Reversal of acute liver failure with N-acetylcysteine and prednisone in a patient with DRESS syndrome: A case report and literature review

Falla hepática aguda en una paciente con síndrome de DRESS que revirtió con N-acetilcisteína y prednisona. Reporte de caso y revisión de la literatura

Drug Rash with Eosinophilia and Systemic Symptoms (DRESS) is an idiosyncratic reaction to medication\(^1\) that is characterized by skin rash, hematologic alterations, and organ involvement.\(^2\) It has been related to the ingestion of phenytoin and other anticonvulsant agents.\(^3\) Skin reactions have been described in up to 19% of patients between 6 and 8 weeks after drug initiation.\(^4\) The mortality rate is above 10% and death is commonly secondary to acute liver failure (ALF).\(^4\)

One month before her hospital admittance, a 46-year-old woman presented with subarachnoid hemorrhage due to a ruptured aneurysm of the posterior left cerebral artery; the affected vessel was clipped and she began adjuvant management with 100 mg of phenytoin every 8 h. She had no past history of herbal medicine or alcohol consumption, or prior use of other medication or hepatotoxic agents. Forty-eight hours after drug initiation, the patient noticed maculopapular lesions on both hands that resolved spontaneously, with no other symptoms. Three weeks later the maculopapular lesions became generalized and turned into an exfoliative dermatitis. She developed unmeasured fever along with pruritus, jaundice, and choloria. Upon hospital admittance the patient presented with dehydration, jaundice, generalized maculopapular lesions with fine flaking (fig. 1), cervical adenomegaly, hepatomegaly of 3 cm under the costal margin, and no hepatic encephalopathy (HE). Her laboratory tests reported: leukocytes 5,000 L\(^{-1}\), eosinophils 1,800 L\(^{-1}\), urea 104 mg/dL, creatinine 5.6 mg/dL, total bilirubin 8.6 mg/dL, albumin 2.9 g/L, alanine aminotransferase 171 U/L, aspartate aminotransferase 333 U/L, alkaline phosphatase 751 U/L, gamma-glutamyl transpeptidase 1,814 U/L, prothrombin time 38%, and international normalized ratio (INR) 1.8. Cultures had no pathogen development, the viral panel was negative for hepatitis A, B, and C, and cytomegalovirus (CMV) and Epstein-Barr virus (EBV) IgM serology were negative. Abdominal ultrasound showed no chronic hepatopathy data, no biliary tract dilation, no vascular thrombosis, and no alterations in either kidney. Management was begun with prednisone 1 g/kg of weight and pentoxifylline 300 mg every 6 h; on her second day in the hospital, the patient presented with stage 2 hepatic encephalopathy characterized by asterixis and bradypsychia. The data indicated: stage 2 HE, INR

\(\text{Please cite this article as: Pérez-Reyes E, Casanova-Lara A, Pérez-Torres E, Córdova J. Falla hepática aguda en una paciente con síndrome de DRESS que revirtió con N-acetilcisteína y prednisona. Reporte de caso y revisión de la literatura. Revista de Gastroenterología de México. 2014;79:208-210.}\)
Clinical symptoms are characterized in addition to ruling out infections. NAC has been proposed as part of the treatment in cases of ALF, due to the ingestion of other medications. The exact incidence of DRESS syndrome is not known; it is estimated at 1 for every 1,500 new users of phenytoin. Its pathophysiology is unknown, but it has been associated with immunologic factors and genetic predisposition. Detoxification defects leading to the formation of reactive metabolites, activated T lymphocytes that release cytokines (IFN-γ and IL-5), exerting cytotoxicity, have been proposed; others have considered it to be a systemic reaction due to reactivation of the human herpes virus 6 and 7, EBV, or CMV. The drugs most associated with the development of DRESS are the anti-epileptic agents (phenobarbital, carbamazepine, phenytoin, sodium valproate, lamotrigine), allopurinol, sulfaalazine, dapsone, and minocycline. Antibiotics or nonsteroidal anti-inflammatory analgesics have rarely been reported. The disorder is independent of the quantity of the drug, having a greater relation to the susceptibility of the individual.

The first step of treatment is the definitive suspension of the medication at fault. Local steroids and antihistamines can be used in cases that are not severe. There are no criteria for the systemic use of steroids, but they are generally used in patients with organ involvement. In cases of ALF, 3 doses of intravenous methylprednisolone at 1 mg/kg of weight, followed by oral steroid, has shown good results. Liver transplantation (LT) is the treatment of choice for patients with ALF that do not respond to high doses of steroids. NAC has been proposed as part of the treatment for patients with ALF secondary to the ingestion of paracetamol, as an antidote to intoxication by that drug, and some studies have shown its usefulness in patients with ALF due to the ingestion of other medications. NAC combined with steroids has been reported mainly in patients with anticonvulsant-induced DRESS, with a favorable response, but the number of studies supporting its use is insufficient.

Our patient’s illness was induced by phenytoin and upon admittance she showed data of ALF and acute renal failure (ARF) (probably due to interstitial nephritis). She was successfully treated with the combination of oral steroid and NAC and the suspension of phenytoin. We believe that early initiation of the double therapy in ALF patients can lead to improvement, averting the need for LT.

Financial disclosure
No financial support was received in relation to this article.

Conflict of interest
The authors declare that there is no conflict of interest.

Bibliografía
3. Oelze L, Pillow T. Phenytoin-induced drug reaction with eosinophilia and systemic symptoms (DRESS) syndrome: A

E. Pérez-Reyes a, A. Casanova-Lara a, E. Pérez-Torres a, J. Córdova b,∗

a Servicio de Gastroenterología, Hospital General de México, México DF, México
b Servicio de Gastroenterología, Instituto Nacional de Ciencia Médicas y Nutrición Salvador Zubirán, México DF, México

∗Corresponding author: Servicio de Gastroenterología, Instituto Nacional de Ciencia Médicas y Nutrición Salvador Zubirán, México DF. C/ Vasco de Quiroga 15 Colonia Sección XVI Delegación Tlalpan CP 14000 México DF. Teléfono: +55 73 34 18.
E-mail address: jacquiemex2@yahoo.com.mx
(J. Córdova).