An infant presenting with Kawasaki disease following immunization for influenza: A case report

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Abstract. Kawasaki disease (KD) is a childhood vascular disorder of unknown etiology. Concerns have recently been raised regarding vaccinations as a potential risk factor for KD. In addition, various forms of vasculitis have been reported as adverse events following administration after various vaccines. Patients exhibiting post vaccination KD have previously been described; however, thus far, to the best of our knowledge, only one patient exhibiting post influenza vaccination KD has been reported in Japan. The present study describes a case of KD 24 h after immunization with influenza in an infant (age, 18 months) following 6 days of high fever, a body rash that had persisted for 2 days and nonsuppurative bilateral conjunctivitis. To the best of the authors' knowledge, this is the first reported case in Korea and the present study reviews various recent studies regarding vasculitis following vaccination and the causal association between them.

Introduction

Kawasaki disease (KD) is one of the most common vasculitides of childhood (1). The cause of KD remains unknown; however, it is hypothesized that the immune system is activated by infectious or environmental triggers in genetically susceptible hosts (1). Previous investigations into vasculitis and KD have reported them as side effects of various vaccinations (2-4). Bonetto et al (5) reviewed the literature from January 1994 to June 2014. Although the majority of the larger, higher quality studies identified no causal association between vaccination and subsequent development of vasculitis, including various studies on KD and Henoch–Schönlein purpura (HSP), the influenza vaccination was ranked first in terms of the number of published articles regarding vasculitis vaccine association (according to vaccine type) and KD was third in terms of the number of published articles regarding vasculitis vaccine association (according to vasculitis type) (5). Cases were reported in which children had received vaccinations, such as those for yellow fever and hepatitis B, and developed KD (2,3). However, a report in 2015 on KD that occurred following influenza vaccination was the only case reported in Japan (4). The present study describes the first case of KD following immunization for influenza in South Korea and reviews various studies regarding vasculitis following vaccination and the causal association between them.

Case report

A male infant, aged 18 months, was admitted to Kyung Hee University Hospital at Gangdong (Seoul, South Korea; 12 October 2015) for evaluation due to a high-grade fever that had started 5 days prior to the visit. The infant had been immunized for influenza 24 h before the onset of fever. He was admitted as he had experienced 6 days of high fever up to 40.0˚C, a body rash that had persisted for 2 days and nonsuppurative bilateral conjunctivitis. Informed consent for the publication of this case report was waived as the infant's next of kin could not be contacted despite numerous attempts to do so.

The body temperature of the infant at the time of admission was 39.6˚C. The physical examination demonstrated bilateral conjunctival injection, cracked red lips, strawberry tongue, erythema of the trunk and the Bacille Calmette-Guérin inoculation site, and erythema and bilateral edema of the hands. However, the cervical lymph nodes appeared normal. The peripheral blood test exhibited leukocytosis (white blood cell count, 12,110 cells/µl), an elevated level of C-reactive protein (15.1 mg/dl) and an elevated quantity of brain natriuretic peptide (403 pg/ml; Table I). Echocardiography demonstrated that the size of the coronary artery was within the normal range, although the left ventricular ejection fraction was somewhat reduced to 50-55%. Based on these findings, KD treatment was initiated with intravenous immunoglobulin (IVIG; 2 g/kg/dose) for 1 day and oral administration of aspirin (50 mg/kg/day). After the end of the IVIG treatment, the fever temporarily improved. However, the infant was injected with...
additional IVIG as a result of a relapse of fever. The second IVIG treatment resulted in the rapid improvement of the KD symptoms, decreased white blood cell count (8,330 cells/µl), level of C-reactive protein (8.6 mg/dl), quantity of brain natriuretic peptide (172 pg/ml) and recovered left ventricular ejection fraction (60%). Therefore, the dose of oral aspirin was reduced to 5 mg/kg/day (Fig. 1). The elevated blood platelet count (625,000 platelets/µl) was then confirmed 7 days after the onset of the disease (Table I).

**Discussion**

Concerns have been raised regarding vaccinations being a potential risk factor leading to KD (6). Various types of vasculitis have been observed and reported in temporal association with the administration of certain vaccines (5). Abrams et al (6) collected and analyzed data of children surveyed from 1996 to 2006 from the Vaccine Safety Datalink (https://www.cdc.gov/vaccinesafety/ensuringsafety/monitoring/vsd/publications.html), and investigated the relevance of childhood vaccines and KD. However, the authors found no evidence that any of the vaccinations increased the risk of KD (6).

Conversely, Zafrir et al (7) demonstrated a possible link between the influenza vaccination and autoimmunity, though the mechanism underlying the development of influenza vaccine-induced vasculitides remains unknown. As an activated immune system following infectious or environmental stimulation contributes to the development of KD, the influenza vaccination may serve as a trigger for the development of KD (4).

Bonetto et al (5) reviewed the literature from 1994 to 2014 and reported that influenza vaccination was ranked first in the number of published articles on vasculitis vaccine association (according to vaccine type) and KD was placed third in the number of published articles on vasculitis vaccine association (according to vasculitis type) (5). Despite the large number of reports, as well as published case reports and case series, suggesting an association between vaccination and vasculitis, the few observational and clinical trials investigating associations between various vaccines and vasculitis subtypes have failed to confirm such an association (5). However, there is increasing evidence establishing various types of infections as potential triggers or causes of different types of vasculitis (5).

Hoffman et al (8) reported a potential association between...
the hepatitis B (HBV) and hepatitis C (HCV) infections and polyarteritis nodosa (PAN) and cryoglobulinemic vasculitis. In France, a successful campaign for vaccination against HBV was followed by a decrease in the incidence of PAN (9). Furthermore, emerging evidence has led to the hypothesis that there is a potential role of infection in KD as well (10). Therefore, the potential of vaccination-induced vasculitis requires consideration in the context of the decreased risk of infection-induced vasculitis (5).

Post-HBV, post-yellow fever and post-influenza vaccination KD have previously been reported (2-4). However, a 2015 report on KD that was performed following vaccination for influenza was, to the best of our knowledge, the only case reported in Japan (4). The present study describes the first case of KD following immunization for influenza in South Korea and reviews previous studies on vasculitis following vaccination and the causal association between them.

In conclusion, it may be hypothesized that influenza vaccination is a trigger of KD when considering the timing of vaccination and onset of KD, although the present study was unable to demonstrate a direct relevance of the influenza vaccine to the onset of KD.

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Availability of data and materials

The analyzed data sets generated during the study are available from the corresponding author on reasonable request.

Authors' contributions

JSW analyzed and interpreted the patient data including the laboratory finding and the pattern of fever. KDH made tables and graph including patient’s information. HMY searched and analyzed references. CSH reviewed and revised about the manuscript. YKL performed an echocardiogram and was a major contributor in writing the manuscript. The final version of the manuscript has been read and approved by all authors.

Ethics approval and consent to participate

Informed consent for the publication of this case report was waived as the infant’s next of kin could not be contacted despite numerous attempts to do so.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References