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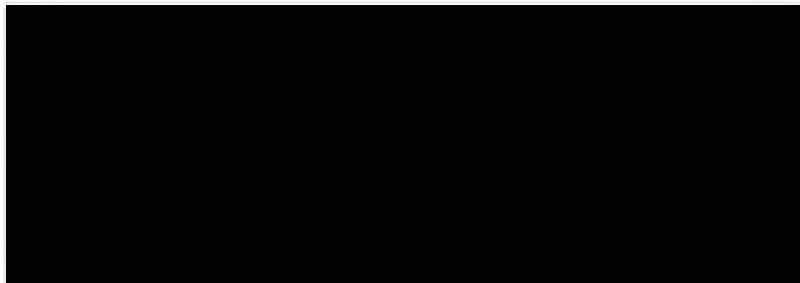
## So this is what Australia signed up for

Inbox

00:09 (8 hours ago)

No consulting. No oversight. No release to public of any information. Every single politician in Australia who voted yes to this legislation needs to be fired... As in fired into the sun!

[WHO New Health Regulations](#)



**WHO New Health Regulations**

Attachments area

[Preview YouTube video WHO New Health Regulations](#)



08:40 (0 minutes ago)

Oh well, I guess someone had to take control and responsibility of de-population, whilst supporting the ongoing expansion of ill health, to support the medical health and foods industries growth of multi trillion-dollar profits. So, WHO better than the CORRUPT WHO, primarily controlled by billionaires and money-making power brokers, BECAUSE, they have TOTAL DIPLOMATIC IMMUNITY, CANNOT BE SUED, like our countries governments and organisations can.

Maybe, I should let it go, because after all, we have overpopulated the world, depleting her resources and are killing the planet, primarily killing mother nature which is the only thing the world survives on. So maybe it's not such a bad thing? But do we all have to go through the extreme pain of all the ill health and pay so dearly from the hip pocket for that luxury?

## **Not so Cheers**

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E-mails monitored by Kasperski and Malwarebytes

**The Long Sad U.S. Corrupt History Of Vaccines and Deaths from the Spanish Flu to Corvid for Money, Power & Control AND since 1986 U.S. Vaccine Companies cannot be sued, so why are we following them?**

**In a recent podcast, Joe ROGAN interviews Robert Francis Kennedy Jr.—the son of former U.S. attorney general Robert F. Kennedy and nephew of former President John F. Kennedy—is an attorney, environmental and anti-vaccine activist, and syndicated talk radio host. He is a co-founder and former president of Waterkeeper Alliance, an organisation monitoring rivers and waterways and suing polluters, including many on the Hudson River in the U.S. which became so polluted, kilometres caught fire and had no fish in them at all, fortunately it has since recovered as a result. He and his organisation have accessed all the studies and all available data and results on everything covered naturally, as if you are going to litigate successfully as he has done against government agencies, big pharma's and large corporations, you have to be right. His litigation success record has brought about, over the last 18 years, there is no one that he has wanted to challenge in a debate that has been prepared to do so. He is now going to run for the next presidency, let's hope he wins and still lives, however no doubt the money-making power brokers won't allow that to happen.**

**I have done my best to summarise the verifiable facts discussed in the podcast below this to make it easier. If you wish to listen to it yourself, you need to have at least the free Spotify version to view the podcast and it is 3 hours long but very worthwhile.**

#### **SUMMARY:**

**1932 Mercury was added to vaccines primarily by Big Pharma's such as ██████████, as a preservative and adjuvant which is a toxic substance to enhanced the vaccines immune response. Mercury is 1,000 times more toxic than lead. Hence, the vaccines have been causing autism and many other life destroying illness and health issues especially for children, crossing the brain barrier**

and causing inflammation. In recent years Mercury has been removed from most of their vaccines but replaced with aluminum, which has the same effect.

Mass independent statistics in the U.S. showed (1:37) One in 37, children had some form of vaccine injury. In the U.S. the vaccinations for youth grew to 72 vaccines.

In Australia it is already up to more than 50. Fortunately, in Australia at least prior to 2021, Australian vaccines were all made the old-fashioned way with an inert form of the virus and/or a non-replicating spike protein, unlike the new mRNA vaccines for Covid, (with exception of ██████ still made the old-fashioned way) with the mRNA spike proteins crossing the blood brain barrier, which enters every organ in the body and lacks the targeting ability to only kill virus cells, hence it can attack healthy cells as well. The worry now is that Australian Government has partnered with ██████ to produce mRNA vaccines for most health conditions and childhood vaccinations, beware. Back to the U.S. info:

In the U.S. the 1986 Vaccine Act, Vaccine Companies cannot be sued. Hence since then Vaccine Companies, did not have to bother about rigidly and thoroughly safety testing their products.

KENNEDY showed a power point video on what they knew about vaccines damage to ██████ (Recent U.S. Chief Health Officer through the Covid pandemic etc) and others. He then challenged ██████ to show any study showing proof of safety testing on any vaccine, prior to licensing. ██████, looked through a stack of books from NIH (National Institute Of Health), but could not find anything. So, KENNEDY and a partner sued NIH. About a year later NIH finally came back with a letter stating that there is none. KENNEDY was involved in many cases of suing large companies and instrumentalities including the huge ██████ organisation re poisons including ██████ and ██████. ██████ ready corn and other legumes, wheats, barleys etc., then being sprayed directly on the crop, cause celiac disease to rapidly increase along with many other illnesses including John Hopkins Lymphoma.

Auto immune diseases have blossomed out of control because of the aluminum (now replacing the mercury) in vaccines, both highly toxic, causing allergies leading to infections and inflammation. Along with many other toxic substances in air, water and products

eaten and used etc., the U.S. has the highest chronic disease levels of any country in the world.

Re the new MRNA Corvid Vaccines. A Cleveland Clinic study showed that it offered some protection for the first two months, but then wanes rapidly into the negative efficacy especially after 7 months, and without bolstering your immune system gives the vaccinated up to 3.5 times the risk of suffering illness than people who are unvaccinated. Even [REDACTED], the then head U.S. Health Officer, in an earlier statement did say that it could make you more susceptible to Corvid and inflammations. Many studies clearly show vitamin deficiency causes metabolic and immune deficiency, ie. Estimated between 70 and 90% of people are vitamin D deficient.

It also covers some of the same ground re the lies about the Chinese Wuhan Institute leak of Corvid supported by U.S. complicit in funding of gain of function with Bat viruses namely Corvid and [REDACTED] lied about it. Together with Interviewer Joe ROGAN, they cover taking anti inflammatories and vitamins to bolster the immune system to deal with corvid, flu etc., instead. It is very hard for a virus or bacterial infections to kill a truly healthy person, but some bacterial infections have a far higher chance than many viruses. The more vaccines you take, the more likely you are to be susceptible to ongoing inflammations.

Other examples were, all the kids in Africa that die are generally malnourished else the disease may not kill them as evidence amongst those not malnourished. In 1963 in the Mississippi delta many [REDACTED] children were dying of measles, but again because they were severely malnourished. The 1918, 1919, Spanish Flu, was not a killer virus and this was even again acknowledged by [REDACTED]. At the time they thought the deaths were as a result of the flu, but after analysis of many samples from thousands of victims, it was established it was a Spanish Flu vaccine (developed in a military research centre in the U.S.) induced flu causing people to die of bacterial pneumonia which could have been prevented with antibiotics. The 1967, 1968 flu pandemic confirmed this.

The NIH, National Institute of Health in America showed a study that Ivermectin was successful in treating adults with moderate Corvid 19, yet the health authorities declared it as ineffective?

Again, Hydroxychloroquine which is used in the treatment of autoimmune conditions such as rheumatoid arthritis and systemic lupus erythematosus, was shown to be effective against Corvid 19. It slows down the progression of the disease and provides relief from pain, swelling, and redness, yet again they declared it as ineffective? Why, because there was a law that said you were not allowed to promote a new drug treatment for a serious illness, if there was another one already available.

Various studies that were organised by the corrupt WHO (World Health Organisation) and paid for by the [REDACTED] [REDACTED] (also one of the largest funders of the WHO and the GVA, The Global Vaccination Alliance), have deliberately conducted studies designed to fail. This included Ivermectin and Hydroxychloroquine (provided by the [REDACTED] [REDACTED] and still successfully given to 100's of people every year in Africa to treat malaria). No body died from taking the recommended dosage, but by giving very large doses of vaccines or medications, instead of the recommended dosage, many people (of which many were in third world countries such as Africa) became very ill and/or died. Hence, they could say that it does not work

On the other hand, another drug, [REDACTED] is a broad-spectrum antiviral medicine. It is used for the treatment of the novel coronavirus disease (COVID-19). It works by inhibiting an RNA dependent enzyme called RNA polymerase that helps the SARS-CoV-2 virus make more copies of itself. This way it decreases the viral load in the body. However, it was used in a study against [REDACTED] where the safety panel pulled it out of use because it was killing more people than [REDACTED] was. So, why would you then allow it's use against a virus that has a serious ill health rate of one percent?

In the Aids crisis, they used [REDACTED], (in full azidothymidine), also called [REDACTED], a drug used to delay development of AIDS (acquired immunodeficiency syndrome) in patients infected with HIV (human immunodeficiency virus). [REDACTED] belongs to a group of drugs known as nucleoside reverse transcriptase inhibitors (NRTIs). It was used as a chemotherapy treatment. It was killing people in a two-week dose virtually faster than cancer was killing



people. They slowed the death rates by giving blood transfusions helping to support their claim that it was reasonably effective in prolonging life.

In the case of █████ when Corvid came along, they had got another vaccine approved elsewhere, which was not available in the U.S. But, having an approved vaccine allowed them to distribute their Corvid 19 vaccine to and collaborate with other government, university, and medical research organisations. They initially promised to do a five-year study, but cut that back to between 4 and 6 months in getting the emergency use approval. They vaccinated 22,000 people and initially used 22,000 people unvaccinated as a control/comparison group, but shortly after it started unblinded it by doing away with the control group, most of which in time got vaccinated, which was a total deception. In the less than 6 months, in the placebo group 1 person died and in the vaccinated group 2 people died, hence they immediately claimed that it was 100 percent effective, because 2 is 100% of 1? Yet when they looked at all-cause mortality in the survey group, the unvaccinated had 17 people die and the vaccine group had 21. From that, 4-5 cardiac arrests occurred in the vaccine group and only 1 in the unvaccinated group?

There was an all-cause mortality deaths records from insurance companies which showed thousands of people including athletes were dying after receiving vaccines. Several records show over 15 million Americans sought medical help after vaccines which has been clearly shown to have been extremely underestimated.

Almost 50% of the FDA's (Food and Drug Administration), budget comes from pharmaceutical companies. The CDC (Centre for Disease Control and Prevention), has a 12 billion budget and almost 5 billion of that goes to buying vaccines in sweetheart deals from the Big 4 Pharma's, and then promoting them to the public. If you are part of the NIH (National Institute of Health) and you work on one of their vaccines or drugs, you can collect royalties on that product, up to U.S. \$150,000 per year, for life, as long as that product continues to sell, i.e. the █████ and █████ vaccines. There are 4-6 close colleges of █████ who are collecting that \$150,000 a year. There are no longer any regulatory authorities left in the U.S. that are not corrupt and monetary driven. Drugs are the

number 3 killer in America and yet America spends more money on health care than any other country in the world, yet their overall health record is less than some third world countries.

Other areas covered:

There are numerous studies showing Wi Fi radiation opens-up the blood brain barrier? KENNEDY has represented many hundreds of people for cancers behind the ear that they favour for their cell phones. He has successfully sued the government bodies responsible for mobile phone Wi Fi.

██████████ in the US water system when given to male frogs caused many to become chemically castrated and others to become female and get pregnant. ██████████ is a pesticide that has been commonly detected in surface and groundwater samples in Australia, usually at less than 1 pg/L<sup>1</sup>. It is one of the most commonly detected pesticides in the Murray-Darling Basin<sup>1</sup>. ██████████ has rarely been found in Australian drinking water supplies, and when detected, it has been below the National Health & Medical Research Council Australian Drinking Water Guideline level of 0.1 pg/L<sup>2</sup>. Australian regulators decided in 2008 to maintain their standard at 40 ppb<sup>3</sup>.

Definitely, Not So Cheers

J.P.

## **Australian Government Initiative Guidelines For Water Quality**

██████████ in freshwater and marine water

## Toxicant default guideline values for protecting aquatic ecosystems

October 2000

Extracted from Section 8.3.7 'Detailed descriptions of chemicals' of the ANZECC & ARMCANZ (2000) guidelines.

The default guideline values (previously known as 'trigger values') and associated information in this technical brief should be used in accordance with the detailed guidance provided in the [Australian and New Zealand Guidelines for Fresh and Marine Water Quality](#).

### Description of chemical

Atrazine (CAS 1912-24-9) is a triazine herbicide developed by Ciba-Geigy, with a selective systemic mode of action, being absorbed mainly through the roots (Tomlin 1994). It inhibits photosynthesis and other enzyme processes (Tomlin 1994). A comprehensive draft review of atrazine by NRA (1997b) indicated that a complete set of data was available.

Its IUPAC name is 6-chloro-N2-ethyl-N4-isopropyl-1,3,5-triazine-2,4-diamine, molecular formula is  $C_8H_{14}ClN_5$  and molecular weight is 215.7. Atrazine is soluble in water at 33 mg/L at 20°C (Tomlin 1994). It has a pKa of 1.7. The current analytical practical quantitation limit (PQL) for atrazine in water is 0.5 µg/L (NSW EPA 2000).

### Uses and environmental fate

Atrazine is used widely against grasses and broad-leaved weeds in a variety of vines, orchards, plantations and crops, particularly maize and sorghum (NRA 1997a). In Australia, atrazine has over 1600 registered uses as 30 registered products. It is often used in conjunction with other herbicides.

Atrazine is relatively stable in water and persists in groundwater with a DT50 greater than 100 days (Tomlin 1994). Atrazine breaks down only slowly in sunlight with a half-life of around 1 year. Its half-life in fresh water systems is around 2 months, or more at lower temperatures, but is more rapid (<1 month) in estuarine systems (NRA 1997b). It does not bioaccumulate significantly. Atrazine is highly mobile and it has been commonly detected in surface and groundwater samples in Australia (Cooper 1996) usually at less than 1 µg/L. In fact, it is one of the most commonly detected pesticides in the Murray–Darling Basin.

### Aquatic toxicology

Freshwater algae: two species, 48 to 96-hour EC50 (growth) of 21 to 377 µg/L. The lowest figure was for *Scenedesmus subspicatus* but another figure of 110 µg/L was also reported for that species.

Freshwater fish: 14 species; 96-hour LC50 between 500 (*Rasbora heteromorpha*, 48-hour LC50) and 71,000 µg/L (*Poecilia reticulata*). A 35-day mortality NOEC of 300 µg/L was obtained for *Brachydanio rerio*; a 21-day mortality NOEC of 60 µg/L for trout *Oncorhynchus mykiss*; a 274-day growth NOEC of 250 µg/L for *Pimephales promelas* but no effect on

reproduction at 2000 µg/L (highest concentration). An acute-to-chronic ratio (ACR) of 300 was reported.

Freshwater crustaceans: five species, 48 to 96-hour EC50 of 5700 to 54,000 µg/L.

Microcosms and mesocosms: NRA (1997b) reviewed a number of aquatic microcosm and mesocosm studies with different composition and test end-points (Moorhead & Kosinski 1986, Pratt et al. 1988, Stay et al. 1989, Neugebauer et al. 1990, Detenbeck et al. 1996, Gruessner & Watkin 1996). Most reported just LOEC values between 50 and 300 µg/L but eight NOEC values were reported; 3.2, 5, 17.5, 20, 20, 20 and 80 µg/L. The lowest figure was a stimulation effect and was not considered. None of these fully satisfied the requirements but they do give added confidence in the trigger value.

Marine fish: two species, 96-hour LC50 between 2000 and 16,200 µg/L.

Marine crustacean: four species, 96-hour LC50 of 94 µg/L (*Acartia tonsa*) to 13,200 µg/L; 8-day NOEC (mortality) between 4200 and 17,500 µg/L for the copepod *Eurytemora affinis*, depending on salinity. Two species of adult crabs were insensitive to atrazine up to its solubility.

Marine diatom: one species, 48 to 72-hour EC50 of 50 to 265 µg/L (PSR, photosynthesis).

## Australian and New Zealand data

The 96-h LC50 to the introduced fish *Gambusia holbrooki* was 18,900 µg/L and to *Hypseleotris gallii* (firedale gudgeon) was 258,000 µg/L, well above the solubility level. The 48-hour LC50 to the water flea *Ceriodaphnia dubia* was 18,300 µg/L. The 72-hour EC50 to the alga *Selenastrum capricornutum* was 359 µg/L. All of those figures were within the range of overseas species, except for the gudgeon, which was particularly insensitive.

## Factors that modify toxicity of atrazine

The NRA (1997b) review did not identify any factors modifying atrazine toxicity. Synergy was not demonstrated with the pyrethroid bifenthrin. Hall et al. (1995) demonstrated varying chronic toxicity of atrazine to an estuarine copepod, *Eurytemora affinis*, under different salinity regimes. It was most sensitive (14.6 mg/L) at 5 ppt salinity and of lowest sensitivity (20.9 mg/L) at 15 ppt. Mixtures of atrazine and metribuzin have additive toxicity to algae (Altenburger et al. 1990).

## Guidelines

A freshwater **moderate reliability** trigger value of 13 µg/L was derived for atrazine using the statistical distribution method with 95% protection and an ACR of 20.2.

Although there are marine data on fish, crustaceans and algae (OECD MPD) it was considered preferable to adopt the freshwater figure as a marine **low reliability** trigger value (13 µg/L). This should be used only as an indicative interim working level.

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