Autoimmune/autoinflammatory syndrome induced by adjuvants (ASIA) after a silicone injection

Victor G Becerra-Gonzalez,1 Amarilys Alarcon-Calderon,1 Garly Saint Croix,1 David De La Zerda2

DESCRIPTION
Autoimmune/autoinflammatory syndrome induced by adjuvants (ASIA) corresponds to a spectrum of immune-mediated diseases triggered by chronic exposure to adjuvants which are substances meant to enhance antigen-specific immune response.1 Silicone implants have been generally considered relatively inert materials; however, they could generate systemic autoimmune reactions in genetically susceptible individuals leading to ASIA. We describe a case of severe ASIA after cosmetic silicone injections. A 49-year-old woman presented to the emergency department with 3 weeks of persistent fever, chills, dysphagia and worsening dyspnoea. She had been previously healthy and had no sick contacts. One month prior to her presenting symptoms, silicone was injected to her breast and gluteus for an augmentation procedure. On physical examination, she had tenderness to palpation in the thyroid area, multiple neck and axillary lymphadenopathies and few non-tender small nodules in the abdominal wall. CT chest and abdomen showed innumerable well circumscribed and calcified lesions involving the chest wall, mediastinum, pericardium, bilateral hilar regions extending to the pleural space and thickened oesophagus (figures 1 and 2). Biopsy of the skin nodule at the site of injection showed silicon granuloma that confirmed the diagnosis of ASIA. She was treated with steroids with resolution of fever and chills; however, dysphagia and dyspnea only improved partially. Over the course of 12 years, she had multiple hospital admission due to worsening dyspnea or dysphagia. Repeating imaging depicted extensive fibrosis involving the pericardium, pleural and gastrointestinal tract. She had esophageal structures requiring multiple dilations and percutaneous feeding tube. Unfortunately, fibrosis was not reversible, and she developed chronic respiratory failure. This diagnosis corresponds to a dysregulation of either the innate or adaptive immune reaction in genetically susceptible individuals triggered by the exposure of an adjuvant (a substance that enhances the immune response to antigens). Some adjuvants have been linked to a severe diffuse inflammatory reaction

Learning points
► Silicone may act as an adjuvant substance that can trigger a dysregulated systemic autoimmune/ inflammatory or autoimmune reaction with multiorgan involvement in susceptible individuals.
► Autoimmune/autoinflammatory syndrome induced by adjuvants is a rare condition and it has been associated with exposure to other inert material (paraffin, acrylamides, hyaluronic acid or methacrylate).
mediated by innate immune pattern recognition receptor activation such as paraffin, silicone, acrylamides, hyaluronic acid or methacrylate. Silicone was previously considered an inert material, but multiple cases of autoimmune diseases have been reported following silicone implants, mainly undifferentiated connective tissue disorder, but also systemic lupus erythematosus (SLE), rheumatoid arthritis (RA), vasculitis and systemic sclerosis. It has been proposed that there is an increase of profibrotic cytokines, immune dysregulation and antisilicone and anticollegen antibodies. Thus far, reported cases have had poor prognosis and outcomes. Currently, therapeutic options are limited. Removal of the offending agent can be helpful in some patients.

Twitter Victor G Becerra-Gonzales @Becerra_MD

Contributors VGB-G, AA-C, GSC and DDLZ conceived the presented idea and contributed in planning, conducting, acquisition of clinical information and interpretation of the case report. VGB-G and AA-C performed the literature review. VGB-G took the lead in writing the manuscript. All authors provided critical feedback and approved the manuscript for publication.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests None declared.

Patient consent for publication Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

ORCID ID
Victor G Becerra-Gonzales http://orcid.org/0000-0001-5092-2773

REFERENCES