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Atypical hemolytic-uremic syndrome due to complement factor I mutation

Almalki AH *et al.* Atypical hemolytic-uremic syndrome

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### Abstract

Atypical hemolytic-uremic syndrome (aHUS) is a rare disease of complement dysregulation leading to thrombotic microangiopathy (TMA). Renal involvement and progression to end-stage renal disease are common in untreated patients. We report a 52-year-old female patient who presented with severe acute kidney injury, microangiopathic hemolytic anemia, and thrombocytopenia. She was

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