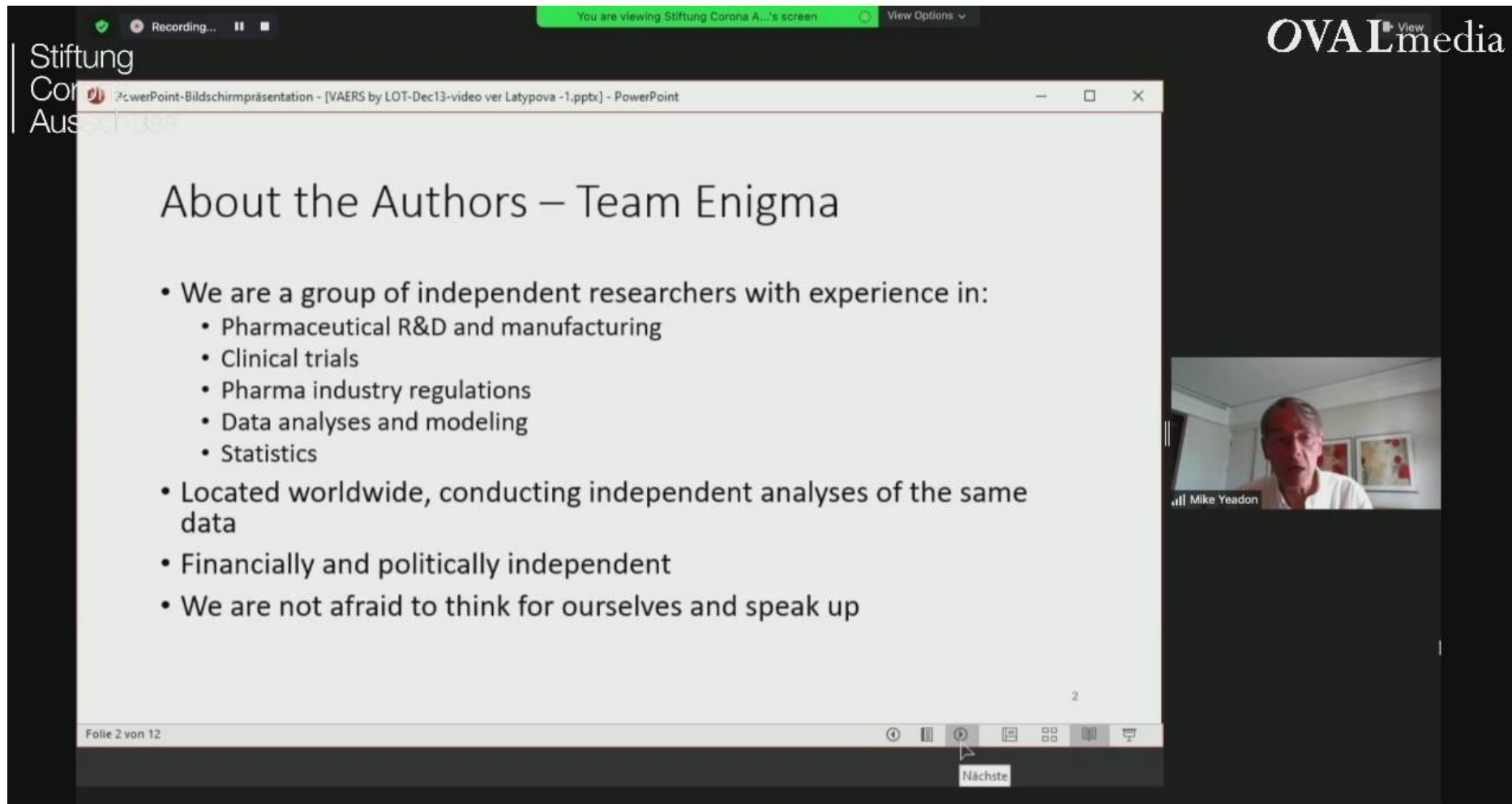


## Team Enigma Presentation

By Dr. Mike Yeadon 7 January 2022

Corona Investigative Committee Session 86, The Fog Is Lifting

<https://odysee.com/@Corona-Investigative-Committee:5/Mike-Session-86-en:0>



Recording... You are viewing Stiftung Corona A...s screen View Options

Stiftung  
Corona  
Aus...

### About the Authors – Team Enigma

- We are a group of independent researchers with experience in:
  - Pharmaceutical R&D and manufacturing
  - Clinical trials
  - Pharma industry regulations
  - Data analyses and modeling
  - Statistics
- Located worldwide, conducting independent analyses of the same data
- Financially and politically independent
- We are not afraid to think for ourselves and speak up

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Nächste

OVA Lmedia

Mike Yeadon

Dr Mike Yeadon: The variability in serious adverse events by vaccine lot “is calibration of a killing weapon”

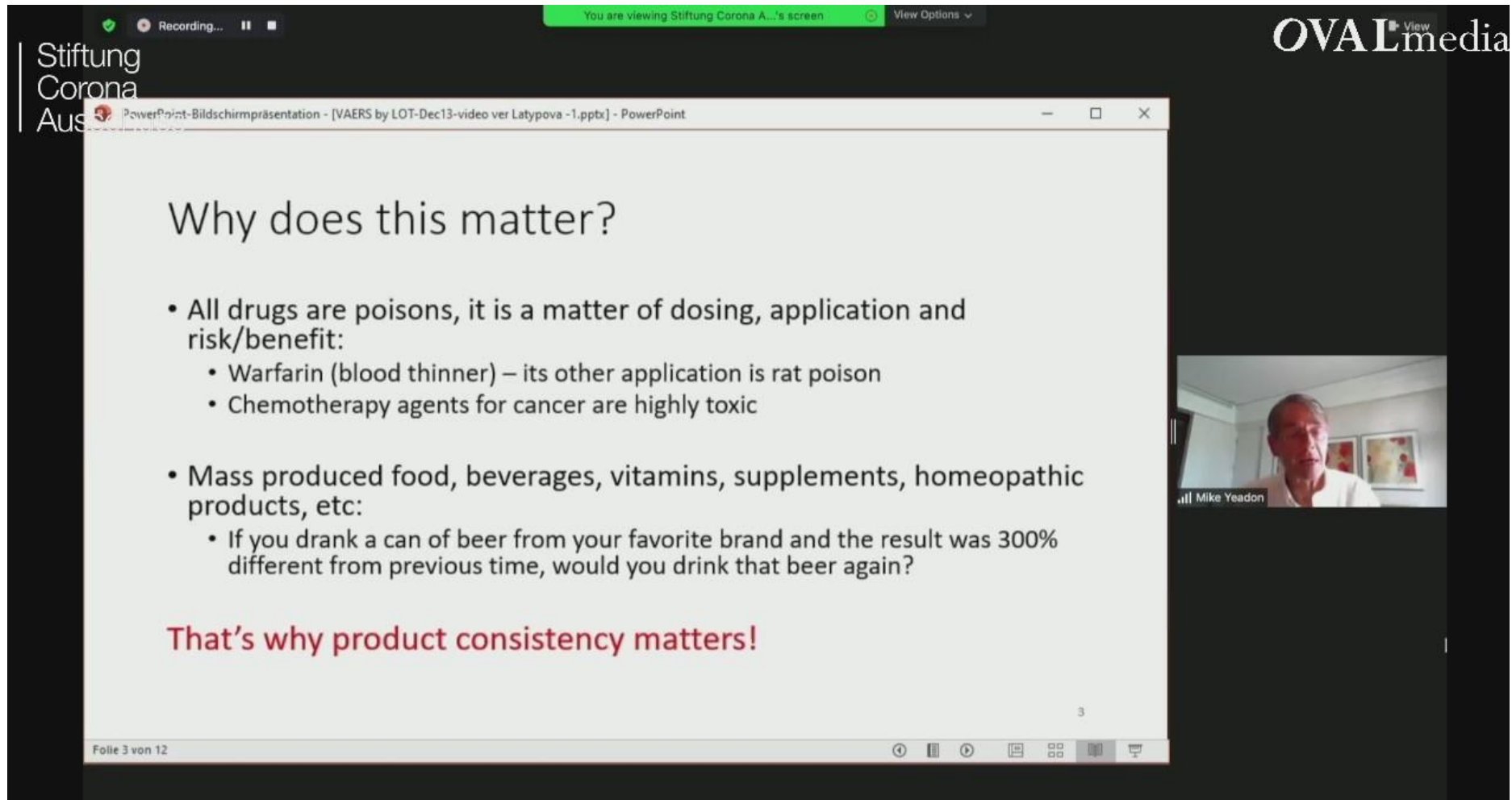
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The screenshot shows a Zoom meeting interface. At the top, there is a green status bar that reads "You are viewing Stiftung Corona A. 's screen" and a "View Options" dropdown. The Zoom logo "OVALmedia" is in the top right corner. The main content is a PowerPoint slide with the following text:

Stiftung Corona  
Aus

### Why does this matter?

- All drugs are poisons, it is a matter of dosing, application and risk/benefit:
  - Warfarin (blood thinner) – its other application is rat poison
  - Chemotherapy agents for cancer are highly toxic
- Mass produced food, beverages, vitamins, supplements, homeopathic products, etc:
  - If you drank a can of beer from your favorite brand and the result was 300% different from previous time, would you drink that beer again?

**That's why product consistency matters!**

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Mike Yeadon

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Stiftung Corona Aus

PowerPoint-Bildschirmpräsentation - [VAERS by LOT-Dec13-video ver Latypova - 1.pptx] - PowerPoint

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

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

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Mike Yeadon

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Stiftung Corona Aus

PowerPoint-Bildschirmpräsentation - [VAERS by LOT-Dec13-video ver Latypova -1.pptx] - PowerPoint

### VAERS Data: Covid 19 mRNA Vaccines vs Seasonal Flu Vaccines

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

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

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Folie 5 von 12

Reiner Fuellmich

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OVALmedia

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

Serious Adverse Event per Lot\* Sorted Alphabetically

Range= 1-37 reports per lot

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

Folie 6 von 12

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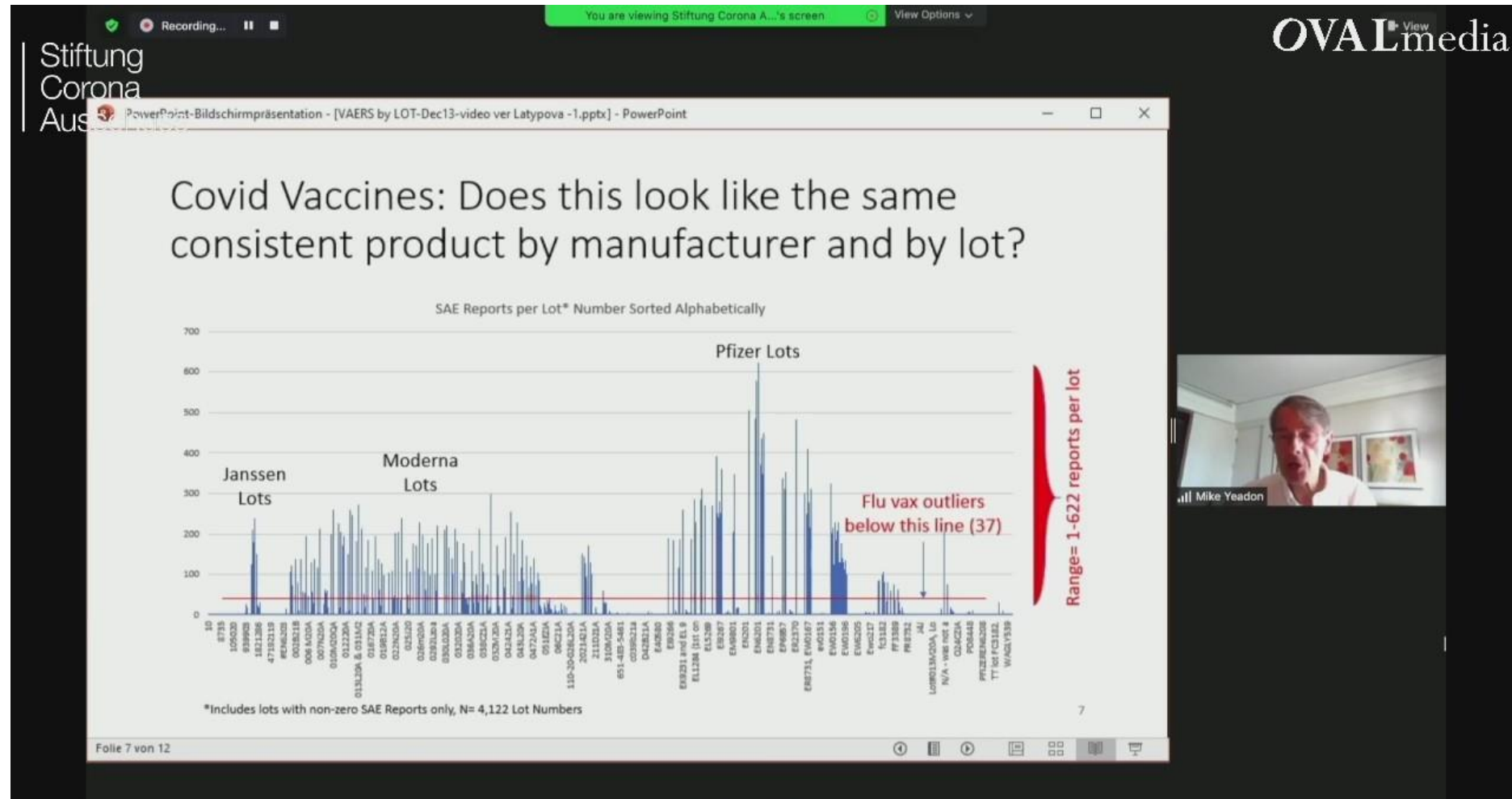
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