. (D) Median travel time by county versus the cumulative population for each geographic region (excluding two outlier counties). Vertical dashed line at 20 minute median travel time. Horizontal dotted lines indicate cumulative population percentage in that region (in parenthesis) residing in counties with <20 minutes median travel time.

the largest database of US testing sites known to the authors. To evaluate completeness (as of 20 April 2020), we identified public testing sites listed in sample areas: 34 in Illinois (https://www.dph.illinois.gov/covid19/covid-19-testing-sites), five in Colorado (https://covid19.colorado.gov/testing-covid-19) and 104 in West Virginia (https://www.wvhealthconnection.com/covid-19). Our database included 169, 85 and 60 sites in each area, respectively. We confirmed our database identified at least one site in every city in Texas operating a drive-through (https://www.dshs.state.tx.us/coronavirus/testing.aspx).

We used published friction-based travel times⁷ between \sim 1 km² gridded cells in the USA, accounting for topography and the most efficient non-air travel method. Median travel times for the shortest path to testing sites across all grid cells in each county (N = 3108) were calculated using the Dijkstra's algorithm.⁸

Generalized linear models (R stats v3.6.2) were used to estimate the correlation of population density, percent minority, percent uninsured and median income on median travel time, by county. We also tested for potential interactions between population density and percent minority or percent uninsured. Influential counties with a Cook's distance measure over 4/N were excluded (up to N = 175).

We collated 6236 SARS-Cov-2 testing sites in the contiguous US states. Testing sites (Supplementary Table 1) were often affiliated with medical centers (43%) and urgent care (47%) and were infrequently drive-through (3%). Testing sites were spatially clustered (Moran's I = 0.037; z = 61.4; $P < 10^{-5}$), around US urban centers (Supplementary Figure 1).

The travel time from each 1 km² grid cell to the nearest US testing site is spatially heterogeneous at the national and state level (Figure 1A–C). Thirty percent of the population live

in a county (N = 1920) with a median travel time over 20 minutes, though with pronounced regional differences (Figure 1D) ranging from 5 to 86%.

Population density, a determinant of population distribution, was associated with a shorter median county-level travel time (Table 1). While controlling for population density as a potential confounder, percent minority was associated with an increase in travel time, as was percent uninsured. These associations remained when also adjusting for median income. We found a significant negative interaction between percent uninsured and population density (P < 0.01) suggesting that the disparity of longer rural travel times is greater in counties where a higher proportion of the population is uninsured. Percent minority and population density did not interact statistically.

Using two large, national datasets of SARS-CoV-2 testing sites paired with estimates of travel times, we demonstrate an uneven distribution of critical public health resources. The testing site distribution recapitulates structural disparities, including inequities among minority, uninsured and rural groups, which may further perpetuate disparities as the pandemic progresses. Differential accessibility to testing may lead to biases in estimation of disease incidence and potentially delay identification of COVID-19 hotspots. In the absence of representative testing, syndromic surveillance tools may provide early warning signals, and augment targeted-testing and other public health interventions.

Despite efforts to ensure comprehensiveness, in some regions, our dataset may be missing testing sites (e.g. West Virginia). While some additional testing sites have been created, given recent difficulties scaling up, we believe our database remains representative. There remains potential for differential missingness of sites in areas with reduced 'webscraping' visibility or sites

Table 1. Generalized linear regression models. Associations between covariates and median travel time in minutes by county in the 48 contiguous US states and DC

	Model 1	Model 2	Model 3	Model 4
Intercept	61.45***	59.99***	56.29***	51.36***
	[59.82, 63.08]	[58.34, 61.64]	[54.17, 58.41]	[47.70, 55.03]
Log of population density	-13.41***	-14.14***	-12.94***	-14.13***
	[-14.02, -12.79]	[-14.76, -13.52]	[-13.56, -12.32]	[-14.78, -13.47]
Percent minority (%)		0.15***		0.13***
		[0.12, 0.18]		[0.10, 0.17]
Percent uninsured (%)			0.41***	0.23**
			[0.30, 0.53]	[0.09, 0.38]
Median income (\$10 000)				2.52***
				[1.46, 3.59]
N	2942	2934	2942	2931
AIC	24 321.32	24 192.59	24 291.55	24 097.61
Pseudo R2	0.38	0.41	0.39	0.41

^{***}P < 0.001; **P < 0.01; *P < 0.05.

specifically placed to address inaccessibility. Nevertheless, this work highlights the need for comprehensive resources and the utility of data sharing during a pandemic.

The travel-time metric used here accounts for the presence of public transportation and routine traffic. Early evidence shows widespread variability in mobility reductions during the epidemic. Our estimates of differential access present a conservative picture of inequality in the USA, which may be worse if public transit closures and private transportation were also modelled. Additionally, our models do not examine other, non-geographic barriers to SARS-CoV-2 testing access (e.g. economic), nor geography for residents in Alaska and Hawaii. Travel time, for example, is shorter for urban uninsured minority groups, and therefore does not explain the below average testing rates in disadvantaged urban areas (e.g. Philadelphia).

In summary, reduced geographic access to SARS-CoV-2 testing sites is associated with sociodemographic factors that, in turn, are linked to poor structural access to care and health outcomes. The location of future testing sites should explicitly account for travel time and sociodemographic predictors, in addition to other public health testing requirements.

Supplementary Data

Supplementary data are available at JTM online.

Author Contributions

B.R., C.M.A., J.S.B. and M.U.G.K. contributed to conceptualization. B.R., K.T.L.S. and K.S. contributed to data acquisition. B.R., C.M.A. and K.T.L.S. contributed to the data analysis. All authors contributed to the interpretation of results and manuscript writing.

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locations were curated and made open-source by volunteers (https://codersagainstcovid.org/about-us) from CodersAgainst-COVID. Testing site data and consultation were also kindly provided by the CarbonHealth team (https://carbonhealth.com/coronavirus).

Data and Code Availability

Median travel time to testing center by county, sociodemographic variables, code and raster for main analysis: https://figshare.com/s/4b2af17d00e4751685c5. For access to the Carbon Health dataset, please email: coviddata@carbonhealth.com. For access to the CodersAgainstCOVID dataset, visit: http://github.com/codersagainstcovid.org.

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Conflict of interest

None declared.

References

- Lipsitch M, Swerdlow DL, Finelli L. Defining the epidemiology of Covid-19—studies needed. N Engl J Med 2020; 382:1194–6. doi: 10.1056/NEJMp2002125.
- Li R, Pei S, Chen B et al. Substantial undocumented infection facilitates the rapid dissemination of novel coronavirus (SARS-CoV2). Science 2020; 3221:eabb3221. doi: 10.1126/science.abb3221.

- Cohen J. The United States badly bungled coronavirus testing—but things may soon improve. Science 2020. doi: 10.1126/science.abb5152. Accessed 21 April 2020.
- Zipfel CM, Bansal S. Health inequities in influenza transmission and surveillance. MedRxiv 2020; 1–16. doi: 10.1101/2020.03.30.20048017.
- Alegana VA, Maina J, Ouma PO et al. National and subnational variation in patterns of febrile case management in Sub-Saharan Africa. Nat Commun 2018; 9:4994. doi: 10.1038/s41467-018-07536-9.
- Yancy CW. COVID-19 and African Americans. JAMA 2020; 60611:19–20. doi: 10.1001/jama.2020.6548.
- Weiss DJ, Nelson A, Gibson HS et al. A global map of travel time to cities to assess inequalities in accessibility in 2015. Nature 2018; 553:333–6. doi: 10.1038/nature25181.

- Hulland EN, Wiens KE, Shirude S et al. Travel time to health facilities in areas of outbreak potential: maps for guiding local preparedness and response. BMC Med 2019; 17:232. doi: 10.1186/s12916-019-1459-6.
- Kaplan S, Thomas K Despite Promises, Testing Delays Leave Americans 'Flying Blin'. 2020 The New York Times. https://www.nytimes.com/2020/04/06/health/coronavirus-testing-us.html. Accessed 21 April 2020.
- 10. Klein B, LaRock T, McCabe S *et al.* Assessing changes in commuting and individual mobility in major metropolitan areas in the United States during the COVID-19 outbreak. 2020:1–29. Available at https://uploads-ssl.webflow.com/5c9104426f6f88ac129ef3 d2/5e8374ee75221201609ab586_Assessing_mobility_changes_in_the_United_States_during_the_COVID_19_outbreak.pdf.