Geroderma osteodysplastica

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Geroderma osteodysplastica. ORPHA:2078

Geroderma osteodysplastica (GO) is characterized by lax and wrinkled skin (especially on the dorsum of the hands and feet and abdomen), progeroid features, hip dislocation, joint laxity, severe short stature/dwarfism, severe osteoporosis, vertebral abnormalities and spontaneous fractures, and developmental delay and mild intellectual deficit.