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Familial hemophagocytic lymphohistiocytosis type 2 in Chinese population: A case report and review of literature

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Abstract

BACKGROUND

Familial hemophagocytic lymphohistiocytosis (FHL2) is a rare genetic disorder presented with fever, hepatosplenomegaly, and pancytopenia secondary to perforin-1 (PRF1) mutation. FLH2 has been described in Chinese but usually presents after one-year-old. We describe a female Chinese neonate with FHL2 sary to compound

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