

Exhibit 456

Report 50: 20% of Post-Jab Strokes Fatal in the 90
Days Following Pfizer COVID mRNA Vaccine
Rollout

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Report 50: 20% of Post-Jab Strokes Fatal in the 90 Days Following Pfizer COVID mRNA Vaccine Rollout

December 26, 2022 • by Barbara Gehrett, MD; Joseph Gehrett, MD; Chris Flowers, MD; and Loree Britt

The War Room/DailyClout Pfizer Documents Analysis Project Post-Marketing Team created the following Stroke System Organ Class (SOC) Review from data in Pfizer document

5.3.6 Cumulative Analysis of Post-Authorization Adverse Event Reports of PF-07302048 (BNT162B2) Received Through 28-FEB-2021 (https://www.phmpt.org/wp-content/uploads/2022/04/reissue_5.3.6-postmarketing-experience.pdf)

(a.k.a., "5.3.6"). The stroke category comprises the diagnoses of strokes attributed to either obstruction of blood flow through arteries to areas of the brain or due to bleeding around or into the brain. Additionally, in this adverse event category, Pfizer included syndromes of diffuse venous clotting in and around the brain and clotting in the venous pools within the skull (cerebral venous sinus thrombosis and cavernous sinus thrombosis). Arterial obstruction blocks oxygen-rich blood delivery, whereas the venous thrombosis prevents drainage of blood from within the head.

Within the stroke data set, there are 275 patients with 300 different events reported; and **20% of the stroke events were fatal.**

It is important to note that the Adverse Events in the 5.3.6 document were reported to Pfizer for *only a 90-day period* starting on December 1, 2020, the date of the United Kingdom's public rollout of Pfizer's COVID-19 experimental mRNA "vaccine" product.

Please read the full report below.



War Room/DailyClout Pfizer Document Analysis

Post-Marketing Team Micro-Report 3:

Stroke System Organ Class (SOC) Review of 5.3.6

SOURCE:
https://www.phmp.org/wp-content/uploads/2022/04/reissue_5.3.6-postmarketing-experience.pdf

5.3.6 AE REPORTING PERIOD:
 "Since the first temporary authorization for emergency supply under Regulation 174 in the UK (01 December 2020) and through 28 February 2021."

ABBREVIATIONS:
 5.3.6 : Pfizer source document

SOC : System Organ Class

AE : Adverse Event

AESI : Adverse Event of Special Interest

EUA : Emergency Use Authorization by FDA

PM : Post-Marketing

BNT162b2 : Pfizer's mRNA COVID-19 vaccine

SEQUELAE: an abnormal condition resulting from a previous disease, injury, or other trauma

AGE GROUPS defined in 5.3.6 (p. 25 footnote) :

Adult	18 - 64
Elderly	≥ 65
Child	2 - 11
Adolescent	12 - < 18
Infant	1 - 23 months



20Dec22

Stroke Search terms: ICD Central nervous system haemorrhages and cerebrovascular accidents	Number of cases: 275 (0.6% of the total PM dataset), of which 180 medically confirmed and 95 non-medically confirmed. Country of incidence: UK (81), US (66), France (23), Germany (21), Norway (14), Netherlands and Spain (11 each), Sweden (9).
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 FDA-CBER-2021-0683-000076

Table 7. AESI Evaluation for BNT162b2

AESI ¹ Category	Post-Marketing Cases Evaluation ² Total Number of Cases (N=260)
<i>(Primary Path) OR ICD Cerebrovascular venous and sinus thrombosis (Primary Path)</i>	Total (9), Italy (3), Belgium (3), Denmark, Finland, Poland and Switzerland (2 each); the remaining 8 cases originated from 8 different countries: • Subjects' gender (n=273): female (182), male (91); • Subjects' age group (n=265): Adult (19), Elderly (205), Child ³ (1); • Number of relevant events: 300, all serious; • Most frequently reported relevant PTs (≥ 1 occurrence) included: o PTs indicative of Ischemic stroke: Cerebrovascular accident (160), Ischemic stroke (41), Cerebral infarction (13), Cerebral ischemia, Cerebral thrombosis, Cerebral venous sinus thrombosis, Ischemic cerebral infarction and Lacunar infarction (3 each), Basal ganglia stroke, Cerebellar infarction and Thrombotic stroke (2 each); o PTs indicative of Haemorrhagic stroke: Cerebral haemorrhage (26), Haemorrhagic stroke (11), Haemorrhage intracranial and Subarachnoid haemorrhage (5 each), Cerebral haematoma (4), Basal ganglia haemorrhage and Cerebellar haemorrhage (2 each); • Relevant event onset latency (n = 241): Range from ~24 hours to 41 days, median 2 days; • Relevant event outcome: fatal and resolved involving (61 each), resolved with sequelae (16), not resolved (83) and unknown (83). Conclusion: This cumulative case review does not raise new safety issues. Surveillance will continue.

https://www.phmp.org/wp-content/uploads/2022/04/reissue_5.3.6-postmarketing-experience.pdf

This category comprises the diagnoses of strokes attributed to either obstruction of blood flow through arteries to areas of the brain or due to bleeding around or into the brain. Additionally, in this adverse event category, Pfizer included syndromes of diffuse venous clotting in and around the brain and clotting in the venous pools within the skull (cerebral venous sinus thrombosis and cavernous sinus thrombosis). Arterial obstruction blocks oxygen-rich blood delivery, whereas the venous thrombosis prevents drainage of blood from within the head.

The occurrence of these events ranged from within the first 24 hours to 41 days post-vaccination, and 50% were within two days.

Outcomes were listed as "resolved or resolving" in 61 events (20%), though the number "resolved" is not independently reported.

- Adverse Events were reported to Pfizer during a 90-day period, following the December 1, 2020, public rollout of its COVID-19 experimental "vaccine" product.
- In the Pfizer 5.3.6 document, these AEs were categorized by System Organ Classes (SOC) – in other words, by systems in the body.
- Within the stroke data set, there are 275 patients with 300 different events reported.
- There were no "non-serious" events. ALL adverse events were categorized as "serious."

95 (32%) did not resolve or "resolved with Sequelae." Thus, those patients were left with health deficits. 83 (28%) unknown outcomes were reported.

The fatal events were 61 (22% of patients; 20% of total stroke events). In the 95 events that did not resolve or "resolved with sequelae," how severely disabled were the survivors? Strokes are life-altering events. Even Pfizer categorized all of the reported stroke adverse events as serious.

An additional observation of note involves the unusual diagnosis of cerebral venous sinus thrombosis. There are three cases reported in this data set. This is an extremely rare diagnosis, but it occurred three times in the first 90 days of the Pfizer mRNA COVID-19 "vaccine" rollout.

Pfizer's Conclusion: "This cumulative case review does not raise new safety issues."

Post-Marketing Team's CONCLUSION:

How many serious ADVERSE EVENTS, UNRESOLVED, and UNKNOWN outcomes does it take? How many DEATHS does it take to RECALL PFIZER'S UNSAFE "VACCINE"?

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<https://dailyclout.io/wp-content/uploads/DC-V3-Post-Marketing-Team-Stroke-micro-report-1-1.pdf> (<https://dailyclout.io/wp-content/uploads/DC-V3-Post-Marketing-Team-Stroke-micro-report-1-1.pdf>)



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Unholy Alliance Between Government, NGOs, and Big Tech Colluded to Smear and Deplatform COVID Dissidents

(<https://dailyclout.io/unholy-alliance-between-government-ngos-and-big-tech-colluded-to-smear-and-deplatform-covid-dissidents/>)

NEXT STORY

Rise of the Fourth Reich: Confronting COVID Fascism with a New Nuremberg Trial

(<https://dailyclout.io/rise-of-the-fourth-reich-confronting-covid-fascism-with-a-new-nuremberg-trial/>)

15 replies added

Scott Kiley

January 10, 2023 Reply

I am active with a Health Freedom group in Collier County Florida. We meet with County Commissioners and push just like you. I reference your site and your work to everyone I know. The Chair wants more information. How do I best obtain, deliver concise, easy to understand information. Scott Kiley Please email so we can begin a conversation.

Scott Kiley

January 10, 2023 Reply

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January 14, 2023 Reply

11B10 | identify as "TRANS-VAXXED!"



Naomi Wolf

January 15, 2023 Reply

Thank you for contacting us!
Best Naomi

Jeanmarie

January 17, 2023 Reply

My very healthy active father of 88 years old (he walked on the treadmill daily and ate blueberries for breakfast etc. Etc.) had been Moderna vaccinated and received a booster around December 2021. After a bout of COVID-19 in January 2022 he then had a brain bleed on March 6, 2023. He is now in a wheelchair and has dementia. He can't walk by himself and he can converse well enough but has lost mental capacity. He knows us. Anyway, I believed it at the time that it was the shot plus contracting COVID-19 subsequently that led to the brain bleed.

Markalete

March 9, 2023 Reply

The more excess deaths like SAD Sudden Adult Death there are, more people will have the chance to wake up. My own family is an anecdotal case study, I think. 74 years old female, (not vaxxed) I caught covid (probably Omicron) December of 2021 and then my husband (73 years old) also got sick with it, both with comorbidities. I'd been taking Quercetin which is in the same family as Hydroxychloroquine as a prophylactic along with supplements etc. recommended to improve immunity, so I didn't get very sick. I just increased the dosage along with other vital supplements including colloidal silver, Oregano EO, etc. The fatigue lasted but I could function to care for myself and my husband. I may have some long covid symptoms of inflammation like a storm (from a reservoir of spike proteins and virus in the epithelial cells of my gut?). My husband was treated with Ivermectin (plus) and is still on it prophylactically. My relatively healthy 87

years old sister contracted covid and survived in her apartment with going to doctor visits and help from family and friends. Whether she has taken any more jabs, I don't know; 85 years old brother, heart attack survivor, (I'm sure vaccinated) developed a cough for several months before it developed into double pneumonia for which he had the most awful medication reserved for such resistant cases and the resultant runaway bacterial diarrhea he also had to be treated for. They sent him home finally to die in his home and passed away in September 2022; another sister with open heart surgery as a child and low ejection numbers and a pacemaker, had a reaction to the booster and tested positive with symptoms for covid(!). She had monoclonal antibody infusion and felt better in 24 hours or so. I didn't tell her that the monoclonal antibodies were produced using a fetal cell line from 1973 "voluntary" abortion. Whether she took any other jabs I don't know. Since then, she developed very painful clots in the epithelial layer of the skin on her legs (microclots) treated with anti-inflammatories, then it was CHF, congestive heart failure and now she is fighting to reduce multiple clots in her lungs on O2 while she decides what to do with custody of her great-granddaughters, her heart is fragile; her son has a diagnosis of chronic lymphatic leukemia and now suddenly has an extremely high white blood count for which he will have to have Chemo; another sister, (stroke survivor) wanted to protect her husband who has a terminal diagnosis so she got vaccinated. Then she developed continual diarrhea, losing a lot of weight and strength while she cares for her husband at home with little help. She had multiple benign polyps removed from her colon. She still has diarrhea. She understands her immune system is compromised from the jab. She and her husband have eaten well for many years so she may be able to reverse the condition and/or control the symptoms with diet; another vaccinated sister reported to me that she thought she was having TIAs (spike proteins crossing the blood/brain barrier producing microclots, my opinion), she now has invited her son to live with her; my two brothers, both vaccinated, one has avoided any adverse effects or covid as far as I know, the other with a heart condition has a cough but not pneumonia, his wife had recent pneumonia and their son with Lyme disease is having trouble with it, probably all vaccinated, following the science, she said.

([HTTPS://DAILYCLOUT.IO/SERIOUS-STROKE-ADVERSE-EVENTS-FOLLOWING-PFIZER-COVID-19-MRNA-VACCINATION/COMMENT-PAGE-1/#COMMENTS](https://dailyclout.io/serious-stroke-adverse-events-following-pfizer-covid-19-mrna-vaccination/comment-page-1/#comments))

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