Satoyoshi syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Satoyoshi syndrome. ORPHA:3130

Satoyoshi syndrome is a rare, multisystemic autoimmune disease mainly characterized by intermittent painful muscle spasms, alopecia (totalis or universalis in most cases) and long-lasting diarrhea that could lead to malnutrition, growth retardation, and amenorrhea. Secondary bone deformities and various endocrine anomalies may also be associated. Antinuclear antibodies are reported in many cases.