

Para: **Añada aquí su Jefe de Estado, ministro de Sanidad, etc. (cartas juntas o separadas).**

Fecha: **25 de noviembre de 2024 (cambie a la fecha en que la envíe).**

Este aviso de extrema preocupación sobre la seguridad y calidad de la vacuna de ARNm modificado COVID-19 se envió el 25 de noviembre de 2024 a los primeros ministros y órganos de gobierno de los países nórdicos y bálticos y del Reino Unido. Representantes de otros países europeos que comparten nuestras preocupaciones se han unido desde entonces a la petición de respuestas.

Se han detectado niveles excesivos de ADN residual en muestras australianas¹, lo que confirma los datos de Francia², Alemania³, Canadá⁴ y Estados Unidos⁵.

La introducción de ADN extraño en las células a través de nanopartículas lipídicas⁶ puede dañar el ADN humano y provocar inestabilidad genómica, cáncer y otras enfermedades muy graves.

Como grupo internacional de políticos y profesionales cualificados, estamos gravemente preocupados por los efectos de las vacunas de ARNm modificado con COVID-19 en nuestras poblaciones y pedimos su suspensión inmediata. Deseamos centrar su atención en tres cuestiones críticas relativas al despliegue de los productos de ARNm modificado con COVID-19 que tienen profundas implicaciones para la salud de los ciudadanos de nuestras naciones.

- Las vacunas contra la COVID-19 nunca fueron probadas para determinar su capacidad de bloquear la transmisión viral⁷. Por ello, los reguladores de productos médicos, así como los gobiernos y los organismos gubernamentales, engañaron a las personas para obligarlas a aceptar estos productos.
- Las vacunas contra la COVID-19 dieron lugar a un nivel sin precedentes de efectos secundarios notificados⁸, incluido muertes⁹. Análisis reproducibles de datos públicos^{10,11,12,13} muestran que fue una lotería qué lote recibió una persona y los efectos secundarios que pudo haber experimentado.
- Los análisis realizados por múltiples científicos independientes evidencian niveles variables y excesivos de ADN plasmídico residual en los viales de los productos de Pfizer y Moderna -este ADN extraño es un subproducto de los procesos de fabricación y nunca debería haber llegado a los viales comerciales.

Los puntos anteriores plantean riesgos críticos, pero no cuantificados para la salud humana que han sido sistemáticamente eludidos por las autoridades reguladoras encargadas de su supervisión. Por lo tanto, en nombre del público pedimos:

- La suspensión inmediata del uso de vacunas de ARNm y el retirada del producto.
- Una investigación independiente y transparente sobre su aprobación y uso.
- Evidencia científica que documente que no existe absolutamente ningún riesgo de daño al ADN humano.

Para información más detallada, consulte el [resumen científico](#) adjunto. Teniendo en cuenta esta información, solicitamos respuestas claras a las siguientes preguntas:



1. ¿Qué se necesita para establecer una investigación forense independiente y transparente sobre los procesos regulatorios que conducen a la aprobación de estos productos?
2. ¿Hay algo que impida al Ministro iniciar y priorizar investigaciones sobre posibles vínculos entre las vacunas de ARNm y el cáncer, la infertilidad u otras enfermedades crónicas a las que se las ha vinculado?

Ante esta amenaza, respaldada científicamente, a la salud humana actual y a la de las generaciones futuras, hemos identificado el riesgo. Le pedimos respetuosamente que actúe en interés de los ciudadanos de su país, por el cual fue elegido.

Con la mayor preocupación y respeto,

en nombre de los miembros del grupo NORTH, lista completa de firmantes a continuación

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