Fanconi anemia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Fanconi anemia. ORPHA:84

Fanconi anemia (FA) is a hereditary DNA repair disorder characterized by progressive pancytopenia with bone marrow failure, variable congenital malformations and predisposition to develop hematological or solid tumors.