

Exhibit 466

Report 66:1,077 Immune-Mediated/Autoimmune
Adverse Events in First 90 Days of Pfizer mRNA
“Vaccine” Rollout, Including 12 Fatalities. Pfizer
Undercounted This Category of Adverse Events by 270
Occurrences.

<https://dailyclout.io/report-66-1077-immune-mediated-autoimmune-adverse-events-in-first-90-days-of-pfizer-mrna-vaccine-rollout-including-12-fatalities-pfizer-undercounted-this-category-of-adverse-events-by-270-occurr/>

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Pfizer Reports

Report 66: 1,077 Immune-Mediated/Autoimmune Adverse Events in First 90 Days of Pfizer mRNA "Vaccine" Rollout, Including 12 Fatalities. Pfizer Undercounted This Category of Adverse Events by 270 Occurrences.

April 13, 2023 • by Joseph Gehrett, MD; Barbara Gehrett, MD; Chris Flowers, MD; and Loree Britt

The War Room/DailyClout Pfizer Documents Analysis Project Post-Marketing Group (Team 1) – Joseph Gehrett, MD; Barbara Gehrett, MD; Chris Flowers, MD; and Loree Britt – wrote an important review of immune-mediated/autoimmune adverse events found 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"). This category is comprised of the numerous diseases resulting from disordered immune attacks against tissues of any of the body's organs.

However, it excludes anaphylaxis which has its own

separate report (<https://dailyclout.io/report-65-in-the-first-three-months-of-pfizers-mrna-vaccine-rollout-nine-patients-died-of-anaphylaxis/>)

, as well as autoimmune diseases attaching nerve tissue (e.g., Guillain-Barré), which have been explored in other 5.3.6 neurological and musculoskeletal reports (<https://dailyclout.io/category/pfizer-reports/>).


It is important to note that the adverse events (AEs) 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.

Highlights of this report include:

- **Twelve immune-mediated/autoimmune adverse events were fatal.**
- Of adverse events with gender specified, **77% (526) were female, and 23% (156) were male** – a female to male ratio of 3.4 to 1.
- Ages of affected patients were **19% elderly, 71% adult, <1% adolescent, and 10% not reported.**
- Time from vaccination to onset was given for 75% of events with a range of 24 hours to 30 days. **Half started within 24 hours of injection.**
- **Only immune-mediated/autoimmune diseases or conditions with over 10 cases are included in Pfizer's reporting, leaving out 270 adverse events of this type.** Thus, adverse events in this category were undercounted by Pfizer.
- Adverse events in this group include:
 - **Hypersensitivity** (which is not further defined, though it accounts for 55% of the AEs.
 - **Peripheral neuropathy** (<https://www.mayoclinic.org/diseases-conditions/peripheral-neuropathy/symptoms-causes/syc-20352061>), which often causes weakness, numbness and pain, usually in the hands and feet, and can also affect other areas and body functions including digestion, urination, and circulation.
 - **Pericarditis** (inflammation of the lining of the heart).
 - **Myocarditis** (immune attack against the heart muscle).
 - **Encephalitis** (brain inflammation disorders).
 - **Diabetes.**
 - **Psoriasis.**
 - **Dermatitis.**
 - **Blistering Skin disorder.**
 - **Autoimmune disorder.**
 - **Raynaud's phenomenon.**
- Immune rejection of transplanted organs was not mentioned whether from there being zero instances of it or, perhaps, because there were 10 or fewer instances.

Pfizer concluded, "This cumulative case review does not raise new safety issues. Surveillance will continue." However, as of the date of this report, 25.5 months after the completion of Pfizer's post-marketing analysis, no additional safety data has

been released to the public.



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War Room/DailyClout Pfizer Document Analysis

Post-Marketing Team Micro-Report 11:
Immune-Mediated/Autoimmune AESIs Review of 5.3.6


SOURCE:
https://www.phmpt.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."

5.3.6 AE CASES/EVENTS:
TOTAL AE CASES: 42,086
TOTAL AE EVENTS: 158,893

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

AUTHORS:
Dr Joseph Gehrett MD
Dr Barbara Gehrett MD
Dr Chris Flowers MD
Loree Britt


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Immune-Mediated/Autoimmune AESIs

Search criteria: Immune-mediated/autoimmune disorders (SMO) (Broad and Narrow) OR Autoimmune disorders FLGT (Primary Path) OR PTs Cytokine release syndrome; Cytokine storm; Hypersensitivity

- Number of cases: 1050 (2.5 % of the total PM dataset), of which 760 medically confirmed and 290 non-medically confirmed;
- Country of incidence (>10 cases): UK (267), US (257), Italy (70), France and Germany (69 each), Mexico (56), Sweden (55), Spain (32), Greece (31), Israel (21), Denmark (18), Portugal (17), Austria and Czech Republic (16 each), Canada (12), Finland (10). The remaining 74 cases were from 24 different countries.
- Subjects' gender (n=682): female (526), male (156).
- Subjects' age group (n=944): Adult (746), Elderly (196), Adolescent (2).
- Number of relevant events: 1077, of which 780 serious, 297 non-serious.
- Most frequently reported relevant PTs (>10 occurrences): Hypersensitivity (596), Neuropathy peripheral (49), Pericarditis (32), Myocarditis (25), Dermatitis (24), Diabetes mellitus and Encephalitis (16 each), Psoriasis (14), Dermatitis Bullous (13), Autoimmune disorder and Reynaud's phenomenon (11 each).
- Relevant event onset latency (n = 807): Range from <24 hours to 30 days, median <24 hours.
- Relevant event outcome: resolved/resolving (517), not resolved (215), fatal (12), resolved with sequelae (22) and unknown (312).

Conclusion: This cumulative case review does not raise new safety issues. Surveillance will continue

https://www.phmpt.org/wpcontent/uploads/2022/04/reissue_5.3.6-postmarketexperience.pdf

This category comprises the numerous diseases resulting from disordered immune attacks against tissues of any of the body's organs. Pfizer has grouped some conditions in a very general way, such as "hypersensitivity." This is defined as an exaggerated response of the normal immune system. However, anaphylaxis, another hypersensitivity reaction, has its own [separate SOC report](#).

Autoimmune diseases attacking nerve tissue (Guillain-Barré, multiple sclerosis, polyneuropathy, and others) are not in this SOC but are found under the [Neurological and Musculoskeletal SOC reports](#). However, dermatitis (skin hypersensitivity) is listed here.

Only diseases or conditions with over 10 cases are described with a diagnosis or symptom. Hypersensitivity had 596 (55%) of the adverse events. The conditions in this large grouping are not further defined in the report. Peripheral neuropathy had 49 events (5%) though, and, as discussed above, other neuropathies are separately listed and presumably different patients. There are **32 diagnosed pericarditis events (3%)** with inflammation of the lining around the heart. **Myocarditis (immune attack against the heart muscle itself) had 25 events (2%).**


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.

Of those whose sex was specified, 526 (77%) were female, and 156 (23%) were male (a ratio of 3.4 to 1).

Age was listed as elderly 196 (19%), adult 746 (71%), adolescent 2 (<1%), and not reported 106 (10%).

There were 1,077 events reported with **780 (72%) serious** and 297 (28%) non-serious.



12 (1.1%) AEs were fatal.

The time from vaccination to adverse event onset was defined in 807 (75%) of the 1,077 events with a range of within 24 hours to 30 days. **Half of the adverse events started within 24 hours of the injection.** 517 (48%) had outcomes reported as "resolved" or "resolving." **215 events (20%) were not resolved, 22 (2%) resolved with sequelae, and 312 (29%) had unknown outcome.**

Further observation regarding autoimmune events:

There was no mention of immune rejection of transplanted organs. It is unknown whether Pfizer found none, or whether there were 10 or fewer and thus not listed explicitly.

There is a VAERS report of a 17-year-old male who was vaccinated 1/19/2021 and was hospitalized mid-May 2021 with heart transplant rejection and heart failure.

<https://dailyclout.io/wp-content/uploads/p1-Post-Marketing-Autoimmune-micro-report.pdf> (<https://dailyclout.io/wp-content/uploads/p1-Post-Marketing-Autoimmune-micro-report.pdf>)

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Post
Marketing
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Encephalitis (brain inflammatory disorders) **occurred 16 times (1.5%)**. There were 16 cases (1.5%) of **diabetes** reported though not specified as insulin dependent or non-insulin dependent.

Skin disorders listed were **psoriasis** 14 events (1.3%), **dermatitis** 24 events (2%) and **blistering skin disorder** 13 events (1.2%). **Autoimmune disorder**, a broad group rather than a specific disease, had 11 events (1%). **Raynaud's phenomenon** (spasm of the small arteries in the extremities) also had 11 events (1%).

Pfizer, by excluding those conditions with fewer than 11 events, has therefore left out explanations of 270 adverse events. Were there instances of Crohn's disease, ulcerative colitis, or glomerulonephritis?

Pfizer's Conclusion:

This cumulative case review does not raise new safety issues.
Surveillance will continue.

NOTE: As of the date of this team report, no additional safety data have been publicly released.



Post-Marketing Team's CONCLUSION
RECALL this unsafe "vaccine."

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