# Exhibit 453

## Report 47: Blood System-Related Adverse Events Following Pfizer COVID-19 mRNA Vaccination

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November 30, 2022 • by Barbara Gehrett, MD; Joseph Gehrett, MD; Chris Flowers, MD; and Loree Britt

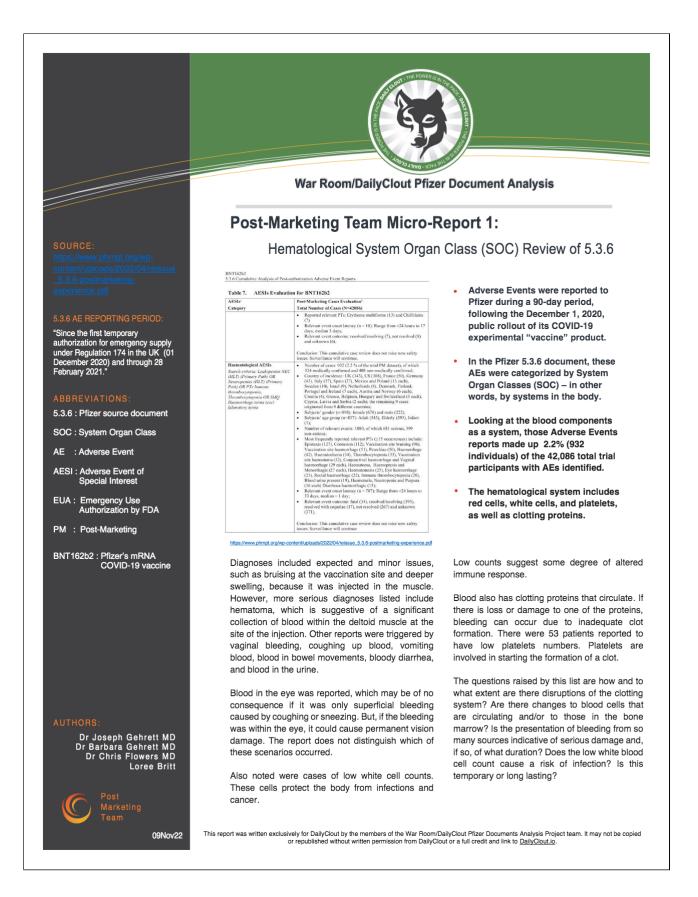
The War Room/DailyClout Pfizer Documents Analysis Project Post-Marketing Team created the following two-page Hematological System Organ Class (SOC) Review from data in Pfizer document

5.3.6 *Cumulative Analysis of Post-Authorization Adverse Event Reports of PF-07302048* (*BNT162B2*) *Received Through 28-FEB-2021* (https://www.phmpt.org/wpcontent/uploads/2022/04/reissue\_5.3.6-postmarketing-experience.pdf)

(a.k.a., "5.3.6"). The hematological system is the human body's blood system and includes red cells, white cells, platelets, and clotting proteins. Viewing the blood components as a system, its Adverse Events (AEs) reports made up 2.2% (932 individuals) of the 42,086 total trial participants with AEs identified.

Fifty percent of the blood-related adverse events reported were noted within 48 hours of Pfizer COVID-19 mRNA vaccination, but there were also cases reported up to 33 days post-injection. In the hematological group of adverse events, **there were 34 deaths and 17 cases of permanent damage**.

It is important to note that the 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.



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https://dailyclout.io/wp-content/uploads/DC-V7-combined-p1-2-Post-Marketing-Hematological-MicroReport.pdf (https://dailyclout.io/wp-content/uploads/DC-V7combined-p1-2-Post-Marketing-Hematological-MicroReport.pdf)

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