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Correspondence to the article by Tabache F. et al “Acute polyarthritis after influenza A (H1N1 immunization”, Joint Bone Spine, 2011, doi:10.1016/j.jbs.2011.02.007:

Primary Sjögren’s syndrome occurring after influenza A H1N1 vaccine administration

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We read with interest the paper published by Tabache et al, reporting a case of acute and transient polyarthritis following influenza A H1H1 vaccination (1). We report here a case of persistent autoimmune disease that developed after the same vaccination.

The patient was a 30-year-old Caucasian woman without medical history. Due to the national H1N1 vaccination program in France, she received one dose of Pandemrix® in December 2009. Seven days later, she presented with arthralgia of shoulders, knees, ankles, wrists and fingers. In March 2010, she still had polyarthralgia and reported symptoms of dry mouth and dry eyes. At physical examination, there were tender joints but no joint swelling. Laboratory investigations showed an erythrocyte sedimentation rate of 24 mm/h, normal C-reactive protein level, positive rheumatoid factors (224 UI/L, nephelometry), positive antinuclear antibodies (1/640, Hep-2 cells), positive antibodies to SSA (211 UI, ELISA), negative anti-citrullinated peptides antibodies, negative SSB and anti-double stranded DNA antibodies. The HLA class II genotyping was: DRB1* 03, *15, DQB1*02, *06. Results from serologic tests for parvovirus B19, Lyme disease, hepatitis B and C virus, human immunodeficiency virus, cytomegalovirus and influenza A and B virus were negative. Shirmer’s test was altered and a lip biopsy showed the presence of lymphocytic infiltrates in the salivary glands. A RT-PCR was also performed on the salivary gland tissue and failed to detect the presence of the influenza A/H1N1 virus. The patient was treated by hydroxychloroquine giving progressive improvement of arthralgia, but she had still mild and persistent symptoms.

Vaccine has been associated to the development of autoimmune diseases for a long time (2,3) and the vaccination causality is mainly suspected by the temporal relationship between these two events. Different mechanisms may explain these autoimmune phenomena: molecular mimicry, epitope spreading, polyclonal activation and bystander activation (3). Our patient had positive antinuclear antibodies and this may be explained by a quick response of the immune system to the vaccine. Alternatively, she could have had these antibodies before the vaccination due to a dormant autoimmune disease and the injected vaccine had revealed the disease. We did not detect the presence of influenza A H1N1 RNA in the labial tissue using PCR method, leading to the conclusion that there was no direct viral presence within the salivary glands. Since the Pandemrix® adjuvant vaccine did not contain viral genetic
particles, we can hypothesize that vaccine antigens could activate the immune system, and immune cells secondarily traffick to the glands, leading to the clinical symptoms of sicca syndrome. Our patient had a predisposing genetic background (DRB1*03, *15) for the development of SS (4). Thus, the vaccine (viral particles and/or the adjuvant or both) (5) may have played a role as a triggering event in this patient with a genetic background for SS. Chronic autoimmune phenomena following vaccine administration (such as influenza A H1N1 vaccine) are important to report in order to better understand the responsibility of this vaccine. However, these cases are infrequent and a coincidental event cannot be ruled out.


