

Pfizer Documents reveal at least 800 people never finished the COVID Vaccine Trial due to Death, Injury or Withdrawn Consent

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By The Exposé

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Pfizer Documents reveal at least 800 people never finished the COVID Vaccine Trial due to Death, Injury or Withdrawn Consent

One of the confidential Pfizer documents that the U.S. Food and Drug Administration (FDA) has been forced to publish by court order reveals that approximately 800 people never completed the phase 1 Pfizer Covid-19 vaccine clinical trial in the USA due to either losing their life, suffering a serious adverse event or suddenly withdrawing their consent.

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The US Food and Drug Administration (FDA) attempted to delay the release of Pfizer's COVID-19 vaccine safety data for 75 years despite approving the injection after only 108 days of safety review on December 11th, 2020.

But in early January 2022, Federal Judge Mark Pittman ordered them to release 55,000 pages per month. They released 12,000 pages by the end of January.

Since then, PHMPT has posted all of the documents on its website. The latest drop happened on 1st June 2022.

One of the documents contained in the latest data dump is '125742_S1_M5_5351_c4591001 fa interim discontinued patients.pdf'.

The document provides a 112-page list of subjects who withdrew from the phase 1 clinical trial of the Pfizer Covid-19 injection, and provides a vague description as to why.

The first 14 pages list 102 subjects who withdrew from the study. This equates to an average of 7.2 subjects per page. So based on a further 93 pages detailing withdrawn subjects this equates to approximately 780 people who withdrew from the first phase of the clinical trial alone. The actual number could be slightly more or slightly less.

Many of the subjects mysteriously revoked consent to continue in the trial due to reasons such as re-reading the consent form and deciding it is not what they had originally agreed to.

					SUBJECT	
C4591001 4444 44441792	1	9	01OCT2020	VACCINATION	WITHDRAWAL BY SUBJECT	Subject, re-reads consent and decided it is not what she'd understood. Decides to revoke consent.
	1	9	01OCT2020	FOLLOW-UP	WITHDRAWAL BY SUBJECT	The subject rejects the terms of consent on a subsequent rereading.

Whilst others withdrew their consent to continue in the study following receipt of dose 1 for unexplained reasons, meaning they did not want to receive the second dose.

					SUBJECT	
C4591001 1254 12541074	2	38	28OCT2020	VACCINATION	WITHDRAWAL BY SUBJECT	subject requested to withdrawal consent
	2	38	28OCT2020	FOLLOW-UP	WITHDRAWAL BY SUBJECT	subject wished to withdrawal consent.
C4591001 1260 12601121	1	20	30SEP2020	VACCINATION	WITHDRAWAL BY SUBJECT	patient withdrew from study prior to vaccine # 2
	1	20	30SEP2020	FOLLOW-UP	WITHDRAWAL BY SUBJECT	withdrawal of consent

But unfortunately, there are several who withdrew their consent to continue the trial due to suffering serious adverse events. Page 110 of the document lists one person who suffered a pulmonary embolism, which is a blood vessel in the lungs blocked by a blood clot. The condition can be life-threatening if not treated quickly.

C4591001 1269 12691124	2	1	29SEP2020	VACCINATION	WITHDRAWAL BY SUBJECT	subject withdrew consent
	2	1	29SEP2020	FOLLOW-UP	WITHDRAWAL BY SUBJECT	Withdrew consent
C4591001 1270 12701057	1	61	02NOV2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/PULMONARY EMBOLISM
C4591001 4444 44441152	1	22	12OCT2020	VACCINATION	WITHDRAWAL BY SUBJECT	The volunteer expresses his desire to withdraw consent to participate in the study.

Page 108 of the document lists one person who suffered a cerebral infarction. The condition is also known as an ischemic stroke and occurs as a result of disrupted blood flow to the brain due to problems with the blood vessels that supply it. A lack of adequate blood supply to brain cells deprives them of oxygen and vital nutrients which can cause parts of the brain to die off.

C4591001 1247 12471066	1	43	10NOV2020	VACCINATION	NO LONGER MEETS ELIGIBILITY CRITERIA	Patient tested positive for COVID 19 and was symptomatic prior to visit 2
C4591001 1247 12471135	1	22	21OCT2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/CEREBRAL INFARCTION

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Page 102 of the document lists a person who suffered a transient ischaemic attack. The condition is also known as a mini-stroke and is again caused by disruption in the blood supply to the brain. Unfortunately, due to suffering the mini-stroke, this person also fell and fractured their ankle at the same time.

C4591001 1224 12241012	1	13	25AUG2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/ANKLE FRACTURE
	1	13	25AUG2020	FOLLOW-UP	ADVERSE EVENT	ADVERSE EVENT/FALL
	1	13	25AUG2020	FOLLOW-UP	ADVERSE EVENT	ADVERSE EVENT/MUSCULAR WEAKNESS
	1	13	25AUG2020	FOLLOW-UP	ADVERSE EVENT	ADVERSE EVENT/TRANSIENT ISCHAEMIC ATTACK

Page 100 of the document lists a person who withdrew consent to continue in the trial due to losing their hearing and going completely deaf in one ear. The page also lists another person who suffered 'syncope' which is a temporary loss of consciousness usually related to insufficient blood flow to the brain.

C4591001 1152 11521476	1	21	15OCT2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/DEAFNESS UNILATERAL
C4591001 1152 11521497	1	31	06NOV2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/SYNCOPE

Page 94 of the document lists a person who withdrew consent due to suffering tachycardia. The condition refers to a heart rate that is too fast and can be caused by poor blood supply to the heart muscle.

	1	30	22SEP2020	FOLLOW-UP	LOST TO FOLLOW-UP	unable to reach subject to return for follow up visit
C4591001	1	2	12AUG2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/TACHYCARDIA
1112						
11121118						

Page 44 of the document lists a person who withdrew consent due to suffering paraparesis. The condition refers to partial paralysis in both legs due to disrupted nerve signals from the brain to the muscles.

C4591001	1	23	13NOV2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/AMNESIA
1140						
11401306						
	1	23	13NOV2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/PARAPARESIS

Sadly, the document also lists several people who were no longer able to take part in the phase 1 clinical trial due to losing their lives. Several deaths can be viewed throughout the document including, but not limited to pages 106/107, 101, 80 and 47.

C4591001	2	16	28SEP2020	VACCINATION	DEATH	DEATH
1231						
12313972						

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16.2.1.4 Listing of Subjects Withdrawn From the Study – All Subjects

Subject	Dose No.	Rel Day ^a	Withdrawal Date	Study Phase	Withdrawal Reason	Comment ^b
	2	16	28SEP2020	FOLLOW-UP	DEATH	DEATH
C4591001	2	30	12OCT2020	VACCINATION	WITHDRAWAL BY SUBJECT	the decision to stop participating is due to personal reasons
1231						
12314003						

Source – Page 106/107

16.2.1.4 Listing of Subjects Withdrawn From the Study – All Subjects

Subject	Dose No.	Rel Day ^a	Withdrawal Date	Study Phase	Withdrawal Reason	Comment ^b
C4591001	1	4	13SEP2020	VACCINATION	DEATH	Cause of Death Atherosclerotic Disease
1162						
11621327						
	1	4	13SEP2020	FOLLOW-UP	DEATH	DEATH

Source – Page 101

C4591001 1006 10061020	1	36	16SEP2020	VACCINATION	ADVERSE EVENT	ADVERSE EVENT/CORONARY ARTERY OCCLUSION
	1	36	16SEP2020	FOLLOW-UP	ADVERSE EVENT	ADVERSE EVENT
C4591001 1007 10071101	2	63	21OCT2020	FOLLOW-UP	DEATH	DEATH

Source – Page 80

16.2.1.4 Listing of Subjects Withdrawn From the Study – All Subjects

Subject	Dose No.	Rel Day ^a	Withdrawal Date	Study Phase	Withdrawal Reason	Comment ^b
C4591001 1152 11521085	1	8	26AUG2020	VACCINATION	DEATH	DEATH
	1	8	26AUG2020	FOLLOW-UP	DEATH	DEATH
C4591001 1156 11561006	1	20	08SEP2020	VACCINATION	NO LONGER MEETS ELIGIBILITY CRITERIA	Exclusion Criteria 5
C4591001 1156 11561015	1	39	28SEP2020	VACCINATION	PREGNANCY	PREGNANCY

Source – Page 47

We know why those who sadly died or were injured didn't complete the trial. But why did hundreds more refuse to continue the phase 1 clinical trial of the Pfizer Covid-19 vaccine in the USA after being eager to participate originally?

And why were the FDA desperate to hide this document among many others for at least 75 years?

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