Open Peer Review on Qeios

Mendelian susceptibility to mycobacterial diseases due to complete IL12RB1 deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Mendelian</u> <u>susceptibility to mycobacterial diseases due to complete IL12RB1 deficiency</u>. <i>ORPHA:319552

Mendelian susceptibility to mycobacterial diseases (MSMD) due to complete interleukin-12 receptor subunit beta-1 (IL12RB1) deficiency is a genetic variant of MSMD (see this term) characterized by mild bacillus Calmette-Guérin (BCG) infections and recurrent Salmonella infections.