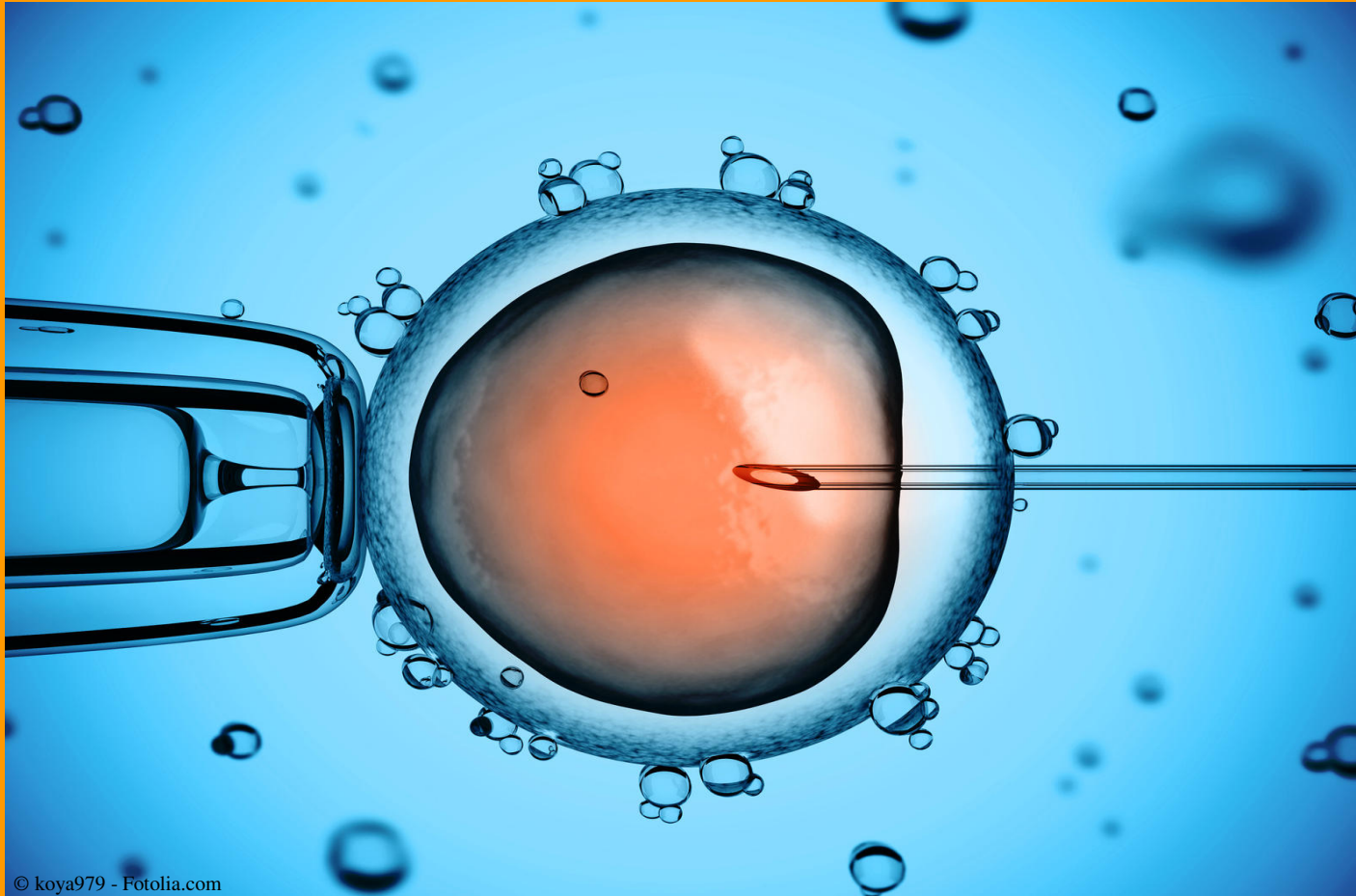


Future of Vaccinations



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Genetics Vaccinations

The Trojan horse of the vaccine manufacturers

For quite some years the use of aluminium as adjuvants in vaccines has been a controversial issue. Since decades top connections between industry and authorities (1) made sure that these aluminum compounds were not removed from vaccines. But behind the scenes research was under way to find other ways of producing vaccines without the help of aluminum. Now an alternative was found: Genetic vaccines with genetically modified viruses, the newest creation of vaccine manufacturing.

Since quite some time mercury was looked upon critically (2). Meanwhile it is prohibited nearly all over and should be outlawed. But vaccines exported to Africa are exempt from this rule. Also Pandemix, a vaccine against swineflu, still contains mercury. Still, most vaccines made with dead or attenuated viruses contain aluminium compounds because without these adjuvants there would be no immune response. No antibodies without aluminium which are the basis for evidence of effect of any vaccine. That is the reason why you cannot completely remove it from vaccines because there just would be no immune response.

During the times of the swine flu and the „blue tongue disease“ of animals, but also during the gulf wars a comprehensive replacement was tested: Squalene for mankind and Saponine for animals. The price for these mass experiments were massive adverse effects which they were able to keep under the lid: narkolepsie, Guillan Barre Syndrome (GBS) and gulf war syndrome for mankind, miscarriages and blood sweating for animals. Thus squalene and saponine could not be used as substitutes for aluminium in vaccines and research had to be continued.

Genetic vaccination (vector vaccines)

First tests on Ebola vaccines were conducted in Lausanne, Switzerland, by the end of October 2014. About 180 volunteers agreed to be injected with this new vaccine. These vaccines have hardly anything in common with the vaccines used up to this point because they are genetic vaccines! „Genetic vaccines“, also called „vektor vaccines“ are produced using a totally new manufacturing process. An article (3) in the German Physician's Magazine about this exact vaccine states: *Some 180 healthy subjects shall be inoculated with a vaccine containing a genetically altered adenovirus which is used as a ferry tool for an Ebola Glykoprotein.* The manufacturers of these „gene ferries“ are GlaxoSmithKline (GSK) and the National Health Institute of the United States (NIH).

An additional article (4) was published in the Pharmaceutical Magazine: *The vaccine contains an attenuated adenovirus of serotype 3 from the chimpanzee, abbreviation ChAd3, (vaccine 2009; 27 (9): 1293). It was gene manipulated by scientists in such a way that it presses out an Ebola-glykoprotein.*

NIH describes their co-produced vaccine as follows: *The investigational vaccine, which was designed by VRC scientists, contains no infectious Ebola virus material. It is a chimpanzee adenovirus vector vaccine into which two Ebola genes have been inserted. This is a non-replicating viral vector, which means the vaccine enters a cell, delivers the gene inserts and does not replicate further. The gene inserts express a protein to which the body makes an immune response. (5)*

With other words: Adenoviruses from monkeys were altered by genetic engineering, parts of DNA from Ebola viruses were integrated into their DNA. These adenoviruses will enter the cells of the vaccinated subjects. Once implanted in the cells the Ebola DNA makes sure that a protein will be produced out of the shell of the alleged Ebola virus; then this whole process produces a heavy immune reaction. How these Ebola proteins are produced exactly – the present publications from GSK and NIH are very vague about it. A search in scientific specialty magazines also did not come up with any consistent results. Sometimes the adenoviruses are producing the Ebola proteins by themselves, sometimes they come out of the viruses, then again these viruses animate the cell to produce the Ebola proteins by themselves. Our impression: They don't know what they are doing but this with all vengeance! One thing you can be sure of: Genetically engineered DNA

smuggled into a cell holds the risk of altering our genetic inheritance. Longterm consequences are not conceivable (6,7,8).

DNA and proteins foreign to the body smuggled into cells – this is an alarm signal for the body. Fact is: these adenoviruses are a Trojan horse which imports foreign gene manipulated DNA into our cells.

Since vaccine studies are not conducted for reasons of ascertaining adverse effects, especially long term ascertainment (9), only to prove general antibody production, a lot will remain to be seen. As nearly all vaccines, these genetically engineered vaccines too, will be uncritically pushed and fast tracked through the licensing process because the whole world is being devastated by this hardship and poverty disease called Ebola, and therefore the fastest possible remedy has to be introduced "vaccinated".

If these Ebola vaccines are licensed – and you can bet on this – these genetically engineered vaccines are going to be the future vaccinations and the justification will be that they brought this terrible disease called Ebola under control so magnificently. Toddlers will not anymore be confronted with these harmful aluminium compounds shortly after birth but they will now be directly gene manipulated. Probably this will make most of the versatile toxic occurrences from the adjuvants disappear which have burdened the organism but soon new gentechnological engineered organisms will take over the part of the „well-tried“ toxic poisons! It is absolutely doubtful if these infantile gene manipulations will be a blessing for mankind and their health. You can see it already in the agricultural sector in which way the genes of plants are mutating. As soon as genetic vaccines are injected, all further body reactions are out of control.

Background on Ebola scare

Ebola dominates the news although the people of Africa have totally different problems: In Sierra Leone every sixth child dies during the first year of life, every third before his or her fifth birthday. Mass extinction tolls – not due to Ebola but because of hunger and polluted water. The media attention of Ebola is not justified in relation to the existing problems in this region.

Additionally you can find doubtful and totally unspecific virus tests, as with every „pandemic“, which generally react to inflammation processes and prove little about a specific infection. Even WHO (10) points out that nearly all of the people tested positively for Ebola don't show the external bleeding typical of Ebola. Why does WHO agree to blow up the effects of poverty and misery to a virus epidemic in the case of Ebola?

WHO is up to 75 % (11) financed by the pharma industry, other health institutions are also dependant on financial contributions from corporations (12). By international regulations (13) and national laws WHO was given the power to declare pandemic stages and enforce relevant provisions for individual states which thus can be or even must be forced upon their people (14). If an international health crisis arises a „protective“ vaccine can be brought to market in a much faster and easier licensing procedure in order to implement mandatory mass vaccination programs under the disguise of „rescuing“ mankind. In this manner the pharma industry is saving millions, if not even billions. Because the less testing has to be done and the faster something is going through the licensing process, the faster the money comes back and the sooner the company stock will rise. This puts us to the question if WHO, governments, authorities and mass media are abusing the people of western Africa to gain financial profits and also thereby convincing the world of the necessity for a new vaccine.

Whoever still believes in philanthropic motivation of the pharma industry does not accept reality. Pharma is operating like mafia organisations (15) and by „buying“ into politics (16,17,18) it accumulates the respective power to have existing laws adjusted to their marketing strategies and sales planings at will.

By inflating the Ebola scare pharma's goal seems to be to establish the use of totally new vac-

cine manufacturing processes to which products no man or woman would agree to have them injected directly into their bodies. This can only be achieved by inventing a deadly and global crisis. Genetic microorganisms injected directly into the human tissue – this would not be feasible without Ebola!

Gene altered viruses that enter cells like pirates are also the replacement for the aluminium compounds! Another benefit for the pharma industry lays in the disappearance of the adverse effects generated by aluminum which were well researched and documented. Adverse effects in genetic constitution developed through gene altered vaccines are not yet known and a lot more difficult and expensive to prove.

Conclusion

Considering the facts one has the impression that the entire Ebola scare was constructed in order to raise the sales volume of the pharma industry. It is doubtful that the Ebola campaign only serves the purpose of financial interest because the goal is mass vaccination which puts our cell nucleus and its DNA into focus. Because of the above and many more reasons we want to warn insistently about genetic vaccination. It's no blessing to mankind but a massive danger to the genetic makeup of mankind whose full potential will probably only show in a future generation.

Gene manufactured vaccines – it's about time to defend our genetic heritage against the pharma industry. As awful as this whole occurrence is, it is also a big chance: A majority of people are opposing genetechnology in food products.

Genetechnological vaccines are a concern for every one of us, there is no way of estimating the potential of damage! If we can prevent the production and implementation of genetechnological vaccines it would be a huge profit for our health. If we fail, the outcome cannot be visualized in our wildest dreams.

The public is visibly better informed and more and more critical to the schemings of the pharma industry. It is our duty to give this information to the public.

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Switzerland, Germany, November 2014

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