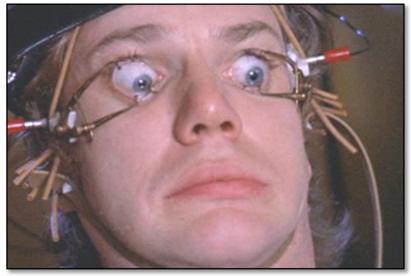
Where do we stand? Four years on.

- Lab Origin.
- Malfeasance Treatments and Evidence of Intent.
- Vaccine Disaster.
- ☐ Why it won't stop:
 - We can't just carry on, without future risks.

This may feel a bit like



but you need to see it.

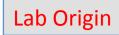


For all the dead, the injured, and the fighters.



There's a novel 19-nucleotide sequence in SARS-CoV-2:

- Not in any eukaryotic/viral organisms pre-2019: https://www.frontiersin.org/articles/10.3389/fviro.2022.834808/full
- Sequence: CTCCTCGGCGGCACGTAG
- If one reverses the sequence and takes the 'DNA compliment', one obtains: CTACGTGCCCGCCGAGGAG
 - Note to make the compliment sequence, swap C<->G and A<->T.
 - Note that <u>reverse transcription</u> uses the <u>complimentary</u> DNA strand when converting mRNA to DNA.



• There's a novel 19-nucleotide sequence in SARS-CoV-2:

- Not in any eukaryotic/viral organisms pre-2019: https://www.frontiersin.org/articles/10.3389/fviro.2022.834808/full
- Sequence: CTCCTCGGCGGCACGTAG
- If one reverses the sequence and takes the 'DNA compliment', one obtains: CTACGTGCCCGCCGAGGAG
 - Note to make the compliment sequence, swap C<->G and A<->T.
 - Note that <u>reverse transcription</u> uses the <u>complimentary</u> DNA strand when converting mRNA to DNA.
- This 100% matches a 2016 Moderna Inc. patent: U.S. Pat. 9587003-B2. (See Seq. ID 11652, nt 2733-2751)
- New sequence Forms a powerful <u>'Furin Cleavage Site'</u>, results in very efficient host-cell entry.
- No other beta-coronavirus even has a Furin Cleavage Site.
- Chance $4^19 = 4x4x4x4x4x4x4... = 1-in-271$ trillion. Divided by two.

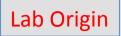


- There's a novel 19-nucleotide sequence in SARS-CoV-2:
 - Not in any eukaryotic/viral organisms pre-2019: https://www.frontiersin.org/articles/10.3389/fviro.2022.834808/full
 - Sequence: CTCCTCGGCGGCACGTAG
 - If one reverses the sequence and takes the 'DNA compliment', one obtains: CTACGTGCCCGCCGAGGAG
 - Note to make the compliment sequence, swap C<->G and A<->T.
 - Note that <u>reverse transcription</u> uses the <u>complimentary</u> DNA strand when converting mRNA to DNA.
 - This 100% matches a 2016 Moderna Inc. patent: U.S. Pat. 9587003-B2. (See Seq. ID 11652, nt 2733-2751)
 - New sequence Forms a powerful <u>'Furin Cleavage Site'</u>, results in very efficient host-cell entry.
 - No other beta-coronavirus even has a Furin Cleavage Site.
 - Chance $4^19 = 4x4x4x4x4x4x4...$ = 1-in-271 trillion. Divided by two.
- <u>DEFUSE proposal</u>: Leaked by Mjr. Murphy & Charles Rixey, both USMC.

 $\underline{https://drasticresearch.files.wordpress.com/2021/09/main-document-preempt-volume-1-no-ess-hr00118s0017-ecohealth-alliance.pdf}$

EcoHealth / Peter Daszak describe adding Cleavage Sites to backbones:
 "we will introduce appropriate human-specific cleavage sites" "viral spike glycoproteins v

"we will introduce appropriate human-specific cleavage sites" ... "viral spike glycoproteins will be synthesized, and those binding to human cell receptor ACE2 will be inserted into SARSr-CoV backbones"



- There's a novel 19-nucleotide sequence in SARS-CoV-2:
 - Not in any eukaryotic/viral organisms pre-2019: https://www.frontiersin.org/articles/10.3389/fviro.2022.834808/full
 - Sequence: CTCCTCGGCGGCACGTAG
 - If one reverses the sequence and takes the 'DNA compliment', one obtains: CTACGTGCCCGCCGAGGAG
 - Note to make the compliment sequence, swap C<->G and A<->T.
 - Note that <u>reverse transcription</u> uses the <u>complimentary</u> DNA strand when converting mRNA to DNA.
 - This 100% matches a 2016 Moderna Inc. patent: U.S. Pat. 9587003-B2. (See Seq. ID 11652, nt 2733-2751)
 - New sequence Forms a powerful 'Furin Cleavage Site', results in very efficient host-cell entry.
 - No other beta-coronavirus even has a Furin Cleavage Site.
 - Chance $4^19 = 4x4x4x4x4x4x4... = 1-in-271$ trillion. Divided by two.
- <u>DEFUSE proposal</u>: Leaked by Mjr. Murphy & Charles Rixey, both USMC.

 $\underline{https://drasticresearch.files.wordpress.com/2021/09/main-document-preempt-volume-1-no-ess-hr00118s0017-ecohealth-alliance.pdf}$

- EcoHealth / Peter Daszak describe adding Cleavage Sites to backbones:
 "we will introduce appropriate human-specific cleavage sites" ... "viral spike glycoproteins will be synthesized, and those binding to human cell receptor ACE2 will be inserted into SARSr-CoV backbones"
- DEFUSE aimed to use 'DC-SIGN' receptor (note: DC-SIGN is used by HIV):
 "we will sequentially introduce clade 2 disrupting residues of SARS-CoV and SHCO14 and evaluate virus growth in Vero cells, non-permissive cells ectopically expressing DC-SIGN and in human monocytes and macrophages anticipating reduced virus growth efficiency"

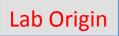


There's a novel 19-nucleotide sequence in SARS-CoV-2:

- Not in any eukaryotic/viral organisms pre-2019: https://www.frontiersin.org/articles/10.3389/fviro.2022.834808/full
- Sequence: CTCCTCGGCGGCACGTAG
- If one reverses the sequence and takes the 'DNA compliment', one obtains: CTACGTGCCCGCCGAGGAG
 - Note to make the compliment sequence, swap C<->G and A<->T.
 - Note that <u>reverse transcription</u> uses the <u>complimentary</u> DNA strand when converting mRNA to DNA.
- This 100% matches a 2016 Moderna Inc. patent: U.S. Pat. 9587003-B2. (See Seq. ID 11652, nt 2733-2751)
- New sequence Forms a powerful <u>'Furin Cleavage Site'</u>, results in very efficient host-cell entry.
- No other beta-coronavirus even has a Furin Cleavage Site.
- Chance $4^19 = 4x4x4x4x4x4x4... = 1-in-271$ trillion. Divided by two.
- <u>DEFUSE proposal</u>: Leaked by Mjr. Murphy & Charles Rixey, both USMC.

https://drasticresearch.files.wordpress.com/2021/09/main-document-preempt-volume-1-no-ess-hr00118s0017-ecohealth-alliance.pdf

- EcoHealth / Peter Daszak describe adding Cleavage Sites to backbones:
 "we will introduce appropriate human-specific cleavage sites" ... "viral spike glycoproteins will be synthesized, and those binding to human cell receptor ACE2 will be inserted into SARSr-CoV backbones"
- DEFUSE aimed to use 'DC-SIGN' receptor (note: DC-SIGN is used by HIV):
 "we will sequentially introduce clade 2 disrupting residues of SARS-CoV and SHCO14 and evaluate virus growth in Vero cells, non-permissive cells ectopically expressing DC-SIGN and in human monocytes and macrophages anticipating reduced virus growth efficiency"
- Restriction ('cut') sites, proposed in DEFUSE for assembly, are found in SARS2:
 https://alexwashburne.substack.com/p/the-nature-of-lab-origin-investigations
 "the pattern of cut/paste sites in the genome of SARS2 is consistent with methods proposed in DEFUSE and it triangulates to the same set of authors (Peter Daszak, Zhi ZhengLi), who were unique in using these specific enzymes"
- Claims 'research was not funded' after much of it was already done, prior to grant proposal.
 Toni Rocke "already developing SARS-CoV spike glycoproteins with respect to the DARPA grant"
 We also do not know what the PLA did, or didn't, fund..

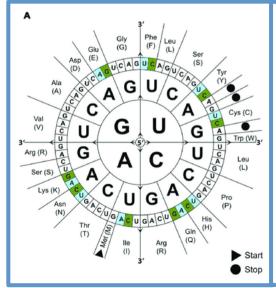


- SARS2 found contaminating Arctic samples from 2019: https://www.researchsquare.com/article/rs-1177047/latest.pdf
 - Variant is a precursor / ancestor of Wuhan-1 strain!
 - Viral host cell DNA was African Green Monkey (obviously not in Arctic, yet is commonly used in labs).
 - Same lab in Shanghai had routinely processed Wuhan lab samples in 2019. Contamination!
 - It means SARS2 existed in Chinese labs during 2019.



- SARS2 found contaminating Arctic samples from 2019: https://www.researchsquare.com/article/rs-1177047/latest.pdf
 - Variant is a precursor / ancestor of Wuhan-1 strain!
 - Viral host cell DNA was African Green Monkey (obviously not in Arctic, yet is commonly used in labs).
 - Same lab in Shanghai had routinely processed Wuhan lab samples in 2019. Contamination!
 - It means SARS2 existed in Chinese labs during 2019.
- Optimised for humans: Furin Cleavage Site 'RRAR'.. Made from codons:





How Amino Acids are formed from 'codons' of 3 nucleotides:

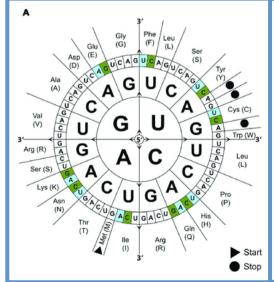
- Work from the centre to the perimeter.
- Choose your nucleotides in order.
- Read off the Amino Acid around the edge of the circle.
- Multiple ways to produce the same Amino Acid.
- Some ways are 'preferred'.
- Note: for mRNA, nucleotide 'U' does not exist; instead uses 'T'.



- SARS2 found contaminating Arctic samples from 2019: https://www.researchsquare.com/article/rs-1177047/latest.pdf
 - Variant is a precursor / ancestor of Wuhan-1 strain!
 - Viral host cell DNA was African Green Monkey (obviously not in Arctic, yet is commonly used in labs).
 - Same lab in Shanghai had routinely processed Wuhan lab samples in 2019. Contamination!
 - It means SARS2 existed in Chinese labs during 2019.
- Optimised for humans: Furin Cleavage Site 'RRAR'.. Made from codons:



- 'RxxR' is a standard Furin Cleavage Site, were 'x' can be anything.
- 'RRxR' is in SARS2, approx. 1000x more efficient.
- Good summary: https://nicholaswade.medium.com/origin-of-covid-following-the-clues-6f03564c038



How Amino Acids are formed from 'codons' of 3 nucleotides:

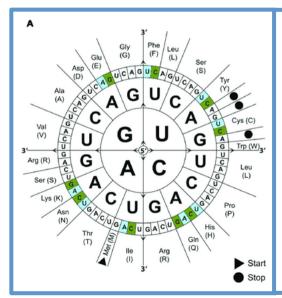
- Work from the centre to the perimeter.
- Choose your nucleotides in order.
- Read off the Amino Acid around the edge of the circle.
- Multiple ways to produce the same Amino Acid.
- Some ways are 'preferred'.
- Note: for mRNA, nucleotide 'U' does not exist; instead uses 'T'.



- SARS2 found contaminating Arctic samples from 2019: https://www.researchsquare.com/article/rs-1177047/latest.pdf
 - Variant is a precursor / ancestor of Wuhan-1 strain!
 - Viral host cell DNA was African Green Monkey (obviously not in Arctic, yet is commonly used in labs).
 - Same lab in Shanghai had routinely processed Wuhan lab samples in 2019. Contamination!
 - It means SARS2 existed in Chinese labs during 2019.
- Optimised for humans: Furin Cleavage Site 'RRAR'.. Made from codons:



- 'RxxR' is a standard Furin Cleavage Site, were 'x' can be anything.
- 'RRxR' is in SARS2, approx. 1000x more efficient.
- Good summary: https://nicholaswade.medium.com/origin-of-covid-following-the-clues-6f03564c038
- Hugely pre-adapted for human-cell infection/transmission 2x'CGG' coding for Arginine is human-preferred, not by virus
- Animal study incl. bats/pangolins found SARS2 binds best to <u>human</u> ACE2: https://arxiv.org/abs/2005.06199 (Piplani et al.)
- SARS2 reproduces poorly in bats direct jump ruled out no evidence of intermediary species.
- SARS2 'out of the box' can use human receptors: ACE2, nAChR, DC-SIGN, along with CD147 ('basigin', note Malaria link)
 NRP1, CD209L, ASGR1, KREMEN1, AXL, SR-B1, GRP78. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC10050803



How Amino Acids are formed from 'codons' of 3 nucleotides:

- Work from the centre to the perimeter.
- Choose your nucleotides in order.
- Read off the Amino Acid around the edge of the circle.
- Multiple ways to produce the same Amino Acid.
- Some ways are 'preferred'.
- Note: for mRNA, nucleotide 'U' does not exist; instead uses 'T'.



- <u>Uncanny matches to HIV</u> (four sequences match the 'gp120' and 'gag' proteins from HIV):
 - Pradhan et al. https://www.biorxiv.org/content/10.1101/2020.01.30.927871v1 (note: this publication was censored)
 - Not a coincidence: spike trimer expresses these in exactly the places they need to be, geometrically, to do damage.
 - Same sequences involved in infection of T-Cells (DC-Sign, CCR4, CXCR5).

Pradhan et al. Uncanny similarity of unique inserts in the 2019-nCoV spike protein to HIV-1 gp120 and Gag bioRxiv preprint doi: https://doi.org/10.1101/2020.01.30.927871

Motifs	Virus Glycoprotein	Motif Alignment	HIV protein and Variable region	HIV Genome Source Country/ subtype	Number of Polar Residues	Total Char ge	pI Valu e
Insert 1	2019- nCoV (GP) HIV1(GP120)	71 76 TNGTKR TNGTKR 404 409	gp120- V4	Thailand */ CRF01_ AE	5 5	2 2	11 11
Insert 2	2019- nCoV (GP) HIV1(GP120)	145 150 HKNNKS HKNNKS 462 467	gp120- V5	Kenya V	6	2 2	10 10
Insert 3	2019- nCoV (GP) HIV1(GP120)	245 256 RSYLTPGDSSSG RTYLFNETRGNSSSG 136 150	gp120- V1	India*/C	8 10	2	10.84 8.75
Insert 4	2019- nCoV (Poly P) HIV1(gag)	676 684 QTNSPRRA QTNSSILMQRSNFKG PRRA 366 384	Gag	India*/C	6 12	2 4	12.00 12.30

Table 1: Aligned sequences of 2019-nCoV and gp120 protein of HIV-1 with their positions in primary sequence of protein. All the inserts have a high density of positively charged residues. The deleted fragments in insert 3 and 4 increase the positive charge to surface area ratio. *please see Supp. Table 1 for accession numbers

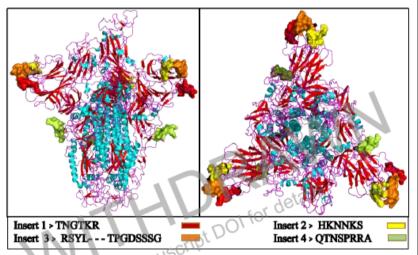


Figure 3. Modelled homo-trimer spike glycoprotein of 2019-nCoV virus. The inserts from HIV envelop protein are shown with colored beads, present at the binding site of the protein.



- <u>Uncanny matches to HIV</u> (four sequences match the 'gp120' and 'gag' proteins from HIV):
 - Pradhan et al. https://www.biorxiv.org/content/10.1101/2020.01.30.927871v1 (note: this publication was censored)
 - Not a coincidence: spike trimer expresses these in exactly the places they need to be, geometrically, to do damage.
 - Same sequences involved in infection of T-Cells (DC-Sign, CCR4, CXCR5).

Pradhan et al. Uncanny similarity of unique inserts in the 2019-nCoV spike protein to HIV-1 gp120 and Gag bioRxiv preprint doi: https://doi.org/10.1101/2020.01.30.927871

Motifs	Virus Glycoprotein	Motif Alignment	HIV protein and Variable region	HIV Genome Source Country/ subtype	Number of Polar Residues	Total Char ge	pI Valu e
Insert 1	2019- nCoV (GP) HIV1(GP120)	71 76 TNGTKR TNGTKR 404 409	gp120- V4	Thailand */ CRF01_ AE	5 5	2 2	11 11
Insert 2	2019- nCoV (GP) HIV1(GP120)	145 150 HKNNKS HKNNKS 462 467	gp120- V5	Kenya V	6	2 2	10 10
Insert 3	2019- nCoV (GP) HIV1(GP120)	245 256 RSYLTPGDSSSG RTYLFNETRGNSSSG 136 150	gp120- V1	India*/C	8 10	2	10.84 8.75
Insert 4	2019- nCoV (Poly P) HIV1(gag)	676 684 QTNSPRRA QTNSSIL.MQRSNFKG PRRA 366 384	Gag	India*/C	6 12	2 4	12.00 12.30

Table 1: Aligned sequences of 2019-nCoV and gp120 protein of HIV-1 with their positions in primary sequence of protein. All the inserts have a high density of positively charged residues. The deleted fragments in insert 3 and 4 increase the positive charge to surface area ratio. *please see Supp. Table 1 for accession numbers

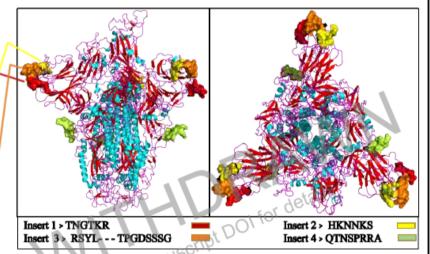


Figure 3. Modelled homo-trimer spike glycoprotein of 2019-nCoV virus. The inserts from HIV envelop protein are shown with colored beads, present at the binding site of the protein.

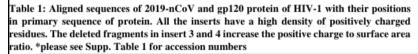
Note: inserts 1,2,3 all join up in the trimer, despite being separated in the sequence by hundreds of amino acids



- Uncanny matches to HIV (four sequences match the 'gp120' and 'gag' proteins from HIV):
 - Pradhan et al. https://www.biorxiv.org/content/10.1101/2020.01.30.927871v1 (note: this publication was censored)
 - Not a coincidence: spike trimer expresses these in exactly the places they need to be, geometrically, to do damage.
 - Same sequences involved in infection of T-Cells (DC-Sign, CCR4, CXCR5).

Pradhan et al. *Uncanny similarity of unique inserts in the 2019-nCoV spike protein to HIV-1 gp120 and Gag* bioRxiv preprint doi: https://doi.org/10.1101/2020.01.30.927871

Motifs	Virus Glycoprotein	Motif Alignment	HIV protein and Variable region	HIV Genome Source Country/ subtype	Number of Polar Residues	Total Char ge	pI Valu e
Insert 1	2019- nCoV (GP) HIV1(GP120)	71 76 TNGTKR TNGTKR 404 409	gp120- V4	Thailand */ CRF01_ AE	5 5	2 2	11 11
Insert 2	2019- nCoV (GP) HIV1(GP120)	145 150 HKNNKS HKNNKS 462 467	gp120- V5	Kenya V	6	2 2	10 10
Insert 3	2019- nCoV (GP) HIV1(GP120)	245 256 RSYLTPGDSSSG RTYLFNETRGNSSSG 136 150	gp120- V1	India*/C	8 10	2	10.84 8.75
Insert 4	2019- nCoV (Poly P) HIV1(gag)	676 684 QTNS	Gag	India*/C	6 12	2 4	12.00 12.30



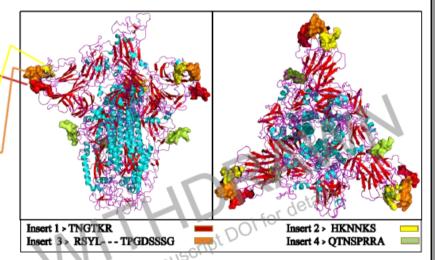


Figure 3. Modelled homo-trimer spike glycoprotein of 2019-nCoV virus. The inserts from HIV envelop protein are shown with colored beads, present at the binding site of the protein.

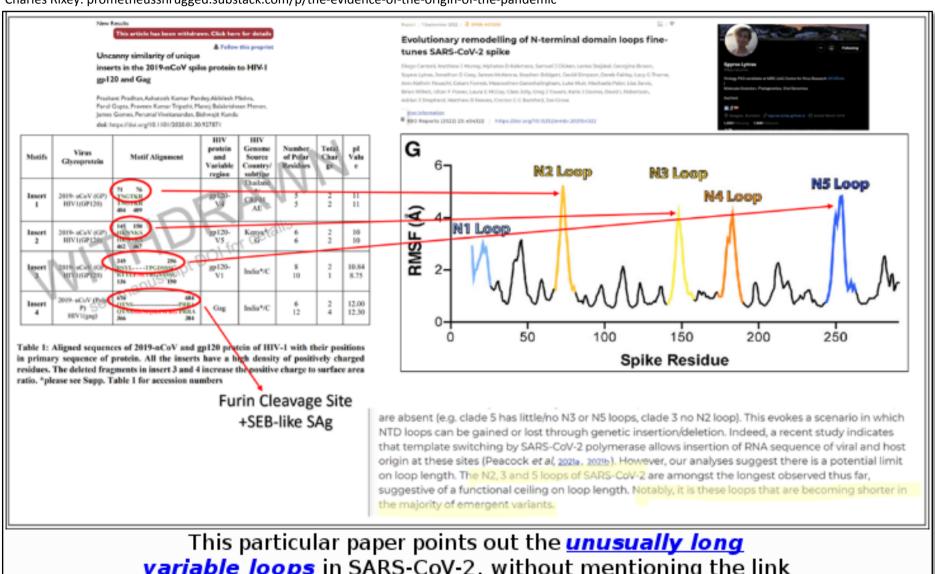
Note: inserts 1,2,3 all join up in the trimer, despite being separated in the sequence by hundreds of amino acids

<u>Weapons research</u> tends to focus on finding the pieces (epitopes) of useful harmful proteins, and splicing them into infectious vehicles.

A Laboratory *did* make this...



Charles Rixey: prometheusshrugged.substack.com/p/the-evidence-of-the-origin-of-the-pandemic

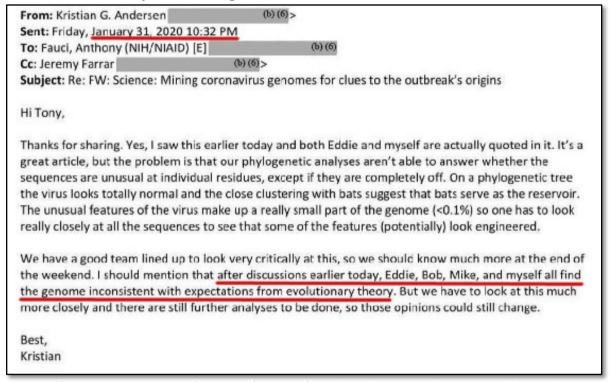


<u>variable loops</u> in SARS-CoV-2, without mentioning the link to the 3 Pradhan et al inserts.



- Duplicity by the authors of 'the natural origin paper': The proximal origin of SARS-CoV-2 & 'Lancet Letter':
 - FOIA'd emails & leaked SLACK messages show the authors believed 'friggin likely' lab origin:
 - Dr. Kristian Andersen: "The lab escape version of this is so friggin likely to have happened because they were already
 doing this type of work & the molecular data is fully consistent with that scenario."

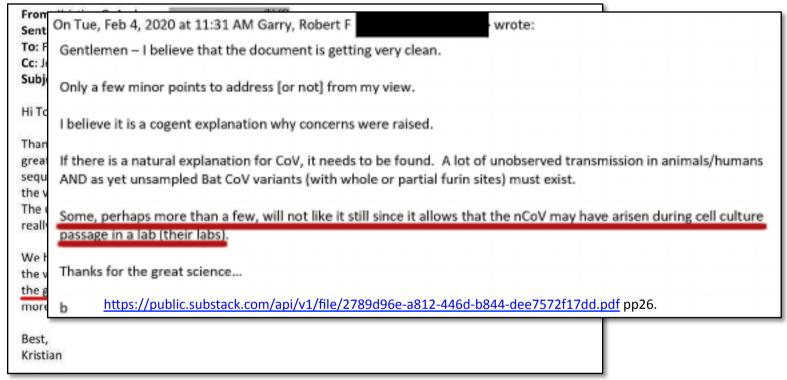
Two weeks before 'proximal origins':





- Duplicity by the authors of 'the natural origin paper': The proximal origin of SARS-CoV-2 & 'Lancet Letter':
 - FOIA'd emails & leaked SLACK messages show the authors believed 'friggin likely' lab origin:
 - Dr. Kristian Andersen: "The lab escape version of this is so friggin likely to have happened because they were already
 doing this type of work & the molecular data is fully consistent with that scenario."

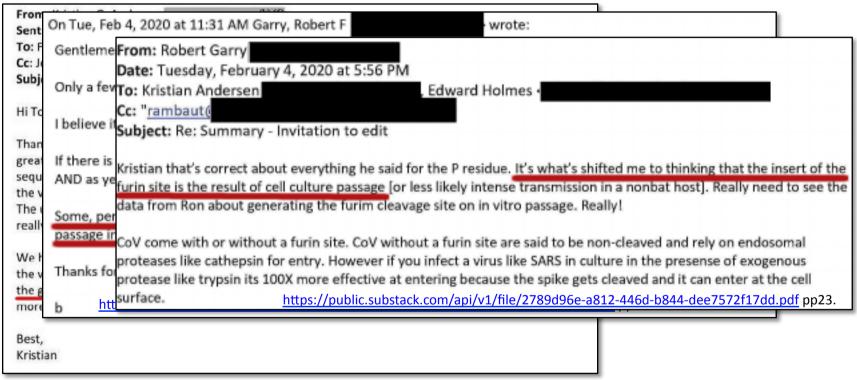
Two weeks before 'proximal origins':





- Duplicity by the authors of 'the natural origin paper': The proximal origin of SARS-CoV-2 & 'Lancet Letter':
 - FOIA'd emails & leaked SLACK messages show the authors believed 'friggin likely' lab origin:
 - Dr. Kristian Andersen: "The lab escape version of this is so friggin likely to have happened because they were already
 doing this type of work & the molecular data is fully consistent with that scenario."

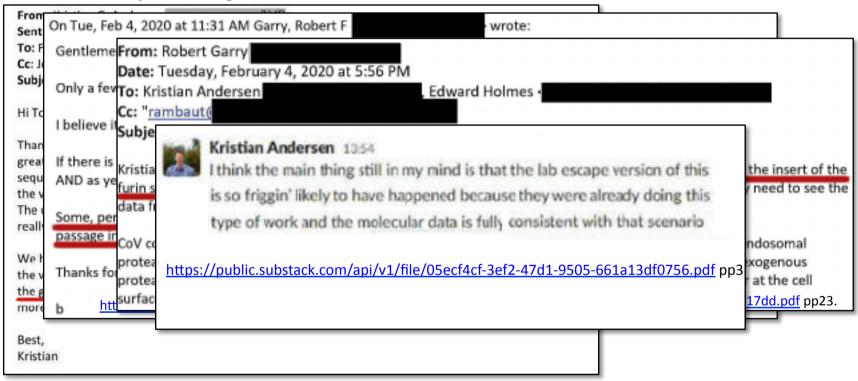
Two weeks before 'proximal origins':





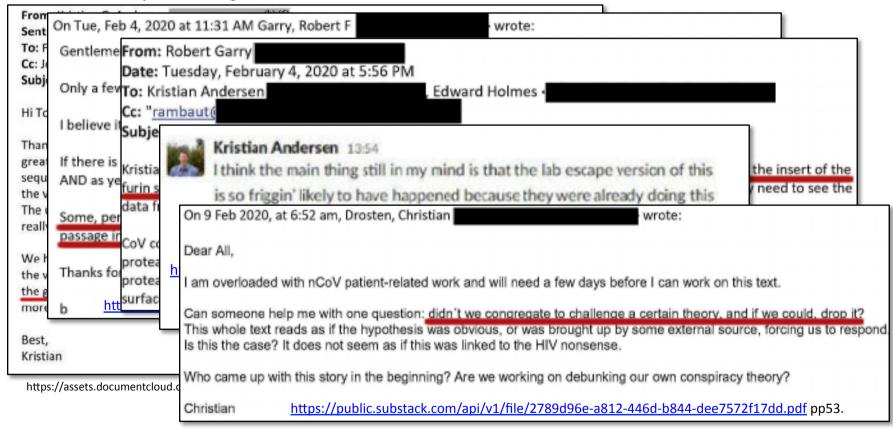
- Duplicity by the authors of 'the natural origin paper': The proximal origin of SARS-CoV-2 & 'Lancet Letter':
 - FOIA'd emails & leaked SLACK messages show the authors believed 'friggin likely' lab origin:
 - Dr. Kristian Andersen: "The lab escape version of this is so friggin likely to have happened because they were already
 doing this type of work & the molecular data is fully consistent with that scenario."

Two weeks before 'proximal origins':



- Duplicity by the authors of 'the natural origin paper': The proximal origin of SARS-CoV-2 & 'Lancet Letter':
 - FOIA'd emails & leaked SLACK messages show the authors believed 'friggin likely' lab origin:
 - Dr. Kristian Andersen: "The lab escape version of this is so friggin likely to have happened because they were already
 doing this type of work & the molecular data is fully consistent with that scenario."

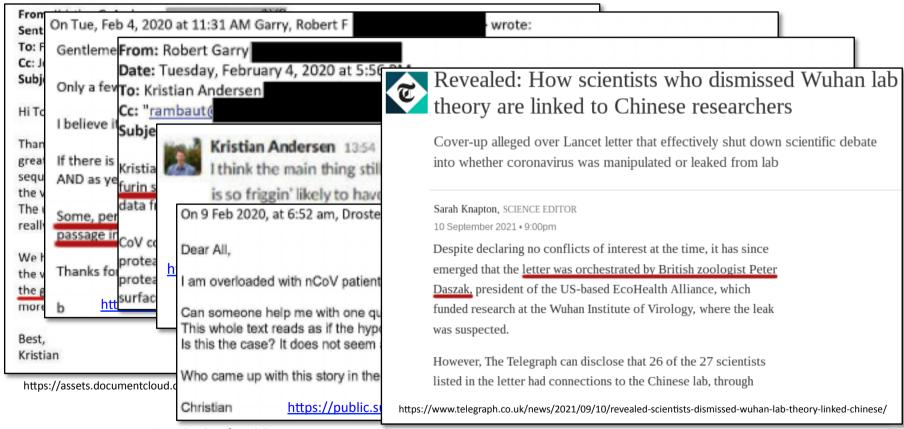
Two weeks before 'proximal origins':





- Duplicity by the authors of 'the natural origin paper': The proximal origin of SARS-CoV-2 & 'Lancet Letter':
 - FOIA'd emails & leaked SLACK messages show the authors believed 'friggin likely' lab origin:
 - Dr. Kristian Andersen: "The lab escape version of this is so friggin likely to have happened because they were already
 doing this type of work & the molecular data is fully consistent with that scenario."

Two weeks before 'proximal origins':

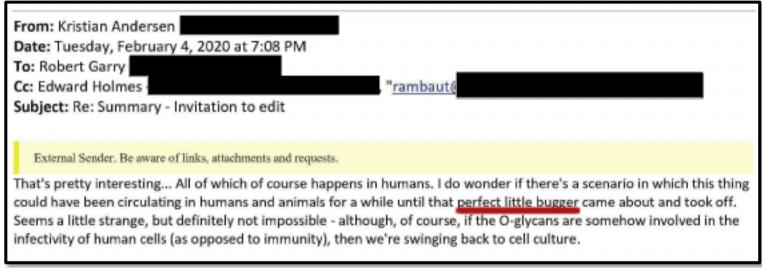


- Lancet Letter organised, drafted by Peter Daszak:
 - CEO EcoHealth, author of DEFUSE proposal, appointed by The Lancet to head COVID commission, whilst taking \$\$ for Coronavirus research in Wuhan.
 - "No competing interests".

Lab origin makes most sense...



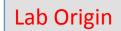
- More duplicity by the authors of 'the natural origin paper': The proximal origin of SARS-CoV-2.
 - The authors also thought SARS2 was 'perfect' and 'pre-adapted for human spread'.
 - "Perfect little bugger"....



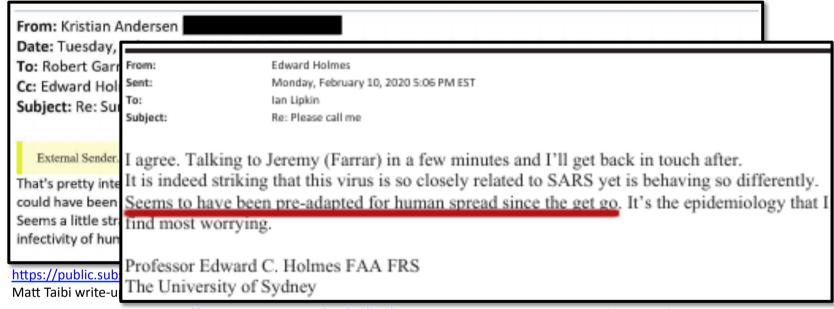
https://public.substack.com/api/v1/file/2789d96e-a812-446d-b844-dee7572f17dd.pdf pp22.

Matt Taibi write-up: https://www.public.news/p/covid-origins-scientist-denounces

Lab origin makes most sense...



- More duplicity by the authors of 'the natural origin paper': The proximal origin of SARS-CoV-2.
 - The authors also thought SARS2 was 'perfect' and 'pre-adapted for human spread'.
 - "Perfect little bugger"....



https://public.substack.com/api/v1/file/2789d96e-a812-446d-b844-dee7572f17dd.pdf pp67.

• Later, Fauci's NIH/NIAID awarded Kristian Andersen's lab \$8.9m (via his Scripps Institute).

https://andersen-lab.com/new-niaid-funded-center-established/

New NIAID-funded center established - Andersen Lab (Aug. 2020)

- Scripps was kicked out of Sierra Leone after an anomalous 'Ebola Zaire' outbreak. Robert F Garry heads up the VHFC in Kenema:
- https://www.independentsciencenews.org/health/did-west-africas-ebola-outbreak-of-2014-have-a-lab-origin/ note "Metabiota".
- https://www.popsci.com/article/science/how-did-deadliest-strain-ebola-travel-central-west-africa/
- https://husseini.substack.com/p/did-west-africas-ebola-outbreak-of VHFC charged Metabiota with culturing Ebola-infected cells.

Lab leaks are not new, or unprecedented..

The Lancet: 21 years of official 'lab leaks' 2000-2021. Includes Ebola, Polio, Prions, Cholera, Anthrax, Zika... https://www.thelancet.com/journals/lanmic/article/PIIS2666-5247(23)00319-1/fulltext

Official Acknowledgement



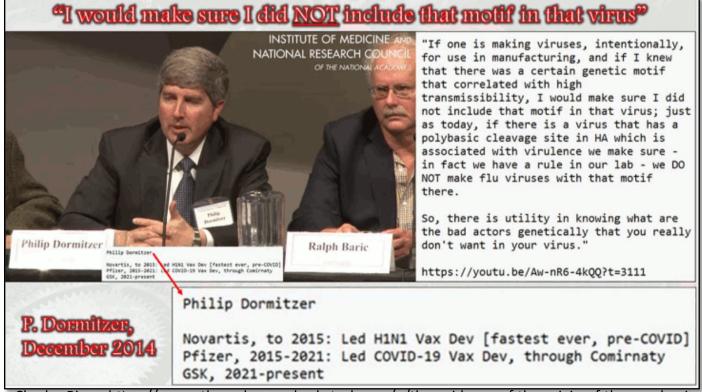
- US Dept. of Energy (DoE):
 - Despite the name, DoE in charge of scientists at CERN, nuclear programmes, defence/weapons focus.
 - Admitted in late Feb. 2023:
 - likely lab leak with low confidence.
 - More to this.. Bette Korber, Lead HIV researcher @DoE behind assessment.
 - Bette removed Furin Cleavage Sites (FCS) from ALL her trial vaccines.
 - Bette would have known about FCS origins, HIV inserts.
- FBI Director Wray:
 - Admitted they believe Chinese Lab release "likely".

Lab Origin

- **Origin**: Was it a Leak, or Intentional?
- **Motive**: Vaccine research? Defence? Weapon?
 - If people had known the lab connections would they have thought twice about vaccination?



- Origin: Was it a Leak, or Intentional?
- Motive: Vaccine research? Defence? Weapon?
 - If people had known the lab connections would they have thought twice about vaccination?
- Option #1: Self-spreading vaccine. The <u>virus</u> could have been a prototype 'self-spreading vaccine'....
 - But the "Dormitzer film" contradicts this hypothesis.. https://www.youtu.be/Aw-nR6-4kQQ?t=3111
 - Strange fact: Apparently "Fusion Inhibitors" EK1C4 / EK1, which can stop a Coronavirus infection, <u>are broken by</u> an epitope in HIV which occurs right after the SARS2 HIV insert.
 - This seems deliberate. Was there a choice made NOT to break Fusion Inhibitors when making SARS2?
 - Those in the loop would have had protection.



Charles Rixey: https://prometheusshrugged.substack.com/p/the-evidence-of-the-origin-of-the-pandemic



- Origin: Was it a Leak, or Intentional?
- Motive: Vaccine research? Defence? Weapon?
 - If people had known the lab connections would they have thought twice about vaccination?
- Option #1: Self-spreading vaccine. The <u>virus</u> could have been a prototype 'self-spreading vaccine'....
 - But the "Dormitzer film" contradicts this hypothesis.. https://www.youtu.be/Aw-nR6-4kQQ?t=3111
 - Strange fact: Apparently "Fusion Inhibitors" EK1C4 / EK1, which can stop a Coronavirus infection, <u>are broken by</u> an epitope in HIV which occurs right after the SARS2 HIV insert.
 - This seems deliberate. Was there a choice made NOT to break Fusion Inhibitors when making SARS2?
 - Those in the loop would have had protection.

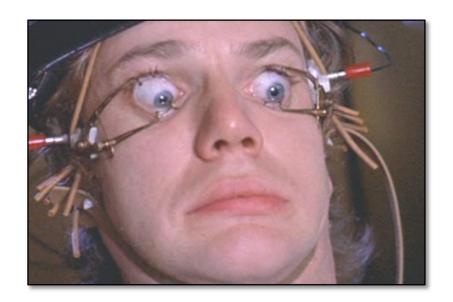


- Origin: Was it a Leak, or Intentional?
- Motive: Vaccine research? Defence? Weapon?
 - If people had known the lab connections would they have thought twice about vaccination?
- Option #1: Self-spreading vaccine. The <u>virus</u> could have been a prototype 'self-spreading vaccine'....
 - But the "Dormitzer film" contradicts this hypothesis.. https://www.youtu.be/Aw-nR6-4kQQ?t=3111
 - Strange fact: Apparently "Fusion Inhibitors" EK1C4 / EK1, which can stop a Coronavirus infection, <u>are broken by</u> an epitope in HIV which occurs right after the SARS2 HIV insert.
 - This seems deliberate. Was there a choice made NOT to break Fusion Inhibitors when making SARS2?
 - Those in the loop would have had protection.
- Option #2: Weapon: There is a lot of evidence suggesting SARS2 is very harmful!
 - CBRN ratings of bioweapon veracity put SARS2 at #4, ahead of Marburg, Ebola, but behind Smallpox.
 - 2018: Agents ranked as bioweapons using 12 criteria, from this paper: https://pubmed.ncbi.nlm.nih.gov/29401327/
 - Beyond the Dirty Dozen: A Proposed Methodology for Assessing Future Bioweapon Threats.
 - 2020: SAME AUTHORS revisited the paper and they evaluated SARS2 as #4 on the list... Above Ebola and Nipah: https://academic.oup.com/milmed/article/188/3-4/531/6586077
 - Analysis of the Virus SARS-CoV-2 as a Potential Bioweapon in Light of International Literature.
- Weapons research revolves around incapacitation; does not break treaties:
 - Immune system damage is one route. (Did I mention that HIV causes dementia, too? "HAND")
 - Weaponisation of Prions has been another long-standing goal. Dementia, Fibrosis.
 - Problem was always effective spread (solved by integrating Prion and HIV epitopes into SARS2).
 - Ideal weapon should be dual-payload, very infectious, mild-moderate initial illness.
 - Incapacitation follows in time.
- MANY OPEN QUESTIONS, but the balance of evidence and 'built-in' harm favours a weapon.

You made it this far...

Malfeasance

- Lab Origin.
- Malfeasance Treatments and Evidence of Intent.
- ☐ Vaccine Disaster.
- ☐ Why it won't stop:
 - We can't just carry on, without future risks.



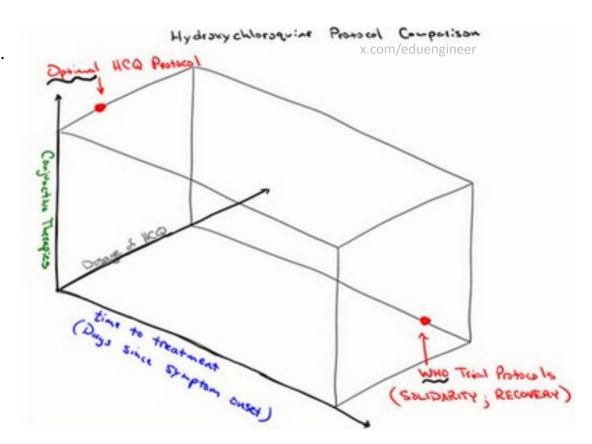
Intent and Malfeasance



- Withholding of treatments:
 - Systematic attempts to discredit HCQ, Ivermectin, Antibiotics ... "#3tablets"
 - Vaccine Emergency Use Authorisations were dependent upon a lack of existing treatments.
- FDA anticipated side-effects from mRNA vaccines, whilst telling you 'safe & effective':
 - Blink and you'd miss it: momentary display of FDA Slide from VRBPAC meeting on October 22, 2020 @2h33m40s.
 - Vaccine biodistribution known, prior to authorisation.
- Regulators asleep, or colluding:
 - We will see a video of June Raine (MHRA, UK) explaining how MHRA became 'an enabler'.
- Midazolam was undoubtedly used to punish and execute people!
- DNA Contamination in the vaccines a danger to human host-cells:
 - Was known about, and has --so far-- been dismssed/covered-up.
 - SV40 promoter used in Pfizer plasmid is very dangerous, and labelling of it was <u>deleted</u> by Pfizer.
- Fraud in the Trials (Pfizer):
 - Not accidental to withhold incidences of illnesses like leukemia or paralysis in trial participants.
 - Retrospective analysis shows more people died in the Pfizer treatment arm, vs. placebo.

HCQ – Treat late, no conjunctives (eg Zinc), ...and overdose

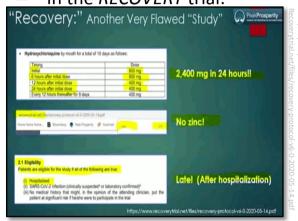
- Trials designed to fail:
 - SOLIDARITY, RECOVERY, TOGETHER.

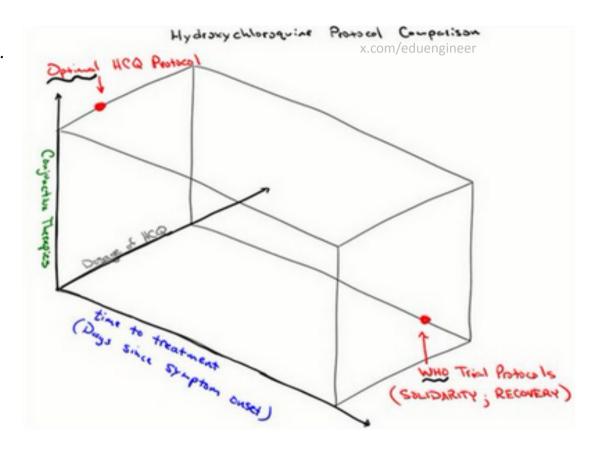


HCQ – Treat late, no conjunctives (eg Zinc), ...and overdose

- Trials designed to fail:
 - SOLIDARITY, RECOVERY, TOGETHER.
- Overdosing of patients happened

– in the RECOVERY trial:



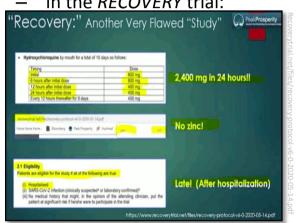


HCQ – Treat late, no conjunctives (eg Zinc), ...and overdose

UCA Protoco

- Trials designed to fail:
 - SOLIDARITY, RECOVERY, TOGETHER.
- Overdosing of patients happened

in the RECOVERY trial:



- Fantastic Documentary:
 - "Epidemic of Fraud" traces the suppression of Quinine over centuries



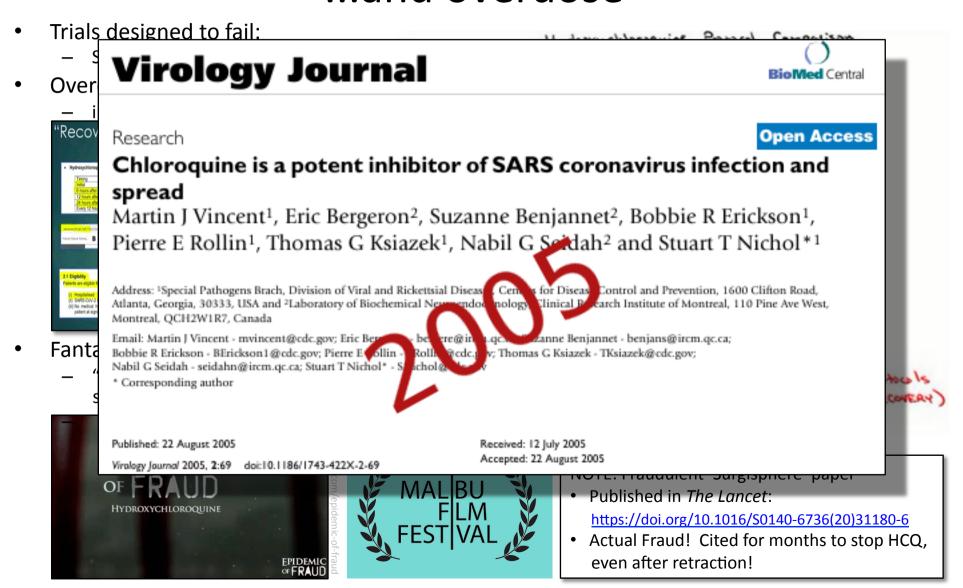
NOTE: Fraudulent 'Surgisphere' paper

Hydroxychloroquine Protocol Comparison

x.com/eduengineer

- Published in *The Lancet*: https://doi.org/10.1016/S0140-6736(20)31180-6
- Actual Fraud! Cited for months to stop HCQ, even after retraction!

HCQ – Treat late, no conjunctives (eg Zinc), ...and overdose



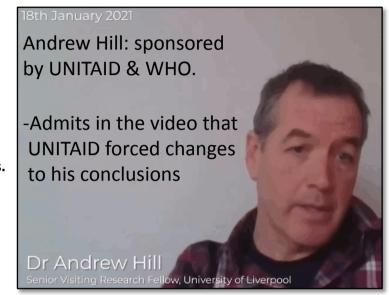
Ivermectin and Andrew Hill

Malfeasance

Over 100 peer-reviewed papers showing efficacy https://c19ivm.org/meta.html

Ivermectin and Andrew Hill

- Over 100 peer-reviewed papers showing efficacy https://c19ivm.org/meta.html
- A meta-analysis by Andrew Hill had its conclusions adjusted shortly before publication.
- Hill, before WHO & UNITAID adjustments (Feb '21):
 - •- "the purpose of this report is to forewarn people..get ready to approve it".
 - •- "can we start to upscale ivermectin as well" [as per vaccines].
- Andrew admitted UNITAID had forced the changes to his conclusions:
 - •- Removed dose-dependent positive treatment effects, among other things.
 - •- No author attribution for the changes.
- In turn, this led the WHO against recommending Ivermectin:
 - •- "not to use ivermectin except in the context of a clinical trial"



oraclefilms.com/alettertoandrewhill

Malfeasance

Ivermectin and Andrew Hill

- Over 100 peer-reviewed papers showing efficacy https://c19ivm.org/meta.html
- A meta-analysis by Andrew Hill had its conclusions adjusted shortly before publication.
- Hill, before WHO & UNITAID adjustments (Feb '21):
 - •- "the purpose of this report is to forewarn people..get ready to approve it".
 - •- "can we start to upscale ivermectin as well" [as per vaccines].
- Andrew admitted UNITAID had forced the changes to his conclusions:
 - •- Removed dose-dependent positive treatment effects, among other things.
 - •- No author attribution for the changes.
- In turn, this led the WHO against recommending Ivermectin:
 - •- "not to use ivermectin except in the context of a clinical trial"



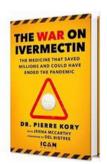
oraclefilms.com/alettertoandrewhill

You could write a book about this aspect alone....

- Dr. Pierre Kory did just that!
- Hill promised Kory that when his contract ended with UNITAID, he would republish, making everything right...he never did.

Chapter 23 of:





Antibiotics Withheld



MYTH: you can't use antibiotics against viruses...

- We know SARS2 has bacteriophage-like activity:
 - https://pubmed.ncbi.nlm.nih.gov/34336189/

Increase of SARS-CoV-2 RNA load in faecal samples prompts for rethinking of SARS-CoV-2 biology and COVID-19 epidemiology

Antibiotics Withheld



MYTH: you can't use antibiotics against viruses...

- We know SARS2 has bacteriophage-like activity:
 - https://pubmed.ncbi.nlm.nih.gov/34336189/
 Increase of SARS-CoV-2 RNA load in faecal samples prompts for rethinking of SARS-CoV-2 biology and COVID-19 epidemiology
- Clinical evidence is vast:
 - The Lancet, Volume 361, Issue 9369, 10 May 2003, pp1615-1617:
 Development of a standard treatment protocol for severe acute respiratory syndrome
 "A series of 31 patients with probable SARS, diagnosed from WHO criteria, were treated according to a

"One patient recovered on **antibacterial treatment alone**, 17 showed rapid and sustained responses, and 13 achieved improvement with step-up or pulsed methylprednisolone."

treatment protocol consisting of antibacterials and a combination of ribavirin and methylprednisolone."

https://pubmed.ncbi.nlm.nih.gov/32972196/
 Clinical evidences on the antiviral properties of macrolide antibiotics in the COVID-19 era and beyond

Antibiotics Withheld

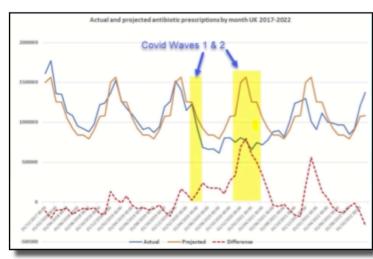


MYTH: you can't use antibiotics against viruses...

- We know SARS2 has bacteriophage-like activity:
 - https://pubmed.ncbi.nlm.nih.gov/34336189/
 Increase of SARS-CoV-2 RNA load in faecal samples prompts for rethinking of SARS-CoV-2 biology and COVID-19 epidemiology
- Clinical evidence is vast:
 - The Lancet, Volume 361, Issue 9369, 10 May 2003, pp1615-1617:
 Development of a standard treatment protocol for severe acute respiratory syndrome
 "A series of 31 patients with probable SARS, diagnosed from WHO criteria, were treated according to a treatment protocol consisting of antibacterials and a combination of ribavirin and methylprednisolone."

"One patient recovered on **antibacterial treatment alone**, 17 showed rapid and sustained responses, and 13 achieved improvement with step-up or pulsed methylprednisolone."

- https://pubmed.ncbi.nlm.nih.gov/32972196/
 Clinical evidences on the antiviral properties of macrolide antibiotics in the COVID-19 era and beyond
- Macrolide antibiotics DO work against viruses.
- Inexplicable blue line; (source: openprescribing.net) _____
 - Would have expected <u>more</u> scripts, not less!
- Secondary infections were allowed to run rampant.
 - Many were killed by bacterial pneumonia.



Malfeasance

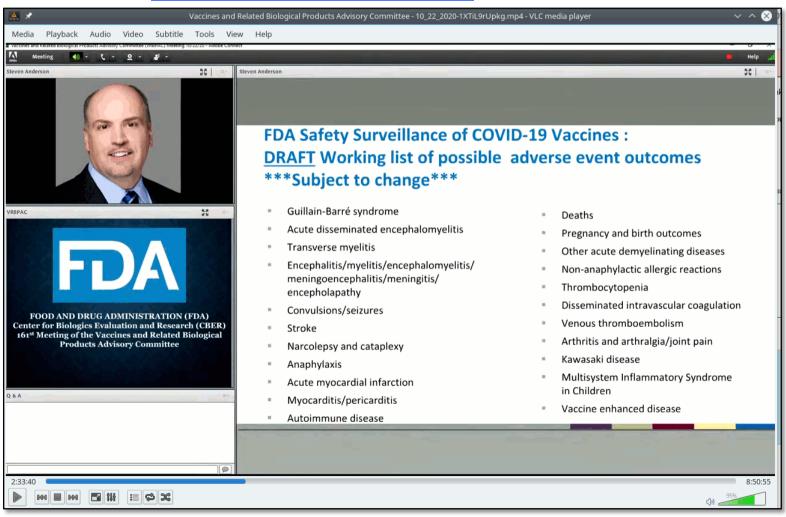
Which Antibiotics Work?

• Petrillo et al.: Increase of SARS-CoV-2 RNA load in faecal samples prompts for rethinking of SARS-CoV-2 biology and COVID-19 epidemiology



Regulators Knew..

- Something very curious happened on October 22, 2020.
- For only a split-second, at the official meeting of the Vaccines and Related Biological Products Advisory Committee (VRBPAC), a slide was shown by the FDA @2h33m40s.... https://www.youtube.com/watch?v=1XTiL9rUpkg



UK's MHRA is Totally Corrupt ... Malfeasance

- The MHRA gets 86% of funding from Pharma/VI's.
 - Other regulators are even worse...

	Australia TGA	Europe EMA	UK MHRA	Japan PMDA	USA FDA	Canada HC
Budgets and fees						
Proportion of budget derived from industry	96%	89%	86%	85%	65%	50.5%
Total annual budget [‡]	AU\$170m (£95m)	€386m (£331m)	£159m	¥29.1bn (£175m)	US\$6.1bn (£5bn)	C\$2.7bn (£1.7bn)
Transparency, COIs, and	data					
Proportion of covid-19 vaccine committee members that declared financial COIs	50%	3%	32%	75%	<10%	0%

https://news.rebekahbarnett.com.au/p/australian-drug-regulator-goes-on

UK's MHRA is Totally Corrupt ...

Malfeasance

- The MHRA gets 86% of funding from Pharma/VI's.
 - Other regulators are even worse...
 - The members of the board have Conflicts of Interest such as declared *investments in firms they regulate*

Last updated: 19th August 2020							
Name & role	Name of company / organisation	Nature of interest	Whether current	Other comments			
Anne-Toni Rodgers Non-Executive Director	AstraZeneca	Shareholder	Yes				
Amanda Calvert Non-Executive Director	Athenex Pharmaceuticals CRUK – digital Experimental Cancer Medicines Team	Consultancy – Fee paying Consultancy – Fee Paying	Yes				
	AstraZeneca Pharmaceuticals	Shareholder	Yes				

https://assets.publishing.service.gov.uk/government/uploads/system/uploads/ attachment_data/file/911028/MHRA_Board_Declarations_of_interest_20_August_2020.pdf

	Australia TGA	Europe EMA	UK MHRA	Japan PMDA	USA FDA	Canada HC
Budgets and fees						
Proportion of budget derived from industry [©]	96%	89%	86%	85%	65%	50.5%
Total annual budget [‡]	AU\$170m (£95m)	€386m (£331m)	£159m	¥29.1bn (£175m)	US\$6.1bn (£5bn)	C\$2.7bn (£1.7bn)
Transparency, COIs, an	data					
Proportion of covid-19 vaccine committee members that declared financial COIs	50%	3%	32%	75%	<10%	0%

https://news.rebekahbarnett.com.au/p/australian-drug-regulator-goes-on

UK's MHRA is Totally Corrupt ...

- The MHRA gets 86% of funding from Pharma/VI's.
 - Other regulators are even worse...
 - The members of the board have Conflicts of Interest such as declared investments in firms they regulate.

Last updated: 19th August 2020							
Name & role	Name of company / organisation	Nature of interest	Whether current	Other comments			
Anne-Toni Rodgers Non-Executive Director	AstraZeneca	Shareholder	Yes				
Amanda Calvert Non-Executive Director	Athenex Pharmaceuticals CRUK – digital Experimental Cancer Medicines Team	Consultancy – Fee paying Consultancy – Fee Paying	Yes				
	AstraZeneca Pharmaceuticals	Shareholder	Yes				

https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/911028/MHRA_Board_Declarations_of_Interest_20_August_2020.pdf

	Australia TGA	Europe EMA	UK MHRA	Japan PMDA	USA FDA	Canada HC
Budgets and fees						
Proportion of budget derived from industry	96%	89%	86%	85%	65%	50.5%
Total annual budget [‡]	AU\$170m (£95m)	€386m (£331m)	£159m	¥29.1bn (£175m)	US\$6.1bn (£5bn)	C\$2.7bn (£1.7bn)
Transparency, COIs, and	data					
Proportion of covid-19 vaccine committee members that declared financial COIs	50%	3%	32%	75%	<10%	0%

Malfeasance

https://news.rebekahbarnett.com.au/p/australian-drug-regulator-goes-on

- June Raine, Chief Executive: MHRA has become 'an enabler'.
 - To me, she literally sounds giddy.



https://www.youtube.com/watch?v=xUQfzTqPUm4 at 31m...

UK's MHRA is Totally Corrupt ...

Malfeasance

- The MHRA gets 86% of funding from Pharma/VI's.
 - Other regulators are even worse...
 - The members of the board have Conflicts of Interest such as declared *investments in firms they regulate*.

Last updated: 19th August 2020							
Name & role	Name of company / organisation	Nature of interest	Whether current	Other comments			
Anne-Toni Rodgers Non-Executive Director	AstraZeneca	Shareholder	Yes				
Amanda Calvert Non-Executive Director	Athenex Pharmaceuticals CRUK – digital Experimental Cancer Medicines Team	Consultancy – Fee paying Consultancy – Fee Paying	Yes				
	AstraZeneca Pharmaceuticals	Shareholder	Yes				

https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/911028/MHRA_Board_Declarations_of_Interest_20_August_2020.pdf

Table 1 How the regulators compare Australia Europe UK USA Canada TGA **EMA** MHRA **PMDA** FDA HC Budgets and fees Proportion of budget 96% 89% 86% 85% 65% 50.5% derived from industry AU\$170m C\$2.7bn €386m ¥29.1bn US\$6.1bn Total annual budget 4 £159m (£95m) (£331m) (£175m) (£5bn) (£1.7bn) Transparency, COIs, and Proportion of covid-19 vaccine committee 50% 32% 75% <10% 0% members that declared financial COIs

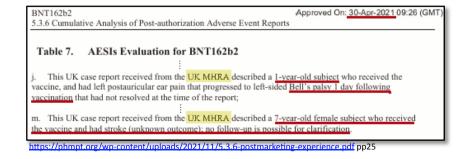
https://news.rebekahbarnett.com.au/p/australian-drug-regulator-goes-on

- June Raine, Chief Executive: MHRA has become 'an enabler'.
 - To me, she literally sounds giddy.

- Pfizer's own trial documents revealed the MHRA asleep at the wheel:
 - A 1yo baby and a 7yo girl. "No follow-up is possible".
 - On a trial written up by April 2021?
 - How does one reliable diagnose just Bell's Palsy in a 1yo?



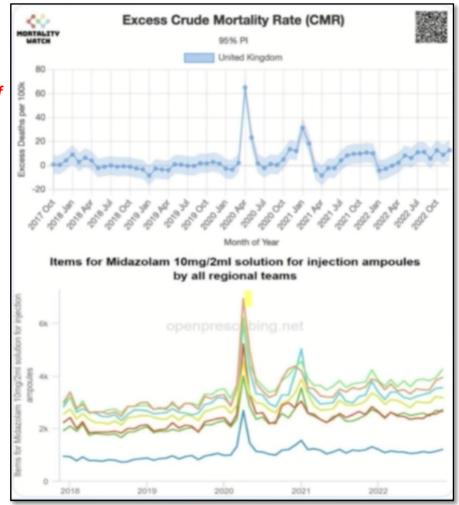
https://www.youtube.com/watch?v=xUQfzTqPUm4 at 31m...



Malfeasance

Midazolam Murders

- Powerful barbiturate (sedative).
- NICE guideline NG163 (no author!)
 - suggested Midazolam+Morphine, powerful respiratory depressants, for 'Managing Breathlessness' in 18+.
 - "Sedation and opioid use should not be withheld because of an inappropriate fear of causing respiratory depression"

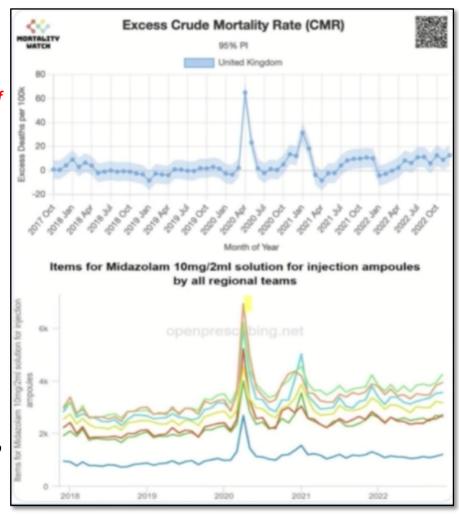


Midazolam Murders

- Powerful barbiturate (sedative).
- NICE guideline NG163 (no author!)
 - suggested Midazolam+Morphine, powerful respiratory depressants, for 'Managing Breathlessness' in 18+.
 - "Sedation and opioid use should not be withheld because of an inappropriate fear of causing respiratory depression"
- Administered outside hospitals:



- "out-of-hospital prescribing of the drug midazolam increased by more than 100% in April compared to previous months."
- "An anti-euthanasia campaigner last night said he suspected that the spike was evidence that many people had been put on end-of-life protocols or 'pathways'."
- "Whistleblowers also claimed to have witnessed misuse of sedatives, with staff told to give them to dementia patients to stop them wandering the corridors."

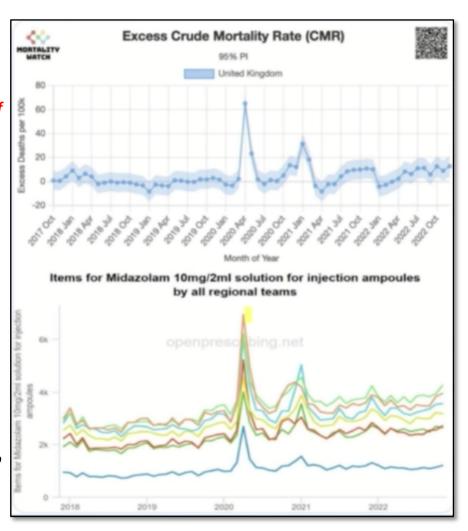


Midazolam Murders

- Powerful barbiturate (sedative).
- NICE guideline NG163 (no author!)
 - suggested Midazolam+Morphine, powerful respiratory depressants, for 'Managing Breathlessness' in 18+.
 - "Sedation and opioid use should not be withheld because of an inappropriate fear of causing respiratory depression"
- Administered outside hospitals:



- "out-of-hospital prescribing of the drug midazolam increased by more than 100% in April compared to previous months."
- "An anti-euthanasia campaigner last night said he suspected that the spike was evidence that many people had been put on end-of-life protocols or 'pathways'."
- "Whistleblowers also claimed to have witnessed misuse of sedatives, with staff told to give them to dementia patients to stop them wandering the corridors."
- Phenomenon only seen in England.
- Often done without consent, or AGAINST wishes.
- Eyewitnesses...



Midazolam "Strongly and causally associated"

In summary, Midazolam was strongly and causally associated with UK excess deaths, particularly in 2020. It was clearly the proximate cause of excess mortality in UK, but it was unlikely to be the primary cause in the chain of causality for deaths, because Midazolam was used mostly for accelerated or assisted dying in euthanasia often to alleviate possible suffering in end-of-life protocols. Midazolam's role based on its pharmaceutics is circumscribed in health policy guidelines.

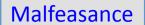
Wilson Sy - Med. Clin Res., Vol. 9, Issue 2 (2024)

Midazolam "Strongly and causally associated"

In summary, Midazolam was strongly and causally associated with UK excess deaths, particularly in 2020. It was clearly the proximate cause of excess mortality in UK, but it was unlikely to be the primary cause in the chain of causality for deaths, because Midazolam was used mostly for accelerated or assisted dying in euthanasia often to alleviate possible suffering in end-of-life protocols. Midazolam's role based on its pharmaceutics is circumscribed in health policy guidelines.

Wilson Sy - Med. Clin Res., Vol. 9, Issue 2 (2024)

- Let's look at the health policy guidelines.
- NICE guideline NG163 (no author!)
 - suggested MZM+Morphine, powerful respiratory depressants, for 'Managing Breathlessness' in 18+.
 - "Sedation and opioid use should not be withheld because of an inappropriate fear of causing respiratory depression"
- Similar documents to NG163 appeared simultaneously in Australia, Canada, etc.
 - More later...



NICE Guideline NG163

England Only?

Z-scores (Std. Dev.) in Excess Deaths (45-64), ONS Table 5 Treatments in the last days and hours of life for managing breathlessness for patients aged 18 years and over Treatment Dosage Morphine sulfate 10 mg over 24 hours via a syringe driver, increasing stepwise to morphine sulfate 30 mg over 24 hours as Opioid required Benzodiazepine if Midazolam 10 mg over 24 hours via the syringe driver, increasing required in addition to stepwise to midazolam 60 mg over 24 hours as required UK (Northern Ireland) opioid Morphine sulfate 2.5 mg to 5 mg subcutaneously as required Add parenteralmorphine Midazolam 2.5 mg subcutaneously as required. or midazolam if required (See BNF for more details on dosages). UK (Scotland) Special considerations Consider concomitant use of an antiemetic (such as haloperidol) and a regular stimulant laxative (such as senna). Continue with non-pharmacological strategies for managing

Notes: At the time of publication (April 2020), opioids and benzodiazepines did not have a UK marketing authorisation for this indication or route of administration (see <u>General Medical Council's guidance on prescribing unlicensed medicines</u> for further information).

breathlessness when starting an opioid.

Sedation and opioid use should not be withheld because of an

inappropriate fear of causing respiratory depression.

NG163 was released in April 2020, was then deleted from the NICE website:

Archived: https://web.archive.org/web/20200409054527/https://www.nice.org.uk/guidance/ng163/resources/covid19-rapid-guideline-managing-symptoms-including-at-the-end-of-life-in-the-community-pdf-66141899069893

UK (Wales)

Original: https://www.nice.org.uk/guidance/ng163/resources/covid19-rapid-guideline-managing-symptoms-including-at-the-end-of-life-in-the-community-pdf-66141899069893

Eyewitnesses: Midazolam only given to really sick people?

Nurse Elena Anamaria

https://odysee.com/@MaajidNawaz:d/Ep6-Radical:9

"On Allegations of Involuntary State Euthanasia Using Midazolam"

- 60m20s: "They put a DNR in place against our wishes"
- 61m05s: "The reason was, he was unvaccinated"
 See also: 70m30s -71m30s "unvaccinated...covid..EOL"
- 63m00s: "Fast-track EOL care" .. Was introduced with COVID and the ONLY requirement was "may be entering terminal phase". Decided by one doctor.
- 65m40s: "admitted 26th, by 27th already on midazolam & morphine, not fed for 11 days. Was crying.. hungry"
- 66m20s: "He tried to escape. They said he is immobile"... Tried four times.. Clothes cut off, catheterised to hinder escape.
- 68m00s: "Post-mortem was refused".
- 69m20s: "Stopped all his usual medication from home"
- 75m30s: "Punished for being unvaccinated".
- 79m00s: Consultant: "Last night, Stuart tried to escape again" and he was overdosed on midazolam
- (Elena asked to take out MZM because night-nurse would overdose &kill him).
- 79m40s: "He put Stuart's midazolam UP"

– 86m30s: Suart died 6th Nov, but records show he received more mzm on 7&8 Nov??

-

Testimony of Nurse Zandra Lewis

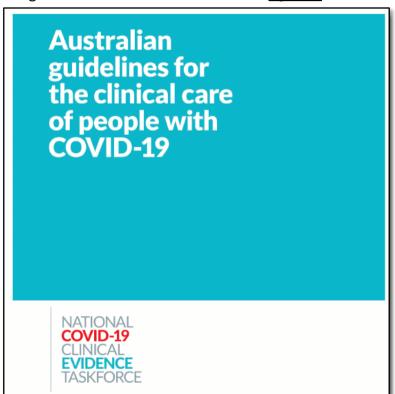
https://www.bitchute.com/video/45ZyErhMuL9c/

- "put all of them apart from two patients on TLC" .. "Liverpool Care Pathway"
- "Rather than try to give fluids and food...they didn't bother".
- "TLC were written up for morphine & midazolam and then they overodosed them"



Other countries had NG163 equivalents!

- "MAGICapp" was the source...
 - In every case, cheap effective repurposed-drug treatments were recommended *against*.
 - The documents have no listed authors.
 - Material is rebadged for every country.
 - Created by Pharma-invested entities.
 - MAGICapp promptly restricted access to UK archived versions.



Other countries had NG163 equivalents!

- "MAGICapp" was the source..
 - In every case, cheap effective repurposed-drug treatments were recommended against.
 - The documents have no listed authors.
 - Material is rebadged for every country.
 - Created by Pharma-invested entities.
 - MAGICapp promptly restricted access to UK archived versions.

Australian guidelines for the clinical care of people with COVID-19

5.1 - Antivirals and other disease modifying treatments

Consensus Recommendation

For patients with COVID-19 illness, only administer antiviral medications or other disease-modifying treatments in the context of clinical trials with appropriate ethical approval.



Other countries had NG163 equivalents!

- "MAGICapp" was the source...
 - In every case, cheap effective repurposed-drug treatments were recommended against.
 - The documents have no listed authors.
 - Material is rebadged for every country.
 - Created by Pharma-invested entities.
 - MAGICapp promptly restricted access to UK archived versions.

Australian guidelines for the clinical care of people with COVID-19

5.1 - Antivirals and other disease modifying treatments

Consensus Recommendation

For patients with COVID-19 illness, only administer antiviral medications or other disease-modifying treatments in the context of clinical trials with appropriate ethical approval.

https://app.magicapp.org/#/guideline/4959

"The MAGICapp will help your admins and authors with planning, authoring and publishing your guidelines according to standards"



- This is why we need to acknowledge the problem.
 - There is a conspiracy underlying it.

ONS Changes Excess Death Methodology. Malfeasance

- mid-crisis.

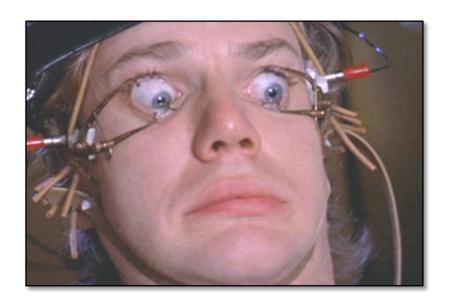
- Why are All-Cause Excess Deaths important?
 - The definition of a 'covid death' is not a factor.
- I have been calculating the All-Cause Excess Deaths based on the original ONS (UK Office For National Statistics) methodology for several years, by age-group.
- ONS always used a 5yr average (the 'average' for each week of the year is made up of 5 datapoints).
 - This changed in 2024.
 - Now, ONS 'bakes in' any excess into the next estimate, so any safety signal gets erased.
 - I have persisted with the old logic, for continuity purposes. Something the ONS strangely did not do!

2024 so far. Many anomalies >3 Std. Dev.

Events >=3SD are supposed to happen -by chance alone- about once in every 300 weeks (~6y). Week Ending Week# 2015-2019 2024 Excess Deaths Excess deaths 2024 Excess Death 2024 Excess Death Colours are indicative only: Standard Deviations calculated using all weeks in 2015-2019 - to obtain smoothest normal distributions 2015-2019 Deaths Fraction "2023 Deaths" / "2015-to-2019 mean" for Age & Week > 1.0 .5 sp Fraction "2023 Deaths" / "2015-to-2019 mean" > 1.0 Plus 0.5xS.D Fraction "2023 Deaths" / "2015-to-2019 mean" > 1.0 Plus 1.0xS.D .5 sp Fraction "2023 Deaths" / "2015-to-2019 mean" > 1.0 Plus 1.5xS.D $_{0}$ SD Fraction "2023 Deaths" / "2015-to-2019 mean" > 1.0 Plus 2.0xS.D 2.5 sp Fraction "2023 Deaths" / "2015-to-2019 mean" > 1.0 Plus 2.5xS.D osp Fraction "2023 Deaths" / "2015-to-2019 mean" > 1.0 Plus 3.0xS.D Fraction "2023 Deaths" / "2015-to-2019 mean" > 1.0 Plus **3.5**xS.D

So many things wrong with Vaccine Disaster asking your body to produce a toxin like SARS2 Spike...

- -Lab Origin.
- -Malfeasance Treatments and Evidence of Intent.
- Vaccine Disaster.
- Why it won't stop:
 - We can't just carry on, without future risks.



⁵⁹ Vaccines: already answered, imo. Vaccine Disaster

- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"
 - Tumor suppressor p53<->Spike interactions: https://pubmed.ncbi.nlm.nih.gov/34696485 (force-retracted, later confirmed by Prof W El-Deiry)
 - Neurodegenerative disease e.g. Korea 558k person study: Alzheimer's up by 23% just 3 months after vaccination: https://academic.oup.com/gimed/advance-article-abstract/doi/10.1093/gimed/hcae103/7684274 Roh et al.

- Auto-immune disease spike has many human protein analogues.
- Genetic damage / genotoxicity: heritable changes? LNPs accumulate in ovaries/testes.
- Inflammation: e.g. myo/pericarditis.
 - Cardiologists:1/35 subclinical after vaccine: https://esc365.escardio.org/presentation/255414 (paywalled) Free:https://docdro.id/0r22Fwj
- Immune system dysfunction: IgG4 'tolerance', Original Antigenic Sin, Imprinting...
- Many, many more.

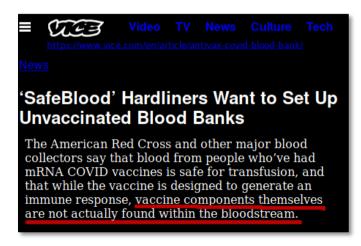
⁶⁰ Vaccines: already answered, imo. Vaccine Disaster

- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"
 - Tumor suppressor p53<->Spike interactions: https://pubmed.ncbi.nlm.nih.gov/34696485 (force-retracted, later confirmed by Prof W El-Deiry)
 - Neurodegenerative disease e.g. Korea 558k person study: Alzheimer's up by 23% just 3 months after vaccination: https://academic.oup.com/gimed/advance-article-abstract/doi/10.1093/gimed/hcae103/7684274 Roh et al.
 - "A potential association between COVID-19 vaccination and development of Alzheimer's Disease"
 - Auto-immune disease spike has many human protein analogues.
 - Genetic damage / genotoxicity: heritable changes? LNPs accumulate in ovaries/testes.
 - Inflammation: e.g. myo/pericarditis.
 - Cardiologists:1/35 subclinical after vaccine: https://esc365.escardio.org/presentation/255414 (paywalled) Free:https://docdro.id/0r22Fwj
 - Immune system dysfunction: IgG4 'tolerance', Original Antigenic Sin, Imprinting...
 - Many, many more.
- Persistence: no 'OFF' switch.
 - mRNA in blood for at least 28 days after vaccination: https://pubmed.ncbi.nlm.nih.gov/36647776/ Castruita et al.
 - Sixty days for vax-spike in lymph nodes: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8786601 Roeltgen et al.
 - Fifteen Months in monocytes: https://www.researchsquare.com/article/rs-1844677/v1 Patterson et al.
 - "SARS-CoV-2 S1 Protein Persistence in SARS-CoV-2 Negative Post-Vaccination Individuals with Long COVID/ PASC-Like Symptoms"

⁶¹ Vaccines: already answered, imo. Vaccine Disaster

- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"
 - Tumor suppressor p53<->Spike interactions: https://pubmed.ncbi.nlm.nih.gov/34696485 (force-retracted, later confirmed by Prof W El-Deiry)
 - Neurodegenerative disease e.g. Korea 558k person study: Alzheimer's up by 23% just 3 months after vaccination: https://academic.oup.com/gimed/advance-article-abstract/doi/10.1093/gimed/hcae103/7684274 Roh et al.

- Auto-immune disease spike has many human protein analogues.
- Genetic damage / genotoxicity: heritable changes? LNPs accumulate in ovaries/testes.
- Inflammation: e.g. myo/pericarditis.
 - Cardiologists:1/35 subclinical after vaccine: https://esc365.escardio.org/presentation/255414 (paywalled) Free:https://docdro.id/0r22Fwj
- Immune system dysfunction: IgG4 'tolerance', Original Antigenic Sin, Imprinting...
- Many, many more.
- Persistence: no 'OFF' switch.
 - mRNA in blood for at least 28 days after vaccination: https://pubmed.ncbi.nlm.nih.gov/36647776/ Castruita et al.
 - Sixty days for vax-spike in lymph nodes: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8786601 Roeltgen et al.
 - Fifteen Months in monocytes: https://www.researchsquare.com/article/rs-1844677/v1 Patterson et al.
 - "SARS-CoV-2 S1 Protein Persistence in SARS-CoV-2 Negative Post-Vaccination Individuals with Long COVID/ PASC-Like Symptoms"



⁶² Vaccines: already answered, imo. Vaccine Disaster

- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"
 - Tumor suppressor p53<->Spike interactions: https://pubmed.ncbi.nlm.nih.gov/34696485 (force-retracted, later confirmed by Prof W El-Deiry)
 - Neurodegenerative disease e.g. Korea 558k person study: Alzheimer's up by 23% just 3 months after vaccination: https://academic.oup.com/gimed/advance-article-abstract/doi/10.1093/gimed/hcae103/7684274 Roh et al.

- Auto-immune disease spike has many human protein analogues.
- Genetic damage / genotoxicity: heritable changes? LNPs accumulate in ovaries/testes.
- Inflammation: e.g. myo/pericarditis.
 - Cardiologists:1/35 subclinical after vaccine: https://esc365.escardio.org/presentation/255414 (paywalled) Free:https://docdro.id/0r22Fwj
- Immune system dysfunction: IgG4 'tolerance', Original Antigenic Sin, Imprinting...
- Many, many more.
- Persistence: no 'OFF' switch.
 - mRNA in blood for at least 28 days after vaccination: https://pubmed.ncbi.nlm.nih.gov/36647776/ Castruita et al.
 - Sixty days for vax-spike in lymph nodes: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8786601 Roeltgen et al.
 - Fifteen Months in monocytes: https://www.researchsquare.com/article/rs-1844677/v1 Patterson et al.
 - "SARS-CoV-2 S1 Protein Persistence in SARS-CoV-2 Negative Post-Vaccination Individuals with Long COVID/ PASC-Like Symptoms"



⁶³ Vaccines: already answered, imo. Vaccine Disaster

- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"
 - Tumor suppressor p53<->Spike interactions: https://pubmed.ncbi.nlm.nih.gov/34696485 (force-retracted, later confirmed by Prof W El-Deiry)
 - Neurodegenerative disease e.g. Korea 558k person study: Alzheimer's up by 23% just 3 months after vaccination: https://academic.oup.com/gimed/advance-article-abstract/doi/10.1093/gimed/hcae103/7684274 Roh et al.

- Auto-immune disease spike has many human protein analogues.
- Genetic damage / genotoxicity: heritable changes? LNPs accumulate in ovaries/testes.
- Inflammation: e.g. myo/pericarditis.
 - Cardiologists:1/35 subclinical after vaccine: https://esc365.escardio.org/presentation/255414 (paywalled) Free:https://docdro.id/0r22Fwj
- Immune system dysfunction: IgG4 'tolerance', Original Antigenic Sin, Imprinting...
- Many, many more.
- Persistence: no 'OFF' switch.
 - mRNA in blood for at least 28 days after vaccination: https://pubmed.ncbi.nlm.nih.gov/36647776/ Castruita et al.
 - Sixty days for vax-spike in lymph nodes: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8786601 Roeltgen et al.
 - Fifteen Months in monocytes: https://www.researchsquare.com/article/rs-1844677/v1 Patterson et al.
 - "SARS-CoV-2 S1 Protein Persistence in SARS-CoV-2 Negative Post-Vaccination Individuals with Long COVID/ PASC-Like Symptoms"

⁶⁴ Vaccines: already answered, imo. Vaccine Disaster

- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"
 - Tumor suppressor p53<->Spike interactions: https://pubmed.ncbi.nlm.nih.gov/34696485 (force-retracted, later confirmed by Prof W El-Deiry)
 - Neurodegenerative disease e.g. Korea 558k person study: Alzheimer's up by 23% just 3 months after vaccination: https://academic.oup.com/gimed/advance-article-abstract/doi/10.1093/gimed/hcae103/7684274 Roh et al.

- Auto-immune disease spike has many human protein analogues.
- Genetic damage / genotoxicity: heritable changes? LNPs accumulate in ovaries/testes.
- Inflammation: e.g. myo/pericarditis.
 - Cardiologists:1/35 subclinical after vaccine: https://esc365.escardio.org/presentation/255414 (paywalled) Free:https://docdro.id/0r22Fwj
- Immune system dysfunction: IgG4 'tolerance', Original Antigenic Sin, Imprinting...
- Many, many more.
- Persistence: no 'OFF' switch.
 - mRNA in blood for at least 28 days after vaccination: https://pubmed.ncbi.nlm.nih.gov/36647776/ Castruita et al.
 - Sixty days for vax-spike in lymph nodes: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8786601 Roeltgen et al.
 - Fifteen Months in monocytes: https://www.researchsquare.com/article/rs-1844677/v1 Patterson et al. "SARS-CoV-2 S1 Protein Persistence in SARS-CoV-2 Negative Post-Vaccination Individuals with Long COVID/ PASC-Like Symptoms"
- **Shedding:** not a myth. Circulating exosomes w/biologically active mRNA & (later) Spike, are found post-vax:
 - Not just mRNA, but Spike shedding too; https://www.jimmunol.org/content/jimmunol/207/10/2405.full.pdf "Circulating Exosomes with COVID Spike Protein Are Induced by BNT162b2"
 - Another one:- Vaccine mRNA Can Be Detected in Blood for 15d: https://www.mdpi.com/2227-9059/10/7/1538 Fertig et al.
 - They knew about, and were concerned regarding the risks of shedding.
 - Pfizer's own documents: https://www.scribd.com/document/484279338/C4591001-Pfizer-Clinical-Protocol-Nov2020
 - JAMA: mRNA Vaccine Shedding in Breast Milk Proven! https://igorchudov.substack.com/p/jama-vaccine-shedding-in-breast-

⁶⁵ Vaccines: already answered, imo.

Vaccine Disaster

- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"

Tu PF-07302048 (BNT162 RNA-Based COVID-19 Vaccines)

Neuro Protocol C4591001

https://www.scribd.com/document/484279338/C4591001-Pfizer-Clinical-Protocol-Nov2020

"A po 8.3.5.1. Exposure During Pregnancy

- An EDP occurs if:
- Genet
- A female participant is found to be pregnant while receiving or after discontinuing study intervention.
- Many.
 A male participant who is receiving or has discontinued study intervention exposes a female partner prior to or around the time of conception.
- Persistend
 - mRNA

https:

Immu

- Sixty of
- Fiftee"SARS
- Shedding:
 - Not ju<u>https:</u>
 - Anoth

- A female is found to be pregnant while being exposed or having been exposed to study intervention due to environmental exposure. Below are examples of environmental exposure during pregnancy:
 - A female family member or healthcare provider reports that she is pregnant after having been exposed to the study intervention by inhalation or skin contact.
 - A male family member or healthcare provider who has been exposed to the study intervention by inhalation or skin contact then exposes his female partner prior to or around the time of conception.
- They knew about, and were concerned regarding the risks of shedding.
- Pfizer's own documents: https://www.scribd.com/document/484279338/C4591001-Pfizer-Clinical-Protocol-Nov2020
- JAMA: mRNA Vaccine Shedding in Breast Milk Proven! https://igorchudov.substack.com/p/jama-vaccine-shedding-in-breast-

led by Prof W El-Deiry)
vaccination:

s://docdro.id/0r22Fwj

ita et al. n et al.

PASC-Like Symptoms"

und post-vax:

7/1538 Fertig et al.

⁶⁶ Vaccines: already answered, imo. Vaccine Disaster

- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"
 - Tumor suppressor p53<->Spike interactions: https://pubmed.ncbi.nlm.nih.gov/34696485 (force-retracted, later confirmed by Prof W El-Deiry)
 - Neurodegenerative disease e.g. Korea 558k person study: Alzheimer's up by 23% just 3 months after vaccination: https://academic.oup.com/gimed/advance-article-abstract/doi/10.1093/gimed/hcae103/7684274 Roh et al.

- Auto-immune disease spike has many human protein analogues.
- Genetic damage / genotoxicity: heritable changes? LNPs accumulate in ovaries/testes.
- Inflammation: e.g. myo/pericarditis.
 - Cardiologists:1/35 subclinical after vaccine: https://esc365.escardio.org/presentation/255414 (paywalled) Free:https://docdro.id/0r22Fwj
- Immune system dysfunction: IgG4 'tolerance', Original Antigenic Sin, Imprinting...
- Many, many more.
- Persistence: no 'OFF' switch.
 - mRNA in blood for at least 28 days after vaccination: https://pubmed.ncbi.nlm.nih.gov/36647776/ Castruita et al.
 - Sixty days for vax-spike in lymph nodes: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8786601 Roeltgen et al.
 - Fifteen Months in monocytes: https://www.researchsquare.com/article/rs-1844677/v1 Patterson et al. "SARS-CoV-2 S1 Protein Persistence in SARS-CoV-2 Negative Post-Vaccination Individuals with Long COVID/ PASC-Like Symptoms"
- **Shedding:** not a myth. Circulating exosomes w/biologically active mRNA & (later) Spike, are found post-vax:
 - Not just mRNA, but Spike shedding too; https://www.jimmunol.org/content/jimmunol/207/10/2405.full.pdf "Circulating Exosomes with COVID Spike Protein Are Induced by BNT162b2"
 - Another one:- Vaccine mRNA Can Be Detected in Blood for 15d: https://www.mdpi.com/2227-9059/10/7/1538 Fertig et al.
 - They knew about, and were concerned regarding the risks of shedding.
 - Pfizer's own documents: https://www.scribd.com/document/484279338/C4591001-Pfizer-Clinical-Protocol-Nov2020
 - JAMA: mRNA Vaccine Shedding in Breast Milk Proven! https://igorchudov.substack.com/p/jama-vaccine-shedding-in-breast-

⁶⁷ Vaccines: already answered, imo. Vaccine Disaster

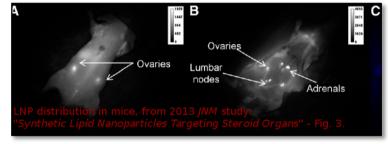
- **Disease-causing:** abundant clinical evidence AND mechanisms of action:
 - Cancer / oncogenesis e.g. at site of injection! https://x.com/ /status/1745616252599464287
 - Using 'pseudo-uridine': https://pubmed.ncbi.nlm.nih.gov/38583833/ "N1-methyl-pseudouridine (m1\P): Friend or foe of cancer?"
 - Tumor suppressor p53<->Spike interactions: https://pubmed.ncbi.nlm.nih.gov/34696485 (force-retracted, later confirmed by Prof W El-Deiry)
 - Neurodegenerative disease e.g. Korea 558k person study: Alzheimer's up by 23% just 3 months after vaccination: https://academic.oup.com/gimed/advance-article-abstract/doi/10.1093/gimed/hcae103/7684274 Roh et al.

- Auto-immune disease spike has many human protein analogues.
- Genetic damage / genotoxicity: heritable changes? LNPs accumulate in ovaries/testes.
- Inflammation: e.g. myo/pericarditis.
 - Cardiologists:1/35 subclinical after vaccine: https://esc365.escardio.org/presentation/255414 (paywalled) Free:https://docdro.id/0r22Fwj
- Immune system dysfunction: IgG4 'tolerance', Original Antigenic Sin, Imprinting...
- Many, many more.
- Persistence: no 'OFF' switch.
 - mRNA in blood for at least 28 days after vaccination: https://pubmed.ncbi.nlm.nih.gov/36647776/ Castruita et al.
 - Sixty days for vax-spike in lymph nodes: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8786601 Roeltgen et al.
 - Fifteen Months in monocytes: https://www.researchsquare.com/article/rs-1844677/v1 Patterson et al.
 - "SARS-CoV-2 S1 Protein Persistence in SARS-CoV-2 Negative Post-Vaccination Individuals with Long COVID/ PASC-Like Symptoms"
- **Shedding:** not a myth. Circulating exosomes w/biologically active mRNA & (later) Spike, are found post-vax:
 - Not just mRNA, but Spike shedding too; https://www.jimmunol.org/content/jimmunol/207/10/2405.full.pdf "Circulating Exosomes with COVID Spike Protein Are Induced by BNT162b2"
 - Another one:- Vaccine mRNA Can Be Detected in Blood for 15d: https://www.mdpi.com/2227-9059/10/7/1538 Fertig et al.
 - They knew about, and were concerned regarding the risks of shedding.
 - Pfizer's own documents: https://www.scribd.com/document/484279338/C4591001-Pfizer-Clinical-Protocol-Nov2020
 - JAMA: mRNA Vaccine Shedding in Breast Milk Proven! https://igorchudov.substack.com/p/jama-vaccine-shedding-in-breast-milk
- **<u>Bio-distribution:</u>** crosses ovarian, testicular, placental & blood-brain barriers, thanks to LNP's.
 - Leaked by Byram Bridle: https://www.naturalnews.com/files/Pfizer-bio-distribution-confidential-document-translated-to-english.pdf

⁶⁸ Biodistribution..They Knew & Lied Vaccine Disaster

Malfeasance

- Bio-distribution of mRNA is far wider than they told you:
 - The fatty Lipid Nanoparticles accumulate in any organ using fats/cholesterol.
 - They knew this principle, and hid it from people taking the vaccines.
 - A 2013 study using LNPs in mice, from the Journal of Nuclear Medicine: https://jnm.snmjournals.org/content/jnumed/54/11/1996.full.pdf Synthetic Lipid Nanoparticles Targeting Steroid Organs (2013).
 - Note: The heart uses fats for energy, thus ensuring some heart-cell transfection from every dose. ALL transfected cells are killed.
 - Heart muscle cells do not re-grow.



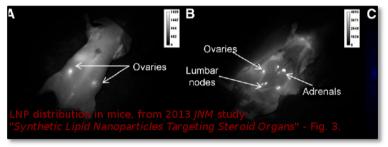
Biodistribution..They Knew & Lied

Vaccine Disaster

Malfeasance

- Bio-distribution of mRNA is far wider than they told you:
 - The fatty Lipid Nanoparticles accumulate in any organ using fats/cholesterol.
 - They knew this principle, and hid it from people taking the vaccines.
 - A 2013 study using LNPs in mice, from the Journal of Nuclear Medicine:
 https://jnm.snmjournals.org/content/jnumed/54/11/1996.full.pdf

 Synthetic Lipid Nanoparticles Targeting Steroid Organs (2013).
 - Note: The heart uses fats for energy, thus ensuring some heart-cell transfection from every dose. ALL transfected cells are killed.
 - Heart muscle cells do not re-grow.



<u>Dendritic Cells were always the target</u>, aside from known accumulation in organs.

This is BioNTech founder Ozlem Tureci giving a talk titled:
 "The Bodyhack: bringing mRNA to the right cells at the right places"

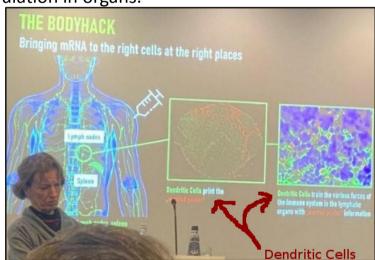


Image tweeted by French MP Virginie Joron

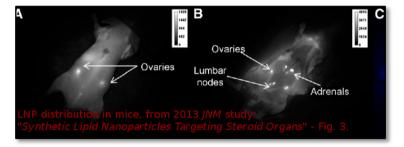
⁷⁰ Biodistribution..They Knew & Lied

Vaccine Disaster

Malfeasance

- Bio-distribution of mRNA is far wider than they told you:
 - The fatty Lipid Nanoparticles accumulate in any organ using fats/cholesterol.
 - They knew this principle, and hid it from people taking the vaccines.
 - A 2013 study using LNPs in mice, from the Journal of Nuclear Medicine:
 https://jnm.snmjournals.org/content/jnumed/54/11/1996.full.pdf

 Synthetic Lipid Nanoparticles Targeting Steroid Organs (2013).
 - Note: The heart uses fats for energy, thus ensuring some heart-cell transfection from every dose. ALL transfected cells are killed.
 - Heart muscle cells do not re-grow.



<u>Dendritic Cells were always the target</u>, aside from known accumulation in organs.

- This is BioNTech founder Ozlem Tureci giving a talk titled:
 "The Bodyhack: bringing mRNA to the right cells at the right places"
- The mRNA & Spike made it to places needed to do severe harm
 - e.g. Brain, Kidneys, Liver, Ovaries, Testes...
 - Means biologically active mRNA/Spike affected babies in the womb.
 - Note Dendritic Cells have DC-SIGN receptors.
 - Dendritic Cells are often found adjacent to nerve fibres/neurons,
 which is bad news for amyloid/prion material reaching the brain.
 - There is considered to be NO SAFE LOWER LIMIT to prions in-brain.
 Just one can spark a chain reaction.

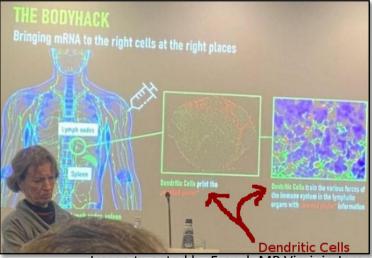


Image tweeted by French MP Virginie Joron

- The wide spread of the material provides potential for harm all over the body.
 - The next few slides will focus on five remaining unanswered questions, in my mind.

Q1:Impact of Vax DNA Contamination? Vaccine Disaster

mRNA is produced using E.coli bacteria, with artificial circular DNA loops called plasmids.

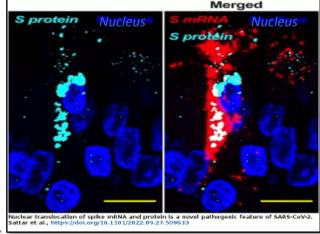
Q1:Impact of Vax DNA Contamination? Vaccine Disaster

- **mRNA** is **produced** using *E.coli* bacteria, with artificial **circular DNA** loops called **plasmids**.
- The Pfizer plasmid is proprietary, but has been sequenced independently. Many surprises:
 - Pfizer used an SV40 mammalian promoter (SV40 is a Simian virus, known to cause cancer).
 - There were Kanamycin & Neomycin antibiotic resistance genes in the plasmid.
 - STOP codons were doubled: they knew there was a risk of 'slippy' transcription when using pseudo-uridine.
 - The plasmid's reverse-strand encoded for a silk-like protein. Silk is known to stabilise mRNA.

- **mRNA** is **produced** using *E.coli* bacteria, with artificial **circular DNA** loops called **plasmids**.
- The Pfizer plasmid is proprietary, but has been sequenced independently. Many surprises:
 - Pfizer used an SV40 mammalian promoter (SV40 is a Simian virus, known to cause cancer).
 - There were Kanamycin & Neomycin antibiotic resistance genes in the plasmid.
 - STOP codons were doubled: they knew there was a risk of 'slippy' transcription when using pseudo-uridine.
 - The plasmid's reverse-strand encoded for a silk-like protein. Silk is known to stabilise mRNA.
- It was found that plasmid DNA was detectable in Pfizer vials at many times the (outdated) supposed limits set by regulators.
 - Speicher et al.: https://osf.io/preprints/osf/mjc97
 - McKernan et al.: https://osf.io/preprints/osf/b9t7m
 - Konig et al.: https://www.mdpi.com/2409-9279/7/3/41#B6-mps-07-00041
 - These limits should be lower (zero!) when LNP-encapsulated DNA is involved!
 - The LNP facilitates entry of the DNA fragments into the cell & nucleus.

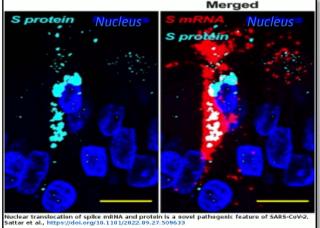
- mRNA is produced using E.coli bacteria, with artificial circular DNA loops called plasmids.
- The Pfizer plasmid is proprietary, but has been sequenced independently. Many surprises:
 - Pfizer used an SV40 mammalian promoter (SV40 is a Simian virus, known to cause cancer).
 - There were Kanamycin & Neomycin antibiotic resistance genes in the plasmid.
 - STOP codons were doubled: they knew there was a risk of 'slippy' transcription when using pseudo-uridine.
 - The plasmid's reverse-strand encoded for a silk-like protein. Silk is known to stabilise mRNA.

- Speicher et al.: https://osf.io/preprints/osf/mjc97
- McKernan et al.: https://osf.io/preprints/osf/b9t7m
- Konig et al.: https://www.mdpi.com/2409-9279/7/3/41#B6-mps-07-00041
- These limits should be lower (zero!) when LNP-encapsulated DNA is involved!
- The LNP facilitates entry of the DNA fragments into the cell & nucleus.
- **DNA fragments in the cell nucleus are a risk** for insertion / mutagenesis.
 - The SV40 promoter is known to traffic DNA/RNA fragments to the nucleus: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4152905 (1999)
 - "The 72bp repeats of the SV40 enhancer facilitate maximal transport"
 - Observation of Spike+mRNA in nucleus:



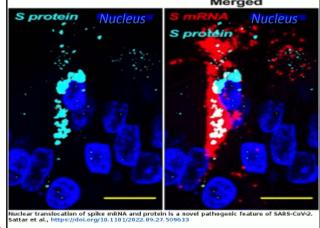
- mRNA is produced using E.coli bacteria, with artificial circular DNA loops called plasmids.
- The Pfizer plasmid is proprietary, but has been sequenced independently. Many surprises:
 - Pfizer used an SV40 mammalian promoter (SV40 is a Simian virus, known to cause cancer).
 - There were Kanamycin & Neomycin antibiotic resistance genes in the plasmid.
 - STOP codons were doubled: they knew there was a risk of 'slippy' transcription when using pseudo-uridine.
 - The plasmid's reverse-strand encoded for a silk-like protein. Silk is known to stabilise mRNA.

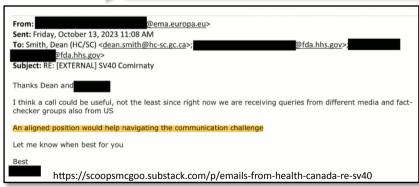
- Speicher et al.: https://osf.io/preprints/osf/mjc97
- McKernan et al.: https://osf.io/preprints/osf/b9t7m
- Konig et al.: https://www.mdpi.com/2409-9279/7/3/41#B6-mps-07-00041
- These limits should be lower (zero!) when LNP-encapsulated DNA is involved!
- The LNP facilitates entry of the DNA fragments into the cell & nucleus.
- **DNA fragments in the cell nucleus are a risk** for insertion / mutagenesis.
 - The SV40 promoter is known to traffic DNA/RNA fragments to the nucleus: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4152905 (1999)
 - "The 72bp repeats of the SV40 enhancer facilitate maximal transport"
 - Observation of Spike+mRNA in nucleus:
- Contamination has been confirmed by Health Canada, the EMA, FDA...
 - There are emails from regulators showing they colluded in their response to this.
 - Pfizer <u>deleted</u> the labelling of their SV40 promoter in the submission to the Euro Medicines Agency, which contravenes rules on mammalian promoter declaration.
 - Should've been a 'slam-dunk' for any regulator: 'adulterated'.



- mRNA is produced using E.coli bacteria, with artificial circular DNA loops called plasmids.
- The Pfizer plasmid is proprietary, but has been sequenced independently. Many surprises:
 - Pfizer used an SV40 mammalian promoter (SV40 is a Simian virus, known to cause cancer).
 - There were Kanamycin & Neomycin antibiotic resistance genes in the plasmid.
 - STOP codons were doubled: they knew there was a risk of 'slippy' transcription when using pseudo-uridine.
 - The plasmid's reverse-strand encoded for a silk-like protein. Silk is known to stabilise mRNA.

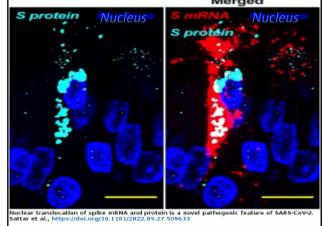
- Speicher et al.: https://osf.io/preprints/osf/mjc97
- McKernan et al.: https://osf.io/preprints/osf/b9t7m
- Konig et al.: https://www.mdpi.com/2409-9279/7/3/41#B6-mps-07-00041
- These limits should be lower (zero!) when LNP-encapsulated DNA is involved!
- The LNP facilitates entry of the DNA fragments into the cell & nucleus.
- **DNA fragments in the cell nucleus are a risk** for insertion / mutagenesis.
 - The SV40 promoter is known to traffic DNA/RNA fragments to the nucleus: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4152905 (1999)
 - "The 72bp repeats of the SV40 enhancer facilitate maximal transport"
 - Observation of Spike+mRNA in nucleus:
- **Contamination has been confirmed** by Health Canada, the EMA, FDA...
 - There are emails from regulators showing they colluded in their response to this.
 - Pfizer deleted the labelling of their SV40 promoter in the submission to the Euro Medicines Agency, which contravenes rules on mammalian promoter declaration.
 - Should've been a 'slam-dunk' for any regulator: 'adulterated'

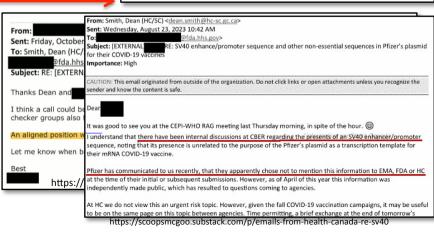




- mRNA is produced using E.coli bacteria, with artificial circular DNA loops called plasmids.
- The Pfizer plasmid is proprietary, but has been sequenced independently. Many surprises:
 - Pfizer used an SV40 mammalian promoter (SV40 is a Simian virus, known to cause cancer).
 - There were Kanamycin & Neomycin antibiotic resistance genes in the plasmid.
 - STOP codons were doubled: they knew there was a risk of 'slippy' transcription when using pseudo-uridine.
 - The plasmid's reverse-strand encoded for a silk-like protein. Silk is known to stabilise mRNA.

- Speicher et al.: https://osf.io/preprints/osf/mjc97
- McKernan et al.: https://osf.jo/preprints/osf/b9t7m
- Konig et al.: https://www.mdpi.com/2409-9279/7/3/41#B6-mps-07-00041
- These limits should be lower (zero!) when LNP-encapsulated DNA is involved!
- The LNP facilitates entry of the DNA fragments into the cell & nucleus.
- **DNA fragments in the cell nucleus are a risk** for insertion / mutagenesis.
 - The SV40 promoter is known to traffic DNA/RNA fragments to the nucleus: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4152905 (1999)
 - "The 72bp repeats of the SV40 enhancer facilitate maximal transport"
 - Observation of Spike+mRNA in nucleus:
- Contamination has been confirmed by Health Canada, the EMA, FDA...
 - There are emails from regulators showing they colluded in their response to this.
 - Pfizer deleted the labelling of their SV40 promoter in the submission to the Euro Medicines Agency, which contravenes rules on mammalian promoter declaration.
 - Should've been a 'slam-dunk' for any regulator: 'adulterated'







This is a technologically difficult thing to confirm!

- Kevin McKernan has designed a test... Long read! https://anandamide.substack.com/p/targeted-enrichment-of-vaccine-dna
- PJ Buckhaults has offered to test patients.
- Buckhaults' testimony to the SC Senate stated that the risk of integration goes up with smaller, more numerous fragments of DNA contamination.
- Note that none of the agencies tasked with public safety are doing (or funding) this work.

- This is a technologically difficult thing to confirm!
 - Kevin McKernan has designed a test... Long read! https://anandamide.substack.com/p/targeted-enrichment-of-vaccine-dna
 - PJ Buckhaults has offered to test patients.
 - Buckhaults' testimony to the SC Senate stated that the risk of integration goes up with smaller, more numerous fragments of DNA contamination.
 - Note that none of the agencies tasked with public safety are doing (or funding) this work.
- **EVIDENCE #1**: Putative integration events in human cancer cell-lines.
 - Kevin McKernan published the discovery: https://anandamide.substack.com/p/vaccine-targeted-qpcr-of-cancer-cell

- This is a technologically difficult thing to confirm!
 - Kevin McKernan has designed a test... Long read! https://anandamide.substack.com/p/targeted-enrichment-of-vaccine-dna
 - PJ Buckhaults has offered to test patients.
 - Buckhaults' testimony to the SC Senate stated that the risk of integration goes up with smaller, more numerous fragments of DNA contamination.
 - Note that <u>none</u> of the agencies tasked with public safety are doing (or funding) this work.
- **EVIDENCE #1**: Putative integration events in human cancer cell-lines.
 - Kevin McKernan published the discovery: https://anandamide.substack.com/p/vaccine-targeted-qpcr-of-cancer-cell
- **EVIDENCE #2:** Prof. Angus Dalgleish (NHS oncologist) describes his colleague's tumor biopsy revealed vaccine mRNA integration into the tumor cell DNA.
 - https://www.youtube.com/watch?v=PnJ5T1Enwq4 at 23m40s...

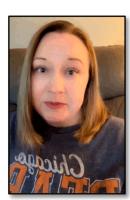


- This is a technologically difficult thing to confirm!
 - Kevin McKernan has designed a test... Long read! https://anandamide.substack.com/p/targeted-enrichment-of-vaccine-dna
 - PJ Buckhaults has offered to test patients.
 - Buckhaults' testimony to the SC Senate stated that the risk of integration goes up with smaller, more numerous fragments of DNA contamination.
 - Note that <u>none</u> of the agencies tasked with public safety are doing (or funding) this work.
- **EVIDENCE #1**: Putative integration events in human cancer cell-lines.
 - Kevin McKernan published the discovery: https://anandamide.substack.com/p/vaccine-targeted-apcr-of-cancer-cell
- <u>EVIDENCE #2</u>: Prof. Angus Dalgleish (NHS oncologist) describes his colleague's tumor biopsy revealed vaccine mRNA integration into the tumor cell DNA.
 - https://www.youtube.com/watch?v=PnJ5T1Enwq4 at 23m40s...



- **EVIDENCE #3**: mRNA->DNA in human liver cell line <u>inside 6h</u>: https://www.mdpi.com/1467-3045/44/3/73
- **EVIDENCE #4**: Adenovirus vectors can integrate: https://ncbi.nlm.nih.gov/pmc/articles/PMC8168329
- **BREAKING:** The first human victm of DNA Integration identified?
 - A lady (I bet she is working with PJ Buckhaults' lab) says she has
 Moderna and Pfizer in her mitochondrial DNA, Lungs, Heart and Liver.
 - "Stay tuned" ... A paper will be on the way, if this is credible.

https://x.com/ericawithac13/status/1819191379232231518



Now at least 12 papers documenting this post-vaccine injury.

Dr. Mikolaj Raszek/ Merogenomics	Farkash etal. 2021	Emmenegger et al. 2022	irrgang et al. 2022	Buhre et al. 2023	Kiszel et al. 2023	Selva et al. 2023	Valketal. 2023	Hartey et al. 2023	Sheehan et al. 2023	Yoshimura et al. 2023	Akhtar et al. 2023	Espino et al. 2024
IgG4 rise only in mRNA vaccinees			X	X	X	X	X	X				
More potent post Moderna than Pfizer				X								
Marginal within month post 2X mRNA	X									X		
Post 2X mRNA after >4 months			X	X					X	X	X	
2X mRNA + >90 days breakthrough infection		X	X			X						
Very common 3X mRNA		X	X		X	X	X	X	X	X	X	X
3X mRNA + breakthrough infection increases IgG4 more		X	X		X	X		X		X		
Persists post 4X, 5X mRNA										X	X	X
Heterologous mRNA combo also induces IgG4 but lesser extent			X	X								
mRNA boosting post other vaccine type induces IgG4						X					X	
Major contributor + can become primary anti-S Ab post 3X mRNA			Х		X		X				X	X
IgG4 rise is accompanied by IL-10 rise											X	
Reduces Fc-mediated effector functions			X			X						
Natural infection prior to mRNA protects				X	X	X						X
Unvaccinated infected do not exhibit IgG4 rise		X			X					X		X
IgG4 rise accompanied by IgG4 memory B cells increase post 2X or subsequent boosters			X					X				
IgG4 rise accompanied by enhanced cross-reactivity and/or shift in targeted epitopes				X				X				X
Observed in blood	X	X	X	X	X	X	X	X	X	X	X	X
Observed in mucosa						X						
Suppressed by immunosuppressive drugs							X					
Does not increase breakthrough infections										X		

- Now at least 12 papers documenting this post-vaccine injury.
- Likely aetiology in cancers (IgG4 essential role in cancer doi: 10.1136/jitc-2020-000661), auto-immunity, fibrosis...
- IgG4 promotes tolerance; similar to reaction with pollen.
 - Except SARS2 is a replicating pathogen!

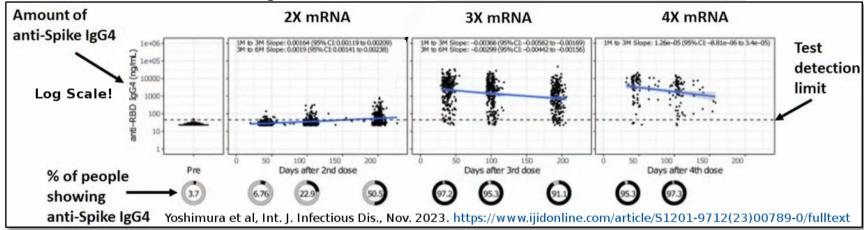


- Now at least 12 papers documenting this post-vaccine injury.
- Likely aetiology in cancers (IgG4 essential role in cancer doi: 10.1136/jitc-2020-000661), auto-immunity, fibrosis...
- IgG4 promotes tolerance; similar to reaction with pollen.
 - Except SARS2 is a replicating pathogen!

https://doi.org/10.3390/vaccines11050991

Review
IgG4 Antibodies Induced by Repeated Vaccination May
Generate Immune Tolerance to the SARS-CoV-2 Spike Protein
Vladimir N. Uversky 1.4®, Elrashdy M. Redwan 2.3®, William Makis 4 and Alberto Rubio-Casillas 5.4®

Studies confirm 3x mRNA induces IgG4 Class-Shift:



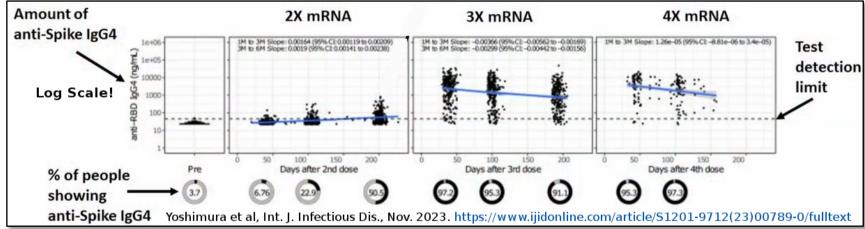
⁻ In Children, just two jabs can induce the class-shift: https://journals.lww.com/pidj/fulltext/9900/delayed induction of noninflammatory sars cov 2.959.aspx

- Now at least 12 papers documenting this post-vaccine injury.
- Likely aetiology in cancers (IgG4 essential role in cancer doi: 10.1136/jitc-2020-000661), auto-immunity, fibrosis...
- IgG4 promotes tolerance; similar to reaction with pollen.
 - Except SARS2 is a replicating pathogen!

https://doi.org/10.3390/vaccines11050991

Review
IgG4 Antibodies Induced by Repeated Vaccination May
Generate Immune Tolerance to the SARS-CoV-2 Spike Protein
Vladimir N. Uversky 1-40, Etrashdy M. Redwan 2-30, William Makis 4 and Alberto Rubio-Casillas 5-40

Studies confirm 3x mRNA induces IgG4 Class-Shift:



- In Children, just two jabs can induce the class-shift: https://journals.lww.com/pidi/fulltext/9900/delayed_induction_of_noninflammatory_sars_cov_2.959.aspx
- IgG4 remains high for at least 2 years: https://www.mdpi.com/1999-4915/16/2/187 Espino et al.
- Vaccination impairs *de novo* immune response to Omicron: a precondition for Original Antigenic Sin https://doi.org/10.21203/rs.3.rs-3579996/v1 *Pusnik et al.*
- Children: sustained decreases in response to viral and bacterial stimulants after Pfizer: https://frontiersin.org/articles/10.3389/fimmu.2023.1242380/full Noe et al.

- Now at least 12 papers documenting this post-vaccine injury.
- Likely aetiology in cancers (IgG4 essential role in cancer doi: 10.1136/jitc-2020-000661), auto-immunity, fibrosis...
- IgG4 promotes tolerance; similar to reaction with pollen.
 - Except SARS2 is a replicating pathogen!

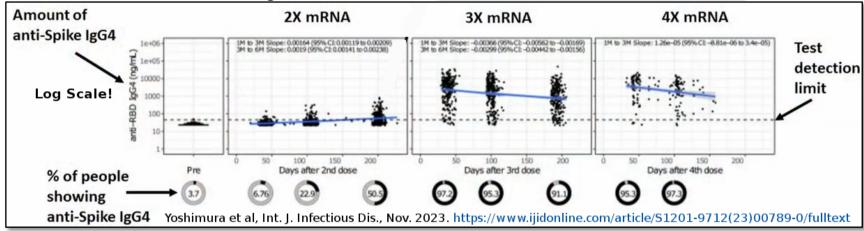
https://doi.org/10.3390/vaccines11050991

Review

IgG4 Antibodies Induced by Repeated Vaccination May
Generate Immune Tolerance to the SARS-CoV-2 Spike Protein

Vladimir N. Uversky 1-0, Etrashdy M. Redwan 2-30, William Makis 4 and Alberto Rubio-Casillas 54-0

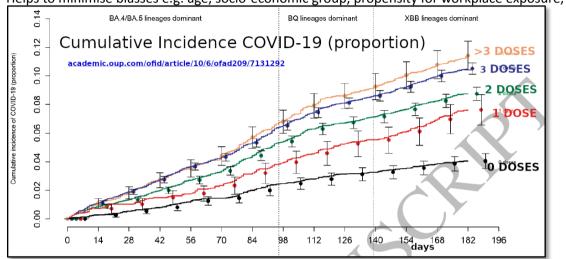
Studies confirm 3x mRNA induces IgG4 Class-Shift:



- In Children, just two jabs can induce the class-shift: https://journals.lww.com/pidi/fulltext/9900/delayed_induction_of_noninflammatory_sars_cov_2.959.aspx
- IgG4 remains high for at least 2 years: https://www.mdpi.com/1999-4915/16/2/187 Espino et al.
- Vaccination impairs *de novo* immune response to Omicron: a precondition for Original Antigenic Sin https://doi.org/10.21203/rs.3.rs-3579996/v1 *Pusnik et al.*
- Children: sustained decreases in response to viral and bacterial stimulants after Pfizer: https://frontiersin.org/articles/10.3389/fimmu.2023.1242380/full Noe et al.
- Notable that HIV's gp120 protein induces IgG4 class-shift.. (remember SARS2 has HIV inserts!) https://journals.aai.org/jimmunol/article/176/7/3931/73688/HIV-1-Envelope-Triggers-Polyclonal-Ig-Class-Switch
 "In the presence of gp120... B-Cells ... undergo CSR from IgM to IgG", "some HIV-1 Ags elicit IgG class switch".

• Cleveland Clinic studied their own staff (n=51k)...



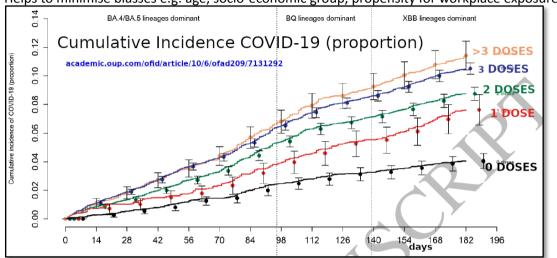


Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)

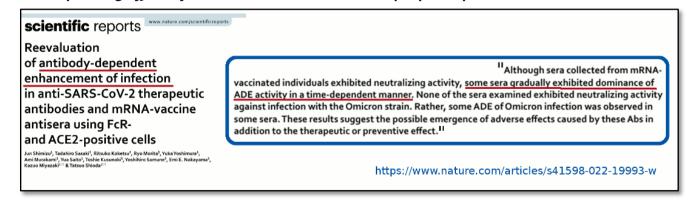
• Cleveland Clinic studied their own staff (n=51k)...





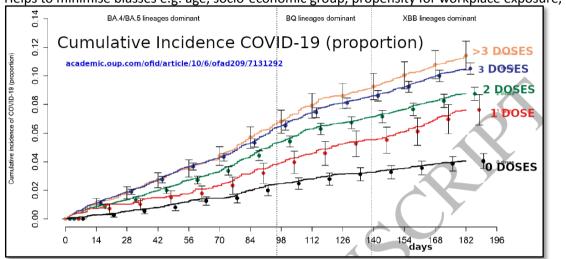
Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)



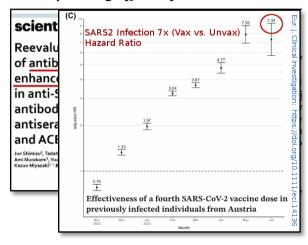
• Cleveland Clinic studied their own staff (n=51k)...





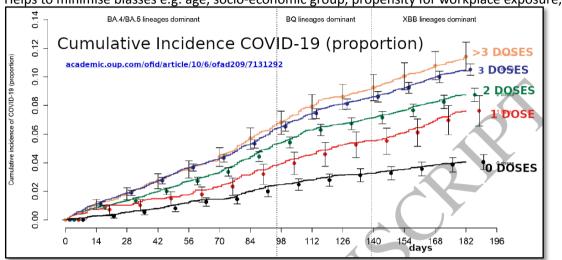
Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)



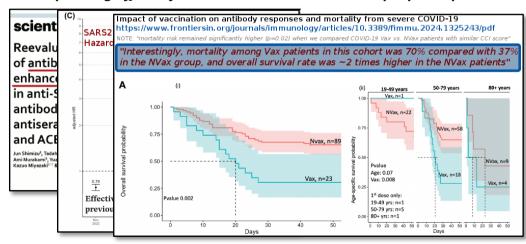
• Cleveland Clinic studied their own staff (n=51k)...





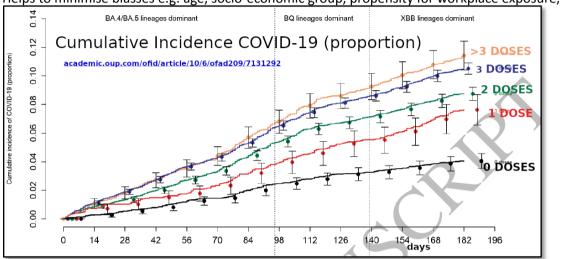
Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)



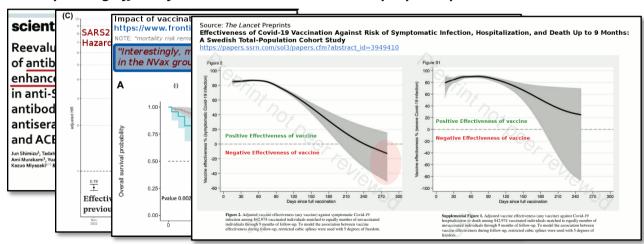
• Cleveland Clinic studied their own staff (n=51k)...



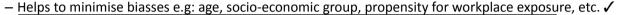


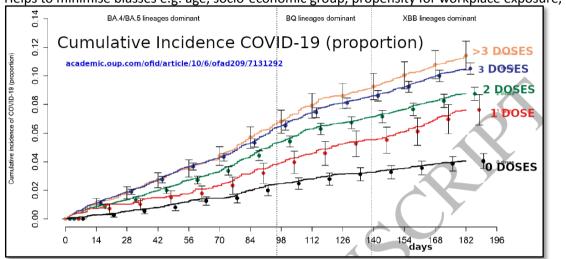
Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)



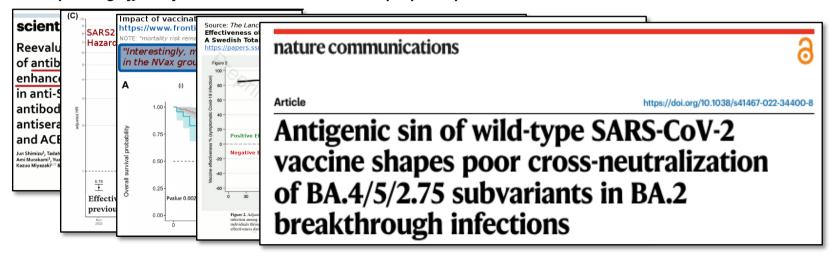
• Cleveland Clinic studied their own staff (n=51k)...





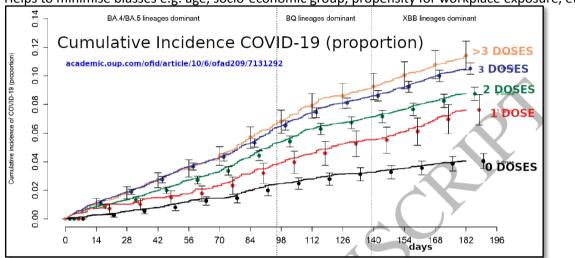
Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)



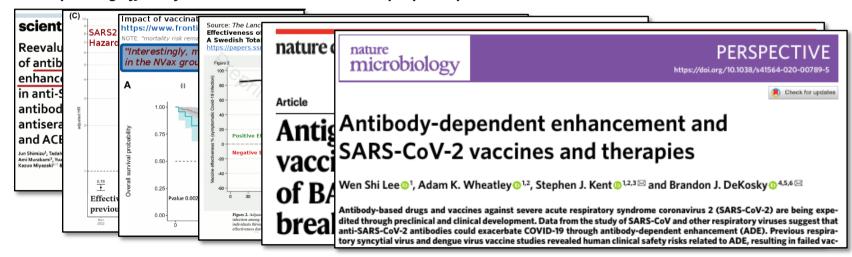
• Cleveland Clinic studied their own staff (n=51k)...





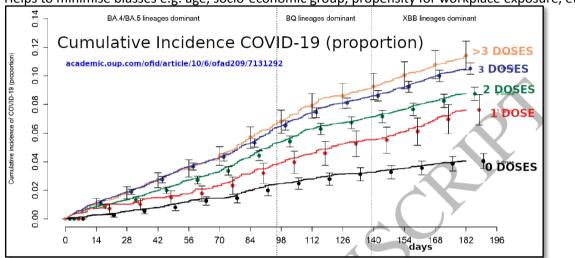
Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)



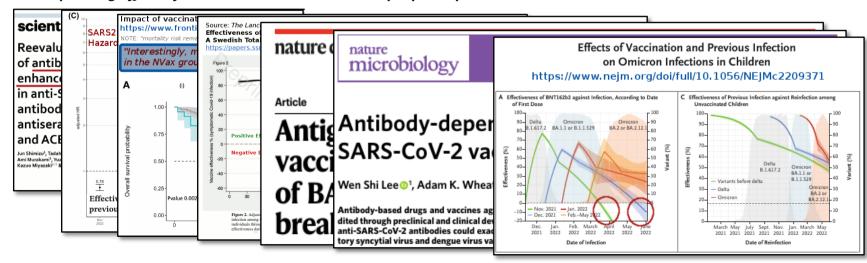
• Cleveland Clinic studied their own staff (n=51k)...



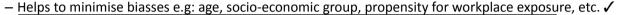


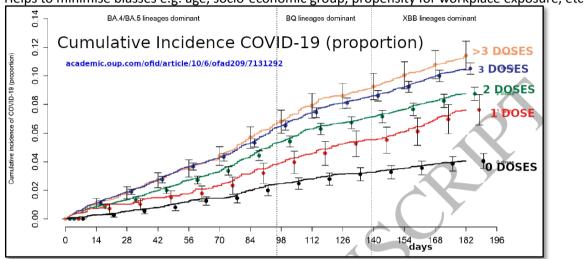
Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)



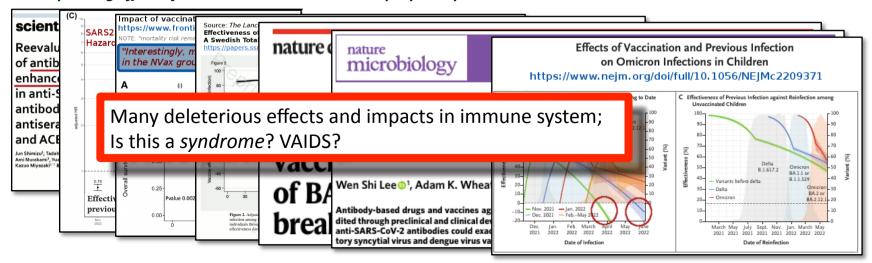
• Cleveland Clinic studied their own staff (n=51k)...





Studies confirm....

https://www.bmj.com/content/381/bmj-2022-074404/rr-0 (includes literature survey)



Q4: Amyloidosis & Prion issues Vaccine Disaster

Basic Amyloid/Prion Biochemistry:

- Amyloid is causative/associated with many diseases (prions cause a form of amyloid):
 - Parkinson's, Alzheimer's, ALS, vCJD, some forms of fibrosis etc., and BSE in cows, CWD in deer.
 - Prions are not destroyed by cooking, bleaching, etc. 1/2000 extracted UK appendices found to contain BSE prions!
 - Prions hijack healthy body proteins and misfold them, causing <u>a chain-reaction</u> leading to death: "uniformly fatal".
 - There is NO SAFE LOWER LIMIT for prions in the brain.
- There is a clear mechanism for amyloid formation, catalysed by SARS2-spike epitopes.
 - IMPORTANT: just specific pieces of Spike are needed- 'epitopes'. Virus not required. Not even whole spike!

Q4: Amyloidosis & Prion issues Vaccine Disaster

Basic Amyloid/Prion Biochemistry:

- Amyloid is causative/associated with many diseases (prions cause a form of amyloid):
 - Parkinson's, Alzheimer's, ALS, vCJD, some forms of fibrosis etc., and BSE in cows, CWD in deer.
 - Prions are not destroyed by cooking, bleaching, etc. 1/2000 extracted UK appendices found to contain BSE prions!
 - Prions hijack healthy body proteins and misfold them, causing a chain-reaction leading to death: "uniformly fatal".
 - There is NO SAFE LOWER LIMIT for prions in the brain.
- There is a clear mechanism for amyloid formation, catalysed by SARS2-spike epitopes.
 - IMPORTANT: just specific pieces of Spike are needed- 'epitopes'. Virus not required. Not even whole spike!

Predictable problems with SARS2-spike 'seeding' amyloid formation:

- Hammerstrom & Nystrom identified dangerous sections of spike, others see it too: Tetz & Tetz, Classen..
 - https://www.biorxiv.org/content/10.1101/2023.09.01.555834v1

"SARS-CoV-2 Spike ... specifically & selectively accelerates amyloid fibril formation of human prion protein and the amyloid-8 peptide"

Q4: Amyloidosis & Prion issues

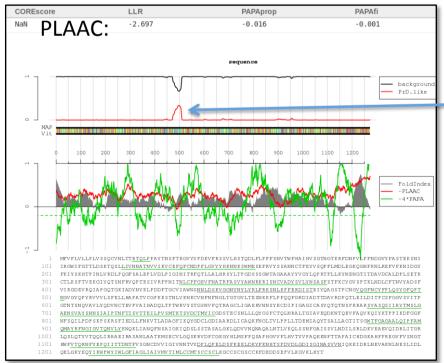
Vaccine Disaster

Basic Amyloid/Prion Biochemistry:

- Amyloid is causative/associated with many diseases (prions cause a form of amyloid):
 - Parkinson's, Alzheimer's, ALS, vCJD, some forms of fibrosis etc., and BSE in cows, CWD in deer.
 - Prions are not destroyed by cooking, bleaching, etc. 1/2000 extracted UK appendices found to contain BSE prions!
 - Prions hijack healthy body proteins and misfold them, causing <u>a chain-reaction</u> leading to death: "uniformly fatal".
 - There is NO SAFE LOWER LIMIT for prions in the brain.
- There is a clear mechanism for amyloid formation, *catalysed by SARS2-spike epitopes*.
 - IMPORTANT: just specific pieces of Spike are needed- 'epitopes'. Virus not required. Not even whole spike!

Predictable problems with SARS2-spike 'seeding' amyloid formation:

- Hammerstrom & Nystrom identified dangerous sections of spike, others see it too: Tetz & Tetz, Classen...
 - https://www.biorxiv.org/content/10.1101/2023.09.01.555834v1
 - "SARS-CoV-2 Spike ...specifically & selectively accelerates amyloid fibril formation of human prion protein and the amyloid-6 peptide"
 - I also identified regions & verified this myself using PLAAC & WALTZ (https://waltzdb.switchlab.org):



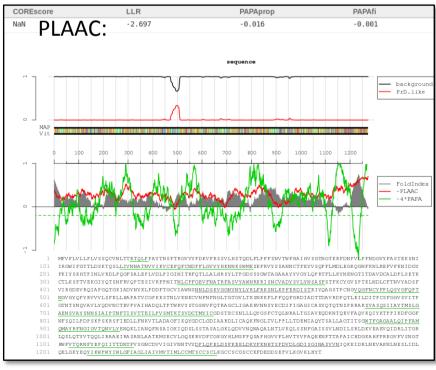
Large 'bump' in prion-ergic tendency around position 500 of the spike sequence.

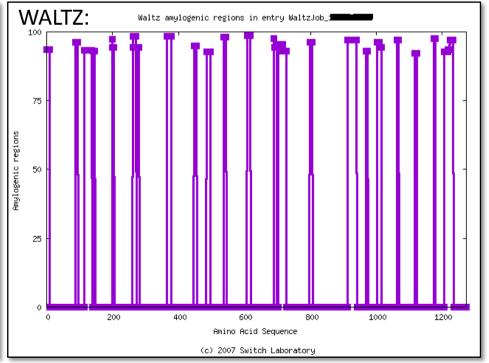
Basic Amyloid/Prion Biochemistry:

- Amyloid is causative/associated with many diseases (prions cause a form of amyloid):
 - Parkinson's, Alzheimer's, ALS, vCJD, some forms of fibrosis etc., and BSE in cows, CWD in deer.
 - Prions are not destroyed by cooking, bleaching, etc. 1/2000 extracted UK appendices found to contain BSE prions!
 - Prions hijack healthy body proteins and misfold them, causing <u>a chain-reaction</u> leading to death: "uniformly fatal".
 - There is NO SAFE LOWER LIMIT for prions in the brain.
- There is a clear mechanism for amyloid formation, *catalysed by SARS2-spike epitopes*.
 - IMPORTANT: just specific pieces of Spike are needed- 'epitopes'. Virus not required. Not even whole spike!

Predictable problems with SARS2-spike 'seeding' amyloid formation:

- Hammerstrom & Nystrom identified dangerous sections of spike, others see it too: Tetz & Tetz, Classen...
 - https://www.biorxiv.org/content/10.1101/2023.09.01.555834v1
 - "SARS-CoV-2 Spike ...specifically & selectively accelerates amyloid fibril formation of human prion protein and the amyloid-8 peptide"
 - I also identified regions & verified this myself using PLAAC & WALTZ (https://waltzdb.switchlab.org):





Q4: Amyloidosis & Prion issues Vaccine Disaster

Basic Amyloid/Prion Biochemistry:

- Amyloid is causative/associated with many diseases (prions cause a form of amyloid):
 - Parkinson's, Alzheimer's, ALS, vCJD, some forms of fibrosis etc., and BSE in cows, CWD in deer.
 - Prions are not destroyed by cooking, bleaching, etc. 1/2000 extracted UK appendices found to contain BSE prions!
 - Prions hijack healthy body proteins and misfold them, causing a chain-reaction leading to death: "uniformly fatal".
 - There is NO SAFE LOWER LIMIT for prions in the brain.
- There is a clear mechanism for amyloid formation, catalysed by SARS2-spike epitopes.
 - IMPORTANT: just specific pieces of Spike are needed- 'epitopes'. Virus not required. Not even whole spike!

Predictable problems with SARS2-spike 'seeding' amyloid formation:

- Hammerstrom & Nystrom identified dangerous sections of spike, others see it too: Tetz & Tetz, Classen..
 - https://www.biorxiv.org/content/10.1101/2023.09.01.555834v1
 - "SARS-CoV-2 Spike ... specifically & selectively accelerates amyloid fibril formation of human prion protein and the amyloid-8 peptide"
 - I also identified regions & verified this myself using PLAAC & WALTZ (https://plaac.wi.mit.edu , https://waltzdb.switchlab.org)

Basic Amyloid/Prion Biochemistry:

- Amyloid is causative/associated with many diseases (prions cause a form of amyloid):
 - Parkinson's, Alzheimer's, ALS, vCJD, some forms of fibrosis etc., and BSE in cows, CWD in deer.
 - Prions are not destroyed by cooking, bleaching, etc. 1/2000 extracted UK appendices found to contain BSE prions!
 - Prions hijack healthy body proteins and misfold them, causing a chain-reaction leading to death: "uniformly fatal".
 - There is NO SAFE LOWER LIMIT for prions in the brain.
- There is a clear mechanism for amyloid formation, catalysed by SARS2-spike epitopes.
 - IMPORTANT: just specific pieces of Spike are needed- 'epitopes'. Virus not required. Not even whole spike!

Predictable problems with SARS2-spike 'seeding' amyloid formation:

- Hammerstrom & Nystrom identified dangerous sections of spike, others see it too: Tetz & Tetz, Classen..
 - https://www.biorxiv.org/content/10.1101/2023.09.01.555834v1
 - "SARS-CoV-2 Spike ... specifically & selectively accelerates amyloid fibril formation of human prion protein and the amyloid-8 peptide"
 - I also identified regions & verified this myself using PLAAC & WALTZ (https://plaac.wi.mit.edu , https://waltzdb.switchlab.org)
- PROBLEM: Lipid Nanoparticles (LNPs) transport the mRNA to the Central Nervous System, crossing the BBB.
- From here on, it's chemistry: amyloid formation is a chain-reaction.
 - Just like BSE prions convert normal human prion to the misfolded form.
- 'Clots' being pulled from people test positive for amyloid.
 - e.g. using birefringence, with stains like Methylene Blue & Congo Red.

Q4: Amyloidosis & Prion issues

Vaccine Disaster

Basic Amyloid/Prion Biochemistry:

- Amyloid is causative/associated with many diseases (prions cause a form of amyloid):
 - Parkinson's, Alzheimer's, ALS, vCJD, some forms of fibrosis etc., and BSE in cows, CWD in deer.
 - Prions are not destroyed by cooking, bleaching, etc. 1/2000 extracted UK appendices found to contain BSE prions!
 - Prions hijack healthy body proteins and misfold them, causing <u>a chain-reaction</u> leading to death: "uniformly fatal".
 - There is NO SAFE LOWER LIMIT for prions in the brain.
- There is a clear mechanism for amyloid formation, <u>catalysed by SARS2-spike epitopes</u>.
 - IMPORTANT: just specific pieces of Spike are needed- 'epitopes'. Virus not required. Not even whole spike!

Predictable problems with SARS2-spike 'seeding' amyloid formation:

- Hammerstrom & Nystrom identified dangerous sections of spike, others see it too: Tetz & Tetz, Classen...
 - https://www.biorxiv.org/content/10.1101/2023.09.01.555834v1
 - "SARS-CoV-2 Spike ...specifically & selectively accelerates amyloid fibril formation of human prion protein and the amyloid-6 peptide"
 - I also identified regions & verified this myself using PLAAC & WALTZ (https://plaac.wi.mit.edu , https://waltzdb.switchlab.org)
- PROBLEM: Lipid Nanoparticles (LNPs) transport the mRNA to the Central Nervous System, crossing the BBB.
- From here on, it's chemistry: amyloid formation is a chain-reaction.
 - Just like BSE prions convert normal human prion to the misfolded form.
- 'Clots' being pulled from people test positive for amyloid.
 - e.g. using birefringence, with stains like Methylene Blue & Congo Red.

Clinical evidence, just one of many:

PET scan sees amyloid at injection site: https://pubmed.ncbi.nlm.nih.gov/34541458



Q4: Amyloidosis & Prion issues

Vaccine Disaster

Basic Amyloid/Prion Biochemistry:

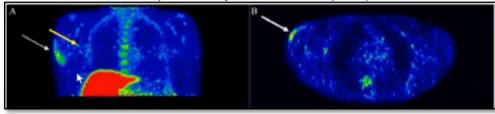
- Amyloid is causative/associated with many diseases (prions cause a form of amyloid):
 - Parkinson's, Alzheimer's, ALS, vCJD, some forms of fibrosis etc., and BSE in cows, CWD in deer.
 - Prions are not destroyed by cooking, bleaching, etc. 1/2000 extracted UK appendices found to contain BSE prions!
 - Prions hijack healthy body proteins and misfold them, causing <u>a chain-reaction</u> leading to death: "uniformly fatal".
 - There is NO SAFE LOWER LIMIT for prions in the brain.
- There is a clear mechanism for amyloid formation, *catalysed by SARS2-spike epitopes*.
 - IMPORTANT: just specific pieces of Spike are needed- 'epitopes'. Virus not required. Not even whole spike!

Predictable problems with SARS2-spike 'seeding' amyloid formation:

- Hammerstrom & Nystrom identified dangerous sections of spike, others see it too: Tetz & Tetz, Classen...
 - https://www.biorxiv.org/content/10.1101/2023.09.01.555834v1
 - "SARS-CoV-2 Spike ...specifically & selectively accelerates amyloid fibril formation of human prion protein and the amyloid-6 peptide"
 - I also identified regions & verified this myself using PLAAC & WALTZ (https://plaac.wi.mit.edu , https://waltzdb.switchlab.org)
- PROBLEM: Lipid Nanoparticles (LNPs) transport the mRNA to the Central Nervous System, crossing the BBB.
- From here on, it's chemistry: amyloid formation is a chain-reaction.
 - Just like BSE prions convert normal human prion to the misfolded form.
- 'Clots' being pulled from people test positive for amyloid.
 - e.g. using birefringence, with stains like Methylene Blue & Congo Red.

Clinical evidence, just one of many:

PET scan sees amyloid at injection site: https://pubmed.ncbi.nlm.nih.gov/34541458



Subcutaneous Uptake on [18F]Florbetaben PET/CT: a Case Report of Possible Amyloid-Beta Immune-Reactivity After COVID-19 Vaccination

Riccardo Laudicella^{1,2,3} 🕒 Irene Andrea Burger^{3,4} 🖰 - Francesco Panasiti^{1,2} - Costanza Longo^{1,2} - Salvatore Scalisi¹ Fabio Minutoli² - Sergio Baldari² 🕒 - Luigi Maria Edoardo Grimaldi⁵ - Pierpaolo Alongi¹ 💍

Renegade science experiments are underway:

- Attempt to determine whether vaccines cause prion / amyloid disease:
 - Vaccines are considered higher risk than SARS2 itself, due to the vaccine biodistribution.

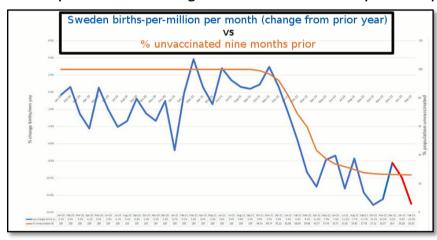


Q5: Birth-Rate drops...

Nine months after vaccines given to pregnant women.

- Observed in multiple countries.
- Sweden: February 2023 birth rates down 13% vs. February 2021: https://boriguagato.substack.com/p/sweden-birthrates-feb-2023-still

"this drop commenced right when vaccination picked up and is not recovering from these really low levels"



- A German Federal Government website:
 - https://www.bib.bund.de/Publikation/2022/Fertility-declines-near-the-end-of-the-COVID-19-pandemic-Evidence-of-the-2022-birth-declines-in-Germany-and-Sweden.html
 - "There is no association of the fertility trends with changes in unemployment, infection rates, or COVID-19 deaths."
 - "However, there is a strong association between the onset of vaccination programmes and the fertility decline nine months after this onset."
- Notably, the UK ONS has not published any 'live births' statistics since 2022, as far as I can see.
 - It is now late 2024.

- It's good to know that these people said these things...
 - What has been unleashed is apparently a *fertility-destroying slow-killer*, which fits with stated goals:



https://www.youtube.com/watch?v=S-un0VLr44c

https://www.youtube.com/watch?v=3rWU VDa1Js

"If you want liberty and more consumption, you have to have fewer people"

"We're now at 7 [billion people], so we have to get back down."

"I hope that this can be slow, and done in a way which is relatively equal"

"That's what lies ahead" (smirking).

Philip: "The growing human population."

"From where we are, there's nothing else"

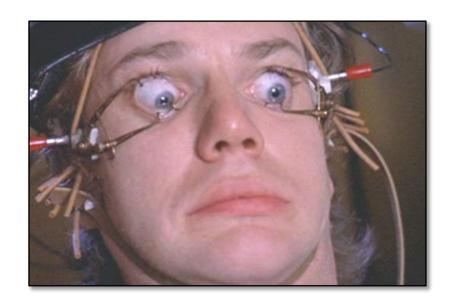
"Do you have views about what should be Interviewer:

"Can't you guess?" Philip:

Might as well keep going. Why it won't stop

because they will, too.

- -Lab Origin.
- -Malfeasance Treatments and Evidence of Intent.
- -Vaccine Disaster.
- Why it won't stop:
 - We can't just carry on, without future risks.



Why it won't stop...

Why it won't stop

• Highwire episode 383:

"There's a pandemic coming, and though [bird flu] can't really spread among humans very well right now, don't worry – they're fixing that in a lab somewhere near you"



• Highwire episode 383:

"There's a pandemic coming, and though [bird flu] can't really spread among humans very well right now, don't worry – they're fixing that in a lab somewhere near you"

 It is now known how to make super-transmissible versions of many pathogens.



Why it won't stop...

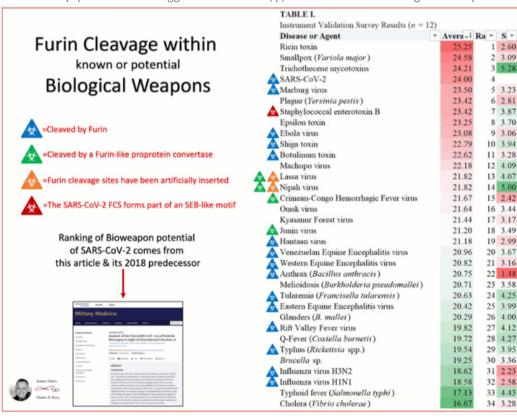
Highwire episode 383:

"There's a pandemic coming, and though [bird flu] can't really spread among humans very well right now, don't worry – they're fixing that in a lab somewhere near you"



Charles Rixey: prometheusshrugged.substack.com/p/the-evidence-of-the-origin-of-the-pandemic

- It is now known how to make super-transmissible versions of many pathogens.
- Charles Rixey, ex-CBRN instructor,
 & US Marine made this chart
 - From the 'dirty dozen' paper revisit (2020).
 - Furin Cleavage can be added.
 - RxxR can be upgraded to RRxR.



On Informed Consent...

Why it won't stop

Informed Consent Rules degraded in USA...

https://www.theepochtimes.com/health/fdas-new-rule-allows-for-medical-research-without-informed-consent-5571962

- U.S. Food and Drug Administration (FDA) has finalized a rule allowing certain clinical trials to operate without obtaining informed consent from participants.
- The hitch? The study cannot pose more than *minimal risk* to humans and must include appropriate safeguards to protect the rights, safety, and welfare of those involved.
- The rule was issued in late December 2023 and went into effect on Jan. 22, 2024.

On Informed Consent...

Why it won't stop

Informed Consent Rules degraded in USA...

https://www.theepochtimes.com/health/fdas-new-rule-allows-for-medical-research-without-informed-consent-5571962

- U.S. Food and Drug Administration (FDA) has finalized a rule allowing certain clinical trials to operate without obtaining informed consent from participants.
- The hitch? The study cannot pose more than *minimal risk* to humans and must include appropriate safeguards to protect the rights, safety, and welfare of those involved.
- The rule was issued in late December 2023 and went into effect on Jan. 22, 2024.
- What is minimal risk? Does mRNA qualify?
- How do "self-spreading vaccines" fit in to new informed consent rules?

Informed Consent Rules degraded in USA...

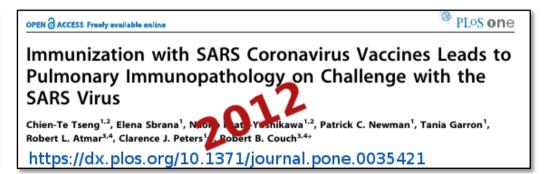
https://www.theepochtimes.com/health/fdas-new-rule-allows-for-medical-research-without-informed-consent-5571962

- U.S. Food and Drug Administration (FDA) has finalized a rule allowing certain clinical trials to operate without obtaining informed consent from participants.
- The hitch? The study cannot pose more than *minimal risk* to humans and must include appropriate safeguards to protect the rights, safety, and welfare of those involved.
- The rule was issued in late December 2023 and went into effect on Jan. 22, 2024.
- What is minimal risk? Does mRNA qualify?
- How do "self-spreading vaccines" fit in to new informed consent rules?
 - Do paid shills like Peter Hotez consider these technologies 'minimal risk'?

Mar 2020: Peter Hotez highlights a big problem with coronavirus vaccines.



https://www.c-span.org/video/?c4861348/user-clip-dr-peter-hotezspeaks-congress-coronavirus-vaccines NOTE: YouTube deleted the video., so c-span used directly.



On Informed Consent..

Why it won't stop

Informed Consent Rules degraded in USA...

https://www.theepochtimes.com/health/fdas-new-rule-allows-for-medical-research-without-informed-consent-5571962

- U.S. Food and Drug Administration (FDA) has finalized a rule allowing certain clinical trials to operate without obtaining informed consent from participants.
- The hitch? The study cannot pose more than minimal risk to humans and must include appropriate safeguards to protect the rights, safety, and welfare of those involved.
- The rule was issued in late December 2023 and went into effect on Jan. 22, 2024.
- What is minimal risk? Does mRNA qualify?

How do "self-spreading vaccines" fit in to new informed consent rules?

Do paid shills like Peter Hotez consider these technologies 'minimal risk'?

Mar 2020: Peter Hotez highlights a big problem with coronavirus vaccines.

June 2020:

2024: Calls for NATO to address "anti-vaccine aggression".



https://www.c-span.org/video/?c4861348/user-clip-dr-peter-hotez speaks-congress-coronavirus-vaccines

NOTE: YouTube deleted the video., so c-span used directly.

https://www.voutube.com/watch?v=QEpp2GbcuEU

The Busine\$\$ Model

Why it won't stop

- Pandemic / Vaccine has now become a business model.
 - No punishment or recognition of wrongdoing
 no dis-incentive.
 - Altering and patenting viruses: US Pat. 7,279,372
 method to produce recombinant Coronavirus (R. Baric).
 - \$\$\$\$\$\$\$ to DARPA, BARDA, PREDICT, EcoHealth, Metabiota, OneHealth, GAVI...

The Busine\$\$ Model

- Pandemic / Vaccine has now become a business model.
 - No punishment or recognition of wrongdoing
 no dis-incentive.
 - Altering and patenting viruses: US Pat. 7,279,372
 method to produce recombinant Coronavirus (R. Baric).
 - \$\$\$\$\$\$\$ to DARPA, BARDA, PREDICT, EcoHealth, Metabiota, OneHealth, GAVI...
- Preparations are being made and lessons learned applied, for future responses.
 - Personally, I felt we were very close to unvaccinated being taken off to camps. Can you imagine how that feels?
 - Some were force-injected <u>against their will</u> during covid.
 - Next time...
 https://www.health.govt.nz/system/files/documents/publications/ pp133
 interim nz pandemic plan v2.pdf



 \mathbb{X}

New Zealand's New "Pandemic Plan" includes Forced injection whilst being restrained by Police

NEW ZEALAND PANDEMIC PLAN

A framework for action - Interim Update July 2024 - Page 125

SPECIAL POWERS

Special powers are authorised by the Minister of Health or by an epidemic notice or apply where an emergency has been declared under the Civil Defence Emergency Management Act 2002.



- The power to detain, isolate or quarantine allows a medical officer of health to 'require persons, places, buildings, ships, vehicles, aircraft, animals, or things to be isolated, quarantined, or disinfected' (section 70(1)(f)).
- The power to prescribe preventive treatment allows a medical officer of health, in respect of any person who has been isolated or quarantined, to require people to remain where they are isolated or quarantined until they have been medically examined and found to be free from infectious disease, and until they have undergone such preventive treatment as the medical officer of health prescribes (section 70(1)(h)).
- Section 71A states that a member of the police may do anything reasonably necessary (including the use of force) to help a medical officer of health or any person authorised by the medical officer of health in the exercise or performance of powers or functions under sections 70 or 71.

'INCLUDING THE USE OF FORCE

link to the document



www.thehealthforumnz.co.nz

The Busine\$\$ Model

3:24 AM · Aug 1, 2024

- Pandemic / Vaccine has now become a business model.
 - No punishment or recognition of wrongdoing
 no dis-incentive.
 - Altering and patenting viruses: US Pat. 7,279,372
 method to produce recombinant Coronavirus (R. Baric).
 - \$\$\$\$\$\$\$ to DARPA, BARDA, PREDICT, EcoHealth, Metabiota, OneHealth, GAVI...
- Preparations are being made and lessons learned applied, for future responses.
 - Personally, I felt we were very close to unvaccinated being taken off to camps. Can you imagine how that feels?
 - Some were force-injected <u>against their will</u> during covid.
 - Next time...
 https://www.health.govt.nz/system/files/documents/publications/ pp133
 interim nz pandemic plan v2.pdf
- If you don't consent, there's the covert option:
 - Fusion inhibitors would protect those 'in the know'.



self-Spreading Vaccines: Self-spreading vaccines are genetically engineered to move through populations like communicable diseases, but rather than causing disease, they confer protection. The vision is that a small number of individuals in a target population could be vaccinated, and the vaccine strain would then circulate in the population much like a pathogenic virus, resulting in rapid, widespread immunity.



Yes, some were forced already.. Why it won't stop

Holding people down and forcing needles into them happened.

In Los Angeles:

https://www.voutube.com/watch?v=QJ8xfO1RMsM



In Berlin:

https://www.bitchute.com/video/hxWbID9xqRrp



And to Children:



Note: deleted from x: https://x.com/ /status/1425818674758365184

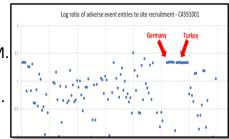
And in India:

https://x.com/ /status/1487926273892618242



Arguably rewarded for fraud (and it's not the first time).

- Aaron Siri / ICAN sued to obtain Pfizer trial documents.
- So many deceptions...
 - 301 participants gone from the trial records.
 - 1258 (vaccinated) & 583 (placebo) participants missing from the 6m report in NEJM.
 - Unblinding of participants and staff during the trial.
 - German & Turkish sites had exactly the same number of adverse events per person.
 - Using "process 2" to make commercial lots NOT what was used in trials.

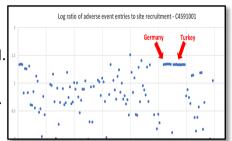


Arguably rewarded for fraud (and it's not the first time).

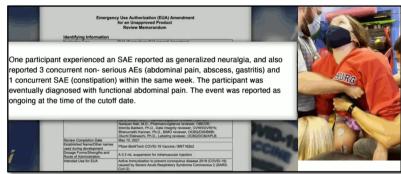
- Aaron Siri / ICAN sued to obtain Pfizer trial documents.
- So many deceptions...
 - 301 participants gone from the trial records.
 - 1258 (vaccinated) & 583 (placebo) participants missing from the 6m report in NEJM.
 - Unblinding of participants and staff during the trial.
 - German & Turkish sites had exactly the same number of adverse events per person.
 - Using "process 2" to make commercial lots NOT what was used in trials.

Several particpants who became ill were taken off the trial:

- Maddie De Garay:
 - 12yo left paralysed.
 - Mis-classified as 'abdominal pain', 'neuralgia'.
 - Was not treated for acute illnesses.
 - Referred to an eating disorder clinic.
 - Had to obtain own wheelchair.
 - Note: Only 1200 children in the trial.



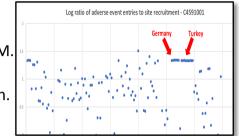
https://x.com/Jikkyleaks/status/1523830222218723328



https://www.bitchute.com/video/DcdbWfuvusaT (see 61m30s)

Arguably rewarded for fraud (and it's not the first time).

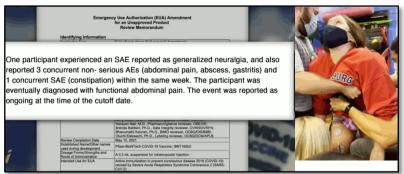
- Aaron Siri / ICAN sued to obtain Pfizer trial documents.
- So many deceptions...
 - 301 participants gone from the trial records.
 - 1258 (vaccinated) & 583 (placebo) participants missing from the 6m report in NEJM.
 - Unblinding of participants and staff during the trial.
 - German & Turkish sites had exactly the same number of adverse events per person.
 - Using "process 2" to make commercial lots NOT what was used in trials.



https://x.com/Jikkyleaks/status/1523830222218723328

Several particpants who became ill were taken off the trial:

- Maddie De Garay:
 - 12yo left paralysed.
 - Mis-classified as 'abdominal pain', 'neuralgia'.
 - Was not treated for acute illnesses.
 - Referred to an eating disorder clinic.
 - Had to obtain own wheelchair.
 - Note: Only 1200 children in the trial.



https://www.bitchute.com/video/DcdbWfuyusaT (see 61m30s)

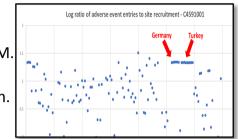
- Augusto Roux (An Argentinian State Prosecutor!) was removed from the trial whilst in hospital.
- Olivia Teseniar Leukemia began on trial:
 - Investigator didn't report to FDA.
 - Her VAERS record was deleted.
 - Olivia secretly filmed trial investigator, Dr Cynthia Strout:



https://x.com/ /status/1610122775200505857 https://www.bitchute.com/video/HdX1AdORTWxV

Arguably rewarded for fraud (and it's not the first time).

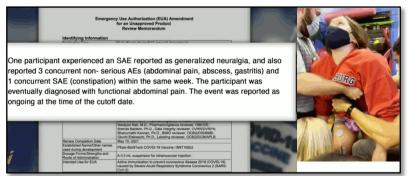
- Aaron Siri / ICAN sued to obtain Pfizer trial documents.
- So many deceptions...
 - 301 participants gone from the trial records.
 - 1258 (vaccinated) & 583 (placebo) participants missing from the 6m report in NEJM.
 - Unblinding of participants and staff during the trial.
 - German & Turkish sites had exactly the same number of adverse events per person.
 - Using "process 2" to make commercial lots NOT what was used in trials.



https://x.com/Jikkyleaks/status/1523830222218723328

Several participants who became ill were taken off the trial:

- Maddie De Garay
 - 12yo left paralysed.
 - Mis-classified as 'abdominal pain', 'neuralgia'.
 - Was not treated for acute illnesses.
 - Referred to an eating disorder clinic
 - Had to obtain own wheelchair.
 - Note: Only 1200 children in the trial.



https://www.bitchute.com/video/DcdbWfuyusaT (see 61m30s)

- Augusto Roux (An Argentinian State Prosecutor!) was removed from the trial whilst in hospital.
- Olivia Teseniar Leukemia began on trial.
 - Investigator didn't report to FDA.
 - Her VAERS record was deleted.
 - Olivia secretly filmed trial investigator, Dr Cynthia Strout:



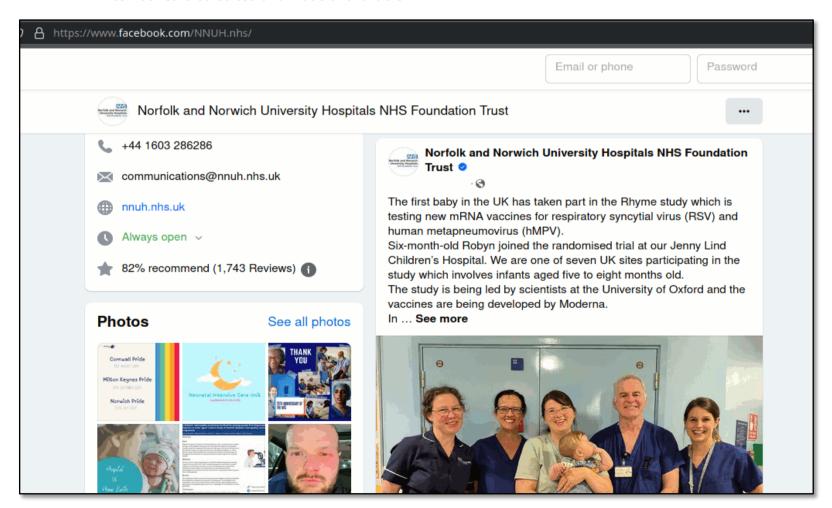
Bear in mind these are TRIAL PARTICIPANTS.

Injuries/events are <u>supposed</u> to be recorded, regardless of aetiology!

https://x.com/ /status/1610122775200505857 https://www.bitchute.com/video/HdX1AdORTWxV

The Pipeline: Depends on Ignorance / Complacency

- MANY mRNA 'vaccines' in the pipelne.
- Poverty of information in the public domain. Same issues...
 - Mother aware of pseudo-uridine danger? (paper: friend or foe of cancer https://pubmed.ncbi.nlm.nih.gov/38583833/)
 - Mother aware of DNA contamination issues? (this is a result of manufacturing process, so must be a risk going forward)
 - Mother aware of bio-distribution data? (transfected cells in heart/brain will not re-grow)
 - No 'OFF' Switch, indeterminate dosing.
 - Alternatives discredited and made unavailable.



Censorship...

- Note that the NNUH NHS Trust tweet I wanted to show on the previous slide about baby Robyn has been deleted. Why?
- Much of the material I saved offline has been systematically removed.
- For a while, a lot of twitter material disappeared when accounts were deleted/banned.
 - Links I had saved were unusable and there has been good improvement since Elon Musk bought the Co.
 - This was industrial-scale, using JIRA, organised by governments against their citizens.
 - They almost got away with it.
- Papers have been 'force-retracted', even though authors disagree with retraction, and many are later corroborated by other papers.
 - Happens when papers highlight topics likely to impair defence/military research budgets.
- In the UK, Ofcom's "material misleadingness" rule has blighted even the few MSM hosts/shows who tried to air both sides of the vaccine debate:
 - Mark Steyn
 - Maajid Nawaz
 - Neil Oliver
 - Ofcom "material misleadingness" rule:
 https://www.ofcom.org.uk/tv-radio-and-on-demand/broadcast-standards/broadcast-standards-and-coronavirus
- Eric Clapton's song(s) censored by MSM.
- I'd like you to think about how much of this information you were aware of.
- They even <u>censored the censorship</u>!
 - Unless you were looking in real-time, you never saw it.

- Bill & Melinda Gates Foundation (BMGF).
 - Large donations, influence peddling, outright corruption.
- UNITAID, WHO (see Astrid Steuckelberger on the takeover of WHO by Gates).
- Brighton Collaboration.
- Regulators: MHRA, FDA, CDC, NIH HC, TGA, EMA,...
 - "Enablers", Conflicted financially (share ownership, biggest funder BMGF, derive income from vaccines)
- UK Statistics Agency, ONS.
 - Stopping publication of key data e.g. live births
 - Changing methodology for Excess Deaths mid-crisis, without parallel running.
- NHS, Surgical/Medical Colleges, GMC:
 - Threatening and firing staff e.g. Dr Sam White, Dr David Cartland
- Biotech and pandemic-related enterprises:
 - EcoHealth, Labyrinth, Metabiota (Ukraine! & Kenema, Sierra Leone where VHFC has a lab and ZEBOV emerged).
 - Note: CEO of EcoHealth (Peter Daszak chaired a closed-door meeting with In-Q-Tel (CIA) and CEO of the Robert Wood Johnson Foundation.
 - Meeting to address media misinformation.
 - I discovered subsequent donations of \$100,000's to FactCheck.org, with nothing prior to the meeting.
- Academic Organisations:
 - The Lancet, Nature, forced retractions of peer-reviewed papers <u>against authors wishes</u>, <u>& without data</u>.
 Findings often confirmed in later papers. E.g. Spike-p53 paper.
 - The Lancet's COVID Commission populated with VI's.
 - Many Universities firing or marginalising academics. Byram Bridle,
- Media: BBC, Sky, Ch4, ...
 - Ofcom "material misleadingness" rule led to media censoring real data, information:
 https://www.ofcom.org.uk/tv-radio-and-on-demand/broadcast-standards/broadcast-standards-and-coronavirus
 - Biassed targetting e.g. Mark Steyn, but not This Morning "100% effective against death" comment by Dr. Sara.

Trustworthy Actors

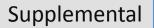
- Building a new system:
 - FLCCC: medical protocols, new fellowships, journal in the works
 - IJVTPR: most articles are of reasonable quality.
 - WCH: Tess Lawrie
 - HART: Clare Craig
 - ★Stars★:- have <u>you</u> heard of any of these people?
 - Kevin McKernan, Dr. Kevin McCairn, Charles Rixey, Dr. Tess Lawrie, Dr. Pierre Kory, Dr. Paul Marik, Prof. Arne Burkhardt, Prof. Angus Dalgleish, Andrew Bridgen, Jikkyleaks, Igor Chudov, Dr. Jessica Rose, Arkmedic, Aussie17, Dr. Steve James, Dr. Dave Cartland, Dr. Sam White, Matt LeTissier, Dr. Harvey Risch, Dr. Ryan Cole, Dr. Joseph Ladapo, Dr. Sabine Hazan, John Paul/ThingsHiddenn, Dr. Peter McCullough, Senator Ron Johnson, Rep. Rand Paul, Joe Rogan, Mark Steyn, Maajid Nawaz, Neil Oliver, Dr. Chris Martenson, Dr. Robert Malone, Aaron Siri, Del Bigtree, Jeffrey Jaxen, Brook Jackson.
- Low-censorship social media:
 - Rumble, X/Twitter, Bitchute, Odysee, Substack.
- Centralised broadcasters:
 - The Highwire, UK Column, ResistanceGB, BrokenTruth.com/John Davidson.

Supplemental

Supplemental #1: Let's talk about Antimicrobial Resistance?

- Meetings were proceeding on AMR, even whilst key antibiotics were being withheld in 2020-21.
- Seemingly relentless pressure to restrict & withhold antibiotics in general, with the stated aim of reducing resistance.
- Has anyone at recent meetings discussed the DNA snippets in the 'vaccines' yet?
 - The Plasmid DNA has Neomycin/Kanamycin resistance genes..
 https://anandamide.substack.com/p/curious-kittens
 - "patient use of Neomycn/Kanamycin after vaccination...could enable selection of Neo/Kan resistant bacteria in the gut"
 - Thankfully, It does appear to be rare to find whole bacterial DNA plasmids in the Pfizer BNT-162b2...
 - Unlike the DNA fragment contamination, which seems ubiquitous!
- This omission casts some doubt in my mind about the real motives regarding antibiotic restrictions.
 - Further doubts because antibiotics withheld from <u>very sick</u> people.

Supplemental #2:

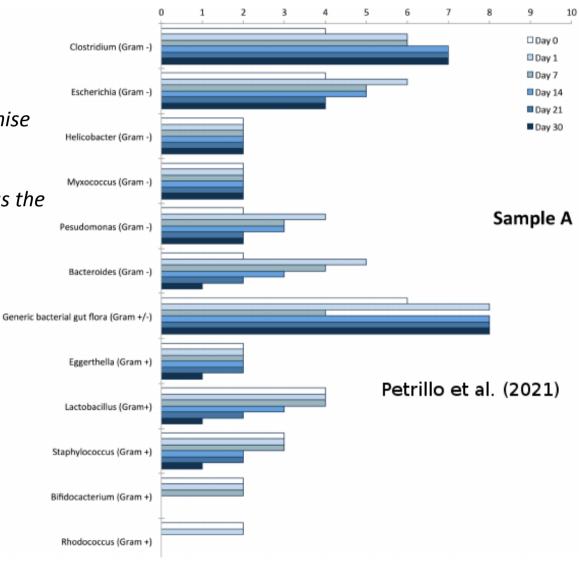


Interesting side-note on *E.coli/C.diff*

Dr. Philip McMillan https://rumble.com/v57vccl-covid-19-infecting-bacteria-mind-blowing-facts-shocking.html

See video above:

- @16m49s.. Groups of gut bacteria present after covid (& 'vaccination').
- @19m03s: "[Spike] seems to weaponise clostridium, ...at the expense of bifidobacteria, Lactobacteria"
- @20m00s "likely to be seeing.. Across the world.. Outbreaks of C. diff".



Thank You.

