Acute-onset central serous retinopathy after immunization with COVID-19 mRNA vaccine

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ARTICLE INFO

Keywords: COVID-19 mRNA vaccine CSR Central serous retinopathy

ABSTRACT

Purpose: We report the case of a 33-year-old male who presented with unilateral central serous retinopathy three days after the injection of a COVID-19 vaccine.

Observations: A 33-year-old healthy Hispanic male referred to the ophthalmology service due to blurry vision and metamorphopsia in the right eye without any flashes, floaters, eye redness or pain. The patient reported that 69 hours prior to presentation he received the first dose of the Pfizer-BioNTech BNT162b2 mRNA COVID-19 vaccine. He denied any past ocular history or pertinent medical history. He does not take any medicines and denies stressful factors in his life. The clinical examination and imaging tests were consistent with central serous retinopathy that resolved in three months.

Conclusions and importance: This is the first report of an ocular complication potentially associated with a COVID-19 vaccination. Our case contributes information of a side effect potentially related to this new vaccine.

1. Introduction

Since the first report of the novel coronavirus SARS-CoV-2 in China in December 2019, COVID-19 has been responsible for more than three million deaths worldwide. The ongoing outbreak of COVID-19 has been declared by the World Health Organization as a public health emergency and a vaccine to control the situation was anxiously expected. Vaccine development occurred in less than one year and thus, long-term data on side effects are still developing.

The most common ocular manifestations of COVID-19 is hyperemia or conjunctival “congestion.”\textsuperscript{1,2} In addition, reports of central retinal vein occlusion,\textsuperscript{3} toxic shock syndrome\textsuperscript{4} and hemorrhages with microinfarcts have been described in COVID-19 patients.\textsuperscript{5} To the best of our knowledge, no intraocular side effects have been reported from any of the COVID-19 vaccinations (a case of Bell’s Palsy was described in a patient who received the Moderna mRNA COVID-19 vaccine).\textsuperscript{6}

We present a unique case of a healthy Hispanic male who presented with unilateral central serous retinopathy (CSR), temporally related to the administration of a COVID-19 vaccine.

2. Case report

A 33-year-old healthy Hispanic male was referred for ophthalmological evaluation due to blurry vision and metamorphopsia in his right eye. He denied flashes, floaters, ocular pain or redness. He did not have any visual symptoms in his left eye.

Past ocular history was negative except for mild hyperopic refractive error. Past medical history was only significant for hip surgery post traumatic injury six years prior to presentation. He denied use of any medications including any remote history of corticosteroids. 69 hours before presentation he received the Pfizer-BioNTech mRNA COVID-19 vaccine, after which, he reported soreness at the injection site and fatigue for 24 hours. Since the inception of the pandemic, the patient has experienced no COVID-19 symptoms and had a negative PCR result in November 2020.

On exam, his best corrected visual acuity (BCVA) was 20/63 and 20/25 in his right and left eye, respectively. Pupils were equally round and reactive to light and accommodation. Intraocular pressure was 10 mmHg in both eyes.

Anterior segment exam was within normal limits. Dilated fundus
examination was normal in the left eye but revealed loss of foveal reflex and swollen appearance of the macula in the right eye. No hemorrhages or vascular abnormalities were noted (Fig. 1A).

Optical coherence tomography (OCT) of the right eye showed a macular serous detachment of the neurosensory retina (Fig. 1B) with a central foveal thickness (CFT) of 457 μm. OCT of the left eye was unremarkable. On fluorescein angiography (FA), a single point of leakage was noted following the classical ink-blot pattern, with progressive expansion of hyperfluorescence emanating from a single point (Fig. 2A). Consistent with previous reports of CSR, OCT angiography (OCTA) showed generally attenuated flow signal in the choriocapillaris that colocalized to the area of serous retinal detachment and foci of increased flow signal (Fig. 2B).

After the initial evaluation, the patient was prescribed spiranolactone 50mg daily and was evaluated two and three months later. At the two-month visit, BCVA improved to 20/40 and CFT decreased to 325 μm. At the three-month visit, BCVA improved to 20/20, CFT decreased to 211 μm, OCT showed complete resolution of subretinal fluid, and the patient was asymptomatic.

3. Discussion

We present a case of acute unilateral CSR that developed shortly after immunization with the Pfizer-BioNTech mRNA COVID-19 vaccine. To the best of our knowledge, this is the first report of an intraocular complication associated with COVID-19 vaccination.

We believe it is prudent to consider the immunization as a potential contributor to disease, given 1) the temporal association between immunization and symptom onset, 2) the relatively low incidence of CSR (9.9 per 100,000 individuals) and 3) our case’s absence of classical risk factors.
factors for CSR development (including a history of exogenous steroid use, recent stressful social history, and type-A personality). While no cases of mRNA COVID-19 vaccine associated CSR have been reported to date, CSR has been associated with vaccinations against influenza, yellow fever, anthrax, and smallpox. Still, these cases are rare. In fact, a search of the terms “central serous”, “central serous retinopathy”, “central serous chorioretinopathy” and “CSR” across all vaccine products by all vaccine manufacturers yielded no results from 1990 to date using the U.S. Centers for Disease Control and Prevention’s Vaccine Adverse Event Reporting System. Nevertheless, vaccines have also been associated with a host of chorioretinal pathologies beyond CSR. For example, vaccinations against influenza, yellow fever, hepatitis B, and Neisseria meningitidis have been associated with uveitis, acute idiopathic maculopathy, acute macular neuroretinopathy, Vogt-Koyanagi-Harada disease, and multiple evanescent white dot syndrome.4–23

The pathophysiology of CSR remains incompletely understood, but the current literature emphasizes the role of a hyperpermeable and thickened choroid, as a result of hydrostatic forces, ischemia, or inflammation.11 Further evidence suggests endogenous and exogenous glucocorticoids contribute to the development of the choroidal vasculopathy.24 The role of retinal pigment epithelium dysfunction remains uncertain. Furthermore, CSR’s difficulty to explain associations with additional risk factors—systemic hypertension,25 alcohol use,25 gastrointestinal reflux disease,26 and sympathomimetic agents27—offer additional questions about the pathophysiology of disease.

There are several possible pathophysiologic mechanisms for the mRNA vaccine to cause CSR. Our hypotheses for possible pathophysiologic mechanisms are divided into three arms, related to increased serum cortisol, free extracellular mRNA, and polyethylene glycol. First, there is an association between CSR and high serum cortisol levels.28 The authors could not find any data related to serum cortisol levels after mRNA immunizations. However in one study of a tetanus toxoid vaccination, serum cortisol increased acutely nearly two-fold after vaccination.29 Therefore, it seems reasonable that mRNA immunizations may similarly trigger endogenous glucocorticoid release. On the other hand, studies have consistently demonstrated that inflammatory cytokines inhibit glucocorticoid receptor expression through decreased receptor translocation and binding affinity.30 Vaccines, including the extensively studied influenza virus vaccine, induce mild inflammatory responses.1–3,11,13–17 Another study showed that pain at injection site and increased body temperature were accompanied by a pro-inflammatory cytokine response.34

Other potential etiologies may arise from the presence of extracellular RNA. Extracellular naked RNA has been shown to increase the permeability of endothelial cells and may thus contribute to leaky choriocapillaris.35 Alternatively, extracellular naked RNA has been shown to promote blood coagulation and thrombus formation.26 Studies of CSR have highlighted that retinal areas with mid-phase inner choroidal staining also have delayed choroidal filling, suggesting choroidal lobular ischemia with associated areas of venous dilation.7,36–38 Interestingly, this phenomenon may explain the attenuated flow signal in the choriocapillaris on OCTA in cases of CSR, including our case.25,29 Further, elevated serum plasminogen activator inhibitor-1, a fibrinolytic, in CSR have led to the suggestion of a thrombotic mechanism for these vascular changes.39

Finally, polyethylene glycol (PEG)-2000 is a lipid ingredient in the vaccine formulation meant to enable the delivery of RNA into host cells. PEGs are the most commonly used hydrophilic polymers used in drug delivery tasked to control pharmacokinetic properties and deliver drug to specific sites. Anaphylactic reactions have been reported in relation to PEG compounds, that are used in bowel preparation regimens and prescription medications (e.g. methylprednisolone acetate).40 In addition, subretinal PEG-8 induces choroidal neovascularization and choroidal vessel thickening through activation of the complement pathway in murine models.41,42

4. Conclusions
Acute CSR may be temporally associated with mRNA Covid-19 immunization.

Patient consent
Informed consent was obtained on this patient at the Instituto “La Raza,” Mexico City, Mexico.

Acknowledgements and Disclosures
The authors have no disclosures.

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