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Research letter

Assessment of geographic access to monoclonal antibodies in the United States

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Neutralizing monoclonal antibodies (mAbs) reduce viral loads in mild to moderate COVID-19 patients¹ and may prevent admission or death in non-hospitalized patients according to the United States (U.S.) Department of Health and Human Services (HHS).2 However, the production of mAbs continues to lag behind daily COVID-19 cases and has led to national shortages and forced rationing.^{1,3} Concerns also exist that rural facilities may have reduced access to these treatments because they lack the capacity for large-scale infusion operations. In September 2021, the Department of Health and Human Services (HHS) announced a new mAb distribution method due to a drastic uptick in COVID-19 cases. HHS calculates incident and hospitalized cases and allocates each state mAbs accordingly, adjusting each week's supply based on previously unused allocations.2 It is reported that existing measures of local disease burden may be an ineffective marker for the distribution of mAbs.¹ In addition, only certain mAbs are an effective form of treatment for the omicron variant of COVID-19.4 Although sociogeographic disparities in accessing COVID-19 resources (e.g. testing, vaccinations have been well documented and recent reports suggest disparities among mAb recipients, 7 to our knowledge there has been no examination of how the current distribution scheme affects geographic access to mAbs across the United

We used previously described methods⁶ to measure the travel time from each populated census tract in the contiguous U.S. $(n=71\,927)$ to the closest facility that had received a delivery of mAbs (n=4240, HHS Data Hub, accessed 6 January 2022) and the closest facility administering a COVID-19 vac-

cine (*n* = 36 259, Vaccines.gov, ⁵ accessed 6 January 2022). We compared the travel time to a vaccine and mAbs to understand the tradeoff in receiving preventive COVID-19 care as opposed to therapeutic COVID-19 care. Travel times were computed by estimating the quickest means of transversal (e.g. car, public transportation) from each 1 km² grid-square of the U.S. to the closest location of interest and averaged across all grid-squares in each tract. Extended travel times (≥30 min) were defined by the Veterans Administration's primary care standard. We also report tract-level 2019 American Community Survey demographics, tract-level collapsed Rural–Urban Commuting Area codes, ¹⁰ county-level COVID-19 vaccination (Covid.CDC.gov, accessed 6 January 2022), and county-level 2020 Presidential voting (ElectionLab.MIT.edu).

Individuals in the mean census tract had to travel twice as long to the nearest mAbs administration site [10.5 min (SD: 14.3)] compared with the nearest COVID-19 vaccine location [4.6 min (SD: 8.9)], although there was high variability especially across the urban–rural spectrum. In addition, a large number of individuals (n = 15252601) live in census tracts with extended travel times to mAbs, but not the COVID-19 vaccine (Figure 1). Extended travel times to mAbs were prevalent in the Mountain division of the U.S. When comparing census tracts under 30 min from mAbs/vaccines to those with extended travel times to mAbs (Table 1), the latter are on average more rural, older (4.3 years), have lower household incomes (\$6389 less), higher percent white (14.5% more), more likely uninsured (0.8% more) and have poorer internet access (6.1% lower). Low-access census tracts were also 16.3 points less likely to be in majority-vaccinated

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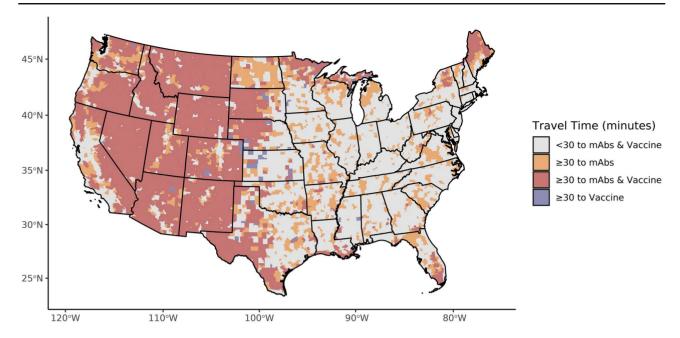


Figure 1. US travel times to monoclonal antibodies and the COVID-19 vaccine. Mean travel times for each census tract in the contiguous United States to the nearest location that previously (as of 6 January 2022) received a delivery of neutralizing mAbs according to the US Department of Health and Human Services and the nearest location distributing the COVID-19 vaccine (as of 6 January 2022) according to Vaccines.gov. Travel times are aggregated at the census tract level which may obscure hyperlocal access measurements, especially in large tracts like those found in the western US.

counties and were 45.0 points more likely to have voted for Donald Trump in 2020.

In this study, we quantified access to two critical pandemic resources and show that travel times to mAbs are double travel times to the COVID-19 vaccine. This result highlights the relative tradeoff and additional cost of receiving mAbs compared with the vaccine. The travel cost is on top of the significant financial and administrative burden of using mAbs,¹ as opposed to a COVID-19 vaccine. The complex logistics of direct administration of mABs (e.g. testing requirements, trained administrators) have resulted in a distribution scheme that has left over 15 million people in the U.S. with extended travel times to this COVID-19 therapy. Our results also highlight profound disparities in access to mAbs and that the areas that are most susceptible to COVID-19 due to low vaccination rates are also less likely to have convenient access to this treatment. These communities also have lower household income and less access to the internet which may exacerbate health service access barriers.

There are important limitations to this study. Our sociode-mographic measures are population aggregates, and our measure of travel time assumes equal access to the quickest means of transportation. Here we use mAB delivery as a proxy for mAB administration which may obscure its complex distribution patterns. Further research should leverage individual-level mAb recipient data to quantify how geographic access differences manifests in outcome disparities. However, this study is the first to use national data to quantify the geographic access of mAbs. Our results show that the urban–rural barriers that are common with accessing health services 10 and other COVID-19 resources 5,6 are also present in accessing mAbs.

Shifting COVID-19 variants and their respective treatments suggest a dynamic landscape where various therapies (e.g.

mAB formula, direct acting antivirals, etc.) may fluctuate in appropriateness. Comprehensive solutions including unconventional mAB delivery technologies such as home administration (e.g. via paramedics) and local community clinic partnerships are essential to remedy access barriers and to ensure equitable availability of this important COVID-19 treatment.

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Conflicts of Interest

The authors have no conflicts of interest to declare.

Data Sharing

All data used in this analysis is accessible from publicly available sources.

Author Contributions

Rader, Whaley, Brownstein and Cantor conceived of the study. All authors were involved in the acquisition, analysis, or interpretation of data. Rader and Cantor wrote the original draft of the manuscript. All authors engaged in critical revision of the manuscript for important intellectual content. Whaley and Brownstein obtained funding and supervised the project. All authors have seen and approve the final text.

Table 1. Characteristics of US census tracts (n=71 927) by travel time to mAbs and the COVID-19 vaccine

	<30 min to mAbs and vaccine ($n = 66781$)	\geq 30 min to mAbs ($n = 3840$)	\geq 30 min to mAbs and vaccine ($n = 1235$)	\geq 30 min to vaccine ($n = 71$)	Total $(n = 71927)$
Rural–Urban designation, n (%)					
Urban	51 100 (99.14%)	398 (0.77%)	43 (0.08%)	0 (0.00%)	51 541 (71.66%)
Suburban	6570 (83.38%)	1067 (13.54%)	238 (3.02%)	5 (0.06%)	7880 (10.96%)
Large rural	5259 (85.04%)	738 (11.93%)	177 (2.86%)	10 (0.16%)	6184 (8.60%)
Small rural	2205 (70.54%)	722 (23.10%)	186 (5.95%)	13 (0.42%)	3126 (4.35%)
Isolated rural	1494 (49.65%)	894 (29.71%)	579 (19.24%)	42 (1.40%)	3009 (4.18%)
No designation	153 (81.82%)	21 (11.23%)	12 (6.42%)	1 (0.53%)	187 (0.26%)
Sex	, ,	,	, ,	,	,
Female, mean % (SD)	50.85 (±4.73)	49.74 (±4.98)	$48.54 (\pm 5.64)$	50.14 (±2.38)	50.75 (±4.77)
Median age, years	,	, ,	, ,	,	,
Mean (SD)	$39.28 (\pm 7.71)$	$43.57 (\pm 7.72)$	$45.18 (\pm 9.40)$	$43.85 (\pm 6.76)$	$39.62 (\pm 7.84)$
Missing, n (%)	23 (0.03%)	1 (0.03%)	0 (0.00%)	0 (0.00%)	24 (0.03%)
Household income, \$1000	, ,	,	,	,	,
Mean (SD)	$33.70 (\pm 14.16)$	$27.31 (\pm 6.98)$	$26.58 (\pm 7.05)$	$26.58 (\pm 5.26)$	$33.23 (\pm 13.88)$
Missing, n (%)	77 (0.12%)	5 (0.13%)	5 (0.40%)	0 (0.00%)	87 (0.12%)
Race, mean % (SD)	, ,	,	,	,	,
White	71.35 (± 25.22)	$85.82 (\pm 16.76)$	$81.35 (\pm 25.67)$	$81.06 (\pm 26.75)$	$72.31 (\pm 25.09)$
Black	14.55 (±22.18)	$7.09 (\pm 14.45)$	$2.87 (\pm 10.12)$	$7.80 \ (\pm 20.79)$	13.95 (±21.80)
Hispanic	17.21 (±21.72)	9.35 (±16.32)	$14.47 (\pm 20.34)$	11.59 (±18.24)	16.74 (±21.52)
American Indian or Alaska	$0.69 (\pm 2.70)$	1.66 (±6.95)	9.71 (±24.07)	6.71 (±19.50)	0.90 (±4.59)
Native ^a	,	, , ,	, , ,	(,	, , , ,
Asian	$5.18 (\pm 9.18)$	$0.79 (\pm 1.39)$	$0.66 (\pm 1.43)$	$0.48 (\pm 0.64)$	$4.86 (\pm 8.93)$
Hawaiian or Pacific Islandera	$0.13 (\pm 0.60)$	0.08 (±0.37)	$0.12\ (\pm0.59)$	$0.07 (\pm 0.20)$	$0.13 (\pm 0.59)$
Two or more races	3.20 (±2.78)	2.43 (±2.49)	$2.55 (\pm 2.73)$	2.19 (±2.15)	$3.15 (\pm 2.77)$
Other race	4.90 (±8.89)	$2.13 (\pm 5.46)$	$2.73 (\pm 6.58)$	1.70 (±5.20)	4.71 (±8.73)
Health insurance, mean % (SD)	, ,	,	,	,	,
Employer	45.65 (±15.03)	$39.00 (\pm 10.72)$	$33.68 (\pm 12.52)$	$36.18 (\pm 8.79)$	$45.08 (\pm 14.94)$
Direct	6.51 (±4.22)	6.68 (±3.87)	$7.72 (\pm 5.48)$	8.32 (±4.64)	6.54 (±4.23)
Medicare	5.36 (±3.13)	6.78 (±2.99)	$7.70 (\pm 3.66)$	6.29 (±2.33)	5.48 (±3.16)
Medicaid	$15.81 (\pm 12.28)$	$16.03~(\pm 8.52)$	17.01 (±11.44)	$16.10 (\pm 10.54)$	15.84 (±12.09)
Veterans administration	$0.28 (\pm 0.51)$	$0.37 (\pm 0.52)$	$0.42 (\pm 0.57)$	0.33 (±0.31)	0.29 (±0.51)
Tricare	$0.96 (\pm 3.86)$	$0.79 (\pm 2.29)$	$0.88 (\pm 3.86)$	$0.48 (\pm 0.83)$	$0.95 (\pm 3.79)$
Two or more	5.96 (±3.25)	6.61 (±2.86)	$6.01 (\pm 3.34)$	6.53 (±2.64)	5.99 (±3.24)
Uninsured	$8.75 (\pm 7.02)$	$9.50 (\pm 6.04)$	$11.34 (\pm 7.42)$	$10.20~(\pm 6.27)$	8.84 (±6.98)
Missing, n (%)	96 (0.14%)	6 (0.16%)	2 (0.16%)	0 (0.00%)	104 (0.14%)
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Mean (SD)	$14.73 (\pm 10.36)$	$20.86 (\pm 9.76)$	23.58 (±14.09)	25.78 (±11.15)	15.22 (±10.56)
Missing, n (%)	167 (0.25%)	8 (0.21%)	4 (0.32%)	0 (0.00%)	179 (0.25%)
Majority of adults fully	92.73 (92.53–92.93)	76.43 (75.05–77.76)	76.36 (73.87–78.68)	76.06 (64.20–85.05)	91.56 (91.36–91.76
vaccinated ^b , % (95% CI)	(= =.00 > =.>0)	(((2.320 00.00)	(, 1.00 / 1.70
Won by Donald Trump in 2020 ^b , N (95% CI)					
Mean (SD)	37.47 (37.10–37.84)	82.50 (81.25-83.68)	75.38 (72.86–77.74)	85.92 (75.16–92.68)	40.57 (40.22–40.93
Missing, n (%)	0 (0.00%)	0 (0.00%)	3 (0.24%)	0 (0.00%)	3 (0.00%)

^aOnly contiguous United States included in analysis. ^bCounty-level statistic (e.g. 40.57% of census tracts are in a county won by Trump and 91.56% of census tracts are in a county where the majority (50%+) of adults (18+) are fully vaccinated against COVID-19)

References

- Goldstein RH, Walensky RP. The challenges ahead with monoclonal antibodies: from authorization to access. *JAMA* 2020; 324:2151.
- U.S. Department of Health & Human Services. Bamlanivimab/etesevimab. Public Health Emergency. https://www.phe.gov/eme rgency/events/COVID19/investigation-MCM/Bamlanivimab-e tesevimab/Pages/default.aspx (4 November 2021, date last accessed).
- Persad G, Peek ME, Shah SK. Fair allocation of scarce therapies for COVID-19. Clin Infect Dis 2021; ciab1039.
- VanBlargan LA, Errico JM, Halfmann PJ et al. An infectious SARS-CoV-2 B.1.1.529 omicron virus escapes neutralization by therapeutic monoclonal antibodies. Nat Med 2022.
- Rader B, Astley CM, Sewalk K et al. Spatial accessibility modeling of vaccine deserts as barriers to controlling SARS-CoV-2. medRxivPublished online June 12 20212021.06.09.21252858. 10.1101/2021.06.09.21252858 preprint: not peer reviewed.
- Rader B, Astley CM, Therese Sy KL et al. Geographic access to United States SARS-CoV-2 testing sites highlights healthcare disparities and may bias transmission estimates. J Travel Med 2020; 2020:1–4.

- 7. Wiltz JL, Feehan AK, Molinari NM et al. Racial and ethnic disparities in receipt of medications for treatment of COVID-19 United States, March 2020–August 2021. MMWR Morb Mortal Wkly Rep 2022; 71:96–102.
- 8. 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.
- Massarweh NN, Itani KMF, Morris MS. The VA MISSION act and the future of veterans' access to quality health care. *JAMA* 2020; 324:343.
- Peek-Asa C, Wallis A, Harland K et al. Rural disparity in domestic violence prevalence and access to resources. J Womens Health 2011; 20:1743–9.