Lietuvos Respublikos Ministrui Pirmininkui Gintautui Paluckui 2024 m. lapkričio 25 d.

Pranešimas Šiaurės ir Baltijos šalių bei Jungtinės Karalystės ministrams pirmininkams ir valdymo organams apie ypatingą susirūpinimą dėl modifikuotos mRNR vakcinos nuo COVID-19 saugumo ir kokybės.

Australijos mėginiuose rastas leistiną ribą viršijantis DNR likučių kiekis 1 – tai patvirtino Prancūzijos, Vokietijos, Kanados ir JAV duomenis. $^{2-5}$

Svetima DNR, su lipidinėmis nanodalelėmis (LNP) patekusi į ląsteles, gali pažeisti žmogaus DNR bei sukelti genomo nestabilumą, vėžį ir kitų labai sunkių ligų.

Mes, tarptautinė politikų ir kvalifikuotų profesionalų grupė, esame labai susirūpinę dėl modifikuotų mRNR vakcinų nuo COVID-19 poveikio mūsų šalių gyventojams ir raginame nedelsiant sustabdyti jų vartojimą. Norime atkreipti Jūsų dėmesį į tris esmines problemas, susijusias su modifikuotų mRNR preparatų nuo COVID-19 vartojimu ir turinčiais didžiulę reikšmę piliečių sveikatai.

- Niekada netirta, ar COVID-19 vakcinos blokuoja apkrėtimą virusu. ⁶ Taigi, vaistų agentūros, vyriausybės ir vyriausybinės įstaigos klaidino žmones, siekdamos priversti juos skiepytis šiais preparatais.
- COVID-19 vakcinos sukėlė precedento neturintį kiekį šalutinių poveikių ⁷, tarp kurių yra mirtys ⁸. Atkartojamos viešai skelbiamų duomenų analizės ⁹⁻¹² rodo, kad tai buvo loterija kokį šalutinį poveikį galėjo patirti žmogus priklausė nuo to, kurios serijos preparato jam buvo suleista.
- Daugelio nepriklausomų mokslininkų analizės rodo, kad Pfizer ir Moderna preparatų buteliukuose yra įvairus, leistiną ribą viršijantis plazmidžių DNR likučių kiekis – ši svetima DNR susidaro gamybos proceso metu kaip šalutinis produktas, kuris niekada neturėjo patekti i teikiamus vartoti buteliukus ¹⁻⁵.

Aukščiau išvardyti dalykai kelia kritinį, tačiau kiekybiškai neįvertintą pavojų žmonių sveikatai, kurį pripažinti nuolat vengia už vaistų priežiūrą atsakingos institucijos. Dėl to visuomenės vardu raginame:

- nedelsiant nutraukti modifikuotos RNR vakcinų nuo COVID-19 vartojimą ir atšaukti jas iš rinkos;
- atlikti nepriklausomą ir skaidrų jų užregistravimo ir vartojimo tyrimą;
- dokumentuotų mokslinių įrodymų, kurie patvirtintų, kad nekyla jokia žmogaus DNR pažeidimo rizika.

Išsamesnės informacijos rasite pridedamoje mokslinėje santraukoje. Atsižvelgdami į šią informaciją, prašome aiškių atsakymų į šiuos klausimus:

- 1. Ko reikia, kad būtų pradėtas nepriklausomas ir skaidrus viešas kriminalinis tyrimas, kaip buvo užregistruoti šie preparatai?
- 2. Ar kas nors trukdo inicijuoti galimo mRNR vakcinų ryšio su vėžiu, nevaisingumu, kitomis ūminėmis, lėtinėmis ir genetinėmis ligų tyrimus, bei teikti šiems tyrimams prioritetą?

Atsižvelgdami į šią moksliškai pagrįstą grėsmę dabartinės ir būsimų kartų sveikatai, mes pagarbiai prašome Jūsų veikti savo šalies piliečių, kurie Jus išrinko, labui.

'Raising awareness about the potential harms posed by modified mRNA therapeutics' https://northgroup.info/

Su didžiausiu rūpesčiu ir pagarba, ŠIAURĖS grupės narių vardu (pasirašiusiųjų sąrašas pateikiamas toliau).

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RGN

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