

To: President Donald Trump, President of the United States

Robert F Kennedy Jr, Secretary of Health and Human Services

Jay Battacharya, Director of National Institute of Health

Marty Makary, Director of Food and Drug Administration

Date: 12th May 2025

This notice of extreme concern about COVID-19 modified mRNA vaccine safety and quality was originally sent on 25<sup>th</sup> November 2024 to prime ministers and governing bodies of the Nordic and Baltic countries and the United Kingdom. Representatives of other European countries and Canada have since joined the call for answers.

Excessive levels of residual DNA were identified in Australian samples<sup>1</sup>, confirming data from France<sup>2</sup>, Germany<sup>3</sup>, Canada<sup>4</sup>, and the USA<sup>5</sup>.

The introduction of foreign DNA into cells via lipid nanoparticles (LNPs)<sup>6</sup> may damage human DNA leading to genomic instability, cancer, and other extremely serious conditions.

As an international group of politicians and qualified professionals, we are gravely concerned about the effects of COVID-19 modified mRNA vaccines on our populations and call for their immediate suspension. We wish to focus your attention on three critical issues regarding the deployment of COVID-19 modified mRNA products that have profound implications for the health of our nations' citizens.

- COVID-19 vaccines were never tested for their ability to block viral transmission<sup>7</sup>. Hence, medical product regulators, as well as governments and governmental bodies, misled people in order to coerce them into accepting these products.
- COVID-19 vaccines resulted in an unprecedented level of reported side effects<sup>8</sup>, including deaths<sup>9</sup>. Reproducible analyses of public data<sup>10,11,12,13</sup> shows that it was a lottery as to which batch a person received and the side effects that they may have experienced.
- Analyses by multiple, independent scientists evidence variable and excessive levels of residual plasmid DNA in vials of Pfizer and Moderna's products this foreign DNA is a by-product of the manufacturing processes and should never have made it into commercial vials.

The above points raise critical but unquantified risks for human health that have been systematically avoided by the regulatory authorities charged with their oversight. Therefore, on behalf of the public we call for:

- An immediate halt to the use of COVID-19 modified mRNA vaccines and a product recall.
- An independent and transparent investigation into their approval and use.
- Scientific evidence that documents that there is absolutely no risk of damage to human DNA.

For more detailed information please see the accompanying <u>scientific summary</u>. Considering this information, we ask for clear answers to the following questions *regarding the regulatory obligations*:

- 1. When will an independent, transparent, public, and forensic investigation into the safety and quality concerns of modified mRNA vaccines against COVID-19 be initiated based on the available evidence?
- 2. When will the causality between mRNA vaccines and cancer, infertility, or other acute, chronic, and genetic diseases be scientifically examined in a sufficient manner?



With this scientifically supported threat to current human health and that of future generations, we have identified the risk. We respectfully ask that you act in the interest of your country's citizens, upon which you were elected.

With the utmost concern and respect,

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#### Signatures from Elected Officials (Elected officials are also listed in their respective countries)

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