CREST syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. CREST syndrome. ORPHA:90290

CREST syndrome is a subtype of limited cutaneous systemic sclerosis (lcSSc; see this term) whose name is an acronym for the cardinal clinical features of the syndrome: calcinosis, Raynaud's phenomenon, esophageal dysmotility, sclerodactyly and telangiectasia.