Lichtenstein syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lichtenstein syndrome. ORPHA:2390

Lichstenstein syndrome is characterised by frequent infections associated with osteoporosis, a tendency for fractures and osseous anomalies. It has been described in two monozygotic twin brothers. Transmission is autosomal recessive.