

Clinical Case Reports

Herpes zoster triggered by ChAdOx1 nCoV-19 coronavirus vaccine (recombinant)

NEETHU MARY GEORGE

ABSTRACT

Herpes zoster is a viral infection caused due to the reactivation of varicella-zoster virus that is localized to a single dermatome unilaterally. The factors responsible for its reactivation are increasing age, immunosuppressive drugs, malignancies, chronic liver and renal diseases. Herpes zoster was found to be one of the cutaneous manifestations of coronavirus disease 2019 (Covid-19). Various skin manifestations post-vaccination are being reported, which include injection site urticarial, maculopapular rash and positive dermatographism. We report a patient of herpes zoster triggered by the viral vector (ChAdOx1 nCoV-19) coronavirus vaccine (recombinant).

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INTRODUCTION

Coronavirus disease 2019 (Covid-19) is a multisystem disease caused by severe acute respiratory syndrome coronavirus 2. Since the emergence of the Covid-19 pandemic, the prevention of this rapidly spreading infection became a priority for all countries. A number of different vaccines were developed based on various mechanisms: RNA vaccines (Pfizer, Moderna), viral vector vaccines (Oxford-AstraZeneca, Sputnik V, Johnson and Johnson), inactivated virus vaccine (Covaxin by Bharat Biotech) and protein-based vaccines (Novavax). Two vaccines were used in India—Covaxin by Bharat Biotech, which is an inactivated virus vaccine and the viral vector vaccine by Oxford-AstraZeneca, which was manufactured locally by Serum Institute of India as Covishield. As this was one of the largest vaccine drives with vaccines getting approved in a short time, vaccine safety was closely assessed. Covid-19 is associated with a wide variety of cutaneous manifestations including urticarial rash, chickenpox-like vesicles, erythematous maculopapular rash, pernio-like lesions, acral erythema, dactylitis, purpuric maculopapules and acro-ischaemic lesions.¹ Limited literature is available on skin manifestations following the Covid-19 vaccine, especially a viral vector vaccine. We report an immunocompetent patient with herpes zoster after the first dose of the viral vector vaccine.

THE CASE

A 24-year-old healthy man with no comorbid conditions started

Mar Sleeva Medicity Palai, Kottayam, Kerala, India; neets.1x@gmail.com

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developing pain over the arm and forearm of the same side, extending to the thumb and index finger, 2 weeks after the first dose of inactivated virus vaccine. Two days later, he noticed small fluid-filled lesions over the arms and forearms in groups and red-coloured rash on the palms (Figs 1 and 2). Examination revealed lesions involving the C6–C7 dermatome. Our final diagnosis was herpes zoster considering the characteristic grouped vesicles associated with pain and dermatomal distribution. He was started on oral valacyclovir thrice daily for 5 days and the lesions subsided over few days with no sequelae or post-herpetic neuralgia. The second dose was taken 4 weeks after the first and the post-vaccine days were uneventful.

DISCUSSION

After a primary infection of chickenpox, the varicella zoster virus remains latent in the dorsal root ganglia or cranial nerve ganglia, which gets reactivated spontaneously or gets triggered by trauma, stress, fever or immunosuppression. Herpes zoster reactivation has been recorded post-vaccination such as inactivated influenza, hepatitis A and rabies vaccines.² Even after Covid-19, many cases of herpes zoster reactivation were reported.³ This is thought to be due to Covid-19-induced lymphopenia and functional impairment of CD4+ T cells and activation with subsequent exhaustion of CD8+ T cells.⁴ Clinicians investigated patients for Covid-19 when they presented with herpes zoster, especially in those without any obvious triggers. After the vaccination drive started, there were many reports of herpes zoster getting triggered by vaccines, especially with mRNA vaccines⁵ and inactivated viral vaccines.⁶ mRNA vaccine is considered to act by stimulation of innate immunity through toll-like receptors (TLRs)³ causing



FIG 1. Erythematous grouped papulo-vesicles over the lateral aspect of the arm



Fig 2. Erythematous macules over first interdigital space

Haemophagocytic lymphohistiocytosis post-ChAdOx1 nCoV-19 vaccine: A rare case

VIKAS MARWAH, ROBIN CHOUDHARY,
TENTU AJAI KUMAR, MANISH SHARMA

ABSTRACT

The severe acute respiratory syndrome coronavirus 2 pandemic started in December 2019, spread like wildfire and took an immense toll on human life. ChAdOx1 nCoV-19 vaccine was used worldwide for the prevention of Covid-19. Covid-19 has been implicated in the causation of severe haemophagocytic lymphohistiocytosis (HLH) syndrome. However, the same has not been reported with ChAdOx1 nCoV-19 vaccine in the literature. We report a young man who developed secondary HLH post-ChAdOx1 nCoV-19 vaccination.

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Army Institute of Cardiothoracic Sciences, Pune, Maharashtra, India
VIKAS MARWAH, ROBIN CHOUDHARY,
TENTU AJAI KUMAR Department of Pulmonary, Critical
Care and Sleep Medicine
MANISH SHARMA Department of Pathology

Correspondence to ROBIN CHOUDHARY; robinch19@gmail.com

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reactivation of herpes zoster. The time of reactivation of herpes zoster post-vaccine varied in different reports—2 weeks in our patients, 5 days in a patient reported by Bostan *et al.* for inactivated virus vaccine,⁶ and 2 days after the first dose in case of mRNA vaccine.⁵ Even though it is not possible to establish a direct relationship between herpes zoster and the Covid-19 vaccine, it alerts clinicians to look for a relationship between vaccination and herpes zoster reactivation. Thus, careful documentation and reporting of cutaneous manifestations associated with the Covid-19 vaccine is important.

Conflicts of interest. None declared

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INTRODUCTION

The severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) primarily affects the lung but can also have severe systemic manifestation in the form of cytokine storm syndrome. Haemophagocytic lymphohistiocytosis (HLH) is a severe, life-threatening inflammatory syndrome associated with intense cytokine release, which has been implicated in Covid-19 disease. Covishield (ChAdOx1 nCoV-19) is a recombinant coronavirus vaccine that consists of recombinant, replication-deficient chimpanzee adenovirus vector encoding the SARS-CoV-2 spike (S) glycoprotein. It was widely used in various countries including India, England, Brazil and South Africa. Hyperferritinaemic HLH has not been reported previously following administration of the vaccine. We describe a patient who developed severe cytokine release-related HLH following administration of the ChAdOx1 nCoV-19 vaccine.

THE CASE

A 40-year-old man, non-smoker, with no known comorbid conditions presented with a history of insidious onset, intermittent fever of 2 weeks to our hospital which had started 48 hours after vaccination with ChAdOx1 nCoV-19 vaccine. There was a history of occasional dry cough for 1 week. On examination, he had fever (103.7 °F), tachycardia (heart rate 110/minute) and a spleen palpable 2 cm below the costal margin. He had anaemia (haemoglobin 10.8 g/dl), lymphopenia (total lymphocyte count 2800/cmm), deranged liver enzymes (aspartate aminotransferase/alanine aminotransferase 171/202 i.u./L), hyponatraemia (121 mEq/L), raised triglyceride (320 mg/dl), serum fibrinogen (410 mg/dl) and lactate dehydrogenase (LDH) levels (926 i.u./L) (Table I). The C-reactive protein and ferritin levels (>1200 ng/ml on two occasions 7 days apart) were raised.