COVID 19 Vaccines by Lot Number

Highly unusual data patterns

Evidence of corporate/government malfeasance

More Information at HOWBAD.INFO

What is this about:

- Not a science debate about VAERS analysis
- Not efficacy or safety debate (although related to safety overall)
- Evidence of malfeasance and willful misconduct by:
  - Pharma manufacturers
  - FDA, CDC other government health agencies
  - Local health agencies
  - Others – media, healthcare establishment, employers and schools, etc
- Vaccine manufacturing must be stopped, investigated and perpetrators prosecuted

C19 “Vaccines” are for NOBODY

- They are not safe – 21,000+ deaths, 1 million+ injuries, under-reported by 10-40x
- Have no efficacy nor positive risk/benefit ratio in any age or risk group, and overall show negative efficacy over time
- There is no notion of a “safe” lot number:
  - If a bag of candy contains some poisoned candies, the whole bag is poisoned
  - Even if a lot shows few adverse events, VAERS contains only short-term reports. Long-term issues such as cancer, prion disease, autoimmune disease, infertility, etc. are not captured
  - Vaccine Lots should be considered adulterated, containing toxic and super toxic substances

FDA Good Manufacturing Practices (GMP): 21CFR210.1

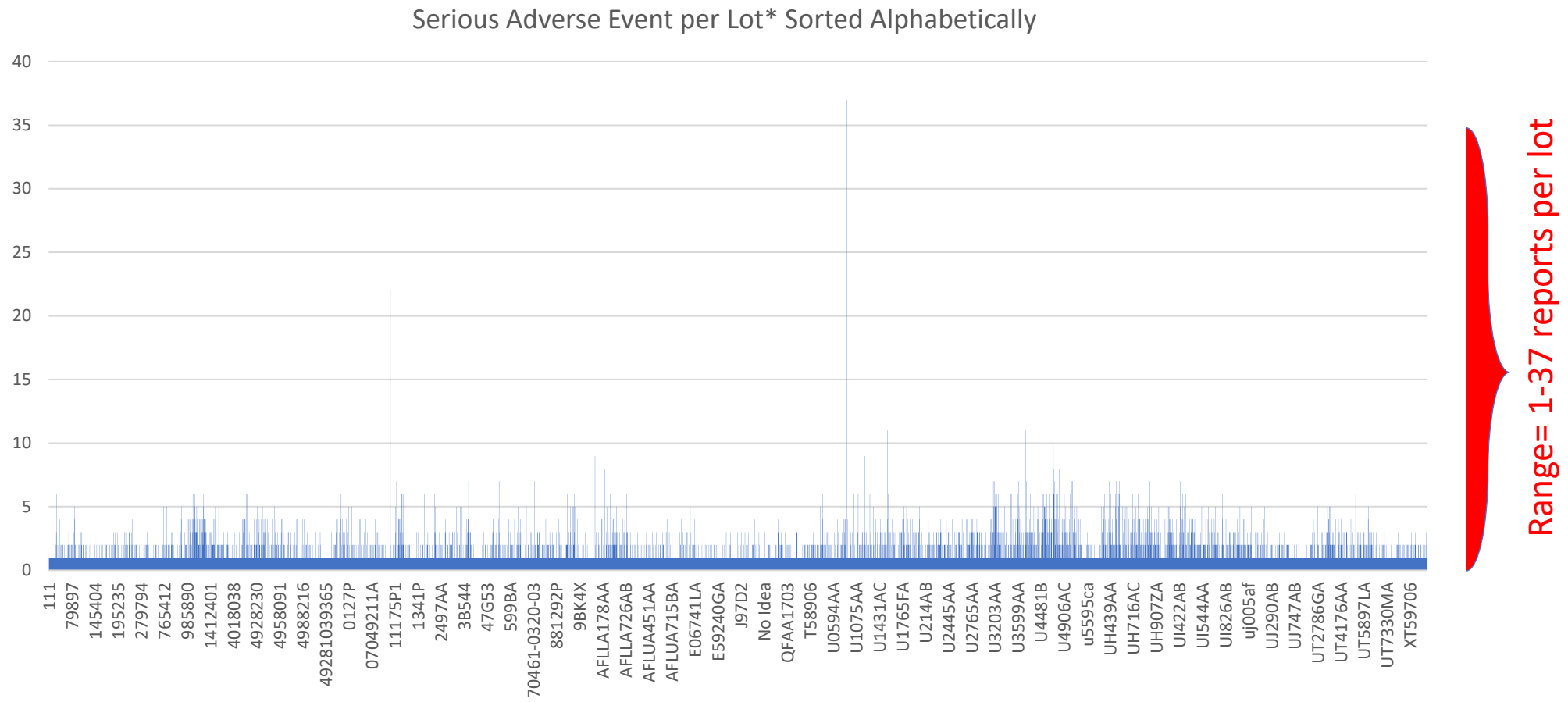
- High quality, consistency and purity standards for drugs/vaccines:
  - Expectation that every new lot/batch is “almost the same” as all previous lots
  - Expectation that vaccines from different manufacturers for a disease indication are “the same” or interchangeable product
- Current FDA GMP regulations developed after several adulterated products poisoned and killed 100s of people (early 1900’s to 1960’s)
- “The failure to comply …shall render such drug to be adulterated …shall be subject to regulatory action”.

VAERS Data: Covid 19 mRNA Vaccines vs Seasonal Flu Vaccines

Seasonal Flu Vaccines	Covid 19 mRNA Vaccines
9 injectable products	3 injectable products
10+ manufacturers	3 manufacturers (Pfizer, Moderna, Janssen)
10+ years of data	<12 months of data
22,334 manufacturing lots*	24,945 manufacturing lots*
9,429 Serious Adverse Events**	47,239 Serious Adverse Events**
906 deaths**	7,335 deaths**

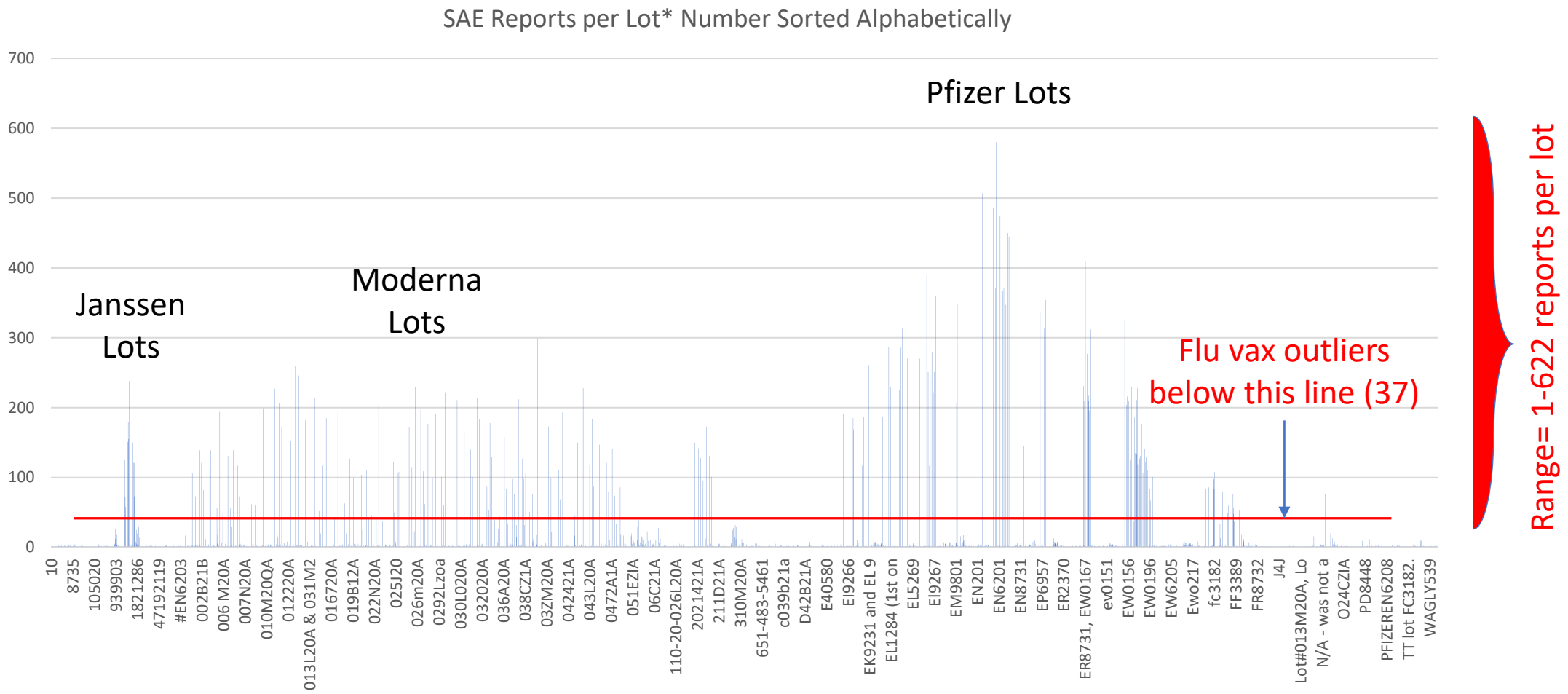
*Unvalidated – i.e. lots that are present in VAERS database, not complete list, contains typos. This does not matter when comparing two datasets that have the same error-prone data.
**Numbers of events reported to VAERS in each category

Flu Vaccines: Consistent product across many manufacturers, lots, years, only 2 outliers



*Includes lots with non-zero SAE Reports only, N= 5,797 Lot Numbers

Covid Vaccines: Does this look like the same consistent product by manufacturer and by lot?



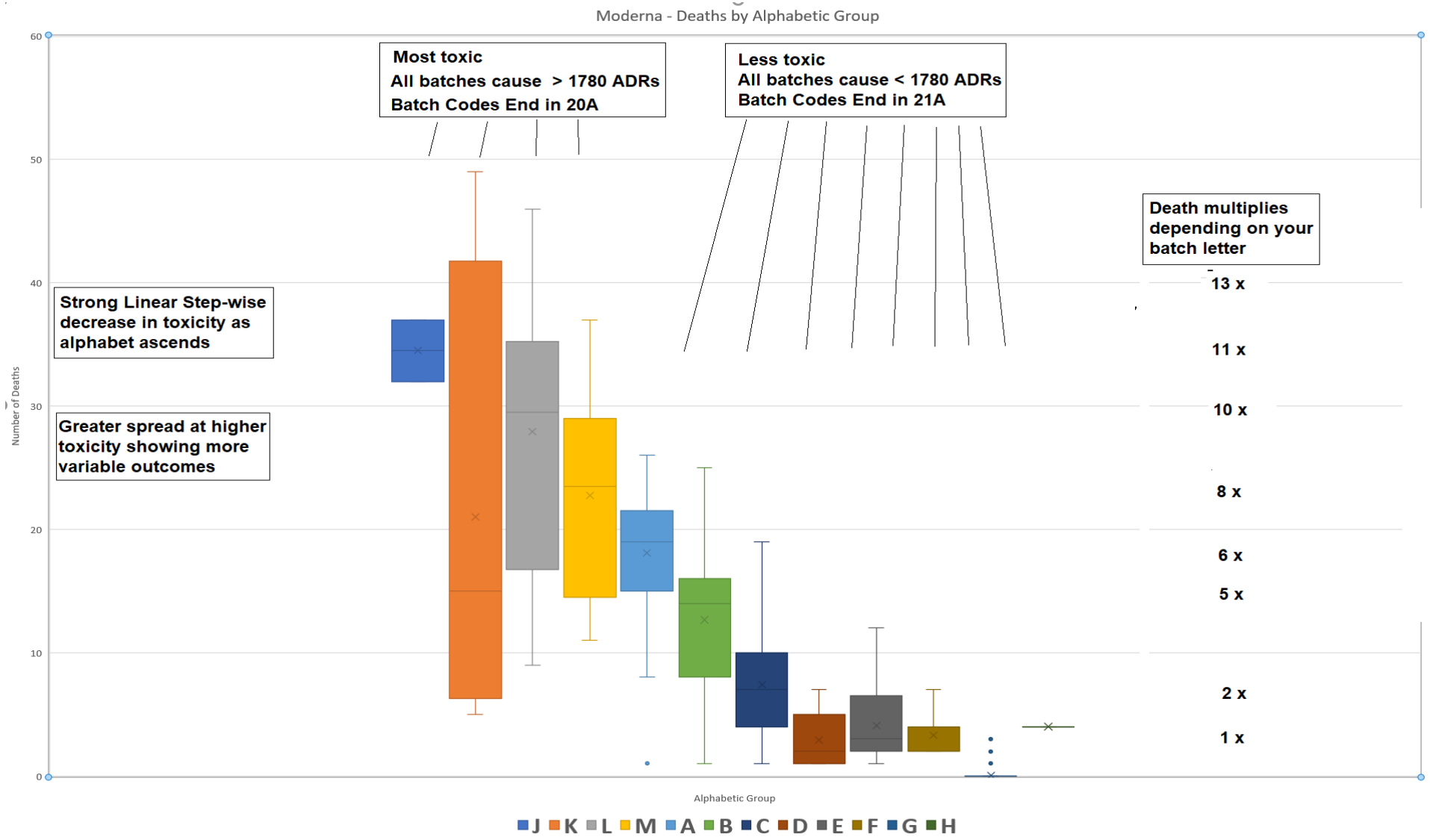
*Includes lots with non-zero SAE Reports only, N= 4,122 Lot Numbers

5-12 Times Higher Variability in C19 Vaccines Lot-to-Lot vs Control Dataset

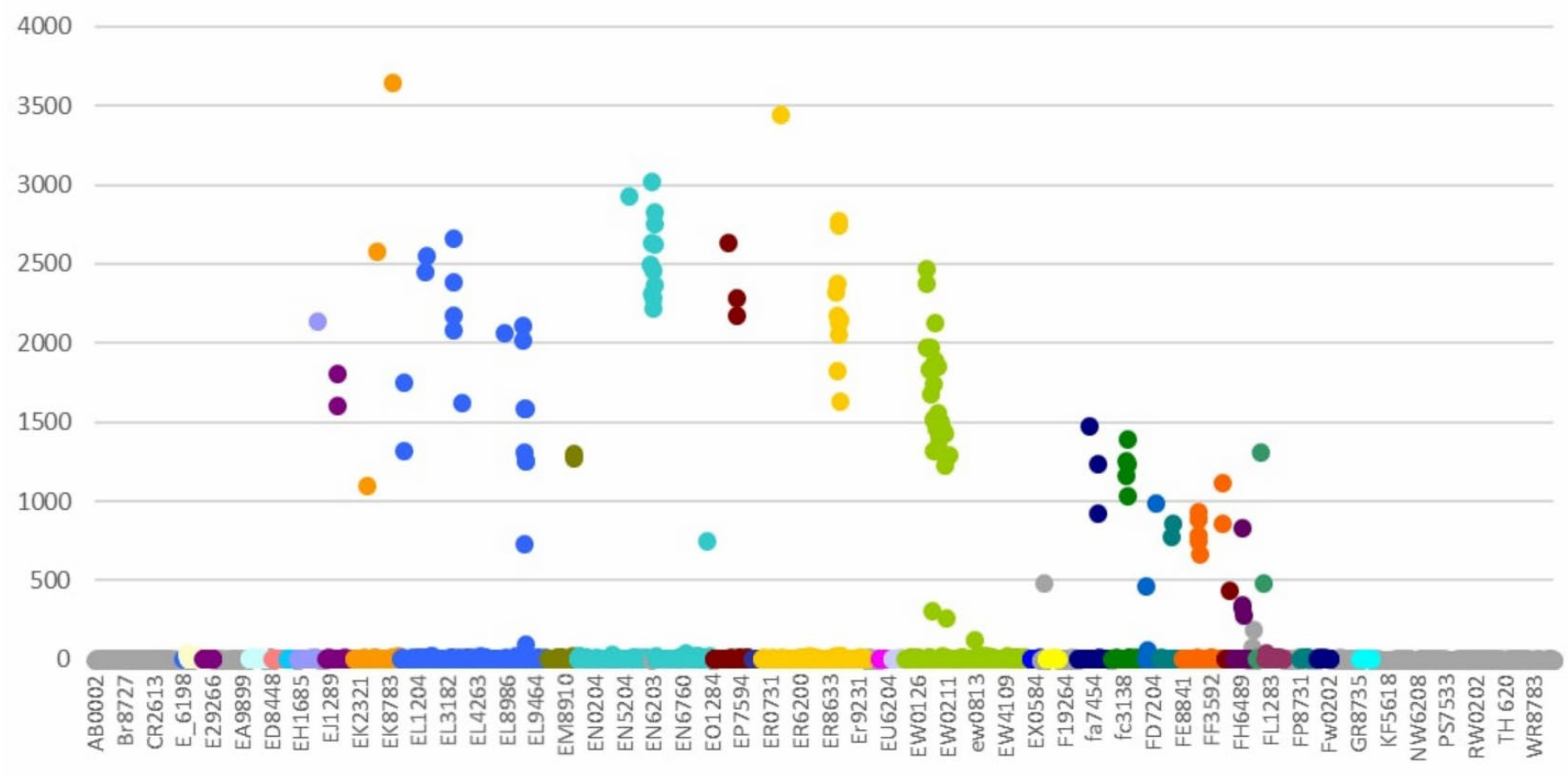
Coefficient of Variation for Each Dataset

Product Type	SAE Reports	Deaths
ALL COVID19 VACCINES	1019%	1195%
MODERNA ONLY	1005%	1159%
ALL FLU VACCINES	224%	99%

Evidence that Lots Are Labeled for Toxicity



Pfizer Batches
When Pfizer batch codes are arranged alpha-numerically along the x-axis, the following pattern appears -
Batches cluster alphanumerically, with the first 2 letters indication a different level of “toxicity”.



Possible Labeling Schema for C19 Vaccines

PFIZER

EN > EP

EP > ER

ER > EW

FA > FC

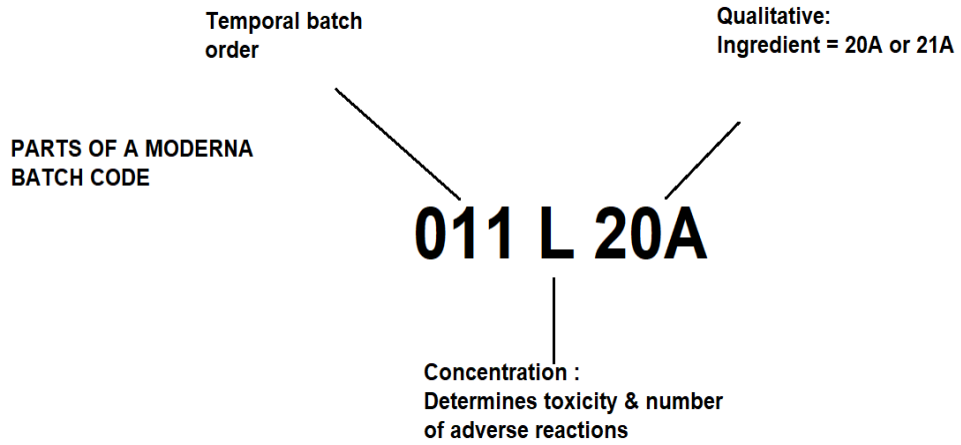
FC > FD

FD > FE

FE > FH

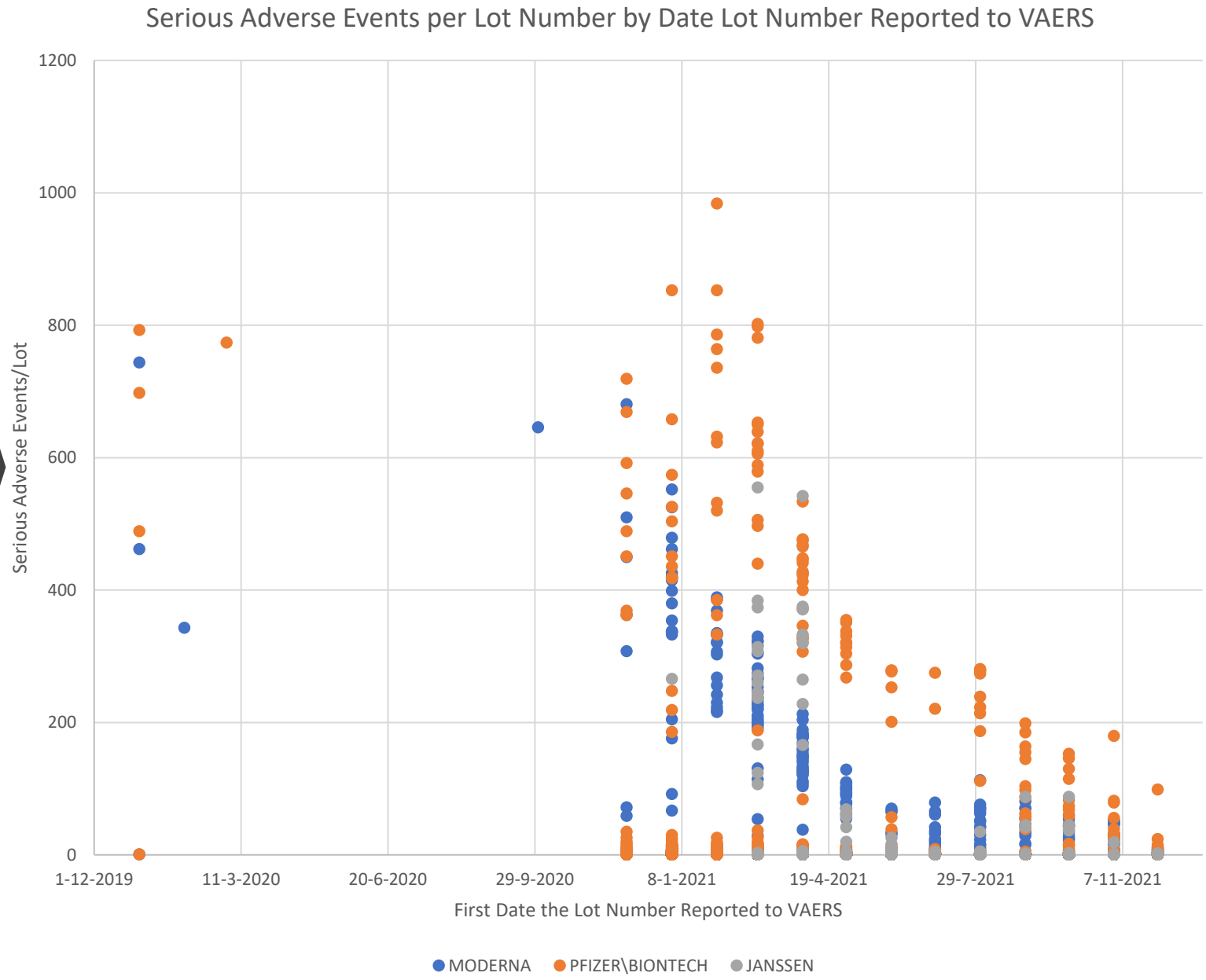
Batches cluster alphanumerically into specific ranges of toxicity

MODERNA

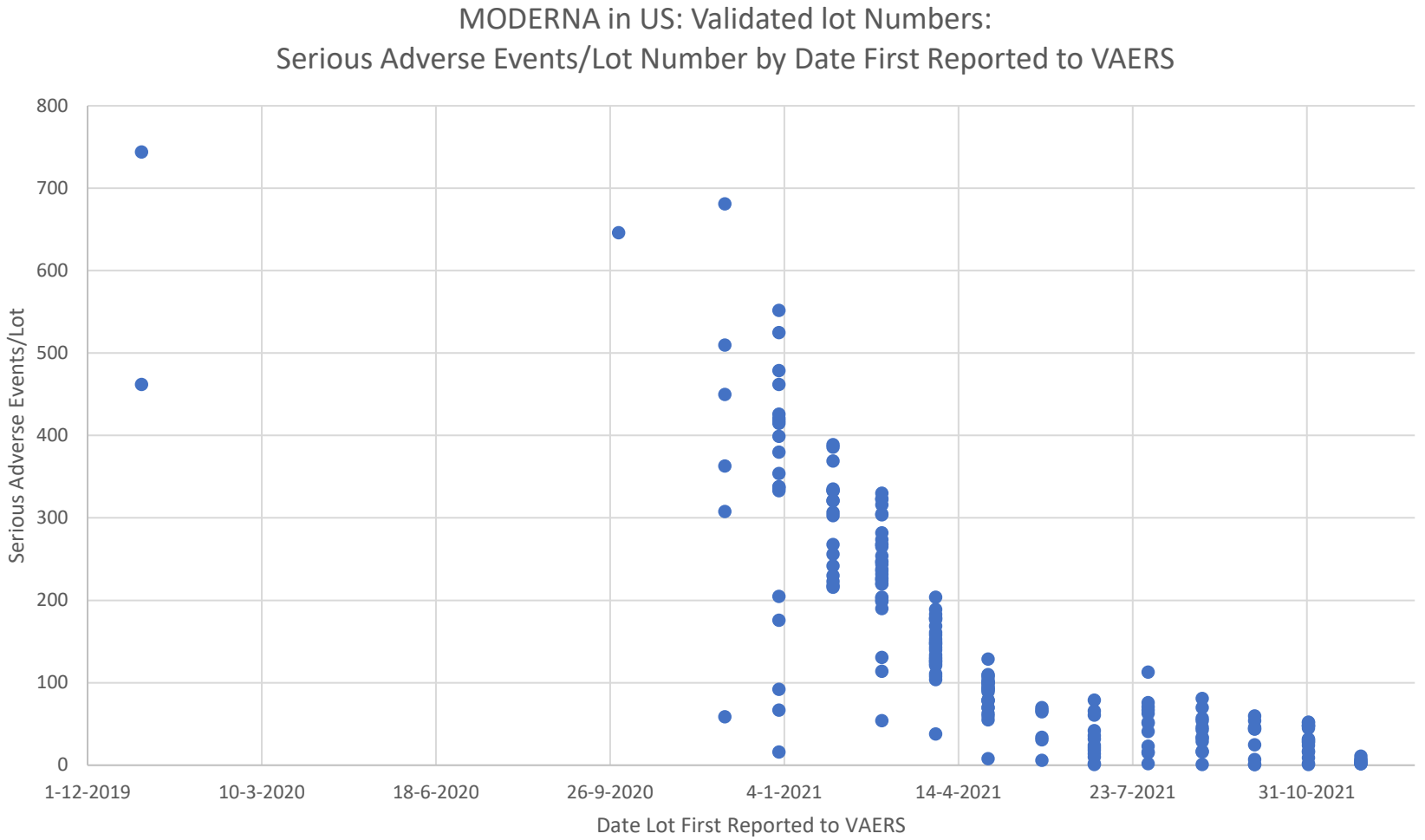


The last three characters of a Moderna batch code are usually either 20A or 21A. When all the Moderna batches are ranked in order of number of adverse reactions, it is found that the top 26 slots (the batches associated with most adverse reactions) are all batches ending in 20A. All of the Moderna batches associated with over 1780 adverse reactions end in 20A.

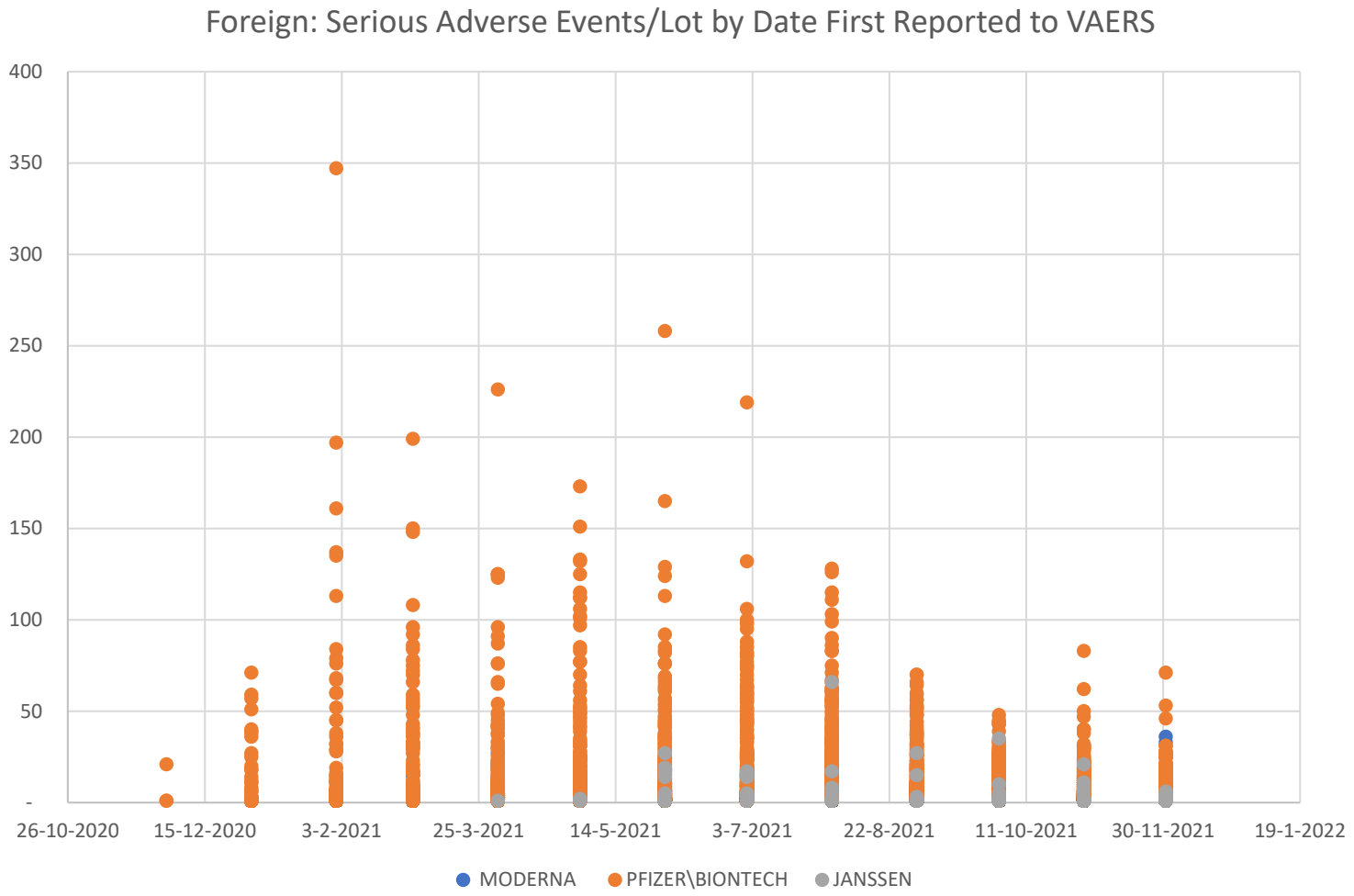
Time sequence of all lots looks unusual and cannot be explained by random variability



Same Pattern When ONLY Validated Lot Numbers are Used – Typos do not matter

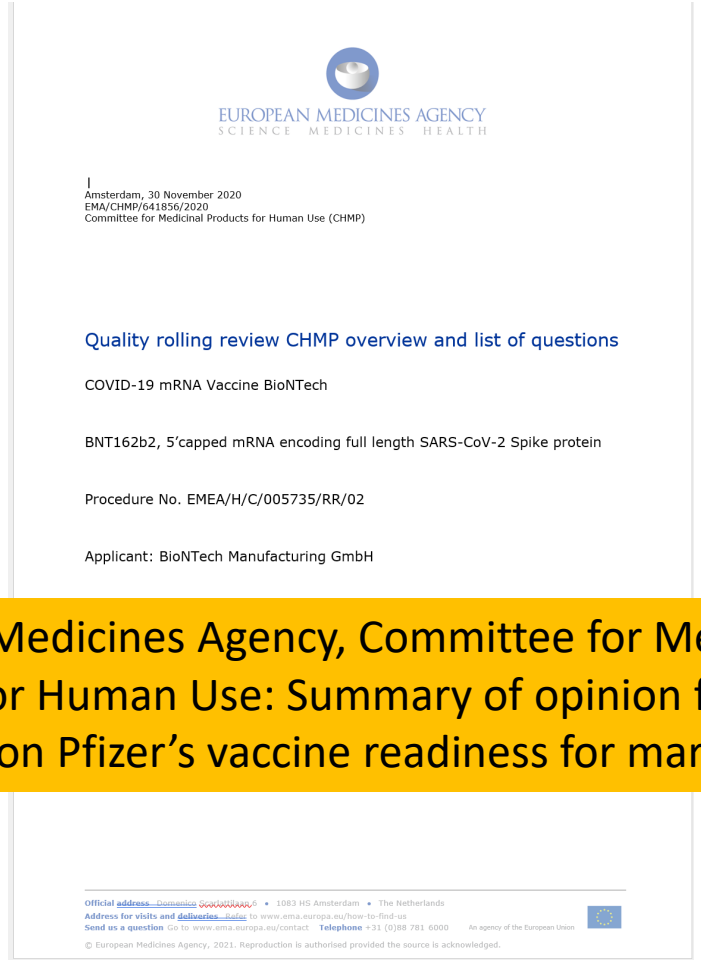


VAERS data ex-US: different pattern, why? Large variability still present.



Is It Willful/Premeditated?

Pfizer, FDA, and EMA Documents from End of November 2020



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Amsterdam, 30 November 2020
EMA/CHMP/641856/2020
Committee for Medicinal Products for Human Use (CHMP)

Quality rolling review CHMP overview and list of questions

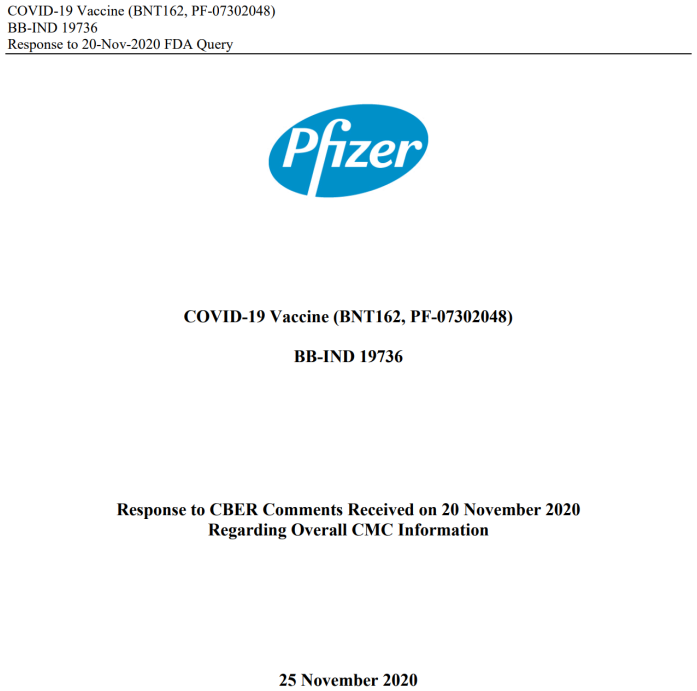
COVID-19 mRNA Vaccine BioNTech

BNT162b2, 5'capped mRNA encoding full length SARS-CoV-2 Spike protein

Procedure No. EMEA/H/C/005735/RR/02

Applicant: BioNTech Manufacturing GmbH

European Medicines Agency, Committee for Medicinal Products for Human Use: Summary of opinion from the regulators on Pfizer's vaccine readiness for market use



COVID-19 Vaccine (BNT162, PF-07302048)
BB-IND 19736
Response to 20-Nov-2020 FDA Query

Pfizer

COVID-19 Vaccine (BNT162, PF-07302048)

BB-IND 19736

Response to CBER Comments Received on 20 November 2020
Regarding Overall CMC Information

25 November 2020

Pfizer's responses to the FDA questions regarding manufacturing quality and compliance with current regulatory requirements

Opinion of the European Regulators:

- Pfizer/BioNTech manufacturing processes (at all sites) were NOT GMP compliant:
  - Not able to produce consistent product
  - No plan/deadline as to when they were going be in compliance
- 117 Major Objections and Concerns listed by the regulators on 30+ pages:
  - For any normal product, all would need to be resolved before authorization would happen

If non-GMP compliant production is allowed, ANY ingredient, process, quality control or packaging step of the vaccine can be manipulated, changed, and interfered with accidentally or on purpose!

33 Pfizer C19 Vaccine Lots for 28 million doses had been produced before EMA Opinion. They were shipped anyway.

Data From VAERS as of 12/2021

Lot (batch)	# Vials	Date manuf	All AE	Perm Disabil	Life Threat	Deaths
ED3938	19,010	16-Jul-20	-	-	-	-
EE3813	30,193	29-Jul-20	-	-	-	-
EE8492	67,665	5-Aug-20	656	123	9	2
EE8493	68,445	5-Aug-20	597	118	14	2
EG5411	201,258	3-Sep-20	-	-	-	-
EH9899	179,400	7-Oct-20	3,630	45	34	23
EH9978	304,869	23-Sep-20	-	-	-	-
EJ0553	164,580	25-Sep-20	476	74	19	20
EJ0701	200,265	26-Sep-20	-	-	-	-
EJ0724						8
EJ1685						4
EJ1686						8
EJ1688						2
EJ1691						-
EJ6795						7
EJ6796						5
EJ6797						3
EK1768						4
EK2808						-
EK4175						7
EK4176	131,625	16-Oct-20	1,339	39	26	34
EK4237	140,985	5-Nov-20	122	18	1	2
EK5730	191,295	22-Oct-20	4,102	37	40	24
EK9231	230,685	4-Nov-20	3,860	63	46	49
EL0140	155,610	29-Oct-20	1,856	24	28	61
EL0141	156,195	29-Oct-20	499	68	12	14
EL0142	138,060	29-Oct-20	1,802	28	36	42
EL0725	272,073	30-Oct-20	919	68	20	50
EL0739	294,239	3-Nov-20	1,023	131	33	19
EL1283	245,895	11-Nov-20	2,492	48	51	60
EL1284	214,305	17-Nov-20	2,790	40	34	45
EL1484	277,608	4-Nov-20	1,478	152	32	37
EL3246	204,360	19-Nov-20	2,417	54	24	43
Total	5,724,844		40,097	1,614	747	1,025

- 40,000+ Injuries
- 1,600+ Permanent Disabilities
- 747 Life Threatening Events (we don't know if later survived)
- 1,025 Deaths
- Includes 1 death of a child (5yo)

Unusual Patterns in the Data Point to Possibly Premeditated Acts

- C19 Vaccines are manufactured in NON-GMP compliant way:
  - C19 Vaccines should be deemed adulterated products, whether accidental or on purpose
  - Non-GMP compliance can be easily used to cover nefarious agenda
  - Evidence of willful misconduct as no recalls or investigations happened so far

- Manufacturers make assurances of product consistency and GMP compliance to FDA/EMA, in SEC filings, to public, Medicare/Medicaid:
  - Are these assurances fraudulent?
  - If manufacturers are exempt from GMP for covid vaccines – which specific aspects are waived?
  - Do patients and providers know this? Do investors and insurers know this?