Berardinelli-Seip congenital lipodystrophy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Berardinelli-Seip congenital lipodystrophy. ORPHA:528

Berardinelli-Seip congenital lipodystrophy (BSCL) is characterized by the association of lipoatrophy, hypertriglyceridemia, hepatomegaly and acromegaloid features. BSCL belongs to the group of extreme insulin resistance syndromes, which also includes leprechaunism, Rabson-Mendenhall syndrome, acquired generalized lipodystrophy, and types A and B insulin resistance (see these terms).