The scandal of aluminum

Statement by Pr Jean-Bernard Fourtillan

Aluminum should never have been present in gastric anti acidic oral drugs or in vaccines. Because it is at the origin of the biggest scandal ever seen in the history of medicine: the imposed "massacre" of all the children born in France since January 1st, 2018.

It's simply the truth!

Professor Jean-Bernard FOURTILLAN (Poitiers, October 23, 2019)

But I come immediately reassure the parents of these children

Thanks to the discovery of the sleep-wake system, and, to the extraordinary and marvelous efficiency of the patches that result, we understood the origin of Parkinson's neurodegenerative diseases, as well as multiple sclerosis, and found their treatment:

1- Administration of organo-silicon compound (the silicium G7 of Dr. Loïc Le Ribault) able to eliminate from the brain the cause of these diseases: Aluminum, but also Mercury and Lead.

2- Suppression of the symptoms of these diseases by administration of mixed transdermal patches of Valentonine and 6-Methoxy-Harmalan

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Who are the responsible, makers of the scandal?

It is not the 4 Pharmaceutical Laboratories that manufacture vaccines with aluminum adjuvants.

The entire responsibility lies with the National Agency for the Safety of Medicines and Health Products, the ANSM, its experts, and the Minister of Health, who have given the marketing authorization.

How did we get there?

The formal prohibition of the presence of aluminum in all drugs for human use, antacid gastric dressings (Maalox®, Phosphalugel®, Smecta®) and vaccines containing aluminum adjuvant, stems from a toxicokinetic study, conducted at the request of the FDA, to evaluate the toxicity of dietary aluminum, that is to say of aluminum administered orally, in animals.

This is the only study in the world conducted to date

This FDA study, taken up by WHO in 1996, states that:

The Minimal Risk Level (or Minimal Toxic Dose) for aluminum is 1 mg per kg of body weight per day.

Thus, this Commandment, above all laws, states that it is strictly forbidden to administer in humans oral doses of aluminum equal, and even higher, to the Minimal Toxic Dose of 1 mg of aluminum per kg of body weight per day.

This study was conducted:

- by Pr Christopher EXLEY, undisputed specialist in aluminum and its toxicity in biological media, such as the brain;
- and by **Dr P. JOUHANNEAU** who performed the only pharmacokinetic study by measuring the amount of aluminum, "bioavailable", which enters the body completely, after oral dosing.
- P. JOUHANNEAU concluded that the oral bioavailability of aluminum is between 0.1 and 1%.

Thus, these two leading aluminum specialists, knowing, according to Christopher EXLEY, that the minimum toxic dose in the body is equal to 0.01 mg / kg, have placed themselves in the worst conditions of aluminum toxicity when given orally, that is to say at its maximum absorption of 1%. This allowed FDA experts to calculate the Minimum Toxic Oral Dose by multiplying the Minimum Toxic Dose in the body, as 0.01 mg / kg, per 100.

Insofar: Minimal Toxic Oral Dose = 0.01 mg / kg x 100 = 1 mg / kg of body weight / day. And they added daily, because this dose corresponds to the aluminum present in the daily diet.

Thus, the Minimum Toxic Dose of aluminum in the diet, that is to say orally, 1 mg of aluminum / kg / day, was calculated from the Minimum Toxic Dose of the aluminum in the body, ie 0.01 mg of aluminum per kg of body weight, when the aluminum is injected in the form of vaccines containing the aluminum adjuvant.

As a result, when injecting 0.85 mg of aluminum into a vaccine to a 5 kg infant, it receives 0.85 mg / 0.05 mg (minimal toxic dose per 5 kg of weight) = 17 times the minimal toxic dose.

Following the "mortifère law", applied from January 1, 2018, in France, it is imposed to all children born since January 1, 2018, to receive 11 vaccines, 8 of which contain aluminum.

Thus, from the age of 2 months, and during their first year of existence, these children receive, at a minimum, 2,775 mg to 3,635 mg of aluminum, which represents for an average weight of 7 kg, during their 1st year, 40 times to 52 times the minimal Toxic Dose.

Which is monstrous!

There is no word to describe the "massacre" of these children who have been sentenced to severe neurological diseases before the age of 5 to 10 years.

It must be emphasized that, to my knowledge, there is no precedent. Indeed, there is no authorized medicine in humans with only a fraction, even a tiny fraction of a minimum toxic dose, promulgated by a health authority.