FAMILY FINANCIAL DISCLOSURE FORM FOR COVID-19 INJECTIONS

~ March 1, 2021 ~

Disclaimer: This form is provided to facilitate effective family due diligence, communication, and planning. It is essential that each person and each family take responsibility to identify and access the information they believe to be most relevant to their situation and decisions, and take responsibility to assess and manage their individual and collective risk as they believe best.
Introduction

The goal of this Family Financial Form is to ensure that an adverse event or death of one family member does not translate into long-lived or permanent financial destruction for the entire family.

This form was created to assist families to communicate regarding and to prepare for the family-wide financial impact of adverse events, if any, resulting from a Covid-19 injection. Examples of adverse events from Covid-19 injections include Covid-19 infection; anaphylaxis; neurological disorders; autoimmune disorders; other long-term chronic diseases; blindness and deafness; infertility, fetal damage, miscarriage, and stillbirth; and death (see Table 1 for examples of each).

Traditionally, informed consent forms for vaccination do not provide disclosure or statistics related to financial costs of possible injury, disability, or death, nor do they discuss the impact on family time, resources, health, and wealth—impacts that may include reduced career potential, divorce, and effects on siblings’ education and future plans.
Consequently, it is essential that prior to receiving a Covid-19 injection, parents and family members with financial responsibility for children and spouses not only perform thorough due diligence—providing adequate disclosures to their families regarding the potential costs to family members of adverse events or death—but also take steps to protect themselves and family members from the material adverse financial consequences of an adverse event or death.

**What Are the Covid-19 Injections?**

The leading Covid-19 injections currently in use in the U.S. and other OECD countries are experimental messenger RNA (mRNA) injections developed by Pfizer (with German partner BioNTech) and by biotechnology company Moderna (in partnership with the National Institute of Allergy and Infectious Diseases). The two experimental products are being distributed through emergency use authorizations (called “conditional marketing authorization” in the EU and “provisional approval” in Australia) granted following abbreviated clinical trials and without long-term safety testing. As yet, neither injection has received full approval or licensure from the FDA or any other national regulatory agency.

Though marketed as “vaccinations,” the Covid-19 mRNA injections are experimental gene therapy. Vaccine developers openly describe the never-before-authorized mRNA approach as a means of “programming a person’s cells”\(^1\) or, using Moderna’s terminology, deploying new “software.”\(^2\) In prior research, mRNA injections have displayed an intrinsic inflammatory component that has made it difficult to establish an “acceptable” risk/benefit profile.\(^3\)

The mRNA approach requires an in-built “gene delivery system” (also called a “carrier system”) to deliver the synthetic mRNA into the cells’ cytoplasm before the mRNA breaks down. The Pfizer and Moderna Covid-19 injections use lipid nanoparticles (LNPs) for this purpose; the LNPs not only shield the mRNA and promote cellular uptake but also function as adjuvants, “revving up” the immune system. Pfizer’s and Moderna’s LNPs are coated with polyethylene glycol (PEG), a synthetic, nondegradable, and controversial polymer associated with adverse immune responses. Moderna acknowledges that its LNPs “could lead to significant adverse events.”\(^4\) The FDA has
identified PEG as the possible culprit responsible for anaphylactic reactions to the Covid-19 injections.\(^5\)

In late February, the FDA authorized a third Covid-19 injection for emergency use in the U.S., manufactured by Johnson & Johnson’s Belgium-based pharmaceutical subsidiary, Janssen. J&J’s injection is an “adenovirus-vectored” vaccine that, like the mRNA injections, is intended to “trick” the cells into making coronavirus spike protein. The injection uses a genetically modified live common cold virus as a Trojan horse to “shuttle” spike protein DNA (genetic instructions) into human cells. In late 2019, the FDA approved an adenovirus-vectored Ebola injection, and the technology has also been featured in experimental—and problematic—Zika and HIV injections.\(^6\) The J&J Covid-19 injection is the first adenovirus-vectored injection to be authorized (on an emergency basis) for general population use.\(^7\)

In Europe, the EU has granted conditional marketing authorization to a different adenovirus-vectored Covid-19 injection—using an adenovirus that usually infects chimpanzees—developed by AstraZeneca and Oxford. AstraZeneca called several time-outs during its Covid-19 vaccine clinical trials because trial participants developed transverse myelitis, a condition that damages the insulating material around nerves\(^8\) and is associated with pain, muscle weakness, paralysis, and bowel and bladder problems; two-thirds of the individuals who experience it remain permanently disabled.

Dozens of other Covid-19 injections are under development, including RNA-based, DNA-based, and viral vector injections as well as injections using other technologies.

The World Health Organization (WHO) has granted emergency authorization to both the Pfizer and AstraZeneca Covid-19 injections, opening the door for the injections to begin rolling out in poorer countries.

**Adverse Event Reporting**

As these Covid-19 injections are new, adverse event reporting is still in its early stages. However, compilations of news accounts\(^9\) and reports submitted to national databases such as the Vaccine Adverse Event Reporting System
(VAERS) in the U.S.\textsuperscript{10,11} already provide representative information about the Covid-19 injections’ potential health impact. See Table 2 for a list of selected public databases that make reports of adverse events available to the public and/or are collecting reports of injuries and deaths following Covid-19 injection.

\textbf{From: [Adult Family Member]}

\textbf{To: [Adult Spouse and Children]}

\textbf{DUE DILIGENCE}

I have completed my due diligence on the Covid-19 injection that I propose to take.

- I have reviewed the available databases provided of material adverse events from Covid-19 injections, including deaths reported to date for people who have received these injections.

- I understand that this Covid-19 injection is being distributed under an emergency use authorization and that it has not been approved by [FDA/national regulatory agency].

- I understand that this Covid-19 injection is made by:
  - Moderna – a company that in 10 years had never brought a single product to market prior to the coronavirus vaccine\textsuperscript{12}
  - Pfizer – a company with a demonstrated history of enforcement settlements for fraudulent marketing\textsuperscript{13}
  - Johnson & Johnson – a company with a demonstrated record of health care fraud\textsuperscript{14}
  - AstraZeneca – a company that paid one of the top 10 pharmaceutical company settlements ever\textsuperscript{15}

- If Pfizer or Moderna: I understand that this Covid-19 injection is an experimental gene therapy.
___ I understand that the injection has only been designed to protect against moderate symptoms of Covid-19 and that it may not protect me from more severe symptoms.16

___ I understand that Covid-19 injections may not protect against transmission.17

___ I understand that by agreeing to this injection, I may be required to take further Covid-19 injections as indicated by the manufacturer’s protocol or requirements, including potential annual “booster shots.”18

___ I understand that this Covid-19 injection is not designed to address mutating versions or additional variants of the coronavirus.

___ I have attached a copy of the manufacturer’s fact sheet [traditionally called a package insert] for the Covid-19 injection, which states that the injections are not FDA-approved, describes the ingredients, outlines potential material adverse events, and acknowledges that not all risks are known. I am willing, able, and available to review and explain the fact sheet to you.

Moderna: https://www.fda.gov/media/144638/download

Pfizer: https://www.fda.gov/media/144414/download


___ I also understand that prior to Covid-19 injection, health care providers are legally required to communicate information “consistent with the fact sheets” to patients and either provide a hard copy or direct patients to the appropriate website.
I understand that because some of the ingredients of these Covid-19 injections are proprietary and may, therefore, be secret, the ingredients listed in the manufacturers’ fact sheets may be incomplete. I also understand that prior research on other vaccines has demonstrated the presence of nanoparticles, heavy metals, fetal tissue, and other substances not disclosed (or not fully disclosed) in “vaccinations.”

[U.S. only] I understand that under the 1986 National Childhood Vaccine Injury Act and the 2005 PREP Act, it will be difficult if not impossible to hold the manufacturer of this Covid-19 injection responsible for any damage to my health or death resulting from this injection.

[Non-U.S.] I have reviewed the policies or agreements in place in my country regarding indemnification and compensation; I understand that depending on these policies or agreements, it may be difficult if not impossible to hold the manufacturer of this Covid-19 injection responsible for any damage to my health or for death resulting from this injection.

I understand that it will be difficult if not impossible to hold the health institutions, doctors, and nurses that distributed this Covid-19 injection to me responsible for any damage to my health or for my death.

I understand that it will be difficult if not impossible to hold federal, state, and local health care officials and regulators responsible for any damage to my health from the Covid-19 injection or for my death.

I understand, therefore, as a practical matter that I and my closest relatives will experience and shoulder the full cost in terms of time and money of any Financial adverse impact of a material adverse event resulting from my taking this Covid-19 injection.

For Families Planning on Having Additional Children

Pfizer and Moderna Covid-19 injections: I understand that this injection has the potential to alter my DNA in ways that no one yet understands and that this injection could alter the DNA of my unborn children or any woman who carries my unborn children.
_____ Pfizer and Moderna Covid-19 injections: I understand that knowledgeable experts have shared serious concerns in a petition filed with the European Medicines Agency that components of the Covid-19 mRNA injections could trigger an immune reaction against syncytin-1, a protein responsible for development of the placenta and essential for a successful pregnancy, resulting in potential infertility.²¹

_____ Johnson & Johnson Covid-19 injection: I understand that this injection is produced in genetically modified human embryonic retinal cells (PER.C6 TetR) and that the presence of fetal DNA fragment contaminants in injections has been linked to autism spectrum disorder.²²

_____ AstraZeneca Covid-19 injection: I understand that this injection is produced in genetically modified human embryonic kidney cells (HEK 293).

_____ My spouse has agreed to assume the risks of such alterations of my DNA and any impact it may have on our ability to have children or on our unborn children.

Material Adverse Events

_____ I understand and have reviewed the material adverse events reported in connection with the Covid-19 injections. Known adverse events include Covid-19 infection; anaphylaxis; neurological disorders; autoimmune disorders; other long-term chronic diseases; blindness and deafness; infertility, fetal damage, miscarriage, and stillbirth; and death (see Table 1 for examples of each; see also endnote #9).

Reasons for Taking Injection

_____ I understand that Covid-19 has a statistical probability of death²³ of 0.003% for youth (ages 0-19), 0.02% for those ages 20-49, 0.5% for individuals aged 50-69, and 5.4% for seniors age 70 and older.²⁴

_____ I also understand that there are multiple, low-cost, non-injection therapeutic drug protocols for early intervention and prophylaxis that have a high rate of success in helping defend against or recover from Covid-19.²⁵
Nevertheless, I want to take these Covid-19 injections. The reason(s) why is (are):

_______________________________________________________________
_______________________________________________________________
_______________________________________________________________

HEALTH CARE

Due to the difficulties of accessing the appropriate care in an emergency, I have identified and arranged health care providers who will be available on a timely basis in the event of a material adverse event from the Covid-19 injection:

Covid-19 Infection

Contact Info:

Anaphylaxis

Contact Info:

Neurological Disorders

Contact Info:

Autoimmune Disorders

Contact Info:

Other Long-term Chronic Diseases

Contact Info:

Blindness and Deafness:

Contact Info:

Infertility, Fetal Damage, Miscarriage, and Stillbirth (women only)

Contact Info:
HEALTH CARE PROXY

____ I have reviewed the Aging with Dignity planning process and filled out the Five Wishes planning form (https://agingwithdignity.org and https://fivewishes.org) and have provided a Health Care Proxy to you along with detailed instructions on resuscitation and extreme measures at end of life.

____ I have reviewed this form with the following people who have authority in my Health Care Proxy and have agreed to assume responsibility in the event of a material adverse event or death resulting from the Covid-19 injection.

[List Here] ____________________ ____________________
____________________

INSURANCE
In the event of a material adverse event from Covid-19 injection, my health care insurance ___ will cover ___ will not cover all health care and hospitalization expenses.

My insurance broker has confirmed that page ___ of my ____ policy states that taking an experimental or emergency-use Covid-19 injection ___ will ___ will not impact my insurance coverage.

In the event that I am unable to work for a period of time or lose my job, profession, or business, my disability insurance will cover the following amounts for ___ months/years:

[__________] 

I have reviewed my decision with my insurance broker and additional health care and disability or other insurance ___ is available ___ is not available to cover any material adverse event from a Covid-19 injection on the following basis:

[_________________________]

In the event of my death from Covid-19 injection, my life insurance will provide the following protection to you:
I have reviewed my decision with my insurance broker and additional life insurance ___ is available ___ is not available to cover any material adverse event or death from Covid-19 injection on the following basis:  

___ I have provided sufficient time and resources for my family and I to arrange for other available insurance that my family members and I believe are prudent.

**FINANCIAL INVESTMENT**

**Loss of Income**

In the event of a material adverse event from Covid-19 injection, the potential range in the loss of income is estimated to be [provide range if unable to work for 1 year, 5 years, or permanently]:

*Covid-19 Infection:*

_________________________________________________

*Anaphylaxis:*

_________________________________________________

*Neurological Disorder:*

_________________________________________________

*Autoimmune Disorder:*

_________________________________________________

*Other Long-term Chronic Disease:*\(^{27}\)

_________________________________________________

*Infertility, Fetal Damage,*

*Miscarriage, Stillbirth* *(women only):*

_________________________________________________
Health Care Expenses

In the event of a material adverse event from Covid-19 injection, the potential range of health care expenses not covered by our health care insurance is estimated to be [estimate potential expenses for 1 year, 5 years, or long-term]:

Covid-19 Infection:

_________________________________________________

Anaphylaxis: _______________________________________

Neurological Disorder:

_________________________________________________

Autoimmune Disorder:

_________________________________________________

Other Long-term Chronic Disease: 28

_________________________________________________

Infertility, Fetal Damage, Miscarriage, Stillbirth
(women only): _______________________________________

Death: ______________________________________________

Chronic Disease: ______________________________________

Blindness & Deafness:

_________________________________________________

Long-Term Care
If a material adverse event from a Covid-19 injection results in the need for long-term care, this is how I propose to arrange such care and fund it:

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**Investment of Family Time**

In the event of a material adverse event from a Covid-19 injection, here is the time I would request from my family or professional caregivers paid by my family to assist me:

*Covid-19 Infection:*

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*Anaphylaxis:*

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*Neurological Disorder:*

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*Autoimmune Disorder:*

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*Other Long-term Chronic Disease:*

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*Blindness & Deafness:*

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*Infertility, Fetal Damage, Miscarriage, Stillbirth* (women only):

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*Death:*

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**Proposed Sources of Financial Support**
If a material adverse event from a Covid-19 injection results in adverse financial events—loss of income and/or increased expenses—these are my estimates of costs and my arrangements to fund them:

*Covid-19 Infection:*
_________________________________________________

*Anaphylaxis:*
_________________________________________________

*Neurological Disorder:*
_________________________________________________

*Autoimmune Disorder:*
_________________________________________________

*Other Long-term Chronic Disease:*
_________________________________________________

*Blindness & Deafness:*
_________________________________________________

*Infertility, Fetal Damage, Miscarriage, Stillbirth (women only):*  
_________________________________________________

*Death:*
_________________________________________________

**DEATH**

____ In the event of my death from a Covid-19 injection, I have finalized an estate plan, have reviewed it with my attorney and executor, and have provided instructions for my funeral and disposition of my remains as follows:
_________________________________________________

Having completed my due diligence on the Covid-19 injection and having made my decision to proceed, I am available to review my findings and arrangements with my family.
I am responsible for my health care choices and am committed to taking responsibility for the true costs of my choices and their impact on those I love and not shifting these costs and risks to them without their full knowledge, due diligence, and consent.

Please let me know when you would like to review and discuss. Your loving [spouse/parent]

Signed

Date:

TABLE 1. Examples of Adverse Events Reported Following Covid-19 Injections

Note: For further examples, see endnote #9.

Covid-19 Infection

Examples:


2. 4,500 people diagnosed with COVID after getting 1st vaccine dose (January 12, 2021)

3. Informed consent disclosure to vaccine trial subjects of risk of COVID-19 vaccines worsening clinical disease


**Anaphylaxis**

*Examples:*

4. VAERS COVID reports

https://www.openvaers.com/covid-data
5. One-third of deaths reported to CDC after COVID vaccines occurred within 48 hours of vaccination (VAERS data through February 12, 2021)

6. COVID vaccine injury reports grow in number, but trends remain consistent (March 5, 2021)
https://childrenshealthdefense.org/defender/vaers-covid-vaccine-injury-reports-increase/


8. Metro healthcare worker describes severe allergic reaction to COVID-19 vaccine (January 14, 2021)

9. California shelves batch of Moderna vaccines over allergy concerns (January 18, 2021)
https://nypost.com/2021/01/18/california-shelves-batch-of-vaccines-over-allergy-concerns/

Neurological Disorders

Examples:

10. I took the Moderna mRNA vaccine without doing my own research (February 19, 2021)
https://www.bitchute.com/video/1lyFggcw9P9c/

11. Louisiana woman convulsing after Pfizer experimental Covid vaccine (January 14, 2021)
http://www.bitchute.com/video/YoJLKtbksyKv/

12. The intertwined history of myelitis and vaccines [AstraZeneca/Oxford] (September 25, 2020)


Autoimmune Disorders

Examples:

13. Professor and molecular genetics expert Dolores Cahill expects autoimmune reactions to result from the mRNA injections “months later,” stating that “this gene therapy...is setting up an autoimmune disease chronically.”

14. Immunologist Bart Classen—who has been issuing warnings for years about chronic, late-occurring adverse events from vaccination—is concerned that the mRNA injections could create new mechanisms of vaccine adverse events. Classen notes that adverse events such as type 1 diabetes “may not occur until 3-4 years after a vaccine is administered.”

15. Are we on the verge of a “super-epidemic” of autoimmune diseases? (February 19, 2021)
16. Vaccination and autoimmune diseases: is prevention of adverse health effects on the horizon? (July 20, 2017)  
https://link.springer.com/article/10.1007/s13167-017-0101-y


**Long-term Chronic Disease**

*Examples:*

18. Immunologist Bart Classen—who has been issuing warnings for years about chronic, late-occurring adverse events from vaccination—is concerned that the mRNA injections could create new mechanisms of vaccine adverse events. Classen notes that adverse events such as type 1 diabetes “may not occur until 3-4 years after a vaccine is administered.”


**Blindness and Deafness**

*Example:*

20. United Kingdom: 12 people deaf, five blind after Pfizer BioNTech mRNA shots (February 11, 2021)  
https://thecovidblog.com/2021/02/16/united-kingdom-12-deaf-five-blind-after-pfizer-mrna-shots/

**Infertility, Fetal Damage, Miscarriage, and Stillbirth (women only)**
Examples:

21. Eight unborn babies dead soon after their mothers received COVID-19 vaccine (as of January 22, 2021)


23. Health officials push pregnant women to get COVID shots, despite known risks (February 23, 2021)

https://childrenshealthdefense.org/defender/health-offcials-push-pregnant-women-covid-vaccine/?itm_term=home

Death

Examples:


28. One-third of deaths reported to CDC after COVID vaccines occurred within 48 hours of vaccination (VAERS reports through February 12, 2021) [https://childrenshealthdefense.org/defender/latest-data-cdc-vaers/](https://childrenshealthdefense.org/defender/latest-data-cdc-vaers/)


30. UK data show 402 reports of deaths following COVID vaccines (March 2, 2021)

[https://childrenshealthdefense.org/defender/uk-data-deaths-following-covid-vaccines/](https://childrenshealthdefense.org/defender/uk-data-deaths-following-covid-vaccines/)

**TABLE 2. Public Databases Compiling Reports of Adverse Events**

**UK Medical Freedom Alliance (UKMFA) (global)**

*See UKMFA Collated Data – Adverse Reactions Summary document*

[https://www.ukmedfreedom.org/resources/covid-19-vaccine-info](https://www.ukmedfreedom.org/resources/covid-19-vaccine-info)

**Vaccine Adverse Event Reporting System (VAERS), U.S. Department of Health and Human Services (United States)**

The Vaccine Adverse Event Reporting System (VAERS) is a highly flawed voluntary reporting system co-administered by the CDC and FDA. Dr. Sherri Tenpenny estimates that reporting of adverse events to VAERS represents approximately 10% of actual adverse events. A Harvard study commissioned in 2010 by the federal government produced an even lower estimate, stating that less than 1% of adverse events get reported: [https://digital.ahrq.gov/sites/default/files/docs/publication/r18hs017045-lazarus-final-report-2011.pdf](https://digital.ahrq.gov/sites/default/files/docs/publication/r18hs017045-lazarus-final-report-2011.pdf).
VAERS reports are accessible through the CDC Wonder search engine or through the more user-friendly MedAlerts search engine maintained by the National Vaccine Information Center (NVIC). MedAlerts also offers more powerful search capabilities and more extensive reporting.
https://vaers.hhs.gov
https://wonder.cdc.gov/vaers.html
https://www.medalerts.org/

VaxxTracker

VaxxTracker describes itself as a “repository for vaccine adverse effects.” The site collects adverse event reports and provides data summaries with monthly counts, top 10 reaction types, outcomes, descriptions of symptoms, and more.
https://www.vaxxtracker.com/VaxDefault.asp

Children’s Health Defense

Children’s Health Defense encourages those who experience vaccine injuries to report the adverse event to VAERS and vaxxtracker.com as well as to Children’s Health Defense. https://childrenshealthdefense.org/defender/injured-by-vaccine-how-to-report-it/

The Informed Consent Action Network (ICAN) also collects reports of Covid-19 vaccine injuries. Recognizing that “Potential safety issues may not be disclosed or fully disclosed to individuals receiving these experimental products,” ICAN can “assist in investigating whether [individuals] have been adequately warned of the potential injury.”

EudraVigilance

The European Medicines Agency maintains a database of suspected adverse drug reactions that is accessible to the general public (see box for “Healthcare professionals, patients and the general public”). Under the letter “C,” Covid-19 reports can be generated for the Moderna, Pfizer, and AstraZeneca injections. As of February 23, 2021, EudraVigilance had received over 71,000 suspected

ENDNOTES:


3 https://www.nature.com/articles/s41541-020-0159-8.


7 An informative description of the FDA’s “lackluster” emergency use authorization proceedings for the J&J injection is available here: https://anthraxvaccine.blogspot.com/2021/02/j-and-js-vaccine-actually-manufactured.html. For a basic comparison of mRNA and adenovirus vector injections, see: https://news.ncsu.edu/2020/12/vaccines-koci-101/.


9 A sample of news reports:

10/26/20, South Korea: Singapore halts use of flu vaccines after 48 die in South Korea https://www.telegraph.co.uk/news/2020/10/26/singapore-halts-use-flu-vaccines-48-die-south-korea/
12/27/20, Germany, Ansbach: Vaccination in Darmstadt-Dieburg district begins in Asbach – Number of Corona deaths in Darmstadt-Dieburg explodes


01/01/21, Israel: 4 people died and 240 got COVID19 in Israel after being injected with Pfizer experimental mRNA vaccine


01/04/21, Iceland: 3 deaths after receiving Covid19 vaccine in Iceland

https://www.mbl.is/frettir/innlent/2021/01/04/thrir_nu_latist_eftir_bolusetningu_her_a_landin/

01/04/21, Portugal: “Perfectly healthy” 41-year-old pediatric assistant dies suddenly after injected with experimental Pfizer COVID vaccine


01/05/21, Israel: Israeli woman diagnosed with Bell’s palsy after taking Covid-19 vaccination


01/05/21, Canada: 27-year-old Canadian healthcare worker faints and suffers multiple seizures after Pfizer experimental COVID vaccine


01/05/21, USA: Hundreds sent to emergency room after receiving COVID-19 vaccines


01/05/21, Mexico: 32-year-old Mexican doctor suffers seizures and is paralyzed after receiving the Pfizer experimental vaccine


01/06/21, Norway: Norway investigating death of two people who received Pfizer’s coronavirus vaccine


01/07/21, USA: “Very healthy 56-year-old” Miami obstetrician dies after being injected with the experimental Pfizer COVID vaccine
01/07/21, USA: CDC reveals at least 21 Americans have suffered life threatening allergic reactions to Pfizer’s COVID vaccine

https://www.dailymail.co.uk/health/article-9119029/At-21-Americans-life-threatening-anaphylaxis-receiving-Pfizers-vaccine-CDC-reveals.html

01/08/21, Norway: 82-year-old resident at Sola nursing home dies one day after being vaccinated against COVID-19


01/10/21, Israel: Young man develops “rare life-threatening syndrome” after Covid-19 vaccine

https://www.israelnationalnews.com/News/News.aspx/294606

01/10/21, Germany: 10 dead in Germany within 4 days of Covid-19 vaccine inoculation; probe ordered


01/11/21, USA: Nurse develops facial paralysis after Covid-19 vaccination


01/11/21, India: 42-year-old Indian dies after vaccination


01/12/21, USA: 24 Corona deaths in New York nursing home after vaccination


01/12/21, Germany: 55-year-old dead 10 days after vaccination

https://static.wixstatic.com/media/252392_5506fea41cdd49bbb714a636b9ddc602~mv2.png/v1/fill/w_600,h_180,al_c,q_85,usm_0.66_1.00_0.01/Bildschirmfoto%202021-01-22%20um%2014_22_08.png.webp

01/13/21, Israel: Corona explosion after vaccination

http://www.wochenblick.at/nach-impfkampagne-explodieren-in-israel-die-corona-zahlen/

01/13/21, Israel: 75-year-old Israeli woman found lifeless hours after second dose of Covid-19 vaccine

01/13/21, Germany: 89-year-old dies about an hour after receiving Corona vaccine


01/14/21, USA: Louisiana woman suffers uncontrollable convulsions after getting experimental Pfizer COVID-19 vaccine

https://greatgameindia.com/uncontrollable-convulsions-pfizer-vaccine/

01/14/21, USA: Moderna victim, convulsions all over body

http://www.bitchute.com/video/P2eQoM0JyVu/?list=notifications&randomize=false

01/14/21, USA: Woman suffers whole body convulsions after taking experimental Moderna COVID-19 vaccine

https://greatgameindia.com/body-convulsions-moderna-vaccine/

01/14/21, Germany: Ten people die after Covid vaccination


01/15/21, France: Severe adverse reactions in 30 cases after Covid-19 vaccination


01/15/21, France: Elderly man in French care home dies two hours after receiving the vaccine


01/15/21, Israel: 4500 Corona infected after vaccination


01/15/21, France, Nice: 50 dead in home after vaccination

https://youtu.be/FB3-RmgaBUc

01/15/21, USA: Public health agency confirms 29 dangerous reactions to CoV vaccine

http://www.wochenblick.at/gesundheitsbehoerde-bestaetigt-29-gefaehrliche-reaktionen-auf-cov-impfung/

01/16/21, Israel: 13 Israelis suffer facial paralysis after taking Pfizer Covid jab


01/16/21, Germany: German nursing home sees Corona outbreak after vaccines


01/16/21, Norway: Scandal in Norway’s nursing homes: 23 deaths after Covid vaccinations


01/17/21, USA: 55 people died after receiving COVID-19 vaccine and 1388 in emergency departments


10 As of Feb 12, 2021, 15,923 adverse events, including 929 deaths had been reported to VAERS. Data may reflect serious delays in reporting. See example:

@ke11ybender: 45 yr old Angelia was hospitalized for seizures following COVID vaccination. She was diagnosed w/ right frontal lobe brain damage. The CDC was delayed in contacted [sic] her due to a month long backlog of adverse reaction reports. https://facebook.com/angelia.griner/posts/10158760300795180
https://www.facebook.com/angelia.griner/posts/101587603007951

Angelia Gipson Desselle, February 18 at 4:19 PM: The CDC finally called me today and have assigned someone to my case. She told me they have a month long back log of people having adverse reactions... that alone is a big eye opener. I am not alone in this. Apparently there’s a lot of us. I personally have connected to 5 people just like me. Although today was a bad one... that call meant so much!

11 Dr. Sherri Tenpenny estimates that reporting of adverse events to VAERS represents approximately 10% of actual adverse events. A Harvard study commissioned in 2010 by the federal government produced an even lower estimate, stating that less than 1% of adverse events get reported: https://digital.ahrq.gov/sites/default/files/docs/publication/r18hs017045-lazarus-final-report-2011.pdf

12 See Motley Fool’s three “red flags”: https://www.fool.com/investing/2020/08/29/3-red-flags-for-modernas-potential-coronavirus-vac/. In the company’s briefing document reviewed by the FDA (https://www.fda.gov/media/144452/download, page 15, tables 7 and 8), Moderna admits to a *23.9%* rate of adverse events. Among the 15,185 people in the study, there were 3,632 adverse events in the first 28 days, statistics that do not include the longer-term effects of DNA modification, infertility, autoimmunity, etc.

13 See “Softball Pfizer vaccine rollout interview goes horribly wrong,” https://www.youtube.com/watch?v=LMAZJtOdYNI.

14 Johnson & Johnson is facing billions in payouts for marketing baby powder known for decades to be tainted with cancer-causing asbestos. See, for example, https://markets.businessinsider.com/news/stocks/johnson-johnson-sets-aside-3-9-bln-for-baby-powder-verdict-settlements-1030109923?op=1. For details

15 AstraZeneca paid the settlement after illegally marketing an anti-psychotic drug to children and the elderly. See https://www.enjuris.com/blog/resources/largest-pharmaceutical-settlements-lawsuits/.


18 https://childrenshealthdefense.org/defender/pfizer-ceo-annual-vaccine-covid/.

19 In 2017, Italian researchers reviewed the ingredients of 44 types of so-called “vaccines.” They discovered heavy metal debris and biological contamination in every human vaccine they tested. The researchers stated, “The quantity of foreign bodies detected and, in some cases, their unusual chemical compositions baffled us.” They then drew the obvious conclusion, namely, that because the micro- and nanocontaminants were “neither biocompatible nor biodegradable,” they were “biopersistent” and could cause inflammatory effects right away—or later. From, “The Injection Fraud: It’s Not a Vaccine” by Catherine Austin Fitts: https://childrenshealthdefense.org/news/editorial/the-injection-fraud-its-not-a-vaccine/. Analyses of aborted fetal cell lines in “vaccines” identified abnormal human DNA, including genes associated with cancer, in all samples analyzed in quantities “up to 300 times higher than the limit imposed by the [European Medicines Agency] for carcinogenic DNA.” From, “New data shows DNA from aborted fetal cell lines in vaccines” by the Corvelva team, https://childrenshealthdefense.org/news/new-data-shows-aborted-fetal-cells-in-vaccines/.

20 In the U.S., individuals injured by Covid-19 injections may apply for compensation from the little-known Countermeasures Injury Compensation Program (CICP). However, the program has a one-year statute of limitations and does not compensate for pain and suffering or attorneys’ fees. Over the past decade, the CICP has rejected 92% of claims submitted. See https://www.concordmonitor.com/The-Race-for-COVID-19-Injury-Benefits- 37737582?fbclid=IwAR0GXi-8fEHelioAQdvnAT08-ZC1hAchHqPndM7Qe8QWodD5wKkVAnOVe8 and https://www.nbcwashington.com/news/local/critics-question-vaccine-injury-compensation-program-readiness-as-covid-19-claims-come-in/ 2573094/.


23 See Table 1 in https://www.cdc.gov/coronavirus/2019-ncov/hcp/planning-scenarios.html.

24 See CDC statistics here: https://www.cdc.gov/nchs/ncvs/vsrr/covid_weekly/index.htm/#SexAndAge. Additional sources of statistics can be found at Swiss Policy Research (https://swprs.org/facts-about-
25 See, for example, “Review of the emerging evidence demonstrating the efficacy of ivermectin in the prophylaxis and treatment of COVID-19”:


28 See endnote 27.


31 See endnote 30.