Autosomal recessive secondary polycythemia not associated with VHL gene

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal recessive secondary polycythemia not associated with VHL gene. ORPHA:247378

A rare, hereditary, hematologic disease characterized by an increase in hemoglobin, hematocrit and erythrocyte mass resulting in plethora or ruddy complexion, headache, dizziness, tinnitus and exertional dyspnea. In some cases, thrombophlebitis and arthralgia have also been reported.