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**Case Report**

**Haemolytic anaemia – Initial presentation of Wilson Disease**

**Dr. Bageshree Seth, Dr. Jeetendra Gavhane, Dr. Revathi N., Dr. Maninder Singh Setia**

Department Of Pediatrics , M.G.M.Medical College and Hospital, Kamothe, Navi Mumbai , India

**Corresponding author:** Dr. Bageshree Seth

**Abstract:**

Wilson disease is an autosomal recessive inborn error of copper metabolism characterized by toxic accumulation of copper in liver, brain, cornea and other tissues. It usually presents with hepatic dysfunction or neuropsychiatric manifestations, Hemolytic anemia is an uncommon presenting manifestation of Wilson disease.We present a case of a child who presented with hemolytic anemia and was subsequently diagnosed as Wilson disease.

**Keywords**: Hemolytic anemia, Wilson disease, first manifestation