Dear Senator/Member,

A DECLARATION AND URGENT DEMANDS
Parliamentary Health Reform Package

The COVID-19 pandemic and the rapidly driven institutional responses to it had many unfortunate and preventable outcomes.

WE HEREBY DECLARE

1. Experimental, inadequately tested medical products were given Provisional Approval. These were not traditional vaccines but new technology with a new mode of action.

   The approval pathway was granted not through their mode of action – *gene therapy* – but through their title – that of *vaccines*. These products lacked data on: safety, full pharmacokinetics, biodistribution, genotoxicity, reproductive toxicity, and carcinogenicity.

2. Despite the Provisional Approval and incomplete data, the public were falsely assured all processes had been followed, and the products were repeatedly pronounced as "Safe and Effective" by Public Health figures and politicians.

3. Effective and cheap off-label drugs for preventing or treating Covid-19 were banned by the TGA. Doctors who used these medicines were subject to legal action and de-registration.

4. No proactive monitoring process to collect safety signals from the injections was put in place in line with the gene therapy "vaccine" roll out.

5. These gene therapy injections did not have the advertised 95% reduction in infection. In fact, there was no significant reduction in infection.

   Consequently, there was no significant reduction in transmission. While data showed the "vaccines" reduced illness severity for a few months, unfortunately recent UK, Danish, and NSW data show that those with more doses – via "boosters" – now tend to have more severe infections and a higher risk of death. Various immunological

mechanisms can explain this finding of reduced efficacy, and even "negative efficacy", as time passes and doses increase.

Australia has never promoted "vaccines" that don't significantly reduce infection or transmission, particularly a "vaccine" with unprecedented side effects including death.

6. State Governments mandated the use of these experimental products as "vaccines" and used financial, legal, and labour penalties to promote them.

7. Serious side effects began to turn up for medical practitioners. Patients have myocarditis/pericarditis, reactivation of cancers, DNA viruses, myocardial infarctions, strokes, and fatigue syndromes. Hundreds more symptoms and injuries are being reported.

8. Doctors speaking for people were dismissed through use of the term "anti-vaxxers".

9. Medical voices were gagged by AHPRA which deregistered Doctors whose public voice did not align to the official "COVID-19 narrative".

10. Informed Consent on COVID-19 injections was lost to the Australian population.

11. The unprecedented number of reports of death, illness, and injury from the "vaccines" in the TGA DAEN system continue to be ignored. Post-mortems were not mandated for these experimental injections. Deaths associated with the "vaccines" continue to be dismissed without post-mortem or pathologic assessment.

12. The Precautionary Principle, where a product had to be proved safe, was replaced with the TGA denying there was enough evidence to prove that it was unsafe, thus inverting the Precautionary Principle.

13. Pfizer's post release Safety data, kept secret by the FDA, was released through FOI. This provided serious, sad, and sobering information on the damage these injections cause, including death. Pfizer and the FDA knew in early 2021 that the mRNA vaccine was unsafe. Subsequently, more recent FOI obtained Pfizer documents and the British Medical Journal publishing of whistleblower testimony, point to data irregularities indicative of scientific fraud by Pfizer.

The attached proposed amendments to the ***Therapeutic Goods Act*** and ***Health Practitioner Regulation National Law*** will reduce these unfortunate and preventable outcomes.

We, the below signatories, being Australian health practitioners and scientists, urgently demand through the enactment of the proposed amendments:

The restoration of Informed Consent to the Australian Community.

The restoration of a Health Practitioner's duty to afford the Australian Community Informed Consent free from interference.

The restoration of the Doctor-Patient relationship free from interference.

The restoration of a Doctor's right to issue a Medical Exemption free from interference.

A safer and transparent process for granting drugs Provisional Approval.

A safer and more reliable process for suspending or cancelling an unsafe drug.

Protections for the Australian Community against mandating provisionally approved drugs.

Protections for the Australian Community against false and misleading messaging about provisionally approved drugs.

Protections for the Australian Community through ensuring Health Practitioners collectively foster and directly determine Health policies, standards, and codes.

SO SAY WE AND DECLARE AND URGENTLY SUBMIT FOR YOUR IMMEDIATE ACTION ON BEHALF OF THE AUSTRALIAN COMMUNITY

Respectfully

We, the undersigned:

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