

LIST OF ARTICLES NOT INCLUDED FOR REVIEW

1. Abdemoula MS. La methemoglobinemie hereditaire recessive de type II. A propos d'une observation. *Revue Maghebaine de Pediatrie* 2002; XII-IV: 207–10.
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13. Nishina T, Miwa S, Hara N, Asakura T. Hereditary methemoglobinemia with mental and growth retardation found in a Japanese boy. With special reference to xylitol, sorbitol metabolism and methemoglobin reduction in the erythrocyte. *Acta Haematol Jpn* 1970;33:455-61.

14. Orsini A, Vovan L, Brusquet Y, Gabriel B, Sebag F, Galtier M. Congenital methemoglobinemia due to NADH (DPNH) dependent methemoglobin reductase deficiency. *Mars Med* 1972; 109: 279–81.
15. Ronconi G, Ferracin G. [Congenital methemoglobinemia of the recessive type due to diaphorase deficit with oligophrenia.]. *Riv Clin Pediatr* 1964; 74: 152–9.
16. Ronconi G, Ferracin G. On a new case of congenital methemoglobinemia caused by deficiency of diaphorase with grave cerebropathy. *Fracastoro* 1968; 61: 121–8.
17. Roussel A, Maestraggi P, Tremoulet M, Marchand .A new case of recessive congenital methemoglobinemia. *Arch Fr Pediatr* 1963; 20: 745–50.
18. Sacerdotti-Favini. Methemoglobinemia costituzionale con cerebropatia e oligofrenia. *Acta pediat Lat* 1948; 11: 255.
19. Shotelersuk V, Tosukhowong P, Chotivitayatarakorn P, Pongpunlert W. A Thai boy with hereditary enzymopenic methemoglobinemia type II. *J Med Assoc Thai* 2000; 83: 1380–6.
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GEOGRAPHIC ORIGIN AMONG PATIENTS ANALYZED FROM LITERATURE REVISION

India: 13 patients

Japan: 5 patients

Turkey, France, Algeria: 3 patients each

Italy, Cuba, Saudi Arabia: 2 patients each

England, South Africa, Thailand, Suriname, Germany, Pakistan, Lebanon, Tunisia, Egypt, Honduras, Australia, USA: 1 patient each

Mixed ethnicity: French-Spanish in one patient

Not clearly reported: 4 patients