



To: AN TAOISEACH, Mr. Micheál Martin

Date: 28<sup>th</sup> 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 and on 29<sup>th</sup> November to Taoiseach, Mr. Simon Harris. Representatives of many 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 [scientific summary](#) and supplementary [letter](#). 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,

**Ms. Una McGurk S.C. Barrister,**  
Senior Counsel

**Professor Francis A. Boyle** Author of the U.S. domestic implementing legislation for the Biological Weapons Convention known as the Biological Weapons Anti-terrorist Act of 1989 that was passed unanimously by both Houses of the United States Congress and signed into law by U.S. President George H.W. Bush with the approval of the U.S. Department of Justice.

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**Nikola Grmoja**

**Nino Raspudić**  
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**Marin Miletić**  
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*'Raising awareness about the potential harms posed by modified mRNA therapeutics'*

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*'Raising awareness about the potential harms posed by modified mRNA therapeutics'*

[NORTHGroup.info](https://NORTHGroup.info), version 3

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*'Raising awareness about the potential harms posed by modified mRNA therapeutics'*

[NORTHGroup.info](https://NORTHGroup.info), version 3

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*'Raising awareness about the potential harms posed by modified mRNA therapeutics'*

[NORTHGroup.info](https://NORTHGroup.info), version 3

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*'Raising awareness about the potential harms posed by modified mRNA therapeutics'*

[NORTHGroup.info](https://northgroup.info), version 3

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