

## Opis choroby \*

### Definicja

A rare systemic or rheumatologic disease characterized by peripheral osteolysis (especially carpal and tarsal bones), interphalangeal joint erosions, subcutaneous fibrocollagenous nodules, facial dysmorphism, and a wide range of associated manifestations.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

MONA spectrum

Spektrum MONA

NAO syndrome

Nodulosis-arthropathy-osteolysis syndrome

Torg-Winchester syndrome

#### Kod ORPHA

371428

#### Kod OMIM

277950

#### Kod ICD10

M89.5

#### Kod ICD11

-

---

#### \*Źródło

orphanet