

## Opis choroby \*

### Definicja

A rare autoinflammatory syndrome characterized by the presence of features of relapsing polychondritis and Behçet's disease in the same individual. This includes cartilage inflammation of the ears, nose, throat, and rib cage, as well as recurrent oral and genital ulcers, respectively. Patients may also present ocular involvement (in particular anterior uveitis or scleritis), arthritis, fever, colitis, thrombophlebitis, central nervous system vasculitis, or, in rare cases, arterial aneurysms. Symptoms of polychondritis occur secondary to those of Behçet's disease in the vast majority of cases.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Mouth and genital ulcers-inflamed cartilage syndrome

Wrzody ust i narządów płciowych z zapaleniem chrząstek

#### Kod ORPHA

324972

#### Kod OMIM

-

#### Kod ICD10

D89.8

#### Kod ICD11

-

---

#### [\\*Źródło](#)

orphanet