Submission and evidence to the COVID-19 Response Inquiry

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Background

I scrutinise Regulators.

From 2008 to ~2011 I paid FSANZ hundreds of dollars of my private money to access the Regulatory Submissions of the GM crop inventors – Monsanto, Bayer, Syngenta, Dow, DuPont and BASF.

It seemed to me that only two people in the whole of Australia scrutinised this data – I was one of them. I suspect I was frequently the only person in Australia doing this. I wrote hundreds of letters to MPs and Senators to tell them about the messes of genetic inventions, about corporate fraud, and about our malfeasant Health regulators.

Eventually I teamed with a virologist and an environmental scientist, put it all down on paper and we published it in the Journal of Biotechnology and Genetic Engineering Reviews. It is a paper cited as often by genetic engineers as it is by precautionary scientists.
https://www.tandfonline.com/doi/full/10.1080/02648725.2017.1357295

GM crops were frequently designed to make "just one protein", but they were messes of many proteins, broken code and innumerable transcripts. Every genetic invention is a mess, and this has been documented. Regulation was co-ordinated and global, and FSANZ never rejected a GM crop application.

It is beyond belief that another Australian Health Regulator, the TGA, without any Guidelines to assess the safety of the Adenoviral and modMRNA inventions, approved the direct injection of the genetic messes, to the most extraordinary harms.

This COVID-19 Response Inquiry

I understand this is a sham Inquiry, incapable of scrutinising all the Federal and State players and their decisions in the COVID-19 Response. We need a Royal Commission.

I am using my private money again to purchase data to show the lives lost to these genetic messes, that should never have been approved, and certainly not mandated.