Relapsing polychondritis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Relapsing polychondritis. ORPHA:728

Relapsing polychondritis (RP) is a rare, clinically heterogeneous, multisystemic inflammatory disease characterized by inflammation of the cartilage and proteoglycan rich structures leading to cartilage damage with joint, ocular and cardiovascular involvement.