Behçet disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Behçet disease. ORPHA:117

Behçet's disease (BD) is a chronic, relapsing, multisystemic vasculitis characterized by mucocutaneous lesions, as well as articular, vascular, ocular and central nervous system manifestations.