80% of the Australian Population has been given COVID injections which should never have been approved for release

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The Australian Federal Court to decide if Pfizer and Moderna have broken the law.

The COVID-19 gene-based synthetic mRNA so-called “vaccines” have been a failure. It is widely recognised and admitted that these injections do not prevent infection or transmission of infection and they have been associated with the highest incidence of serious adverse events, ongoing disability and death of any therapeutic agent ever released. The ongoing non-COVID all cause mortality around the world has skyrocketed since these injections were rolled out. In Australia alone, data suggests more than 30,000 Australian non-COVID deaths have been associated with the injections. All data suggests the unexplained non-COVID injections are to blame for the deaths. Governments cannot provide a credible alternative explanation as to why this is happening and they even refuse to investigate. Our drug regulator, the Australian Therapeutic Goods Administration (TGA), waived the normal requirements for proof of safety and we are all suffering because of this reckless approach to the release of these experimental gene-based injections which have never been used on such a massive scale and used in healthy people, children and pregnant women.

But were these COVID injections legally approved for release? That is a question that may soon be answered. AMPS has outlined the Federal Court case which has now been lodged. See below:

Amps Australian Federal Court Gmo Case O...
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You see, the COVID-19 “vaccines” produced by Pfizer and Moderna satisfy the Australian legal definition for being properly deemed Genetically Modified Organisms (GMOs), pursuant to Section 10 of the Gene technology Act 2000. The Act defines an “Organism” as any biological entity that is: “viable; or capable of reproduction; or capable of transferring genetic material”.

The lipid nanoparticle modified RNA or the lipid nanoparticle complex which encapsulates the modified RNA (“modified RNA” because it is not natural mRNA) fulfill the criteria as being “any biological entity” and the modified RNA is “genetic material”. Also, the lipid nanoparticle modified RNA and the lipid nanoparticle complex containing the modified RNA clearly are “capable of transferring genetic material” throughout the body. Furthermore, modified RNA in the injections is also “capable of reproduction” in that it can self-replicate independently within human cells. So, clearly the gene-based injections satisfy the definition of a GMO.

Now, it is a criminal “Offence” for a person to knowingly “deal” with a GMO (like the TGA) without a licence because if the risk to life. It is alleged that both Pfizer and Moderna avoided their legal obligation to properly seek a GMO Licence from the Office of Gene Technology Regulator (OGTR) which is the Australian legal entity responsible for the issuance of such licences.

The full legal brief is below:

Ag Information 27 September 2023
566KB · PDF file

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This is serious business because it is now known that the COVID-19 injections contain synthetic modified genetic material which can enter the nucleus of cells and potentially integrate irreversibly into a person's DNA and this altered DNA could be passed on to future generations. In addition, it has now been shown that the COVID-19 injections contain massive amounts of dangerous plasmid DNA and endotoxins derived from the manufacturing process (Process 2, CLICK HERE to see my Substack of 5 Oct.) which was not present in clinical trial batches which utilised a cleaner manufacturing process based on real-time PCR manufacture.

It could reasonably be assumed that Pfizer and Moderna were aware of these dangers prior to application to market these injections and did not adequately advise the OGTR or the TGA. The Australian public was also not made aware of these dangers and senior public health "experts" declared the injections "safe" without any qualification. Health professionals who administered the injections also failed to properly provide informed consent under both international and national guidelines (CLICK HERE to see my Substack of 20 Sept).

To his enormous credit, Senator Gerard Rennick has written to the Attorney General to bring to his attention the need to act quickly. See his letter below.

Sen Rennick Letter Re Gmo Investigation
431KB · PDF file

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Where will this lead? I will leave that to the legal eagles amongst us. But what can be said is that our drug regulatory system has failed to inform and protect us from harm. There continues to be a veil of secrecy surrounding the nature of the COVID-19 injections including their real nature and potential harm. This is why a Royal Commission is so necessary. This is why Big Pharma indemnity from prosecution for harm caused by "vaccines" must be abolished. We must understand how the system failed us so we avoid any repetition of this tragic saga.