# Neurofibromatosis type 1

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Neurofibromatosis type 1. ORPHA:636*

Neurofibromatosis type 1 (NF1) is a clinically heterogeneous, neurocutaneous genetic disorder characterized by café-au-lait spots, iris Lisch nodules, axillary and inguinal freckling, and multiple neurofibromas.