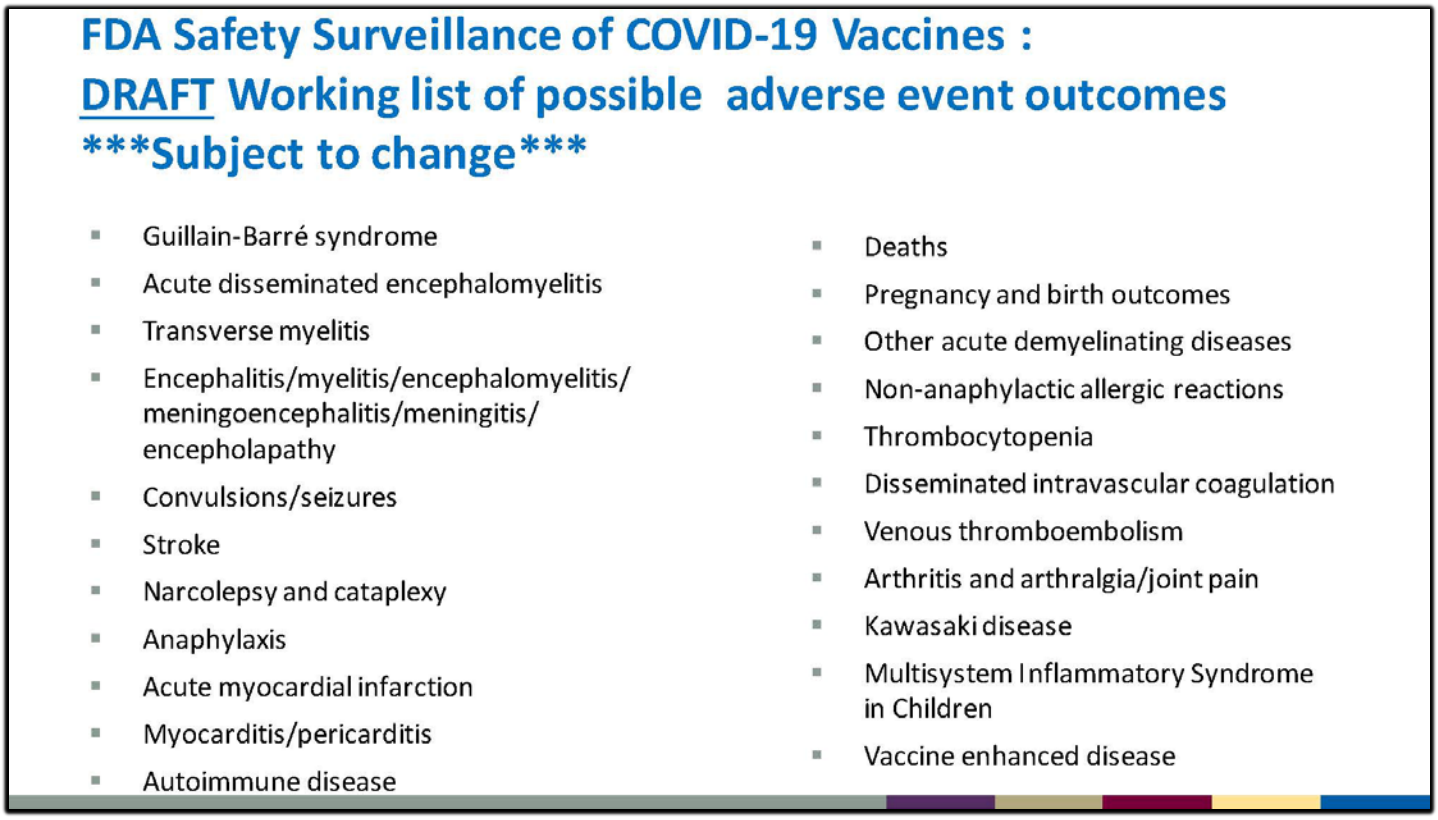
Observation memo from Paul V. Sheridan

Excerpted below, from the secret FDA meeting of 22 October 2020, Page 17 of 27:



FDA Safety Surveillance of COVID-19 Vaccines :
DRAFT Working list of possible adverse event outcomes
*****Subject to change*****

- Guillain-Barré syndrome
- Acute disseminated encephalomyelitis
- Transverse myelitis
- Encephalitis/myelitis/encephalomyelitis/ meningoencephalitis/meningitis/ encepholapathy
- Convulsions/seizures
- Stroke
- Narcolepsy and cataplexy
- Anaphylaxis
- Acute myocardial infarction
- Myocarditis/pericarditis
- Autoimmune disease
- Deaths
- Pregnancy and birth outcomes
- Other acute demyelinating diseases
- Non-anaphylactic allergic reactions
- Thrombocytopenia
- Disseminated intravascular coagulation
- Venous thromboembolism
- Arthritis and arthralgia/joint pain
- Kawasaki disease
- Multisystem Inflammatory Syndrome in Children
- Vaccine enhanced disease

Unbeknownst to the layperson, the above contains a fundamental but purposeful deception; a deception that was well-known by vested interests. This deception was well-known to Dr. Anthony Fauci, as well as Ivy League life science professors and immunological researchers:

The initial deception is in the titled use of the phrase: "COVID-19 Vaccines."

It was well-known that the needles from Moderna and Pfizer did not contain a vaccine in any historical, reasonable, or "standard art definition" use of that term. Prior-to and after release by the FDA (under their Emergency Use Authorization), the term "vaccine" was promoted from the highest members of government, academia, and corporate board rooms, **but its use was a lie.**

Concealed from the general public, the so-called *"COVID-19 vaccine,"* which was injected into billions worldwide, was known to contain an experimental modified RNA or self-amplifying RNA technology that had never been successful in *any* mammal, let-alone humans. Even the phrase *"mRNA vaccine"* is a lie. **But for emphasis and contrast, this memo asks the following *simple* question; looking at the FDA excerpt above:**

When did the flu vaccine, used for respiratory illness (ala COVID-19), over decades of use, ever have listed as a *"possible adverse event"* the outcome of :

Venous Thromboembolism ?!

In case members of the US Congress and US Senate missed it, lying is a form of "willful misconduct" . . .

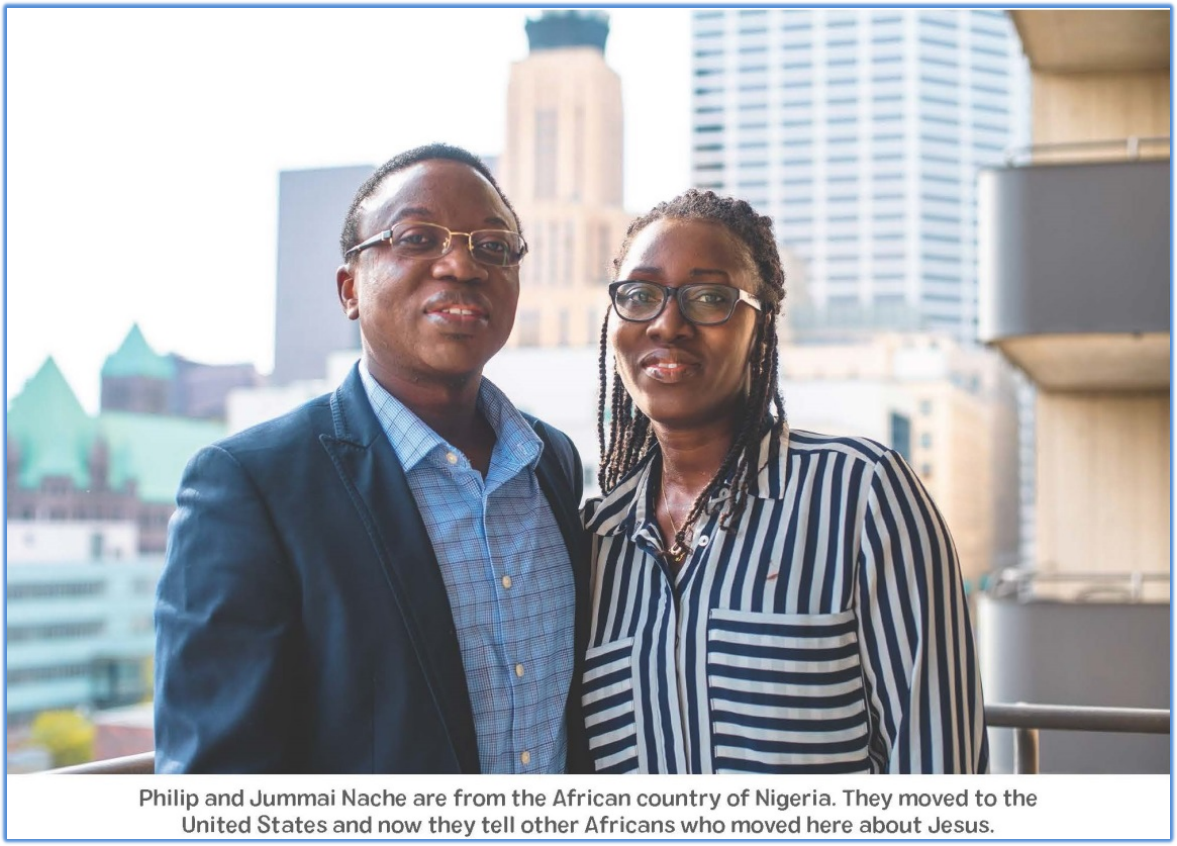
Subject : **Mrs. Jummai Nache and her Family**

Current Medical Status Post Employer Mandated Pfizer modRNA Injections

Mrs. Jummai Nache and devoted husband Philip; learning how to walk stairs on prosthetic limbs. Amputation of her limbs caused by modRNA injection mandated by her former employer, the University of Minnesota (UMinn).

A non-smoker, a non-drinker, a model of health prior to Pfizer needle; amputations in pictorial review caused by modRNA-induced venous and arterial thromboembolism. Such dangers were not merely foreseeable but discussed in-detail at secret FDA meeting of 22 October 2020. That meeting occurred prior to Emergency Use Authorization (EUA) meeting of 11 December 2020, where these "COVID-19 vaccine" dangers were ignored. **The EUA allowed Pfizer, hospital administrators, university officials, medical doctors, clinicians, nurses, et al. to inject the modRNA poison under the secretive protections of LIABILITY IMMUNITY.**

Subject : Mrs. Jummai Nache and her Family



Philip and Jummai Nache are from the African country of Nigeria. They moved to the United States and now they tell other Africans who moved here about Jesus.

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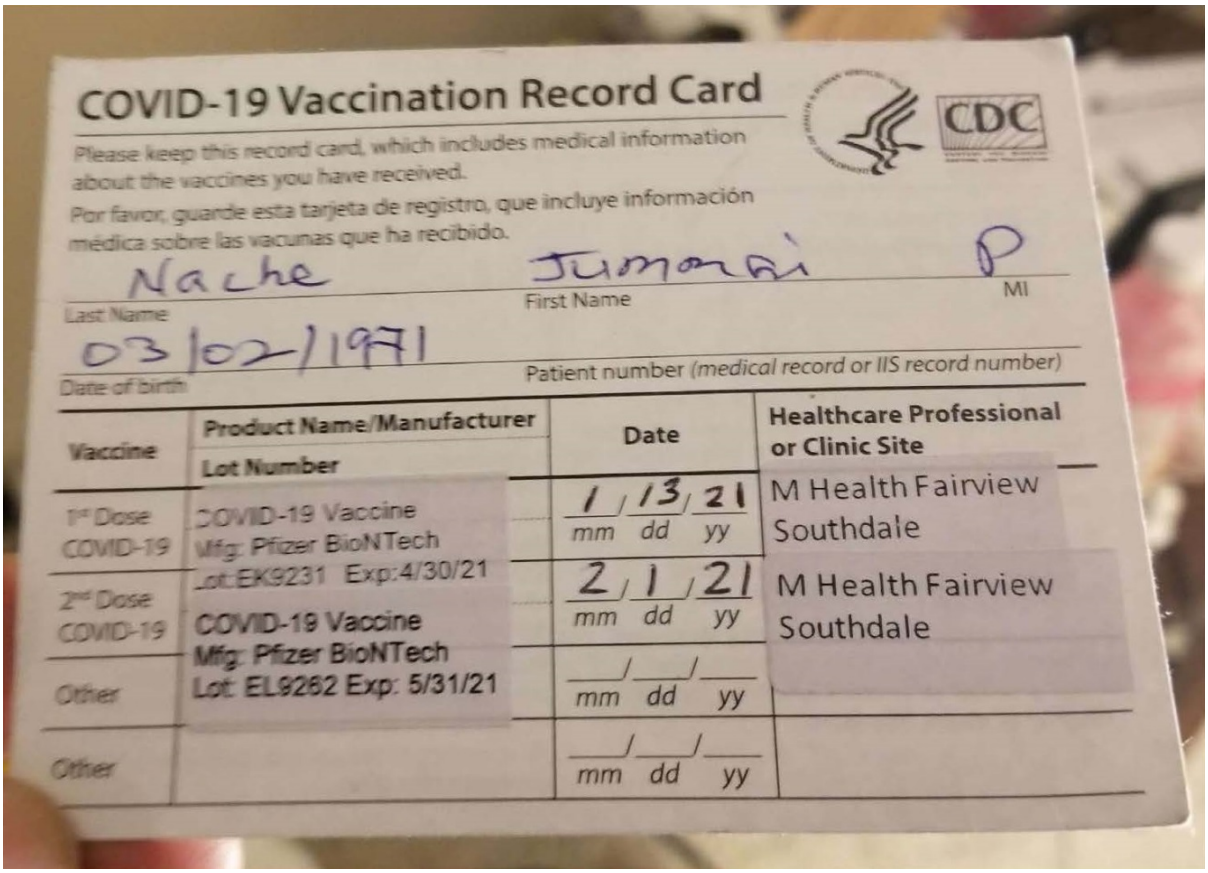


COVID-19 Vaccination Record Card

Please keep this record card, which includes medical information about the vaccines you have received.

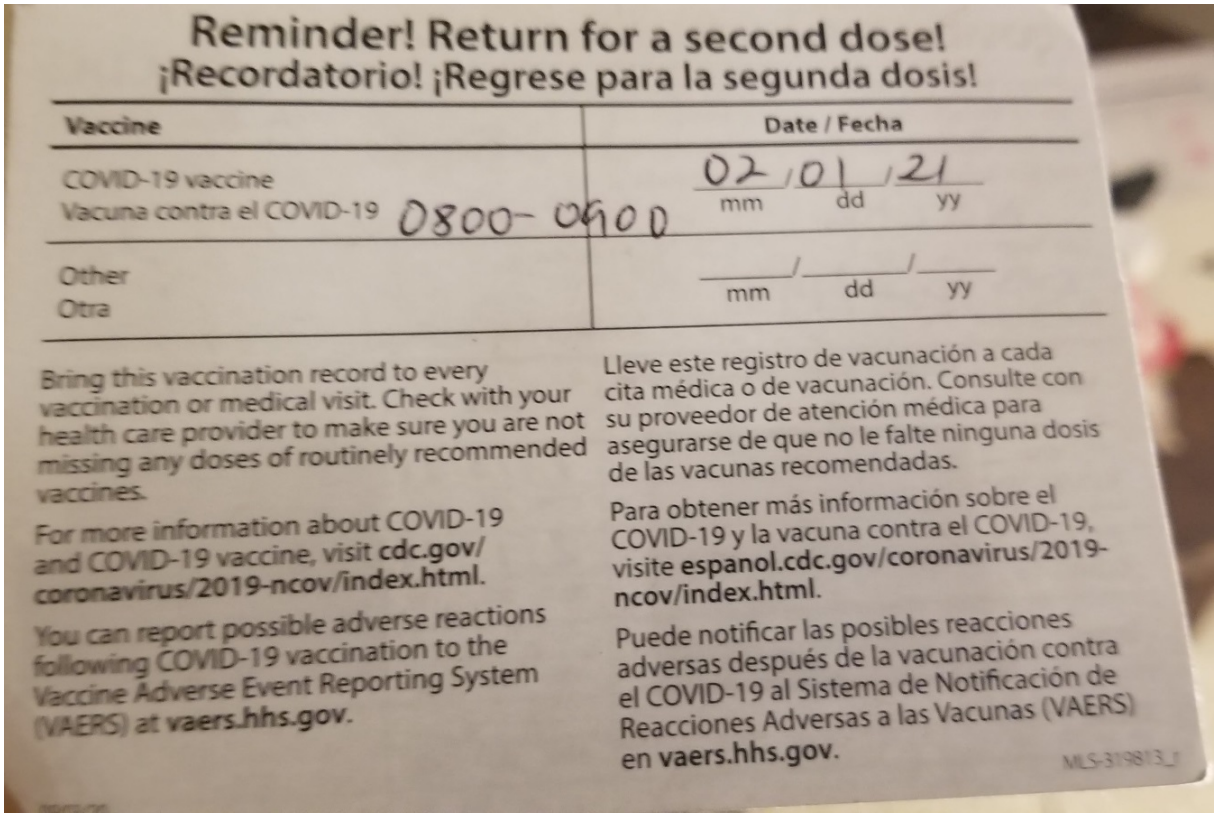
Por favor, guarde esta tarjeta de registro, que incluye información médica sobre las vacunas que ha recibido.

Nache (Last Name) Jummai (First Name) P (MI)

03/02/1971 (Date of birth)

Patient number (medical record or IIS record number)

Vaccine	Product Name/Manufacturer Lot Number	Date	Healthcare Professional or Clinic Site
1st Dose COVID-19	COVID-19 Vaccine Mfg: Pfizer BioNTech Lot:EK9231 Exp:4/30/21	1/13/21 (mm/dd/yy)	M Health Fairview Southdale
2nd Dose COVID-19	COVID-19 Vaccine Mfg: Pfizer BioNTech Lot: EL9262 Exp: 5/31/21	2/1/21 (mm/dd/yy)	M Health Fairview Southdale
Other		___/___/___ (mm/dd/yy)	
Other		___/___/___ (mm/dd/yy)	



Reminder! Return for a second dose!

¡Recordatorio! ¡Regrese para la segunda dosis!

Vaccine	Date / Fecha
COVID-19 vaccine Vacuna contra el COVID-19 0800-0900	02/01/21 (mm/dd/yy)
Other Otra	___/___/___ (mm/dd/yy)

Bring this vaccination record to every vaccination or medical visit. Check with your health care provider to make sure you are not missing any doses of routinely recommended vaccines.

For more information about COVID-19 and COVID-19 vaccine, visit **cdc.gov/coronavirus/2019-ncov/index.html**.

You can report possible adverse reactions following COVID-19 vaccination to the Vaccine Adverse Event Reporting System (VAERS) at **vaers.hhs.gov**.

Lleve este registro de vacunación a cada cita médica o de vacunación. Consulte con su proveedor de atención médica para asegurarse de que no le falte ninguna dosis de las vacunas recomendadas.

Para obtener más información sobre el COVID-19 y la vacuna contra el COVID-19, visite **espanol.cdc.gov/coronavirus/2019-ncov/index.html**.

Puede notificar las posibles reacciones adversas después de la vacunación contra el COVID-19 al Sistema de Notificación de Reacciones Adversas a las Vacunas (VAERS) en **vaers.hhs.gov**.

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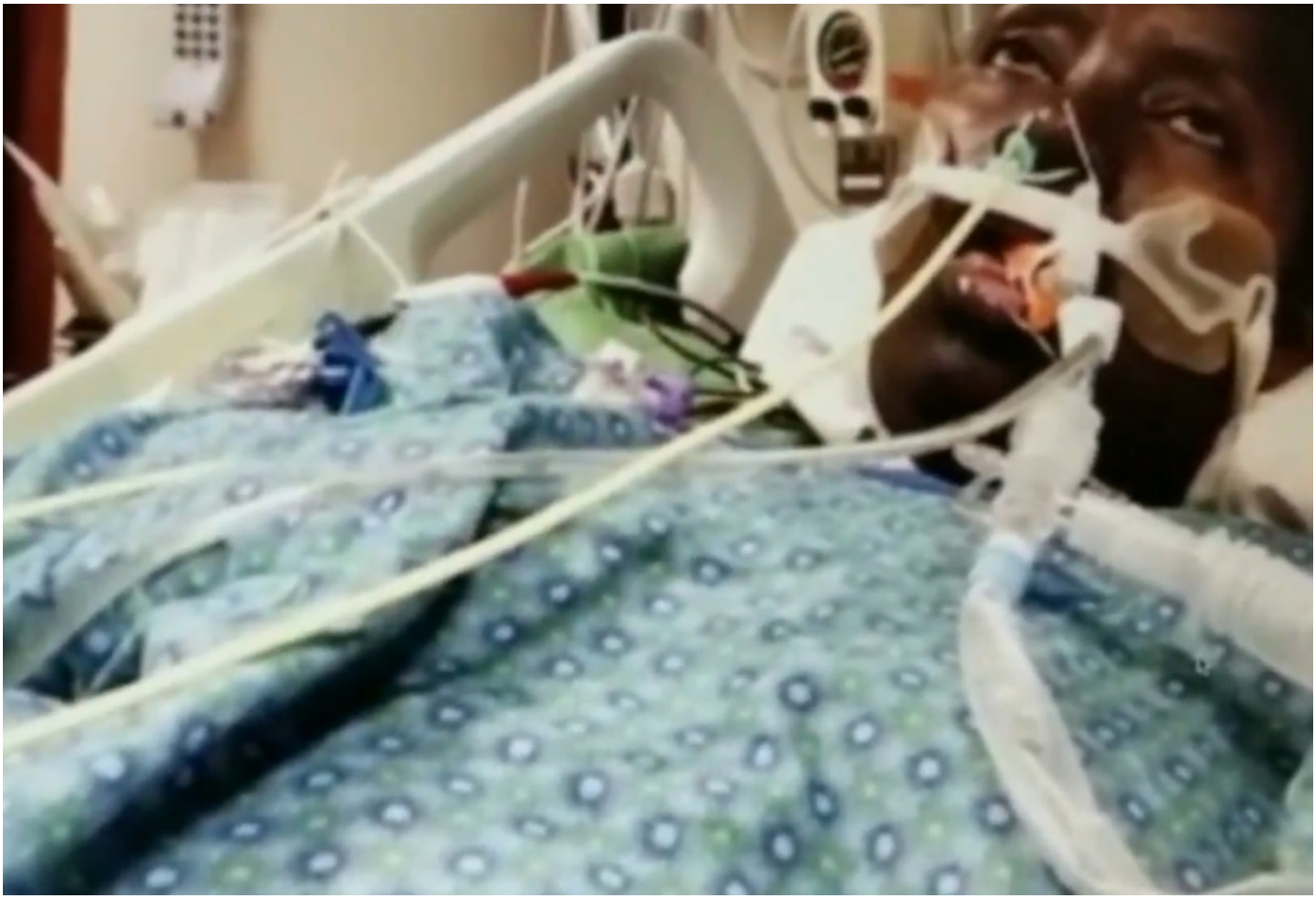

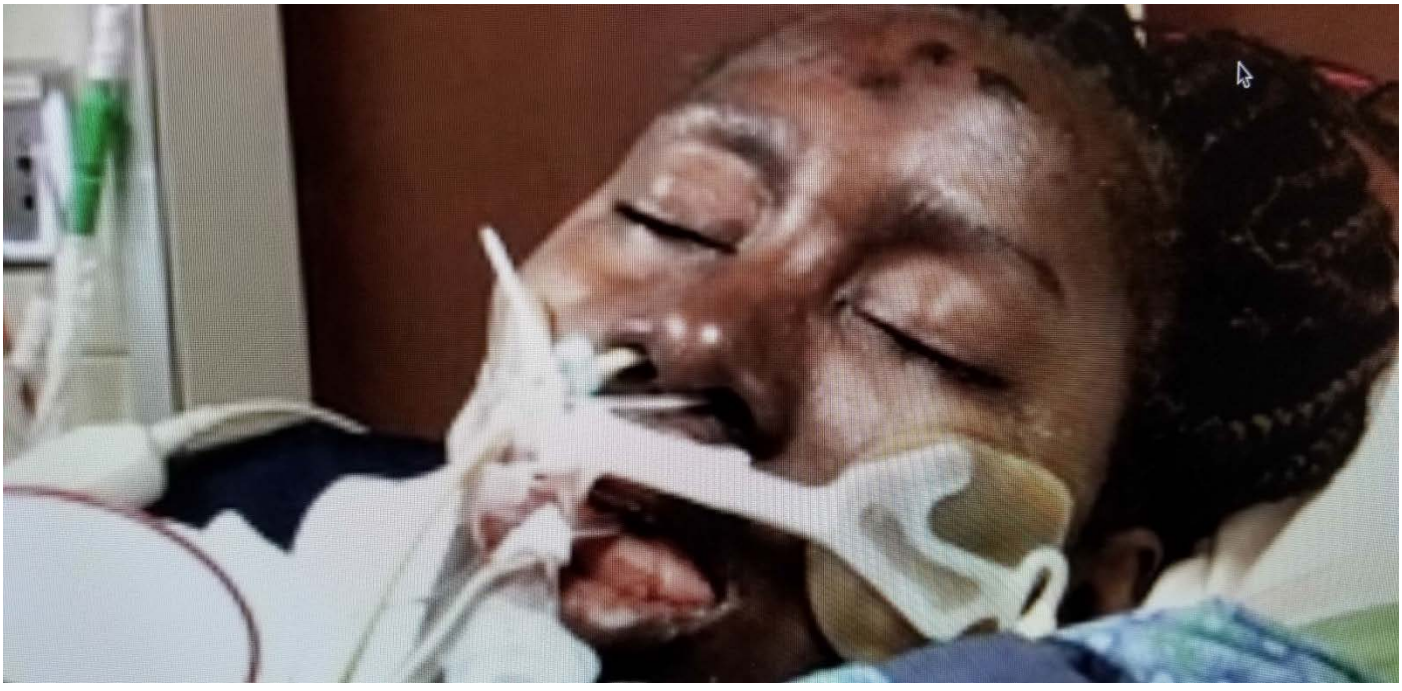

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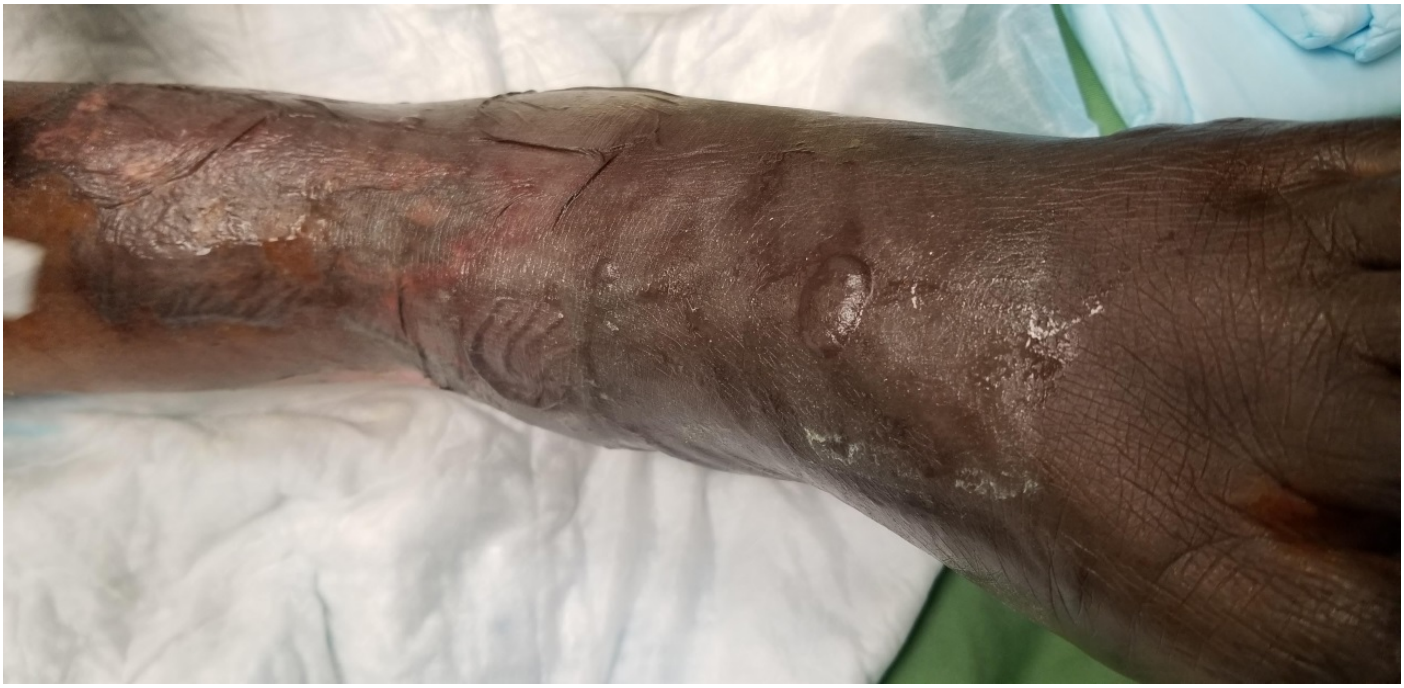

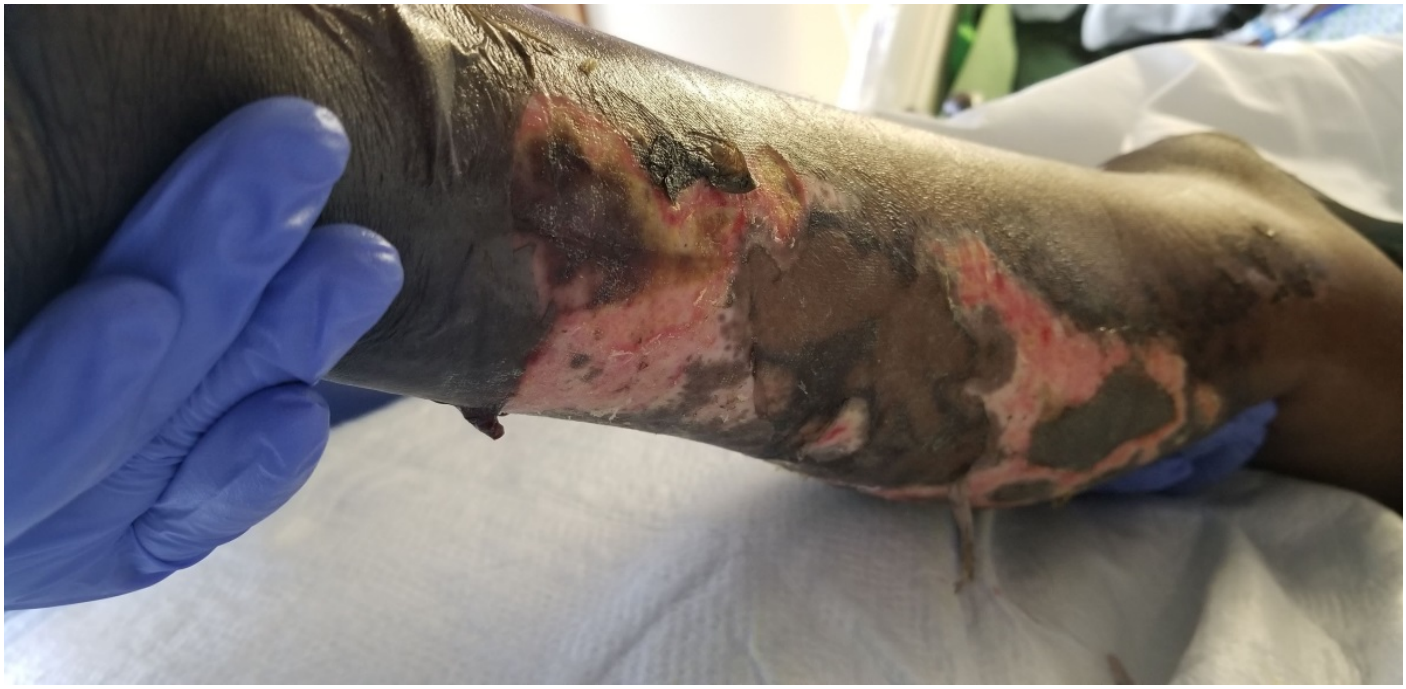

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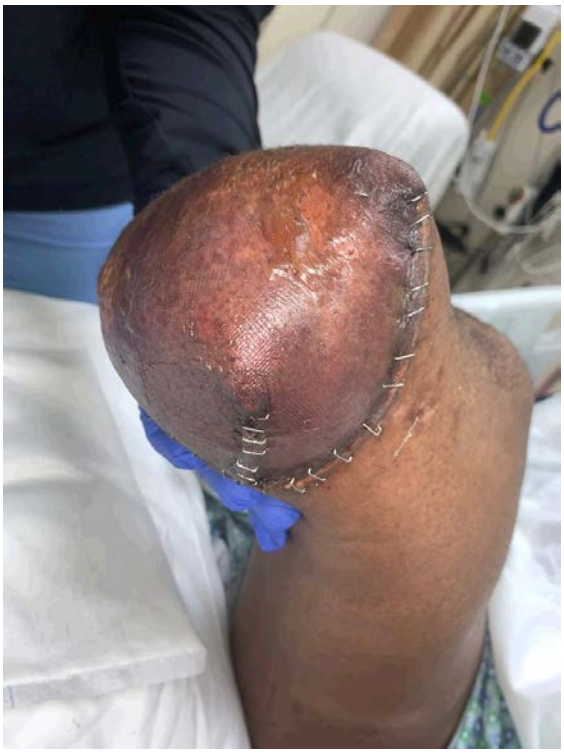

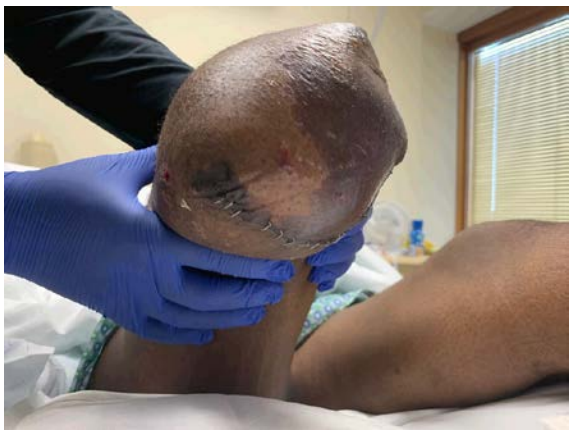

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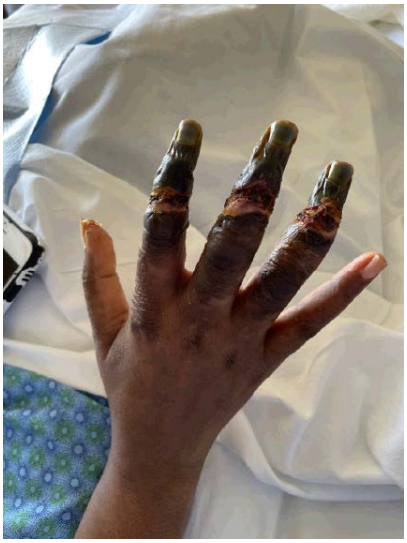

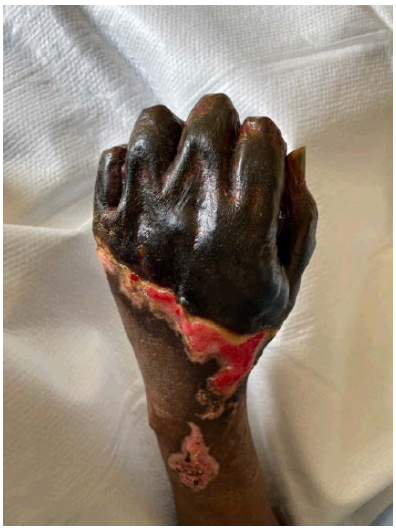

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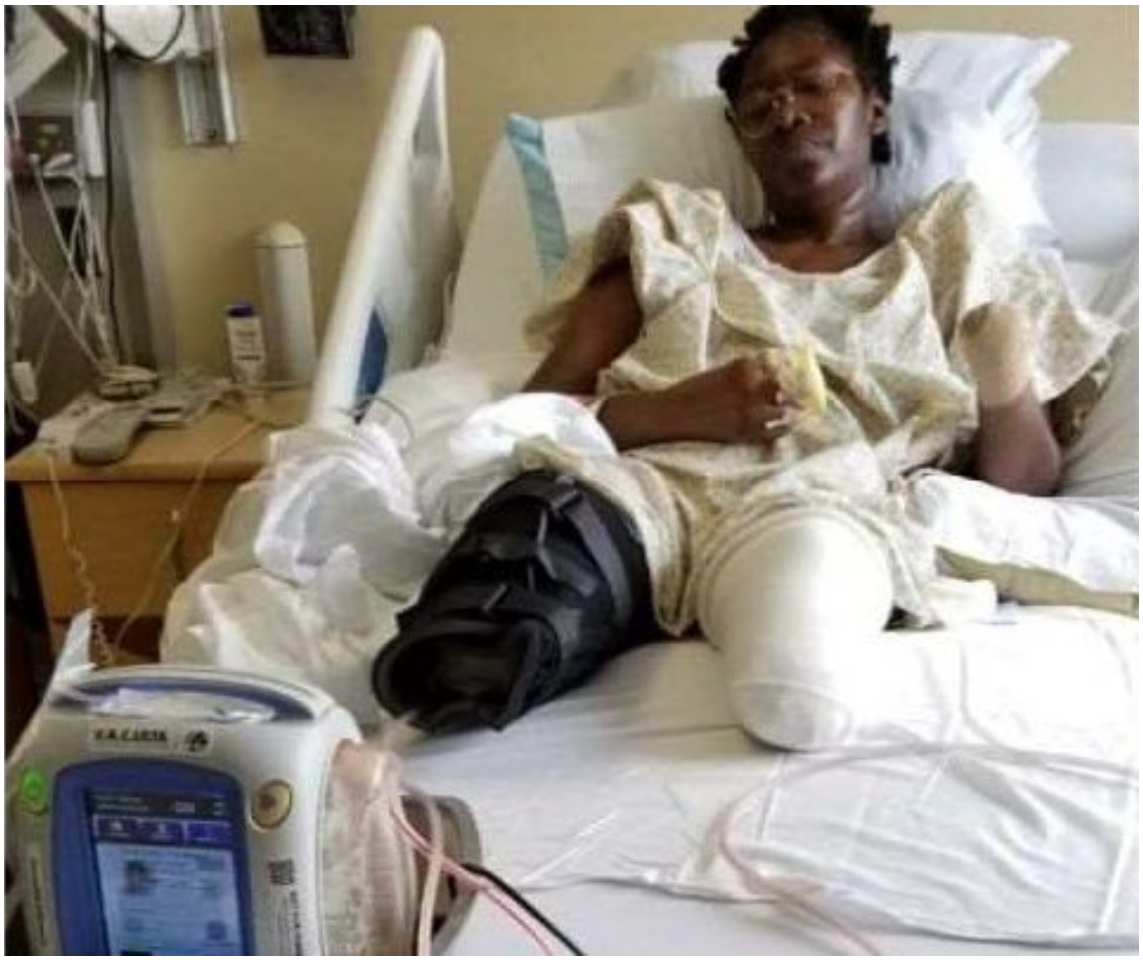

Excerpt Page 3 of 22: ***" I am confident that AG Ellison, and the other eleven AG signatories . . . acted in-cooperation with the Biden White House. "***

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A non-smoker, a non-drinker, a model of health **prior** to Pfizer needle; amputations in pictorial review (Tab 5 below) caused by modRNA-induced venous and arterial thromboembolism; not merely foreseeable but a known "mRNA" danger discussed in-detail at the secret FDA meeting of 22 October 2020, which occurred **prior** to their Emergency Use Authorization (EUA) of 11 December 2020. The EUA allowed Pfizer, hospital administrators, university officials, medical doctors, clinicians, nurses, et al. to inject the modRNA poison under the secretive protections of **LIABILITY IMMUNITY.** *(Please see CONCLUSION Part A, Page 15 of 22).*