POEMS syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. POEMS syndrome. ORPHA:2905

POEMS syndrome is a paraneoplastic syndrome characterized by polyradiculoneuropathy (P), organomegaly (O), endocrinopathy (E), clonal plasma cell disorder (M), and skin changes (S). Other features include papilledema, extravascular volume overload, sclerotic bone lesions, thrombocytosis/erythrocytosis, and elevated VEGF levels.