

Letters

RESEARCH LETTER

Asymptomatic and Symptomatic SARS-CoV-2 Infections After BNT162b2 Vaccination in a Routinely Screened Workforce

A 2-dose regimen of the BNT162b2 vaccine (Pfizer-BioNTech) against SARS-CoV-2 was authorized in December 2020 based on reported 94.8% efficacy.¹ Although an association between vaccination and a reduction in symptomatic disease has been well described, an association with asymptomatic infection remains unclear.^{2,3}

Methods | In March 2020, St Jude Children’s Research Hospital initiated routine, test-based screening of asymptomatic workers and targeted testing for symptomatic individuals and those with known exposure. Polymerase chain reaction-based testing of midturbinate samples from asymptomatic employees was performed at least weekly. On December 17, 2020, vaccination with BNT162b2 was initiated. “Vaccine-eligible” workers were individuals meeting state vaccination guidelines.⁴ Vaccinated employees receiving BNT162b2 were followed up from their first dose date. Unvaccinated employees were followed up from December 17, 2020, or their first asymptomatic screen result, whichever was later. The end of surveillance was March 20, 2021, employment termination, a positive test result, or receipt of other vaccines, whichever



Related article

was earlier. No person contributed to both groups. Individuals with prior COVID-19 exposure were excluded. When asymptomatic infections were analyzed, symptomatic and known exposure cases were treated as competing risks; when symptomatic infections were analyzed, positive results from asymptomatic screening were treated as competing risks.

The incidence rate ratio (IRR), the ratio of confirmed COVID-19 cases per person-days of follow-up in vaccinated compared with unvaccinated groups, with 95% CIs,⁵ was used as a measure of association between vaccination and infection. An analysis by time after dose 1 and 2 was also performed. Cumulative incidence curves were estimated with the Kaplan-Meier estimator. Analyses were performed in R version 4.0.3.

The study was determined by the St Jude institutional review board to be exempt (secondary use of data), for which participant informed consent is not required.

Results | Between December 17, 2020, and March 20, 2021, 5217 workers met vaccination criteria, 3052 (58.5%) received at least 1 BNT162b2 dose, and 2776 (53.2%) received 2 doses; 2165 (41.5%) were unvaccinated. Median follow-up was 81 days in the unvaccinated group and 72 days among vaccinated employees. In the vaccinated group, 66.0% were women, 60.3% White individuals, 19.4% Black individuals, 88.7% younger than 65 years, and 47.2% health care personnel⁶; in the unvaccinated group, 58.3% were women, 40.3% White individuals, 24.6%

Table. Estimated Incidence Rate Ratio Against Any SARS-CoV-2 Infection and Asymptomatic or Symptomatic/Contact SARS-CoV-2 Infection^a

Vaccination status	Follow-up time, person-days (No. at risk)	Any positive test result		Asymptomatic screening positive test result		Screening positive test result based on the presence of symptoms or known COVID-19 exposure	
		No.	IRR (95% CI) ^b	No.	IRR (95% CI)	No.	IRR (95% CI)
Unvaccinated total follow-up ^c	149 718 (2165)	185		79		106	
Vaccinated total follow-up	198 480 (3052)	51	0.21 (0.15-0.28)	29	0.28 (0.18-0.42)	22	0.16 (0.10-0.25)
Vaccinated periods							
0-11 d after dose 1	32 807 (3052)	24	0.59 (0.39-0.91)	10	0.58 (0.30-1.12)	14	0.60 (0.35-1.05)
≥12 d after dose 1 and before dose 2	32 481 (2942)	17	0.42 (0.26-0.70)	10	0.58 (0.30-1.13)	7	0.30 (0.14-0.65)
0-6 d after dose 2	16 492 (2776)	4	0.20 (0.07-0.53)	3	0.35 (0.11-1.09)	1	0.09 (0.01-0.61)
≥7 d after dose 2	116 700 (2724)	6	0.04 (0.02-0.09)	6	0.10 (0.04-0.22)	0	0 ^d

Abbreviation: IRR, incidence rate ratio.

^a Follow-up periods for unvaccinated employees began on December 17, 2020, or on their first asymptomatic screening date, whichever was later. Individuals who received vaccines other than BNT162b2 were censored on vaccination. Follow-up periods for vaccinated workers began when they received their first dose. Workers who remained SARS-CoV-2 negative during follow-up were censored on March 20, 2021, or on the employment termination date,

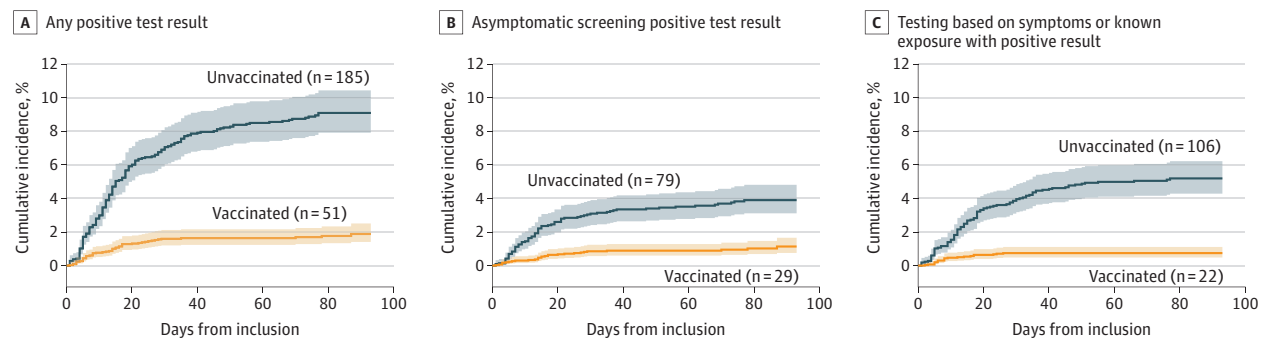
whichever was earlier. No person contributed to both groups. Individuals with prior COVID-19 exposure were excluded.

^b Incidence rate ratio is the ratio of confirmed COVID-19 cases per person-days of follow-up in vaccinated compared with unvaccinated groups.

^c The unvaccinated group was treated as the reference group for all calculations.

^d 95% CI does not apply.

Figure. Cumulative Incidence of COVID-19 Against SARS-CoV-2 Infections After the First Dose



A total of 2165 unvaccinated employees and 3052 vaccinated employees were included. A, Any SARS-CoV-2 infection among St Jude employees during follow-up. B, Asymptomatic infections identified through routine asymptomatic screening; SARS-CoV-2 cases through testing based on the presence of symptoms or known COVID-19 exposure were treated as

competing risks. C, Positive results via testing based on the presence of symptoms or known COVID-19 exposure; positive results from asymptomatic screening were treated as competing risks. Shaded areas are 95% CIs.

Black individuals, 84.8% younger than 65 years, and 25.7% health care personnel.

Among vaccinated employees, 51 tested positive for SARS-CoV-2 during follow-up (41 before and 10 after the second dose); 29 (56.9%) were diagnosed through asymptomatic screening. Among unvaccinated employees, 185 tested positive and 79 (42.7%) were asymptomatic. The IRR was 0.21 (95% CI, 0.15-0.28) for any SARS-CoV-2 infection, 0.28 (95% CI, 0.18-0.42) for asymptomatic screen results, and 0.16 (95% CI, 0.10-0.25) for symptomatic or known exposure cases (Table). The IRR within the first 11 days after the first dose was 0.58 to 0.60 for all 3 outcomes. The IRR for positive results via asymptomatic screening from 12 days after the first vaccine dose until the second dose (median interval between doses, 21 days [range, 11-49 days]) was 0.58 (95% CI, 0.30-1.13), within 7 days after the second dose, 0.35 (95% CI, 0.11-1.09), and 7 days or more after the second dose, was 0.10 (95% CI, 0.04-0.22). There were no positive symptomatic or known exposure cases more than 7 days after the second dose. Unvaccinated employees had higher cumulative incidence of a positive test result than vaccinated employees, and higher incidences of positive test results via asymptomatic screening, for symptoms, or for known exposure (Figure).

Discussion | This study found an association between vaccination with BNT162b2 in hospital employees and a decreased risk of symptomatic and asymptomatic infections with SARS-CoV-2. Limitations include the observational design; short follow-up time; small cohort size, which led to an inability to match the 2 groups and unequal follow-up; differential temporal risk during the surveillance; and that the group choosing not to be vaccinated may have been more prone to higher-risk behavior. The unequal follow-up time and the latter 2 limitations may have biased the results in favor of vaccination. Further research is needed to determine whether a reduction in risk of asymptomatic infection leads to reduced transmission.

Li Tang, PhD
Diego R. Hijano, MD, MSc
Aditya H. Gaur, MD, MBBS
Terrence L. Geiger, MD, PhD
Ellis J. Neufeld, MD, PhD
James M. Hoffman, PharmD, MS
Randall T. Hayden, MD

Author Affiliations: Department of Biostatistics, St Jude Children's Research Hospital, Memphis, Tennessee (Tang); Department of Infectious Diseases, St Jude Children's Research Hospital, Memphis, Tennessee (Hijano, Gaur); Department of Pathology, St Jude Children's Research Hospital, Memphis, Tennessee (Geiger, Hayden); Department of Hematology, St Jude Children's Research Hospital, Memphis, Tennessee (Neufeld); Department of Pharmaceutical Sciences, St Jude Children's Research Hospital, Memphis, Tennessee (Hoffman).

Corresponding Author: Li Tang, PhD, Department of Biostatistics, St Jude Children's Research Hospital, 262 Danny Thomas Pl, Mail Stop 768, Memphis, TN 38105 (li.tang@stjude.org).

Accepted for Publication: April 12, 2021.

Published Online: May 6, 2021. doi:10.1001/jama.2021.6564

Author Contributions: Dr Tang had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. Drs Tang and Hijano are co-first authors. Drs Hoffman and Hayden are co-senior authors.

Concept and design: Tang, Hijano, Gaur, Geiger, Hoffman, Hayden.

Acquisition, analysis, or interpretation of data: Tang, Hijano, Gaur, Neufeld, Hoffman, Hayden.

Drafting of the manuscript: Tang, Hijano, Gaur, Hoffman.

Critical revision of the manuscript for important intellectual content: All authors.

Statistical analysis: Tang.

Administrative, technical, or material support: Geiger, Hoffman, Hayden.

Supervision: Hijano, Gaur, Geiger, Neufeld, Hoffman.

Conflict of Interest Disclosures: Dr Neufeld reports serving as a consultant to Pfizer outside the submitted work. Dr Hayden reports serving on advisory boards for Roche Molecular, Quidel Corporation, and Inflammatrix outside the submitted work. No other disclosures were reported.

Funding/Support: This work was supported in part by the American Lebanese Syrian Associated Charities (ALSAC).

Role of the Funder/Sponsor: The funder had no role in design and conduct of the study; collection, management, analysis, and interpretation of the data; preparation, review, or approval of the manuscript; and decision to submit the manuscript for publication.

Additional Contributions: We acknowledge the following members from St Jude Children's Research Hospital for their contributions to this work: data

aggregation and analysis efforts of Yilun Sun, MS, Biostatistics, and Sandra Dennis, BS, Human Resources; the insight and guidance of Richard Webby, PhD, Infectious Diseases, Greg Armstrong, MD, Epidemiology and Cancer Control, and Motomi Mori, PhD, Biostatistics; and the commitment and support of Hana Hakim, MD, Infectious Diseases, Kari Lahmon, BSN, Nursing Administration, and Sri Suganda, MT, MB (ASCP), Pathology. No one received financial compensation for their contributions.

1. Polack FP, Thomas SJ, Kitchin N, et al; C4591001 Clinical Trial Group. Safety and efficacy of the BNT162b2 mRNA Covid-19 vaccine. *N Engl J Med*. 2020;383(27):2603-2615. doi:10.1056/NEJMoa2034577
2. Amit S, Regev-Yochay G, Afek A, Kreiss Y, Leshem E. Early rate reductions of SARS-CoV-2 infection and COVID-19 in BNT162b2 vaccine recipients. *Lancet*. 2021;397(10277):875-877. doi:10.1016/S0140-6736(21)00448-7

3. Dagan N, Barda N, Kepten E, et al. BNT162b2 mRNA Covid-19 vaccine in a nationwide mass vaccination setting. *N Engl J Med*. 2021;384(15):1412-1423. doi:10.1056/NEJMoa2101765

4. State of Tennessee Health Department. COVID-19 vaccination plan. Posted March 8, 2021. Accessed March 21, 2021. https://www.tn.gov/content/dam/tn/health/documents/cedep/novel-coronavirus/COVID-19_Vaccination_Plan.pdf

5. Hightower AW, Orenstein WA, Martin SM. Recommendations for the use of Taylor series confidence intervals for estimates of vaccine efficacy. *Bull World Health Organ*. 1988;66(1):99-105.

6. Centers for Disease Control and Prevention. What healthcare personnel need to know about COVID-19 vaccines. Updated April 30, 2021. <https://www.cdc.gov/coronavirus/2019-ncov/vaccines/recommendations/hcp.html>