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*Internationale Initiative zur Gewährleistung der Arzneimittelsicherheit
von modifizierten mRNA-Impfstoffen*

An:

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- 3) Nationalratspräsidentin Maja Riniker, Parlamentsgebäude, Generalsekretariat, 3003
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Liestal, 12. Juli 2025

Betreff: Dringender Appell zur Ausübung der Aufsichtspflichten betreffend Arzneimittelsicherheit aufgrund gravierender Sicherheits- und Qualitätsbedenken von Comirnaty (Biontech/Pfizer) & Spikevax (Moderna)

Sehr geehrte Frau Bundesrätin Baume-Schneider
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Sehr geehrte Frau Nationalratspräsidentin Riniker

Das Einbringen fremder DNA in Zellen über Lipid-Nanopartikel (LNP) kann die menschliche DNA schädigen, was unter anderem zu genomischer Instabilität, Krebs und anderen schwerwiegenden Erkrankungen führen [kann](#)¹.

In [australischen Proben](#)² wurde eine übermäßige Menge an Rest-DNA festgestellt, wie auch zuvor in [Frankreich](#)³, [Deutschland](#)⁴, [Kanada](#)⁵ und den [USA](#)⁶ bestätigt.

Diese Schreiben ist Teil der internationalen Initiative *The North Group* zur Gewährleistung der Arzneimittelsicherheit von mRNA modifizierten Impfstoffen. Die unterstützenden Politiker und qualifizierte Fachleute (siehe Anhang I) fordern aufgrund der potenziell negativen gesundheitlichen Auswirkungen von COVID-19 modifizierten mRNA-Impfstoffen auf unsere Bevölkerungen die sofortige Aussetzung der Zulassungen von den zuständigen Verantwortlichen.

Wir möchten Ihre Aufmerksamkeit auf drei kritische Punkte im Zusammenhang mit dem Einsatz von COVID-19 modifizierten mRNA-Produkten lenken, die potenziell negative Auswirkungen auf die menschliche Gesundheit haben können.

- Die COVID-19-Impfstoffe wurden nie auf ihre Fähigkeit getestet, die [Übertragung von Viren zu verhindern](#).⁷ Nichtsdestotrotz haben die Regulierungsbehörden für Medizinprodukte sowie Regierungen und staatliche Behörden die Menschen in die Irre geführt, um sie zu zwingen, diese Produkte zu akzeptieren.

- Die COVID-19-Impfstoffe führten zu einer noch nie dagewesenen [Anzahl gemeldeter Nebenwirkungen](#),⁸ darunter auch [Todesfälle](#).⁹ [Reproduzierbare Analysen öffentlicher Daten](#) bestätigen, dass einzelne Chargen/Batches mit besonders hohen Nebenwirkungsraten korrelieren.^{10,11,12,13}
- Analysen mehrerer unabhängiger Wissenschaftler belegen, dass in den Fläschchen der Produkte von Pfizer und Moderna unterschiedliche und übermäßige Mengen an Restplasmid-DNA vorhanden sind. Diese Fremd-DNA ist ein Nebenprodukt des Herstellungsprozesses und hätte niemals in die kommerziellen Fläschchen gelangen dürfen.

Die oben genannte Evidenz deutet auf unabsehbare Gesundheitsrisiken hin, aufgrund derer die staatlichen Aufsichtsbehörden über Arzneimitteln umgehend zum Tätigwerden gemäß den nationalen Bestimmungen des Heilmittelgesetzes Art. 66 Abs. 3 HMG bzw. nach Art. 59 HMG verpflichtet sind.

Daher fordern wir im Namen der Öffentlichkeit Folgendes:

- Sofortiger Stopp der Verwendung von modifizierten mRNA-Impfstoffen gegen Covid-19 und Rückruf des Produkts.
- Eine unabhängige und transparente Untersuchung ihrer Zulassung und Anwendung.
- Neue, transparent durchgeführte wissenschaftliche Studien, um die Frage zu klären, ob ein Risiko einer Schädigung der menschlichen DNA besteht oder nicht.

Ausführlichere Informationen finden Sie in dem Begleitschreiben ‚Wissenschaftliche Zusammenfassung‘ in Anhang II.

In Anbetracht dieser Informationen bitten wir im Hinblick auf die aufsichtsbehördlichen Pflichten um Beantwortung folgender Fragen:

1. *Wann wird auf Basis der vorliegenden Beweise eine unabhängige, transparente, öffentliche Untersuchung über die Sicherheits- und Qualitätsbedenken von modifizierten mRNA Impfstoffen gegen Covid veranlasst?*
2. *Wann wird die Kausalität zwischen mRNA-Impfstoffen und Krebs, Unfruchtbarkeit oder anderen akuten, chronischen und genetischen Krankheiten wissenschaftlich hinreichend untersucht?*

Die Gesundheit der Bevölkerung einschließlich zukünftiger Generationen ist durch die fortgesetzte Verabreichung der mRNA-Impfstoffe akut bedroht.

Wir ersuchen Sie daher höflich, im Interesse der Bürger unseres Landes ihren gesetzlichen Auftrag zur Gewährleistung der Arzneimittelsicherheit gewissenhaft zu erfüllen und umgehend die Aussetzung bzw. den Widerruf der Marktzulassungen von Comirnaty und Spikevax zu verfügen.



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