

Have IgE and IgG4 antibodies become vaccine injury biomarkers?

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We know that IgE and IgG4 are involved naturally in helminth defense and now in allergies. IgE dominates in mild helminth infections. IgE mediated histamine release, itching and mucus generation are strategies to physically remove helminths/parasites and prevent major infections. If infection intensifies, an IgE mediated reaction can be dangerous leading to anaphylaxis. So naturally the body downregulates IgE and switches to an IgG4 dominated defense state. A chronic low intensity battle ensues with the helminth to avoid immunopathology of an IgE dominated aggressive defense (1).

We know that any injected protein will result in IgE mediated sensitization to that protein (2,3). Vaccines containing gelatin induce IgE mediated sensitization directed against gelatin (4,5). Egg protein containing vaccines induce IgE mediated sensitization directed against egg proteins (6,7). Toxoid containing vaccines induce IgE mediated sensitization directed against the toxoids (8). Influenza vaccines cause IgE mediated sensitization directed against influenza viral proteins (9–12). Bovine serum albumin (BSA) containing vaccines induce IgE mediated sensitization directed against BSA in humans(13), horses (14) and dogs (15). The Institute of Medicine evaluated the entire research from 1950 to 2011 and concluded that viral/bacterial/food/animal proteins in vaccines do induce IgE mediated sensitization (16).

We know from allergy and helminth infection research that IgE is the first stage of the immune response to an injected antigen. Upon continuing exposure to the antigen, the immune system synthesizes IgG4 directed against the antigen thus resulting in “desensitization” (17–23). Many researchers and allergists including the European Academy of Allergy and Clinical Immunology (EAACI) confuse desensitization with tolerance (24–26). Desensitization is a disease state and corresponds to a chronic parasite infection. There is nothing normal about it. It is useful only because it is better than an IgE dominated defense state that involves anaphylaxis risk.

Vaccines contain numerous proteins including food proteins, fungal proteins, animal proteins and aeroallergens. With the exception of food to some extent, people have no control of exposure to these proteins. Once they develop vaccine-induced IgE mediated sensitization to any of these proteins, unless they suffer a severe allergic reaction, they will continue to be exposed and begin synthesizing IgG4 directed against the antigens. So one can expect IgG4 directed against food, fungal, animal and aeroallergens. Since in developed countries parasite infections are rare, parasite directed IgE/IgG4 are also rare. Therefore in developed countries, IgE/IgG4 are likely to be reliable vaccine injury biomarkers.

IgE/IgG4 directed against food such as milk results in food allergy/eosinophilic esophagitis (27,28). IgG4 directed against bovine folate receptor alpha proteins (a milk protein) results in autism and cerebral folate deficiency (29,30).

IgE/IgG4 directed against bovine serum albumin (BSA) causes BSA allergy/membranous nephropathy (31,32).

IgG4 directed against bovine insulin (a protein in cow’s milk) causes type 1 diabetes (33,34).

IgG4 directed against vaccine antigens can bind to human self proteins due to molecular mimicry thus resulting in numerous autoimmune disorders (35) and IgG4 related diseases (IgG4-RD).

Immunotherapy for vaccine-induced allergies is another major cause of IgG4 synthesis.

The explosion of IgG4-RD is basically proof of an explosion of vaccine injuries. IgE mediated sensitization caused by vaccines is the first iatrogenic disease of the cascade. IgG4-RD due in part to immunotherapy is the next iatrogenic disease.

Allergists and allergy organizations such as the EAACI discourage people from having IgG4 tests done because they (allergists) don't know how to interpret the results (26). These allergists and allergy organizations use corrupted science (36–39) to deny vaccine injury so they are of course incapable of interpreting the results of IgG4 testing.

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