

# If Truth Be Told About Aluminum (Aluminium), A Neurotoxin In Vaccines

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Aluminium is neurotoxic. Its free ion,  $Al^{3+}_{(aq)}$ , is highly biologically reactive and uniquely equipped to do damage to essential cellular (neuronal) biochemistry. This unequivocal fact must be the starting point in examining the risk posed by aluminium as a neurotoxin in humans. *Apr 30, 2014*

Source: [What is the risk of aluminium as a neurotoxin?](#)

Nothing explains the aluminum free ion  $Al^{3+}$  more intelligently than the following:

**The likely principal antagonist in all such events is  $Al^{3+}_{(aq)}$  and its mechanism of action will involve numbers of different agents or intermediates.** For example, we know that *aluminium is a potent pro-oxidant, its interaction with the superoxide radical anion establishing, fuelling and sustaining redox cycles.* The potency of these effects are all the more significant in that the enhanced formation of reactive oxygen species may be accelerated at sites which are distinct and divorced from locations housing the cell's anti-oxidant machinery. For example, aluminium sinks such as the extracellular senile plaques of  $A\beta_{42}$  and the intracellular chromatin of neuronal nuclei are both likely targets of aluminium-driven oxidative damage. Aluminium is

an excitotoxin and a number of mechanisms have been described, whereby aluminium induces elevated and sustained levels of intracellular Ca<sup>2+</sup> with significant implications not only for cellular energy metabolism, but also uncontrolled phosphorylation of biomolecules. The presence of biologically reactive aluminium imposes an immediate energy requirement upon a neuron, whether simply because of the need to produce more Ca<sup>2+</sup>-buffering proteins or because of the requirement to clean-up the consequences of hyperphosphorylation, for example, through autophagosomal activities. Aluminium is a mutagen and the phosphate-rich environment of the nucleus predisposes it to the accumulation of aluminium and subsequent alterations in the expression of genetic materials. The latter may be subtle but sufficient to bring about significant alterations in neuronal physiology over extended time periods. Aluminium is, of course, a powerful immunogen, being the preferred adjuvant in vaccination and immunotherapy. This activity as an adjuvant, and concomitantly as an antigen, at injection sites in skin or muscle must also be considered for focal accumulations of aluminium within the CNS and such reactivity may underlie aluminium's suggested roles in autoimmunity. [3-5]

[CJF emphasis]

Source: [Taylor & Francis Online](#)

**And yet, consensus-vaccinology-science doesn't get it! Why?**

There has to be another designated reason, *plus rationale*, for poisoning the CNS (central nervous system) of a human starting in infancy with mandated hyper-containing-aluminum-salts (as adjuvants) in any of 3 formulations, i.e., *Aluminum hydroxide, aluminum phosphate, and potassium aluminum sulfate (alum)*:

Aluminum adjuvants are used in vaccines such as [hepatitis A](#), [hepatitis B](#), [diphtheria-tetanus-containing vaccines](#),

Haemophilus influenzae type b, and pneumococcal vaccines, but they are not used in the live, viral vaccines, such as measles, mumps, rubella, varicella and rotavirus.

Source: [Children's Hospital of Philadelphia](#)

## **Proving the above**

To check the quantities of aluminum in any vaccine on the CDC/FDA schedule for infants, children and adults, please search each vaccine using the following resource: [Vaccine Package Inserts](#). *Scroll to Section 11 Description, which lists the type of aluminum and the number of micrograms, plus other ingredients.*

## **Medical Care**

*The most important part of emergency medical treatment is the recognition of possible aluminum toxicity based on risks (eg, renal insufficiency, aluminum exposure) and symptoms (eg, altered mental status, anemia, osteoporosis).*

*Treatment of aluminum toxicity includes elimination of aluminum from the diet, TPN, dialysate, medications, antiperspirants, and an attempt at the elimination and chelation of the element from the body's stores.*

**Source: Aluminum Toxicity Treatment & Management** (July 10, 2017)

<https://emedicine.medscape.com/article/165315-treatment>

## **Research**

*Research at Keele University in Staffordshire, UK, has shown for the first time that an individual who was exposed to aluminum at work and died of Alzheimer's disease had high levels of aluminum in the brain.*

[...]

*Professor Chris Exley, of The Birchall Centre, at Keele University, said: "The results showed unequivocally that **the frontal lobe contained an average aluminum content which was at least four times higher than might be expected for an age-matched control brain.***

Source: Elevated brain aluminium, early onset Alzheimer's disease in an individual occupationally exposed to aluminium  
<https://www.sciencedaily.com/releases/2014/02/140212093300.htm>

Dr. Christopher Exley, PhD, premiere aluminum researcher for close to 30 years, recently has had his funding in aluminum research halted!

<https://www.thetimes.co.uk/article/funding-halted-for-professor-chris-exley-linking-vaccines-to-autism-8xvwp0g8p>

## **Medications**

The medication, deferoxamine mesylate, may be given to help eliminate aluminum from your body. This substance works through a procedure known as chelation, which helps the body remove poisonous materials.

**Source: Aluminum Toxicity**

<http://www.winchesterhospital.org/health-library/article?id=164929>

**Notation should be made, and even emphasized,** that the allopathic medical paradigm disapproved of chelation as a "quack-medicine" modality when used by holistic and integrative physicians, and actually prosecuted some physicians for using the modality! The above indicates just how screwed up allopathic medical science actually is, since allopathy now incorporates what it previously condemned. Go figure!

Thus, the mechanism of Al toxicity appears to be different in the two cell lines. It is possible that the principal

*neurotoxic target of the metal is glial and when these cells are in a compromised state, this may secondarily impact the neuronal population and thus eventually lead to [neurodegeneration](#).*

Source: **Differential Toxicity of Aluminum Salts in Human Cell Lines of Neural Origin: Implications for Neurodegeneration**  
<https://www.sciencedirect.com/science/article/pii/S0161813X00000073>

**The purported 'science' behind vaccines must be investigated.**

Why? Most annual flu vaccines are not subject to the testing as other vaccines. The reason: Because they are formulated from a '*sophisticated guessing game formula*' based upon what strains appeared prevalent in the previous Australian winter flu season!

However, ***veterinary medicine*** has taken a different look at animal vaccines, which is discussed in **Vital Vaccination Antibody Titers Versus Vaccination**  
<https://todaysveterinarypractice.com/vital-vaccination-series-antibody-titers-versus-vaccination/>

All the above indicates the need for independent – *no Big Pharma* – involvement in assaying the neurotoxic effects of aluminum in vaccines given to infants as early as 24 hours old, especially in view of the Autism epidemic in the USA, i.e., one in 34 in New Jersey!, and spreading globally.

**Congress, why aren't you performing your Oversight duties?**

**Resources:**

**Aluminum Toxicity**

<https://myersdetox.com/aluminum-toxicity/>

*8 Ways to Protect Yourself from Aluminum Poisoning*

<https://universityhealthnews.com/daily/nutrition/8-ways-to-protect-yourself-from-aluminum-poisoning/>

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