

May 13, 2022

No Vaccine Safety Data for Pregnant Women

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#	Exclusion Number	Criterion Description	Cr
2.A	1	Other medical or psychiatric condition (incl. recent (within past year) or active suicidal ideation/behavior) lab abnormal by that may increase the risk of study participation	
2.B	2	Known infection with human immunodeficiency virus (HIV), hepatitis C virus (HCV), or hepatitis B virus (HBV)	
2.A	3	History of severe adverse reaction associated with a vaccine and/or severe allergic reaction (eg, anaphylaxis) to any component of the study intervention(s)	
2.A	4	Receipt of medical care intended to prevent COVID-19	
2.A	5	Stages 1 and 2 only: Previous clinical or serological diagnosis of COVID-19	
2.F	6	Screened participants in Stage 1 only: Individuals at high risk for severe COVID-19 (full details in protocol)	
2.A	7	Screened participants in Stage 1 only: Individuals currently working in occupations with high risk of exposure to SARS-CoV-2 (eg, healthcare worker, emergency response personnel)	
2.A	8	Immunocompromised individuals with known or suspected immunodeficiency, as determined by history and/or laboratory/physical assessment on	
2.I	9	Screened participants in Stage 1 only: Individuals with a history of autoimmune disease or an active autoimmune disease requiring therapeutic intervention	
2.I	10	Bleeding diatheses or condition associated with prolonged bleeding that would, in the opinion of the investigator, contraindicate intramuscular injection	
2.A	11	Women who are pregnant or breastfeeding	
2.I	12	Previous vaccination with any coronavirus vaccine	
2.A	13	Subjects who receive immunosuppressive therapy, such as cytotoxic agents or systemic corticosteroids	
2.A	14	Screened participants in Stage 1 only: Regular receipt of inhaled/injected corticosteroids	
2.A	15	Receipt of biologic/plasma products or immunoglobulins, from 60 days before study intervention administered on or planned receipt throughout the study	
2.A	16	Participation in other studies involving study intervention within 28 days prior to study entry and/or during study participation	
2.A	17	Previous participation in other studies involving study intervention containing IgG replacement	
2.F	18	Screened participants in Stage 1 only: Positive serological test for SARS-CoV-2 IgM and/or IgG antibodies at the screening visit	
2.A	19	Screened participants in Stage 1 only: Screening hematologic/clinical chemistry lab >=Grade 1 abnormality. Except bilirubin, other stable Grade 1 abnormalities may be considered eligible by investigator	
2.F	20	Screened participants in Stage 1 only: Positive test for HIV, hepatitis B surface antigen	

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2.A	21	Screened participants in Stage 1 only: SARS-CoV-2 NAb1-positive nasal swab within 24 hours before receipt of study intervention	
2.A	22	Investigator is to staff or Pfizer employees directly involved in the conduct of the study, site staff otherwise supervised by the investigator, and their respective family members	

The confidential documents, which Pfizer wanted to slowly release over a period of 75 years are being rapidly released every month by court order. The latest 80,000 pages were released the first week of May. One of the amazing findings within these documents is that 22 groups of people were excluded from the clinical trials including pregnant women and breastfeeding women.

Thus, there are no safety data on the use of the Pfizer vaccine in pregnant or breastfeeding women.

Pregnant women have nonetheless been encouraged to get the shot. Meanwhile, Department of Defense data show a significant increase in congenital malformations, from nearly 11,000 in 2020 to nearly 19,000 in the first half of 2021.

“Interview with Dr. Naomi Wolf,” War Room, Real America’s Voice, May 3, 2022: <https://bit.ly/3KQlrwW>

“Annotated Study Book for Study Design: C4591001 (Study Design Version: 11.0), Sponsor Pfizer, October 12 2020: <https://bit.ly/3P1W2Vo>

Presented daily by Twila Brase, President and Co-founder
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