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

An:

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- 2) Dr. med. Raimund Bruhin, Swissmedic, Hallerstrasse 7, 3012 Bern
- 3) Nationalratspräsidentin Maja Riniker, Parlamentsgebäude, Generalsekretariat, 3003
Bern

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
Sehr geehrter Herr Dr. Bruhin
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 kann¹.

In australischen Proben² wurde eine übermäßige Menge an Rest-DNA festgestellt,
wie auch zuvor in Frankreich³, Deutschland⁴, Kanada⁵ und den USA⁶ 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.



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- 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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‘Warnung vor potenziellen Schäden durch modifizierte mRNA-Therapeutika’ [NORTHGroup.info](https://northgroup.info) Version 3

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