C-19 Injections: Massive Regulatory and Manufacturing Fraud

Article by Tessa Lena

November 22, 2022

Video link

STORY AT-A-GLANCE

- Alexandra Latypova is an ex-pharmaceutical industry and biotech executive with a lot of experience in the areas of drug safety and clinical trials
- Early on, she discovered that, based on the number of reported deaths and adverse events per jab, there was unprecedented veritability in the history of the product
- According to Alexandra, the mRNA shots do not conform to their label specifications, and "in practice, both 'shots' and 'lethal' trials and anything is between is produced"
- Having analyzed massive amounts of publicly available data, as well as documents that became available as a result of FOIA requests and other sources, she has found strong evidence of manufacturing and regulatory fraud

I recently had the pleasure of interviewing Alexandra Latypova, an ex-pharmaceutical industry and biotech executive, who has been investigating and exposing manufacturing and regulatory fraud related to COVID injections.

We talked about the industry standards that were not adhered to during the clinical trials and the manufacturing of those injections, about the vial content quality testing procedures that had not been put in place, about the "hot batches" and their geographical distribution, about signs of fraud at every stage of testing and manufacturing the product, and about the general condition of living in a world run by a mob.

The latter was the lightest part of our conversation — evoking a lot of dark Eastern European humor — since both of us are Soviet expats, and in 2020, neither of us expected a whole lot of imagination to embrace the existential possibility of living in a world run by a mob. We had seen it in the past without a disguise — and when something looks like a duck, walks like a duck, and quacks like a duck, maybe it's just a duck!

Alexandra Latypova's Background

Alexandra grew up in Soviet Ukraine and immigrated to America in the late 1990s. She received her MBA from Fairmount College and the spent about twenty-five years in pharmaceutical industry and biotech (including in the areas of drug safety and clinical trials).

Alexandra has had a very gratifying entrepreneurial career. She has founded a number of successful startups, sold them — all before COVID — and retired, hoping to focus on enjoying her life and especially painting, which she does masterfully.

When 2020 knocked on the door with a whole bag of ugly and weird "new normal" treats, Sasha smelled the rat right away. Initially, she became alarmed by the abominations in "COVID response," including the conspicuous campaign to prevent effective treatment of COVID.

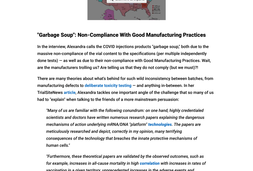
Compelled to understand what was going on, Alexandra got to work. She looked at VAERS and discovered huge discrepancies between the lots, where some batches had just a few reported severe adverse events, and some had over 1500 (she later learned from FDA documents that lot sizes were in a relatively similar range, and thus the discrepancies could not be explained by the lot size).

And when it comes to VAERS, let's not forget the 2010 Harvard Pilgrim study showing that VAERS was severely underreported — NOT overreported — capturing less than 1% of adverse events.

"Hot Batches"

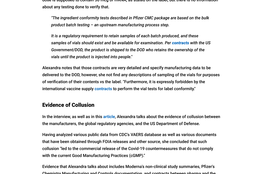
Early on, Alexandra discovered the existence of "hot batches." She is one of team members behind the famous "How Bad Is My Batch?" webpage where people can look up the number of severe adverse events reported to VAERS associated with a COVID injection batch. Other fearless members of the team are Dr Mike Yeadon, ex-head of Pfizer Respiratory Research, Jessica Rose with her blowing Psaschkovsian researcher, and Walter Kirkpatrick.

The slide below show the unprecedented veritability of serious adverse events and deaths in the U.S. per batch. Note the comparison to the veritability of the flu vaccine lots.



In the interview, Alexandra also mentioned the uneven distribution of deaths per a hundred thousand doses from batch to batch in the U.S. The coasts did much better than some of the Midwestern states that showed a very high numbers of reported deaths per a hundred thousand doses. The worst state is South Dakota (30+ reported deaths per 100,000 doses).

The areas that did even worse, according to Alexandra, were some of the U.S. territories with high percentage of indigenous population. (The latter data became available after the presentation was created, not reflected in the slide.)



"Garbage Swap" Non-Compliance With Good Manufacturing Practices

In the interview, Alexandra calls the COVID injections products "garbage swap," both due to the massive non-compliance of the vial content to the specifications (per multiple independently done tests) — as well as due to their non-compliance with Good Manufacturing Practices. Well, are the manufacturers trolling us? Are selling us that they do not comply (but we must)?!

There are many theories about what's behind for such wild inconsistency between batches, from manufacturing defects to deliberate toxicity testing — and anything in between. In her TrialSitenews article, Alexandra tackles one important angle of the challenge that so many of us had to "explain" when talking to the friends of a more mainstream persuasion:

> *Many of us are familiar with the toxicity conundrum: on one hand, highly credentialed scientists and doctors have written numerous research papers explaining the dangerous mechanisms of action underlying mRNA/DNA "platform" technologies. The papers are meticulously researched and depict, correctly in my opinion, the very real consequences of the technology that breaches the innate protective mechanisms of human cells.*
>
> *Furthermore, these theoretical papers are validated by the observed outcomes, such as for example, increases in all-cause mortality in high correlation with increases in rates of vaccination in a given territory, unprecedented increases in the adverse events and deaths recorded by various passive reporting systems, astonishingly high reports of the adverse events and deaths from the pharmacovigilance systems, and autopsy findings in vaccinated post-mortem showing the mechanisms of mRNA technology damage in histopathologic evaluations.*
>
> *On the other hand, many who have received the injections report no adverse effects and deem the points above a 'crazy conspiracy'.*
>
> *It is a regulatory requirement to retain samples of each batch produced, and these samples of vials should exist and be available for examination. The contract with the US Government/DOD, the product is shipped to the DOD who retains the ownership of the vials until the product is injected into people.*

Alexandra notes that these contracts are very detailed and specify manufacturing data to be delivered to the DOD; however, she not find any descriptions of sampling of the vials for purposes of verification of their contents in the label. Furthermore, it is expressly forbidden by the international vaccine supply contracts to perform the vial tests for label conformity!

Evidence of Collusion

In the interview, as well as in this article, Alexandra talks about the evidence of collusion between the manufacturers, the global regulatory agencies, and the US Department of Defense.

Having analyzed various public data from CDC's VAERS database as well as various documents that have been obtained through FOIA releases and other sources, Alexandra stated that data collusion "led to the commercial release of the Covid-19 countermeasures that do not comply with the current Good Manufacturing Practices (cGMP)."

Evidence that Alexandra talks about includes Moderna's non-clinical study summaries, Pfizer's Chemistry Manufacturing and Controls documentation, and contracts between pharma and the DOD for supply of the mRNA/DNA products. According to her, "it reveals disregard for established safety rules, regulations, and safety practices throughout the development, manufacture, and distribution of these products."

Moderna Bad Flags

As reported by Children's Health Defense, Alexandra reviewed 750 pages of documents that Moderna submitted to the FDA as part of its application process and obtained via a Freedom of Information Act request.

And according to her, "out of nearly 750 pages, about 400 pages are irrelevant studies that Moderna repeated multiple times. Moderna also submitted three versions of a single module, she said. And one module contained only narrative summaries of Moderna's studies but no actual study results." Alexandra's conclusion is that we are missing a large number of results, such as full reports that would support their narrative.

"The FDA (obviously did not object to any of this, she said. That's evidence of collusion to me with the manufacturer."

Other "abominations" that Alexandra highlighted both in the interview and in the Children's Health Defense article, were Moderna's clinical trials timeline and the fact that their product has two — not one — Investigational New Drug (IND) number.

Normally there is one IND application for one product. In this case, however, there are two IND applications — one belonging to Moderna, and one belonging to the National Institutes of Health, which partnered with Moderna on its COVID-19 vaccine.

"The Investigational New Drug (IND) application meeting is supposed to occur with the FDA when the company initiates human clinical trials. Moderna and the FDA had a pre-IND meeting on Feb. 19, 2020, and the IND application was formally opened the next day. The global pandemic was declared on March 11, 2020."

In the words of Alexandra, "Somehow these visionaries could predict the future with such certainty that they opened a clinical trial for the vaccine, for which a pandemic was announced a month later."

Pfizer Bad Flags

As Alexandra notes in her article titled, "Did Pfizer Perform Adequate Safety Testing for its Covid-19 mRNA Vaccine in Preclinical Studies? Evidence of Scientific and Regulatory Fraud," both the manufacturer and the regulators behaved in a highly dishonest manner and conspired to push an entirely novel technology and product on millions of people without carrying out a single well designed safety assessment."

For example, she points out that a review of clinical studies released by FDA uncovered that at least a different variants of active ingredient were included in the single Investigational New Drug application by Pfizer INDi19736:

- BNT162b1 — Uninoculated mRNA (uRNA, variant RBL063.3)
- BNT162b1 — Methylpseudouridine-modified mRNA (modRNA, variant RBP020.3)
- BNT162b2 — Methylpseudouridine-modified RNA (modRNA, variant RBP020.2)
- BNT162b2 — Self-amplifying uninoculated mRNA (saRNA, variant RBP020.1)

Alexandra writes that while the use of multiple versions of a product in the early stages of development is often inevitable, each chemical or biological entity is nevertheless deemed legally distinct for the purpose of product approval.

"Therefore, studies conducted with versions of the product that don't conform to the exact specification of the final version may serve only as supporting information for the approval of the latter, but they should never be deemed definitive and sufficient tests for claims of safety or efficacy pertaining to the final product."

She further mentions that in September 2021, the FDA issued a draft guidance entitled "Studying Multiple Versions of a Cellular or Gene Therapy Product in an Early-Phase Clinical Trial," which states that each version of product requires a separate IND application.

However, stunningly, "a footnote in this guidance exempts 'vaccines intended to prevent infectious diseases' from this requirement. No explanation is given as to why this exemption is made, and no convincing scientific or legal basis exists for this exemption, other than that the FDA had already arbitrarily abused this unprecedented deviation from the regulatory standard and later needed to cover their tracks.

In fact, arguably this regulatory 'exception' does not even apply to Pfizer's COVID-19 'vaccine' since the product does not prevent infection or transmission of the disease, but rather to prevent illness alone a sufficient condition? After all, every new drug is intended to do something like preventing an illness, but only few succeedfully do so."

Alexandra's article is very detailed, and I highly recommend reading it in full. You can also find Alexandra on TrialSitenews and on her Bitchute channel. To familiarize her take on Pfizer, she make the following points:

> *Pfizer's program did not include a comprehensive end-to-end test of all components of the final approved product (the mRNA COVID-19 vaccine). Instead, the studies included in the document package submitted to the FDA employed several variants and analogues of the product, whose comparability to the actual COVID-19 vaccine was not demonstrated or evaluated.*
>
> *Thus, no comprehensive assessment of product safety can be made on the basis of these studies.*
>
> *A key determinant of a drug's toxicity is its distribution within the body. However, with the mRNA active ingredient of Pfizer's COVID-19 vaccine, this crucial aspect was never studied!*
>
> *Pfizer claimed absence of potential for "vaccine-elicited disease enhancement" based on studies of an animal species that does not get sick from SARS-CoV-2.*
>
> *The CDC, the FDA and Pfizer all lied about "vaccine staying at the injection site," they knew all along that distribution of the vaccine throughout the body had to be expected.*
>
> *Pfizer skipped major categories of safety testing altogether.*
>
> *Pfizer used dishonest and self-serving interpretation of regulatory guidelines to justify the shortcuts it took in native safety testing.*
>
> *Both FDA and Pfizer knew about major toxicities associated with gene-therapy medicines in general, and they therefore cannot claim lack of anticipatory knowledge of these risks with the particular gene therapy medicine that is Pfizer's COVID-19 vaccine. This points to intentional fraud and collusion between Pfizer and the regulators, who conspired to push this untested dangerous product on the market.*

Yes, They Are Trolling Us — But We Are Not Helpless

Even though it is rather disheartening to know we live in a world that run by a mob, the challenge is centuries old, and remembering it can bring us much needed perspective and patience. The novel and "sudden" part of the challenge is that is happening to us, here and now, in broad daylight. That's shocking! But throughout history, many of our ancestors had to deal with pirats, and today, it is our turn to be brave. May our brave ancestors be our inspiration.

I would like to end this story with a short quote from my earlier article titled, "Is Our World Run Like a Mafia? So What Do We Do?"

"Good news: As the mafia bosses do their predatory thing, something mysterious is happening the hearts of those of us who resist or love. Under pressure, we are forced to remember that we are not theirs." We are not theirs. It is true.

About the Author

To find more of Tessa Lena's work, be sure to check out her bio, Tessa Fights Robots.

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Thank you Tessa Lena for your excellent research exposing that Alexandra Latypova found strong evidence of manufacturing and regulatory fraud on mRNA injections. A total fraud on the efficacy of "vaccines" and on their composition and adulot toxicity. The Fiss, Safe and Effective. A Second Opinion,"focuses primarily on the UK, but according to Lilcoke News, "one of its most powerful contributors is based in the United States and has contributed to numerous documentaries and articles, which have been distributed to the public regarding the effectiveness and potential harms of COVID vaccs." The documentary includes experts who can show how data from vaccine trials was flawed and how "vaccine" makers used figures, "very very misleading" to push injections.

The film seem ex to highlight the "shocking accusations" by clinical trial research specialist Alexandra Latypova, who studied Pfizer that contained "hot batches" and "cold batches" of the jab. Pfizer's Chemistry Manufacturing and Controls

www.youtube.com/watch?v=ZuaTbbbbbbc — Their findings include the following:— Pfizer completely skipped major categories of safety testing.— The active mRNA ingredient in COVID-19 vaccines was never studied for toxicity.— The FDA and Pfizer were aware of the major toxicities associated with class of gene therapy drugs.— The CDC, FDA, and Pfizer lied about "vaccines remaining at the injection site.— My review of leaked Moderna documents also revealed that vaccine-induced antibody-enhanced disease was identified as a serious risk. www.rfkjuniorsays.com/opinion/brilliant-new-documentary-exposes-hard-hitting

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In this article the German Working Group for the Analysis of Vaccines against COVID and international network of more than sixty scientists, doctors, mathematicians, lawyers and others who have committed themselves to carrying out in-depth laboratory analysis of so-called "vaccines" against covid 19 and to report their findings to the public present interesting information. Among the many topics they have covered are the contamination of the vials with DNA, the presence of SV40 promoters, the presence of Luciferase and Luciferase and the breakdown of people who have been injected with them. They have used scanning electron microscopy (SEM), energy dispersive X-ray spectroscopy (EDX), mass spectrometry (MS), inductively coupled plasma (ICP) analysis, lyophilized microscopy (LM), field microscopy (DFM) and Live Blood Diagnostics, as well as image analysis using Artificial Intelligence.

In the UK a conference of over 100 doctors MORE THAN 60 SCIENTISTS COLLABORATING BETWEEN MULTIPLE LABORATORIES IN

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SEVERAL NATIONS REVEAL WHAT IS REALLY IN THE INJECTIONS AND WHAT IS HAPPENING TO THE INJECTION VICTIMS' BLOOD. drtrozzi.org/.../whats-really-in-the-covid-vaccines (09/23/2022) A group of independent German scientists also found toxic components, mostly metals, in all the COVID vaccine samples they analyzed, "without exception", using modern medical and physical measurement techniques.

The COVID Vaccine Testing Task Force says some of the toxic elements found inside vials of vaccines from AstraZeneca, Pfizer and Moderna were not listed on manufacturers' ingredient lists. The findings "build on the work of other researchers in the international community who have reported similar findings, including Dr. Young, Dr. Nagase, Dr. Botha, Dr. Flemming, Dr. Robert Wakeling, and Dr. Noak," Dr. Janci Lindsay, Ph.D., a toxicologist who was not involved in the study, told The Epoch Times.

Mark as Spam · Posted On 11/22/2022 2:02:12 AM

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"The number and consistency of contamination allegations alone, coupled with the disturbing silence from global safety and regulatory agencies, is troubling and disconcerting in terms of 'transparency' and the continuing accusations by these agencies that vaccines genetics are 'safe'". Lindsey added. In the link the metallic elements found.
www.globalresearch.ca/unusual-toxic-components-found-covid-vaccines-wi... (08/22/2022).------- Dr. Jane Ruby told the public that large pharmaceutical companies Pfizer, Moderna, and Johnson & Johnson are rolling out toxic batches of Wuhan coronavirus vaccines. "There is evidence that these companies are intentionally rolling out toxic batches of the COVID-19 vaccine.

19 Ruby said she was told by a group of analysts that all three drug companies are actively testing live human subjects with these mass vaccination campaigns. She said analysts have identified the toxic batches by lot numbers, and it's an emergency to get the information out to people around the world. The team of analysts said these companies are conducting what are called "lethal dose studies," www.brighteon.com/cc47526e-aadb-46db-adc7-7372f2dcdc1f Ben Armstrong spoke with former Pfizer scientist Mike Yeadon on the January 10 episode of "The Ben Armstrong Show" about vaccine manufacturing.

They discussed how a large portion of reported adverse effects from Wuhan coronavirus vaccines came from less than 10 percent of released vaccine batches. Armstrong pointed out that pharmaceutical companies may have been using placebos and other things, and giving different servings or formulas for distribution of their vaccines because Israel, the UK and the US have worse adverse effects compared to other countries that are also using the vaccines. notaskhirzaman.com/6433
.-------www.brighteon.com/878da098-1f94-4e6a-815b-f2ffd9c304a9

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They are not injecting the same substance into each person. 5 percent of batches of each brand are responsible for 95 percent of deaths and injuries. A website allows you to check the toxicity of each batch according to official data.
Diariodevallarta.com/es/la-lotteria-de-vacunas-craig-paardekooper-estudio...
theresiouspandemic.net/2022/01/05/breaking-deadly-vax-lot-numbers-ident... .----- --
diariodevallarta.com/en/the-vaccine-lottery-craig-paardekooper-estudio... .---- One in two hundred batches has much higher levels of adverse reactions (hospitalizations, disabilities, miscarriages, birth defects, infertility, death, etc., ad nauseam.) By connecting with your link, the most satanic eugenics program is being implemented in the history of the human race.In this sense, British researchers say that the "vaccines" against Covid-19 do not provide any real protection against the latest "variant" of the disease, which the establishment calls "Omicron" A new paper published in the journal Science reveals that both antibody and T-cell protection are nowhere to be found after injection.

Meanwhile, the injections leave mysterious substances inside a person's body, often leading to chronic illness or early death. Another thing the article found is that the "fully vaccinated" are suffering from so-called "breakthrough" infections with Moronic, as their T-cell response is now artificially tuned toward earlier variants of the disease like Delta."when vaccinated people but previously uninfected suffer from Omicron infections, their T-cell response is biased towards earlier versions of Sars-Cov-2, not the Omicron variant that actually infects them alexberenson.substack.com/.../dont- read-this-if-youre-vaccinated," writes Alex Berenson on his Substack. alexberenson.substack.com/.../dont- read-this-if-youre-vaccinated (07/16/2022)

Mark as Spam · Posted On 11/22/2022 2:05:08 AM

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versatile
Joined On 10/16/2007 4:23:09 PM
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Frankly, we have better quality control mechanisms for lettuce, egg salad sandwiches, and pizza, than we have for big pharma products. If someone gets sick from a sandwich, it happens. But as soon as a number of people get sick, the epidemiologists step in and shut down the manufacturer, or the producer. The entire production chain is investigated, until the problem is addressed. However, when someone dies from a medical treatment, "Well, people die.' Cause of death? "They had a disease.' Investigation? "The product is patented. Intellectual property. It was scientifically tested. No investigation is needed."

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Jackaroni
Joined On 4/4/2009 8:32:35 AM
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I find it interesting that the most toxic batches of "vaccine" were concentrated in the more conservative areas of the country, places where people are more likely to be resistant to mandates. I strongly suspect that this is not by accident, which would then seem to indicate that not only was it known that the vaccines varied from lot to lot, but which lots were the most toxic and then to distribute them to the most "uncooperative" parts of the country. More proof that our government is trying to kill us. Truly frightening.

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brianallen1
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Fraud has been revealed. Now what?

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dejure
Joined On 8/23/2009 3:38:11 PM
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Keep in mind, unlike many other crimes, there is no statute of limitations on fraud.

Mark as Spam · Posted On 11/22/2022 5:17:20 PM

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MMaster
Joined On 8/14/2021 1:37:23 AM
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South Dakota with a high death count? Does that have something to do with REVENGE? Pharmaceutical companies going after the state that refused to lock down??? Sending the worst and most deadly batches there?

Reply · Mark as Spam · Posted On 11/22/2022 7:55:45 PM

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Kitera
Joined On 7/26/2007 6:24:19 PM
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As the narrative falls apart the true believers just double down and close their eyes to any evidence. They know in their hearts that authorities would never harm them. So the latest from FDA is that they didn't advertise Invermectin was bad and a house dewormer and needed to be banned - they merely didn't recommend it. And that is likely true. My guess is that their PR companies did the advertising and big pharma can not be blamed if health authorities can't tell the difference. If the health authorities are that stupid can big pharma be blamed? I can only comment on Canada. We paid a great deal of money for our health authorities and got "a ship of fools".

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san5129
Joined On 5/16/2021 4:10:10 PM
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Unfortunately, the most recent batch of releases from Pfizer (phmpt.org/pfizers-documents) uses a .xpt file, which cannot be easily opened or read, and I am a scientist! ...and they also changed ALL of the previous docs to the same file. ..If you didn't download the documents when they first came out, you will have a challenge now. I know some of these heroes are keeping tabs of what these companies are doing, but it is really sad that lay persons may not be able to open the files...for instance, the 'post-marketing' document with NINE pages of 'adverse events' listed in sentence form (AE 1, AE 2, AE 3...for nine pages.) Gui, you seem to be the wizardess at documents; let us know what you find.

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dividingcricker
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Lou321
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Label specification?....in the documentary film "Died Suddenly" that can be watched on Bitchute , there is no label specification brochure on the poisonous inyection, just a blank piece of paper.

Reply · Mark as Spam · Posted On 11/22/2022 5:41:30 AM

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san5129
Joined On 5/16/2021 4:10:10 PM
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howbadismybatch.info is where you will find info on the lots!

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