

Lill: **Kapijiet tal-Istat**

Data: 17 ta' Diċembru 2024

Dan l-avviz ta' thassib serju dwar is-sigurtà u l-kwalità tal-vaċċin Covid-19 bil-mRNA modifikat ntbagħat fil-25 ta' Novembru 2024 lil prim ministri u l-entitajiet governattivi tal-pajjiżi Nordiċi u Baltiċi u tar-Renju Unit. Minn dak inhar ingħaqdu pajjiżi oħra, fosthom ċittadini ta' Malta li jaqsmu magħhom dan it-thassib u li bħalhom qegħdin jitolbu tweġibiet.

Livelli eċċessivi ta' fdal ta' DNA instabu f'kampjuni fl-Awstralja¹ li kkonfermaw id-dejta li kienet ġejja minn Franza², Ġermanja³, Kanada⁴ u l-Amerka⁵.

L-introduzzjoni ta' DNA barrani fiċ-ċelluli permezz ta' lipid nanoparticles (LNPs)⁶ jista' jkun ta' hsara għad-DNA u jwassal għal instabilità ġenomika, kanċer u kondizzjonijiet oħra serji hafna.

Bħala grupp internazzjonali ta' političi, ta' professjonisti kwalifikati u organizzazzjonijiet li għandhom l-interess fis-saħħa u s-sigurtà taċ-ċittadini, ninsabu mhassbin dwar l-effetti tal-Vaċċini tal-Covid-19 bil-mRNA modifikat fuq il-popolazzjonijiet tagħna u qegħdin nitolbu biex jiġu sospizi immedjatament.

Nitolbukom tqisu dawn it-tlett punti serji dwar l-użu tal-prodotti tal-Covid-19 bil-mRNA modifikat li għandhom implikazzjonijiet serji għas-saħħa taċ-ċittadini tan-nazzjonijiet tagħna:

- Il-Vaċċini tal-Covid-19 qatt ma ġew ittestjati għall-abilità li jimpedixxu it-trasmissjoni tal-Virus⁷. Għal din ir-raġuni, ir-regolaturi tal-prodotti tal-medicina, kif ukoll gvernijiet u entitajiet tal-Gvern gidbu lin-nies biex iġieghluhom jaċċettaw dawn il-prodotti.
- Bil-Vaċċini tal-Covid-19 ġew rappurtati numru kbir ta' effetti ħżiena⁸ mingħajr preċedenti li jinkludu mwiet⁹. Analizi riproduttivi ta' dejta pubblika^{10 11 12 13} turi li l-għotja tal-vaċċin kienet lotterija skond mill-liema lott il-persuna rċeviet il-vaċċin u l-effetti sekondarji li effettwawhom.
- Analizi minn diversi xjentisti indipendenti pprezentaw evidenza ta' livelli varjati u eċċessivi ta' fdalijiet ta' plasmid DNA fil-kunjeti (vials) tal-prodotti tal-Pfizer u Moderna; dawn huma traċċi sekondarji tal-proċess ta' manifattura u qatt ma kellhom jinsabu f'kunjeti kummerċjali.

Il-punti msemmija hawn fuq iqajmu riskji kritiċi iżda mhux kwantifikabbli għas-saħħa tal-bniedem li ġew sistematikament evitati mill-awtoritajiet regolatorji li kienu inkarigati mis-sorveljanza tagħhom. Għaldaqstant f'isem il-pubbliku nappellaw għal:

- Waqfien immedjat tal-użu ta' Vaċċini tal-Covid-19 b' mRNA modifikat u s-sejha lura tal-prodotti.
- Investigazzjoni indipendenti u trasparenti dwar l-approvazzjoni u l-użu tagħhom.
- Evidenza xjentifika u dokumentata li m'hemm assolutament l-ebda riskju ta' hsara lid-DNA tal-bniedem.

Għal aktar informazzjoni dettaljata jekk jogħġbok ara s-sommarji xjentifiċi meħmuża ma din l-ittra. Wara li eżaminajna din l-informazzjoni, nitolbu tweġibiet ċari għall-mistoqsijiet li ġejjin:

1. X'inhu meħtieġ biex titwaqqaf inkjesta pubblika u forensika, indipendenti u trasparenti dwar il-proċessi regolatorji li wasslu għall-approvazzjoni ta' dawn il-prodotti?
2. X'hemm li jzomm lil ministru biex iniedi u jagħti prijorità lir-riċerka dwar rabtiet potenzjali bejn il-Vaċċini mRNA u l-Kanċer, l-infertilità, jew mard ieħor akut, kroniku u ġenetiku.



B'din it-theddida approvata mix-xjenza għas-saħħa tal-bniedem fil-preżent u għal generazzjonijiet futuri, aħna identifikkaw ir-riskju. Bir-rispett kollu nitolbukom ta'għaxxu fl-interess ta'ċittadini ta'pajjiżkom li għalih ġejtu eletti.

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