Data Sharing Statement

Walter EB, Talaat KR, Sabharwal C, et al. Evaluation of the BNT162b2 Covid-19 Vaccine in Children 5 to 11 Years of Age. N Engl J Med. DOI: 10.1056/NEJMoa2116298.

Question	Authors' Response
Will the data collected for your study	Yes
be made available to others?	
Would you like to offer context for	_
your decision?	
Which data?	Other (eg, partial data sets) — please describe
Additional information about data	—
How or where can the data be obtained?	Upon request, and subject to certain criteria, conditions and exceptions (see https://www.pfizer.com/science/clinical-trials/trial- data-and-results for more information), Pfizer will provide access to individual de-identified participant data from Pfizer-sponsored global interventional clinical studies conducted for medicines, vaccines and medical devices (1) for indications that have been approved in the US and/or EU or (2) in programs that have been terminated (i.e., development for all indications has been discontinued). Pfizer will also consider requests for the protocol, data dictionary, and statistical analysis plan. Data may be requested from Pfizer trials 24 months after study completion. The de-identified participant data will be made available to researchers whose proposals meet the research criteria and other conditions, and for which an exception does not apply, via a secure portal. To gain access, data requestors must enter into a data access agreement with Pfizer.
When will data availability begin?	June 15, 2023
When will data availability end?	
Will any supporting documents be available?	—
Which supporting documents?	_
Additional information about	—
supporting documents	

How or where can supporting documents be obtained?	_
When will supporting documents availability begin?	_
When will supporting documents availability end?	—
To whom will data be available?	—
For what type of analysis or purpose?	—
By what mechanism?	—
Any other restrictions?	_
Additional information	—

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