MEDICAL VERITAS

THE JOURNAL OF TRUTH IN HEALTH SCIENCE

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NEWS RELEASE

Release: No. 9-Pfizer/Moderna Lawsuit
Date Mailed: February 22, 2021

For Immediate Release

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Pfizer/Moderna Acts to Hide Genetic Safety Science from Vaccine Discovery in Federal Court

Las Vegas, NV (MedicalVeritas.org)—Pfizer and Moderna companies, along with vaccine distributor, Henry Schein, Inc., have filed to hide discovery of genetic safety science in favor of billions made in governmental sales according to a federal lawsuit brought by whistleblower and former AIDS science Schein advisor, Dr. Leonard G. Horowitz.

Horowitz claims the companies are guilty of unfair competition and deceptive advertising risking civilization's extinction through vaccine-induced viral mutations.

Multiple mutants are now emerging globally as accurately predicted by the doctor in the wake of the defendants' "novel" mRNA vaccines. The first mutants were reported less than two weeks after Pfizer's mass vaccination campaign began in the UK.

These "genetic therapies" deliver their "payloads" by multiplying massive numbers of AIDS-science-discovered spike proteins, the doctor's lawsuit explains. The manufactured "antigens" attach to human cells like keys unlocking the door to DNA function. The corrupted DNA then makes more of the same key entry-proteins. These spike protein poisons, evidenced to deliver AIDS virus genes, are now spreading throughout populations to presumably produce protective antibodies and 'herd immunity.' But at the same time the genetic-poisons' shed from vaccinated people, and may be spreading, now worsening risks of mutating and multiplying the pandemic.

On February 21, 2021, a new mutant reportedly emerged in Russia combining the coronavirus with the avian flu, H5N8.

"This indicates the risk to civilization is skyrocketing," Horowitz said. "As vaccinated people shed or spread the airborne viral antigens these genetic risks will merge with other viruses, such as herpes, H5N1, or even Ebola. That will 'hyper-weaponize' an increasing number of more deadly strains."

The South African strain contains just such a spike protein. This mutation has been reported to be more deadly.

Meanwhile, the doctor alleges, the companies have not only "neglected crucial genetic safety studies and falsely advertised their vaccines' safety," but Pfizer and Moderna have filed in federal court to block such discovery altogether.

"Vaccination hesitancy" may actually be preventing such risks to humanity," Dr. Horowitz explained. "As these companies spend fortunes, delay discoveries, and conceal here and elsewhere, vaccination hesitancy increases contrary to officials' intentions."

Aside from the genetic safety concerns raised by Horowitz, the efficacy of Pfizer and Moderna's mRNA products has been solidly discredited by the associate editor of the *British Medical Journal*, Peter Doshi, in two hotly-contested articles.

According to Amazon and *Wikipedia*, Horowitz is a Harvard-trained public health expert who authored the best-selling book in the field of communicable diseases, *Emerging Viruses: AIDS & Ebola—Nature, Accident or Intentional?* He is also an award-winning filmmaker and pioneer of the competing anti-viral and vaccine alternative branded OxySilver™—a NASA science advancement. He was subsequently smeared, censored, and deprived of his career in professional education and humanitarian medicine.

Viewed as a pharmaceutical industry competitor, Dr. Horowitz filed this lawsuit after discovering the defendants had falsely advertised the COVID vaccines were "safe" while neglecting and concealing genetic risks. Schein had similarly concealed AIDS science when Horowitz had advanced it and objected to their officials' alleged cover-up.

The questionable mRNA vaccines were approved by the FDA despite 23% of expert commissioners voting in opposition.

As a vaccine risk analyst and industry competitor, the doctor argues that the FDA violated its Emergency Use Authorization ("EUA") requirements and the public's trust. "The 'fast-tracked' vaccines breached the EUA because alternatives, such as anti-oxidants, including OxySilver™ , were available. Much like hydroxychloroquine, OxySilver™ was neglected, unfairly dismissed, and maliciously maligned."

"These are monopolistic practices," Horowitz said. "They are illegal. 'Regulatory capture' by these drug companies best explains the FDA's actions."

Defendants Pfizer, Moderna and Schein filed to inform federal case magistrate, Nicholas P. Mizell, that they intend to file motions to stay discovery of the genetic science requested by Horowitz, pending a motion to dismiss the case to conceal the evidence and their liability.

If the enterprise's motions are granted, the urgent information will remain hidden, and irreparable harm to society will compound according to the complaint.

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