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Clinics in Dermatology

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A definite case of Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS) induced by administration of the Pfizer/BioNTech BNT162b2 vaccine for SARS-CoV2

Jan Walter Schroeder, MD¹, Chiara Gamba, PhD¹, Andrea Toniato, MD^{1,*}, COVID-19 Study Group^{1,2,#}, Franco Rongioletti, MD³

 ¹ From the Unit of Allergology and Immunology, ASST Grande Ospedale Metropolitano Niguarda, Milan, Italy, allergologia@ospedaleniguarda.it
² From the Unit of Pathological Anatomy and Cytogenetics, ASST Grande Ospedale Metropolitano Niguarda, Milan, Italy, anatpat@ospedaleniguarda.it

³ From the Dermatology Clinic, Vita-Salute S. Raffaele University, Milan, Italy

Abstract At the end of December 2020, the anti-SARS-CoV2 vaccination campaign began in Italy. As the number of vaccinated subjects in the general population has increased, several adverse reactions have been observed and reported. Severe cutaneous adverse reactions (SCARs) induced by drugs or vaccines are rare but distinguished by high mortality and include DRESS syndrome or drug induced hypersensitivity syndrome (DiHS), a condition characterized by skin rash, eosinophilia, fever, lymphadenopathy, and involvement of one or more internal organs. Here we present a definite case of DRESS that occured following the administration of Pfizer/BioNTech COVID 19 vaccine. He required hospitalization and was managed with supportive care, antihistamines, and intravenous steroid. © 2022 Elsevier Inc. All rights reserved.

Introduction

At the end of December 2020, the anti-SARS-CoV2 vaccination campaign began in Italy¹. As the number of vaccinated subjects in the general population has increased, several adverse reactions have been observed and reported, some more frequent, such as hyperpyrexia, lymphadenopathy, headache, or muscular-articular pain, and others rarer².

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^{*} Corresponding author: Andrea Toniato, M.D., Unit of Allergy and Immunology, ASST Grande Ospedale Metropolitano Niguarda, Piazza Ospedale Maggiore, 3, 20162 Milan, Italy. Tel. +390264442751.

E-mail address: andrea.toniato@ospedaleniguarda.it (A. Toniato).

[#] COVID-19 Study Group: Valentina Caputo, MD², Luca Giuseppe Balossi, MD¹, Marta Piantanida, MD¹, Linda Borgonovo, MD¹, Joseph Scibilia, MD¹, Camilla Rosa Gentile, MD¹, Martina Liuzzi, MD¹, Selene Nicolosi, MD¹, Alessandro Toscano, MD1, Scarlett Valeria Bernal Ortiz, MD¹, Paolo Calzari, MD¹, Carole Diletta Bagnato, MD¹, Yuri Alessandro Arguello, MD¹, Giovanni Lasagni, MD¹, Jakub Filip Moll, MD¹, Alice Botta, MD¹, Claudia Cordini, MD¹.

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Fig. 1 Pomphoid lesions and diffuse skin edema on admission to our Allergy.

In this regard, the Italian pharmacovigilance agency (AIFA) plays a key role in monitoring adverse reactions to these vaccines². Among the interregional pharmacovigilance projects promoted by AIFA, GRESIF (multiregional pharmacovigilance project "Severe systemic drug hypersensitivity reactions") has the task of detecting and analyzing severe cutaneous adverse reactions (SCARs) induced by drugs or vaccines³. These delayed drug reactions are rare but distinguished by high mortality and include DRESS syndrome or drug induced hypersensitivity syndrome, Stevens-Johnson syndrome, toxic epidermal necrolysis, and acute generalized exanthematous pustulosis (AGEP). In particular, DRESS is a condition characterized by skin rash, eosinophilia, fever, lymphadenopathy, and involvement of one or more internal organs⁴. Resolution of the reaction generally occurs with high-dose steroid therapy. The symptoms usually begin between 3 and 8 weeks of therapy with the culprit drug, but rare cases of this reaction with a much shorter latency following vaccination have been described⁵, including a recent case of possible DRESS/AGEP overlap induced by Janssen vaccine⁶ and another case of DRESS induced by AstraZeneca vaccine7.



Fig. 2 Immunology Unit, during steroid therapy.

Here we present a case of DRESS that occurred following the administration of Pfizer/BioNTech COVID 19 vaccine.

Case report

A 47-year-old man of Asian ethnicity, suffering from ulcerative rectocolitis and allergic asthma, presented to the Emergency Department of the ASST Grande Ospedale Metropolitano Niguarda, in Milan, for high fever ($> 38.5^{\circ}$ C), persistent cough, headache and a diffuse erythematous itchy rash since about a week. The symptoms had occurred after the second dose of Pfizer/BioNtech COVID 19 vaccine that has been given 3 weeks after the first one. The patient reported, after vaccination, the onset of myalgia, abdominal and chest pain associated with a productive cough followed after 5 days by fever, persistent cough and skin rash. The only known drug allergy was angioedema after acetylsalicylic acid with a negative history of allergic reactions to previous vaccinations. The patient denied taking other medications concurrently with the vaccine and in the preceding period except for home therapy consisting of inhalation therapy for asthma and mesalazine.

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Table 1 RegiSCAR score system. Table modified from Kardaun et al. ⁸						
Score	-1	0	1	2	Min	Max
Fever ≥ 38.5°C	No/U	Yes			-1	0
Lymphadenopathy		No/U	Yes		0	1
Eosinophilia					0	2
Eosinophils			700-1499/µl	$\geq 1500/\mu l$		
Eosinophils,			10-19.9%	$\geq 20\%$		
if leukocytes < 4000						
Atypical lymphocytes		No/U	Yes		0	1
Skin involvement					-2	2
Rash extent (> 50% BSA)		No/U	Yes			
Rash suggesting DRESS	No	U	Yes			
Biopsy suggesting DRESS	No	Yes/U				
Organ involvement*		No/U	Yes		0	2
Liver, Kidney, Lung,						
Muscle/heart, pancreas, other						
organ(s)						
Resolution \geq 15 days	No/U	Yes			-1	0
Evaluation other potential			Yes		0	1
causes						
ANA, blood culture, serology for						
HVA/HBV/HCV, C./M.						
pneumoniae, other serology/PCR						
if none positive and > 3 of above						
negative						
TOTAL SCORE					-4	9
U = unknown/unclassifiable; BSA = body surface area; PCR = polymerase chain reaction; C./M. = Chlamydia/Mycoplasma. *after exclusion of other						

explanations: 1 = 1 organ; $2 = \ge 2$ organs. Final score: < 2 = no case; 2-3 = possible case; <math>4-5 = probable case; > 5 = definite case.

Physical examination revealed an erythematous, itchy, maculo-papular skin rash involving the abdomen, chest and upper limbs associated with facial edema and a generalized lymphadenopathy. Laboratory tests performed at the emergency room showed leukocytosis with severe eosinophilia (eosinophils $13,450/\mu$), elevation of alanine aminotransferase (230 U/L) and total bilirubin 1.87 mg/dL, elevation of creatinine 1.37 mg/dL and slight rise in C-reactive protein (CRP) to 2.2 mg/dL. Peripheral blood smear confirmed the leukocyte formula with 70% of cells represented by eosinophils. Skin biopsy could not be performed. The patient was then admitted initially to Emergency Medicine and then to the Department of Allergology and Immunology. At this time, his skin rash exhibited an erythematous and urticarial appearance (Figs. 1 and 2). Abdominal ultrasonography and echocardiogram were negative, while abdominal MRI showed inflammation of the intrahepatic biliary tract, which the hepatologist interpreted as organ involvement related to eosinophilia. The autoimmunity panel (including hepatic autoimmunity), blood cultures, parasitology and viral serologies for HHV-6, HHV-8, Parvovirus B19, Coxsackievirus, CMV, EBV and HIV were also negative. The patient was treated with antihistamine and intravenous steroid (methylprednisone) at a dosage of 1 mg/kg/day with a slow but progressive improvement of the rash associated also with a normalization of both hepato-renal function and eosinophilia. The patient was discharged in good general condition after 2 weeks of hospitalization. At a follow-up of 2 months from the discharge, the patient was doing well both from a clinical and laboratory point of view.

On the basis of the clinical presentation and the results of the investigations performed, the reaction was interpreted as a vaccine-induced SCAR and, in particular, a DRESS syndrome. This diagnosis was confirmed not only by the clinical-laboratory context but also by the application of the RegiSCAR score (see Table 1), which confirmed a definite case.

Discussion

SCARs are rare conditions but associated with high mortality and morbidity. These include DRESS, a delayed drug reaction that usually occurs 3-8 weeks after initiating therapy with the responsible drug. The diagnosis is clinical and is based on diagnostic criteria such as those of the RegiS-CAR score⁸. The most widely used criteria to confirm or exclude the diagnosis of DRESS are those included in the Reg-

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istry of Severe Cutaneous Adverse Reactions (RegiSCAR) scoring system, based on the main clinical manifestations of DRESS⁴: fever >101.3°F or >38.5°C (core) or >100.4°F or >38°C (axillar), enlarged lymph nodes in at least two different body areas, eosinophilia, atypical lymphocytes, skin involvement (extent, rash suggestive of DRESS, biopsy), organ involvement (eg, at least twofold elevation of liver enzymes on at least two different days), and resolution >15 days. A value between -1 and 2 is assigned to each feature (Table 1). The cumulative score ranges from -4 to 9 and defines four levels of certainty regarding the diagnosis of DRESS: excluded, possible, probable, and definite. As some of the variables included in the RegiSCAR DRESS score may not be available when the patient is first evaluated, the score is most useful as a retrospective validation of suspected cases.

Potentially all drugs can cause this syndrome, but the most frequent are aromatic anticonvulsivants and allopurinol⁹, a fact that has also emerged from one of our recent studies¹⁰. However, the literature also describes rare cases associated with vaccinations, which have a shorter onset latency than other drugs⁵. Our patient represents a definite case of DRESS in close temporal association with Pfizer/BioNtech COVID 19 vaccine. Among the COVID-19 vaccine reactions reported, Lospinoso K. et al. described a case of overlapping AGEP-DRESS from Janssen vaccine⁶ and, more recently, O'Connor T. and colleagues presented a case of DRESS following administration of AstraZeneca vaccine⁷. To our knowledge, our case represents the first definite case of DRESS induced by Pfizer/BioNtech COVID 19 vaccine.

As for pathogenesis, it has been suggested that an inefficient detoxification of the drug leads to accumulation of reactive metabolites, which cause an autoimmune reaction in skin and some internal organs in vulnerable patients who have genetic predisposition.

Conclusions

We want to draw attention to the possible occurrence of this type of reactions after vaccination against SARS-CoV2 which, although rare, have significant implications on the health of patients at a time when vaccination for COVID 19 is of primary importance. We also wish to emphasize the importance of a continuous updating by physicians in recognizing this type of reactions so that they can diagnose SCARs early and treat patients in the most appropriate and efficient way.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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