



Atypical Hemolytic Uremic Syndrome in a One-YearOld Male Patient, Successfully Treated with Eculizumab

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Abstract:

Atypical hemolytic uremic syndrome is a rare form of thrombotic microangiopathies resulting from various genetic mutations of the complement factors. Thrombotic microangiopathies include Thrombotic thrombocytopenic purpura, primary HUS further divided into typical HUS or Shiga-toxin related HUS, atypical HUS and secondary forms of HUS. The most common clinical features of the Hemolytic Uremic Syndrome are: kidney failure, thrombocytopenia resulting in hemorrhagic phenomena and also intravascular hemolytic anemia. Most commonly HUS is a diagnosis made from excluding all other plausible causes. HUS predominantly affects pediatric ages. Here we present the case of a 1-year old male patient that was admitted to the pediatric intensive care unit with; a generalized severe condition, vomiting and diarrhea. The infant is agitated. Gross hematuria is noted. The skin and mucosa are pale. Periorbital and pitting edema in the lower extremities. BP of 140/90 mm/Hg.

Biography:

Henri is a medical student currently in the last semester. He has joined voluntarily the Albanian Centre for Medical Emergency amid the COVID-19 outbreak. After one month of training as a volunteer he now works at



the centre of coordination for Medical Emergencies at the COVID-19 department. Henri has participated with a team of 4, in the International Medical Tournament held in Novosibirsk Russia where the team got ranked third. He is a cofounder and vice-president for Internal Affairs at "Albanian Committee of Medical Students" and is the former head of delegation for the integration process in IFMSA (International Federation of Medical Students Associations) a position he held during the IFMSA August Meeting in Taipei Taiwan, where Henri presented the Albanian candidacy. ACMS is now a member of IFMSA. Has two publications at ACTA SCIENTIFIC MEDICAL SCIENCES.

7th International conference on Medical and Nursing education, July 17, 2020, Vienna, Austria

Citation: Henri Ardian Fero, Atypical Hemolytic Uremic Syndrome in a One-YearOld Male Patient, Successfully Treated with Eculizumab, 7th International conference on medical and nursing education , July 17, 2020, Vienna, Austria.