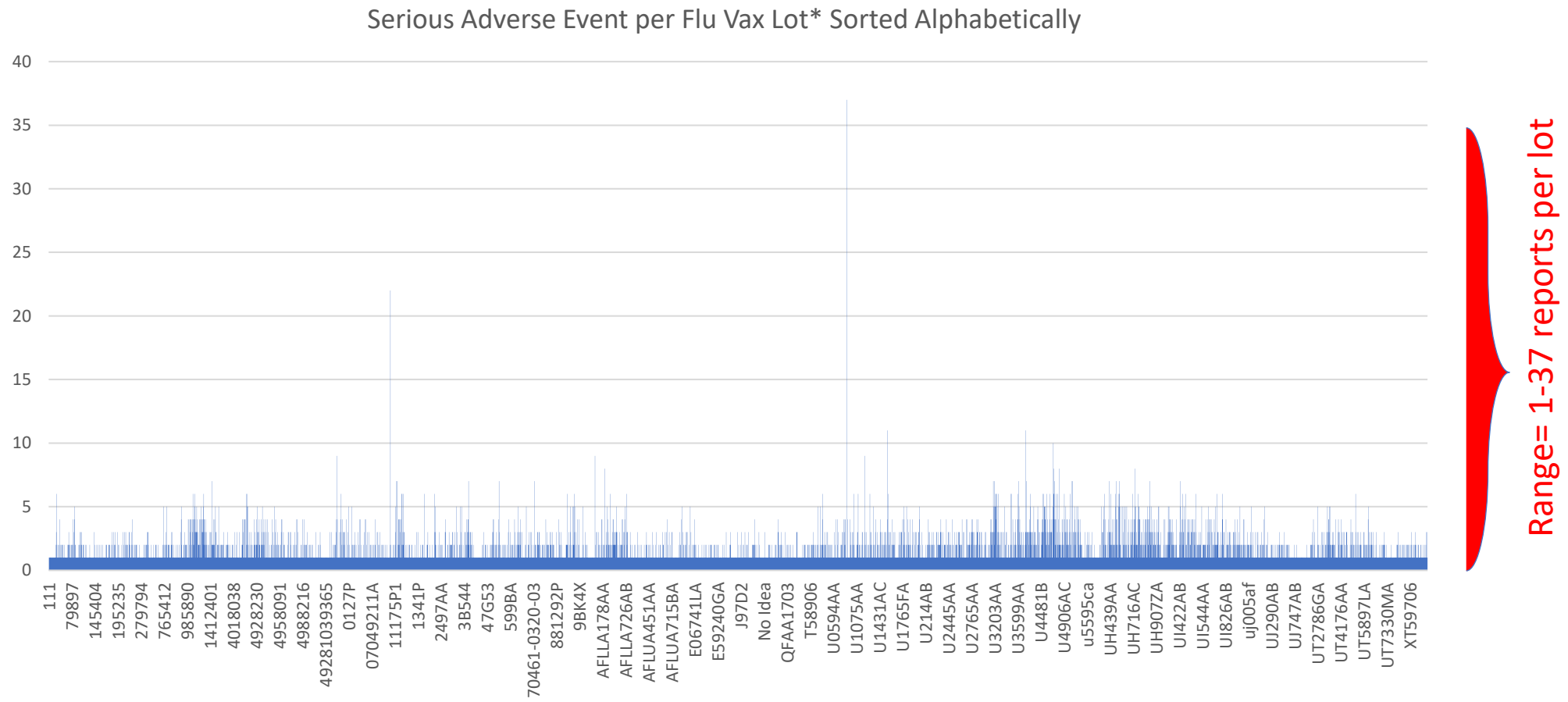
Weaponized “Healthcare”

Sasha Latypova

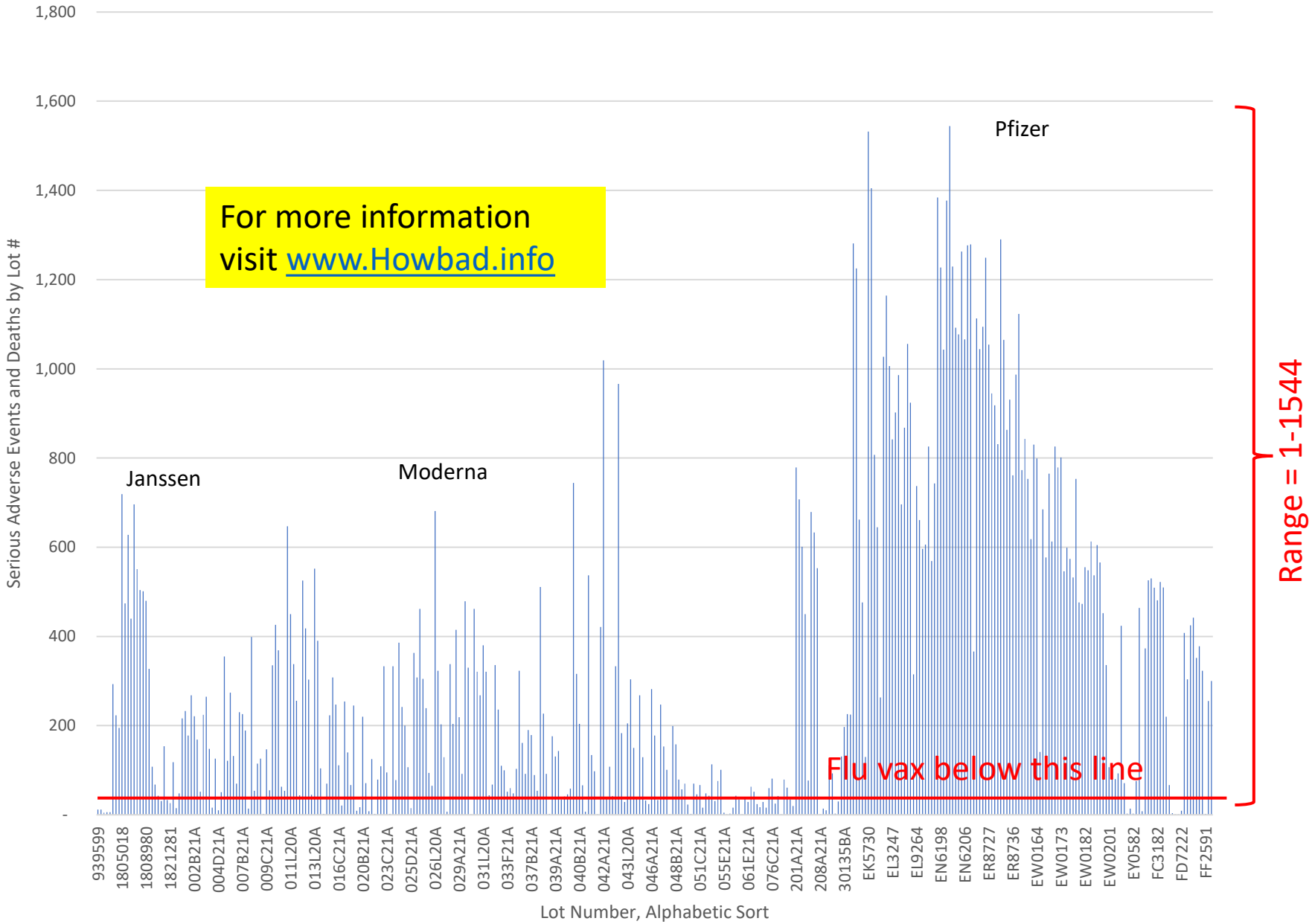
Sashalatypova.substack.com

30 years of VAERS data for all flu vaccines by batch number



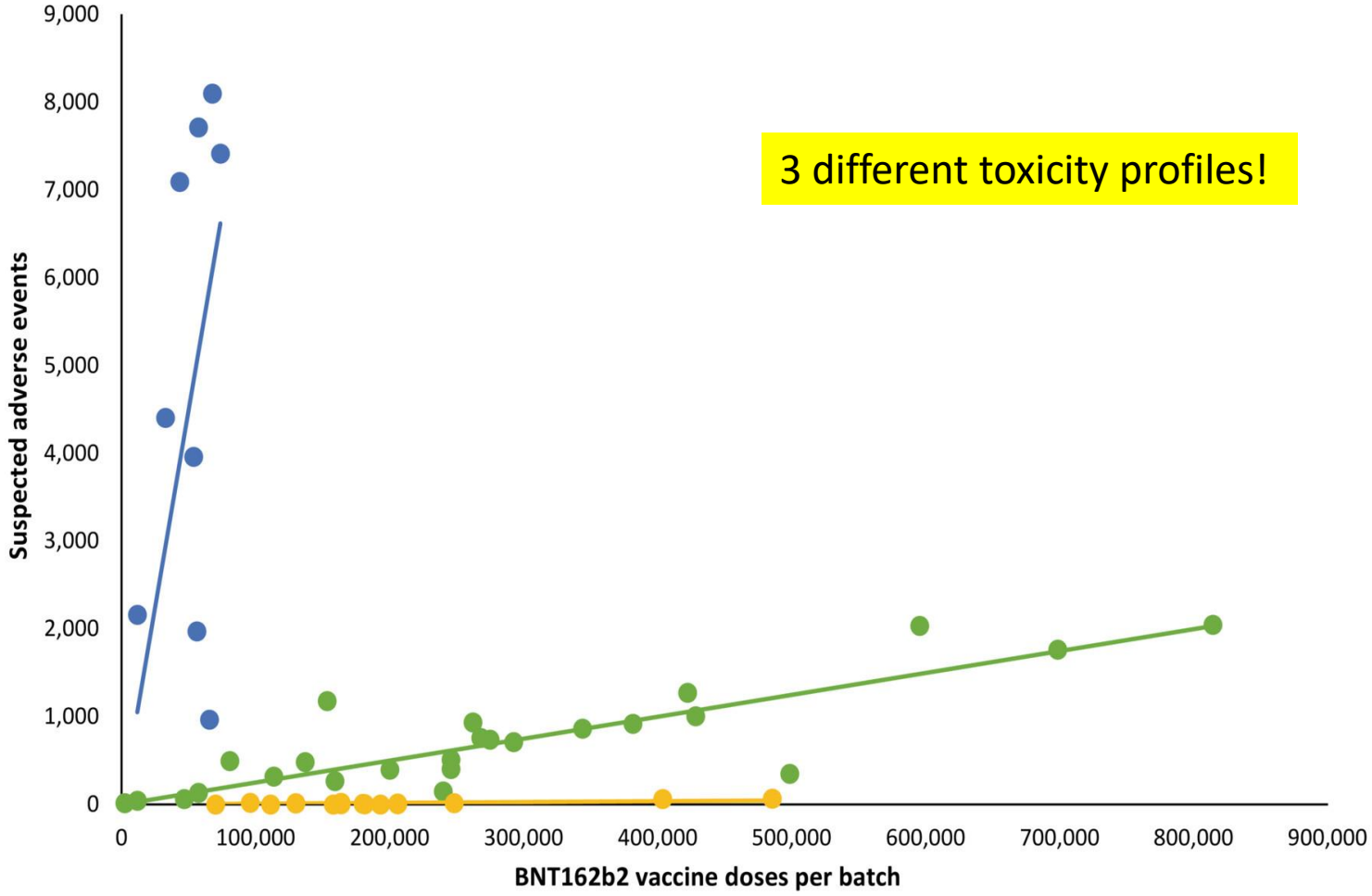
*Includes lots with non-zero SAE Reports only

C19 Vaccines Serious AE/Lot Number*, Sorted Alphabetically



*Includes US CDC lot-matched data only, Jan-Dec 2021

Dose Adjusted Lot-to-Lot Variability in Danish Vax AE Data, Pfizer Lots

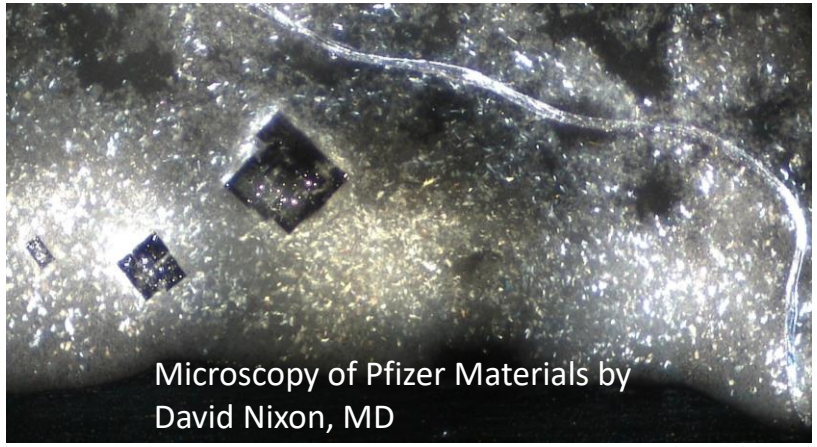


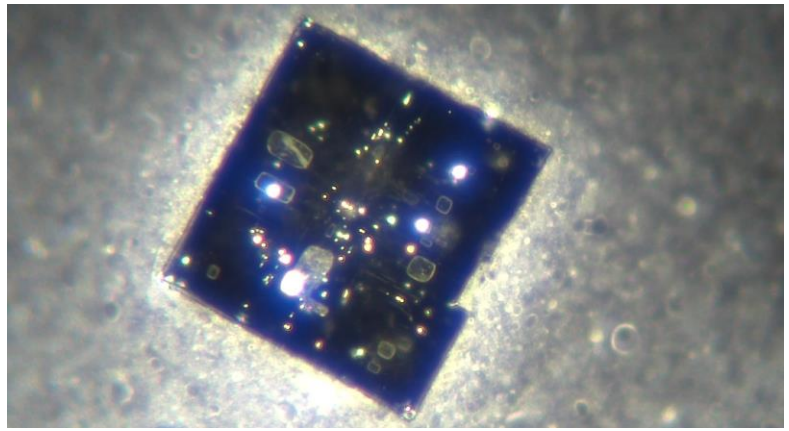
3 different toxicity profiles!

Schmeling M et al. Eur J Clin Investigation, 30 March 2023, DOI: (10.1111/eci.13998)

Product does not conform to labels*

- RNA non-conforming to label
- DNA and protein impurities in massive quantities
- Toxic metals of unknown origin or purpose:
  - Caesium, potassium, calcium, barium, cobalt, iron, chromium, titanium, cerium, gadolinium, aluminum, silicon, sulfur, thulium, antimony.
- Hydrogel (DARPA hydrogel?)
- Graphene oxide?
- Objects/structures: blobs, particles, crystals, square shapes, fibers, ribbons
- Many other yet unexplained materials/phenomena





* David Hughes, PhD https://ijvtpr.com/index.php/IJVTPR/article/view/52

#Plasmidgate: Pfizer and Moderna bivalent vial testing by Kevin McKernan (USA)*

- dsDNA plasmid contaminants from engineered e.coli (~10-100x over acceptable limit):
  - Can transfect human and microbiome cells
  - Contain antibiotic resistant genes
  - SV40 promoter (purpose – transfection of the cell nucleus**) – not declared in regulatory filings
  - Coded for spike protein
- Deliver a combination of:
  - Synthetic chemical mRNA + broken pieces (miRNAs)
  - DNA plasmids - replicate inside human cells/ bacterial cells of gut microbiome
  - Sepsis? Dysbiosis? Cancer? Myocarditis? Sterility? Birth Defects? More covid illness?

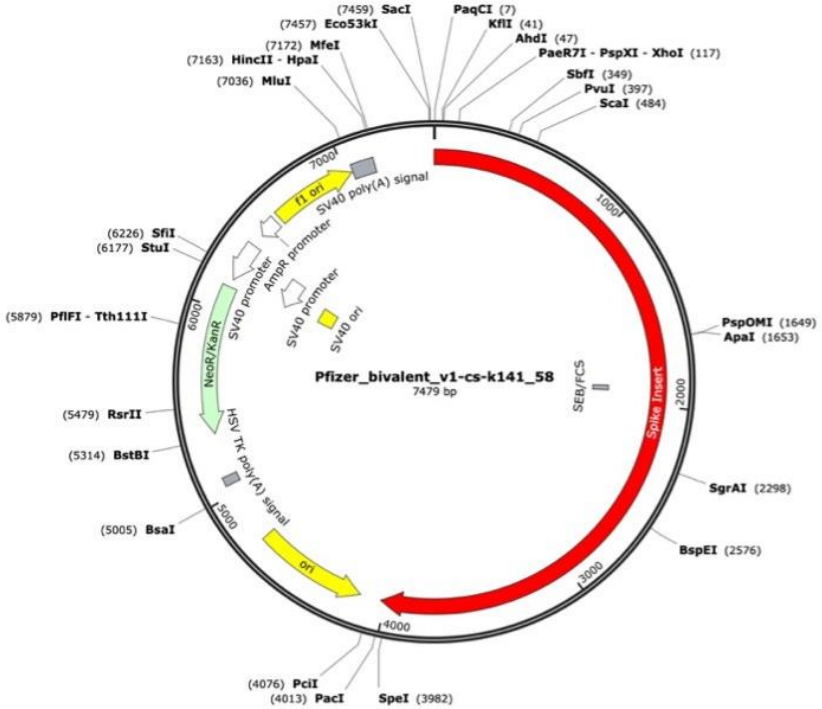


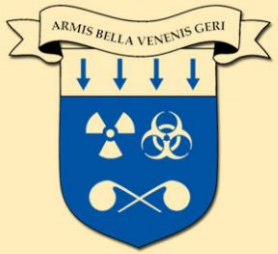
Figure 2. Pfizer bivalent vaccine assembly of the RNA-seq library. Annotated with SEB/FCS, spike insert (red), bacterial origin of replication (yellow), Neo/Kan resistance gene(green), F1 origin (yellow) and an SV40 promoter (yellow and white).

*Preprint DOI 10.31219/osf.io/b9t7m

**University of Rochester, Dean Lab

Next Generation Bioweapons: Genetic Engineering and BW

Michael J. Ainscough



US Air Force
Counterproliferation Center
Future Warfare Series
No. 14

14

the contributions of scientists to problems of national security and public benefit." Their meeting concentrated on the near-term future threat of biological warfare, specifically on genetically engineered pathogens and weapons.

The JASON Group that met in 1997 grouped potential genetically engineered pathogens into six broad groups of potential futuristic threats.[85]

- *Binary biological weapons*
- *Designer genes*
- *Gene therapy as a weapon*
- *Stealth viruses*
- *Host-swapping diseases*
- *Designer diseases*

3) <u>Gene Therapy as a Weapon</u>:[96] Gene therapy will revolutionize the treatment of human genetic diseases. The goal is to effect a permanent change in the genetic composition of a person by repairing or replacing a faulty gene. Genes have already been spliced into bacteria to produce "human" insulin in large quantities.[97] The eventual goal is to splice a gene that codes for the production of insulin into human pancreatic tissue to cure diabetes. Similar research is progressing on adding in the missing gene to prevent the symptoms of cystic fibrosis. However, the same technology could be subverted to insert pathogenic genes.

TGA regulators knew nothing about the genetic technology platform they were supposedly regulating.

Australian Government

Department of Health
Therapeutic Goods Administration

TRIM Ref: D22-5167274

[redacted]
By email: [redacted]

Dear [redacted],

FREEDOM OF INFORMATION REQUEST FOI 3604
Notice of Decision

1. I refer to your request dated 5 February 2022 under the *Freedom of Information Act 1982* (the FOI Act) for access to the following documents:

   "the following documents relating to the provisional approval of the Pfizer-BionTech BNT162b2 vaccine in January 2021:
   1. *"All documents relating to the TGA's assessment of the risk of and/or presence of **micro-RNA sequences (miRNA)** comprised within the Comirnaty mRNA active ingredient (mRNA genomic sequence).*
   2. *All documents relating to the TGA's assessment of the risk of and/or presence of **Oncomirs** (oncogenic miRNA - microRNA) comprised within the Comirnaty mRNA active ingredient (mRNA genomic sequence).*
   3. *All documents relating to the TGA's assessment of the risk of and/or presence of **Stop Codon read-through** (suppression of stop codon activity) arising as a result of the use of pseudouridine in the Comirnaty miRNA active ingredient (mRNA genomic sequence).*
   4. *Any document showing that the TGA has assessed the composition of the **final protein product** (molecular weight and amino acid sequence) produced following injection of the Comirnaty mRNA product in human subjects.*
   5. *All documents relating to the TGA's assessment of the risk of the use of the **AES-mtRNR1 3' untranslated region** of the Comirnaty mRNA product in human subjects."*

Decision Maker

2. I am the Therapeutic Goods Administration (TGA) officer authorised to make this decision under section 23 of the FOI Act. What follows is my decision under the FOI Act.

Decision

3. Unfortunately, I am unable to continue to process your request because the documents you have requested do not exist.

Opinion of the European Regulators (EMA) Nov 30, 2020:

- Pfizer/BioNTech manufacturing processes (at all sites) were NOT GMP compliant:
  - Not able to produce consistent product
  - Not validated, proprietary “black box” process, missing information
  - No plan/deadline as to when they were going be in compliance
- 117 Major Objections and Concerns listed by the regulators on 30+ pages:
  - For any normal product, even fraction of this would halt the approval
  - All objections would need to be resolved before authorization
- 2 weeks later the product is “authorized” worldwide…

Regulatory “approval” was a sham:
Neither the regulators, nor vaccinators, nor pharma employees know what is in the vials.

They are just following orders…
Whose orders??

If the Government is allowed to violate the Constitution in an emergency, they will create the permanent emergency.

The US Constitution is NOT in Force

Step 1: Announce Public Health Emergency:

- Title 42, Public Health Service Act, at §247d, “Public health emergencies” and at §300hh “National All-Hazards Preparedness for PHE”
- Legal state = announcement of war => suspends the Constitution

Step 2: Deploy “countermeasures”:

- PREP Act declarations for covid and marburg in 2020, now include flu, RSV and anything “potential”
- Legal equivalent of announcing use of specific type of weapons and extending liability protection to those who operate them as long as they “follow orders”

Congress transferred — to the HHS Secretary — unilateral, unreviewable power to declare and maintain public health emergency status and direct biochemical “countermeasures” attacks on the American people camouflaged as “vaccination” programs.

Countermeasures are “medicines” without any regulatory safeguards = i.e., poisons/weapons

- 21 USC 360bbb-3(k): **use** of EUA-covered medical countermeasure (MCM) products, once designated as such by the Secretary of Health and Human Services “**shall not be considered to constitute a clinical investigation.**”

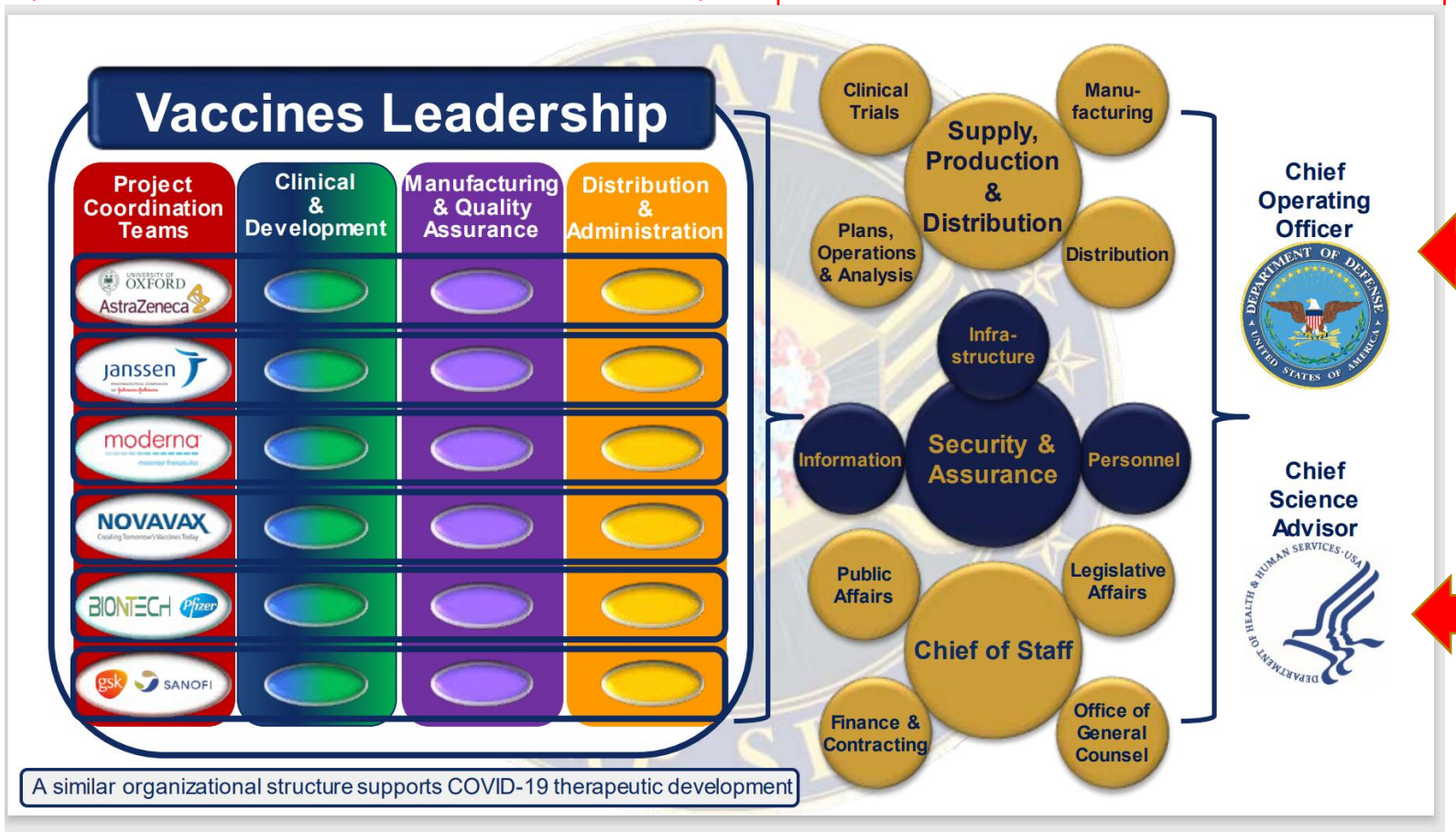
- Countermeasures deployed at sole discretion of the HHS Sec during HHS-declared PHE:
  - No initiation criteria, no data needed, no judicial review allowed, no stopping criteria!

Bio-chemical Weapons Renamed into "Medical Prototype Countermeasures"

- 10 USC 2371b/10 USC 4022 Other Transaction Authority (OTA) program "legalized" DOD contracting with pharmas to produce bio-chemical weapons, in violation of federal and international laws prohibiting same:
  - 10 USC 4022(a)(1) - "[] carry out prototype projects that are directly relevant to enhancing the mission effectiveness of military personnel and the supporting platforms, systems, components, or materials proposed to be acquired or developed by the Department of Defense, […]."
- OTA program renamed bio/chem/radio/nuclear-weapons as "qualified countermeasures, medical countermeasures and security countermeasures".
- BARDA (not FDA) owns and "regulates" countermeasures outside of pharmaceutical manufacturing and distribution laws.

Not in charge: Pharma companies ($$$$)

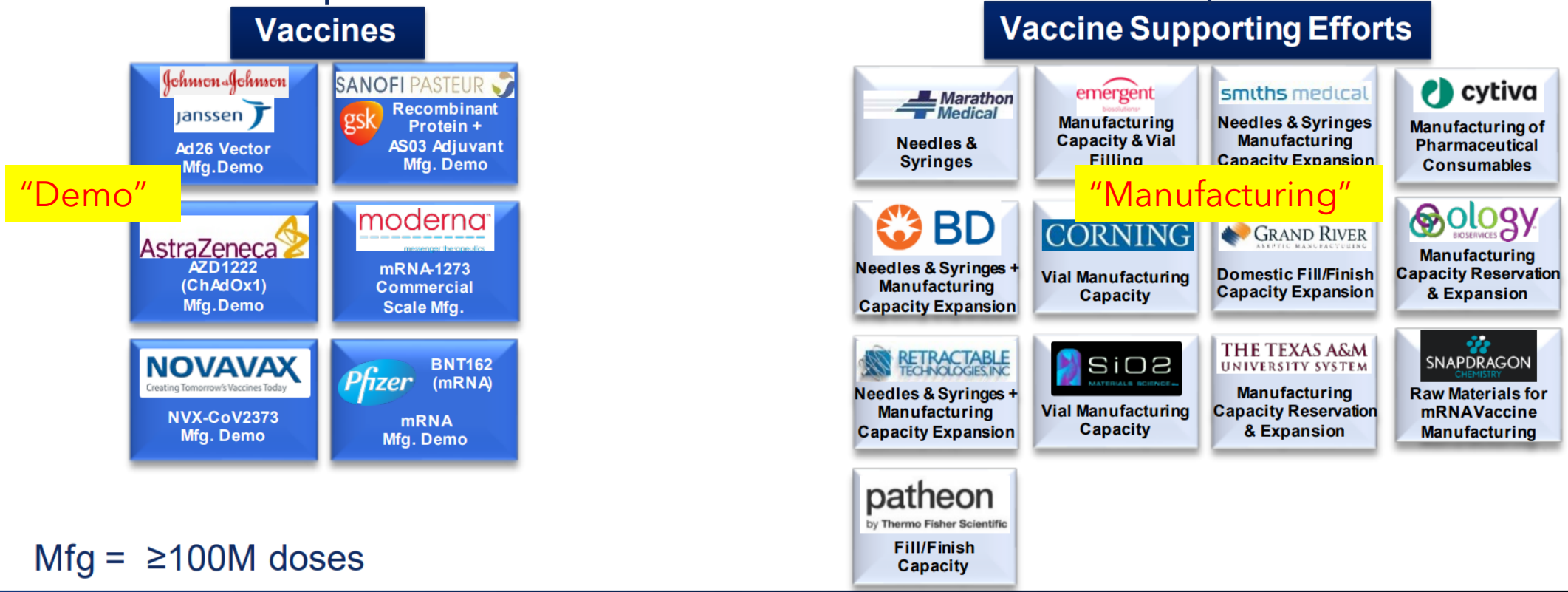
In charge: NSC, DOD, BARDA



Vaccine development and approval was a farse – military campaign

As of October 20, 2020

OWS/BARDA Vaccine Manufacturing Portfolio



Mfg = ≥100M doses

Pfizer Contract

1.2 Scope

The scope of this prototype project is the demonstration by Pfizer of the supply and logistics capability to manufacture and distribute to the Government of 100M doses of a novel mRNA-based vaccine that has received FDA-approval or authorization based on demonstration of efficacy (hereafter FDA-approved or authorized). The criteria for successful Emergency Use Authorization (EUA) are described in *Emergency Use Authorization of Medical Products and Related Authorities: Guidance for Industry and Other Stakeholders, January 2017*; and *Development and Licensure of Vaccine to Prevent COVID-19: Guidance for Industry June 2020*. The successful provision of these doses shall establish the effectiveness of a technology capable of potentially providing immediate and long-term solutions to coronavirus infections. While pre-clinical, clinical, and chemistry/manufacturing/controls (CMC) activities are described in the Background section of this Statement of Work, the Parties acknowledge and agree that such activities not related to the large-scale manufacturing demonstration are out-of-scope for this prototype project as Pfizer and BioNTech have and will continue to fund these activities, without the use of Government funding.

Pfizer Contract

DEPARTMENT OF THE ARMY
U.S. ARMY CONTRACTING COMMAND – NEW JERSEY
PICATINNY ARSENAL, NEW JERSEY 07806-5000

REPLY TO
ATTENTION OF

21 July 2020

Army Contracting Command – New Jersey
ACC-NJ, Building 9
Picatinny Arsenal, NJ 07806

SUBJECT: Technical Direction Letter for Medical CRBN Defense Consortium (MCDC), Request for Prototype Proposals (RPP) 20-11, Objective PRE-20-11 for "COVID-19 Pandemic – Large Scale Vaccine Manufacturing Demonstration" (Pfizer, Inc.)

REF: Prizer Request for Technical Direction Letter, RPP 20-11 under OTA W15QKN-16-9-1002 for Objective PRE-20-11, dated 20 July 2020

Advanced Technology International
ATTN: (b) (6), Sr. Contracts Manager
315 Sigma Drive
Summerville, SC 29486

Dear (b) (6),

The Army Contracting Command – New Jersey (ACC-NJ), in supporting the Joint Project Manager – Medical Countermeasure Systems (JPM-MCS), issued MCDC RPP 20-11 on 09 June 2020. Members of the MCDC submitted proposals in accordance with this RPP. The Government received and evaluated all proposal(s) submitted and a Basis of Selection has been executed, selecting Pfizer, Inc. as the awardee. The Government requests that a Firm-Fixed-Price Project Agreement be issued to Pfizer, Inc. to award this proposal under Other Transaction Agreement W15QKN-16-9-1002, to be performed in accordance with the attached Government Statement of Work (SOW).

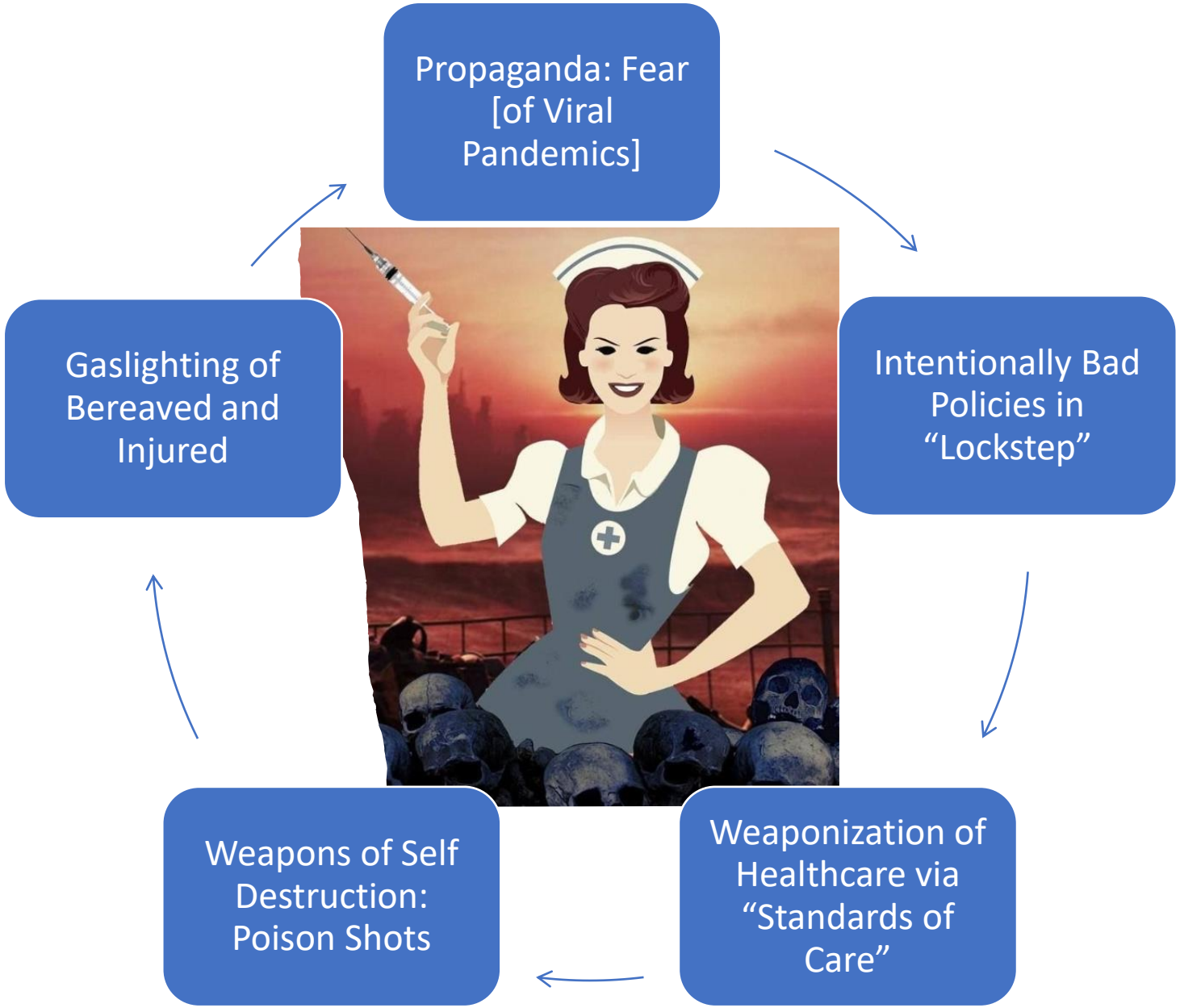
Based upon the acceptable update of Pfizer, Inc.'s proposal for "COVID-19 Pandemic – Large Scale Vaccine Manufacturing Demonstration" and 1) The Project Agreement Recipient's concurrence with the requirements included in the Government SOW; 2) An acceptable milestone schedule that meets SOW requirements, and; 3) The price proposed that has been analyzed by the Government, you are hereby directed to issue a Project Agreement to Pfizer, Inc. for the subject project. The total project value has been determined fair and reasonable and Pfizer, Inc.'s proposal has been selected IAW the above referenced Basis of Selection.

Already invoked in the Motion to Dismiss Brook Jackson's case (Apr 2022)

"Vaccine Development and Approval"- a performance art to convince the public

- The word "demonstration" (i.e. fake) in DOD contracts for vaccines
- Clinical trials were not ordered by DOD/HHS - not possible for countermeasures
- cGMP compliance was not ordered - not possible
- Legally there were no clinical trial subjects or investigators, and no informed consent

FDA leadership are impersonating the regulators and lying to the public – they have no authority to regulate countermeasures



US DoD Dictionary of Military and Associated Terms:

"Kill box = A three-dimensional area reference that enables timely, effective coordination and control, and facilitates rapid attacks."

THE WALL STREET JOURNAL.

THE WEALTH REPORT

Psychopaths

Try to Shrink World's Population, Report Says

By Robert Frank

May 26, 2009 11:57 am ET

Last week's meeting of the Great and the Good (or the Richest and Richer) was bound to draw criticism.

The New York meeting of billionaires Bill Gates, Warren Buffett, David Rockefeller, Eli Broad, George Soros, Ted Turner, Oprah, Michael Bloomberg and others was described by the Chronicle of Philanthropy as an informal gathering aimed at encouraging philanthropy. Just a few billionaires getting together for drinks and dinner and a friendly chat about how to promote charitable giving.

Associated Press

"…there was "nothing as crude as a vote" but a consensus emerged that they would back a strategy in which ***population growth would be tackled as a potentially disastrous environmental, social and industrial threat.****"*

Questions?