



North.info Initiative internationale pour garantir la sécurité des vaccins à ARNm modifiés

À :

- 1) Madame la Conseillère fédérale Elisabeth Baume-Schneider, Département fédéral de l'intérieur, Inselgasse 1, 3003 Berne
- 2) Dr Raimund Bruhin, Swissmedic, Hallerstrasse 7, 3012 Berne
- 3) Madame la Présidente du Conseil national Maja Riniker, Bâtiment du Parlement, Secrétariat général, 3003 Berne

Liestal, le 12 juillet 2025

Objet : Appel urgent à l'exercice des obligations de surveillance concernant la sécurité des médicaments en raison de graves préoccupations de sécurité et de qualité de Comirnaty (Biontech/Pfizer) & Spikevax (Moderna)

Chère Conseillère fédérale Baume-Schneider,
Cher Dr Bruhin,
Chère présidente du Conseil national Maja Riniker

L'introduction d'ADN étranger dans les cellules via des nanoparticules lipidiques (LNP) peut endommager l'ADN humain, ce qui peut notamment entraîner une instabilité génomique, un cancer et d'autres maladies graves.¹

Une quantité excessive d'ADN résiduel a été détectée dans des échantillons australiens,² comme cela a été confirmé auparavant en France,³ en Allemagne,⁴ au Canada⁵ et aux USA.⁶

Cette lettre s'inscrit dans le cadre de l'initiative internationale *The North Group* qui vise à garantir la sécurité des vaccins à ARNm modifié. Les politiciens qui la soutiennent et les experts qualifiés (cf. annexe I) demandent la suspension immédiate des autorisations des vaccins à ARNm modifiés COVID-19 de la part des responsables compétents, en raison des effets potentiellement négatifs sur la santé de nos populations.

Nous souhaitons attirer votre attention sur trois points critiques des effets potentiellement négatifs sur la santé humaine liés à l'utilisation de produits à base d'ARNm modifiés contre le COVID-19.

- Les vaccins COVID-19 n'ont jamais été testés pour leur capacité à empêcher la transmission du virus.⁷ Néanmoins, les autorités de réglementation

des dispositifs médicaux ainsi que les gouvernements et les agences gouvernementales ont induit les gens en erreur pour les forcer à accepter ces produits.

- Les vaccins COVID-19 ont entraîné un nombre sans précédent [d'effets indésirables signalés](#)⁸ y compris des [décès](#).⁹ [Des analyses reproductibles des données publiques](#) confirment que des lots individuels sont en corrélation avec des taux d'effets indésirables particulièrement élevés.^{10,11,12,13}
- Les analyses de plusieurs scientifiques indépendants montrent que des quantités excessives variables d'ADN plasmidique résiduel sont présentes dans les flacons des produits Pfizer et Moderna. Cet ADN étranger est un sous-produit du processus de fabrication qui n'aurait jamais dû se retrouver dans les flacons commercialisés.

Les preuves susmentionnées révèlent des risques imprévisibles pour la santé, qui obligent les autorités nationales de contrôle des médicaments à agir immédiatement conformément aux dispositions nationales de la loi sur les produits thérapeutiques (LPTh), art. 66 al. 3, ou de l'art. 59 LPTh.

Par conséquent, nous réclamons ce qui suit au nom du public :

- L'arrêt immédiat de l'utilisation des vaccins ARNm modifiés contre le Covid-19 et le rappel du produit.
- Une enquête indépendante et transparente sur leur autorisation et leur utilisation.
- De nouvelles études scientifiques menées en toute transparence pour déterminer s'il existe ou non un risque de dommages à l'ADN humain.

Pour plus d'informations, veuillez consulter la lettre d'accompagnement « Résumé scientifique » à l'annexe II.

Compte tenu de ces informations, nous vous prions de bien vouloir répondre aux questions suivantes concernant vos obligations en matière de contrôle :

1. *Quand une enquête publique indépendante et transparente sur les problèmes de sécurité et de qualité des vaccins à ARNm modifiés sera-t-elle lancée au sujet du Covid, sur la base des preuves disponibles ?*
2. *Quand le lien de causalité entre les vaccins ARNm et le cancer, l'infertilité ou d'autres maladies aiguës, chroniques et génétiques sera-t-il suffisamment étudié scientifiquement ?*

La santé publique, y compris celle des générations futures, est gravement menacée par la poursuite de l'administration des vaccins à ARNm.



Nous vous prions donc instamment, dans l'intérêt des citoyens de notre pays, de remplir consciencieusement votre mission légale de garantie de la sécurité des médicaments et d'ordonner immédiatement la suspension ou le retrait des autorisations de mise sur le marché du Comirnaty et du Spikevax.

Veuillez agréer, Mesdames, Monsieur, l'assurance de notre considération distinguée

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