Polyarteritis nodosa

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Polyarteritis nodosa. ORPHA:767

Polyarteritis nodosa (PAN) is a rare, clinically heterogeneous, rheumatologic disease characterized by necrotizing inflammatory lesions affecting small- and medium-sized blood vessels. PAN most commonly affects skin, joints, peripheral nerves, the gut, and the kidney.