DISEASE:
Acrodermatitis continua of Hallopeau

NAME:	Acrodermatitis continua of Hallopeau
DESCRIPTION:	A rare, genetic, chronic, recurrent, slowly progressive, epidermal disease characterized by small, sterile, pustular eruptions, involving the nails and surrounding skin of the fingers and/or toes, which coalesce and burst, leaving erythematous, atrophic skin where new pustules develop. Onychodystrophy is frequently associated and anonychia and osteolysis are reported in severe cases. Local expansion (to involve the hands, forearms and/or feet) and involvement of mucosal surfaces (e.g. conjunctiva, tongue, urethra) may be observed.
ORPHACODE:	163931
XREF(S):	Orphanet ICD-10
ANALYTE(S):	IL36RN AP1S3
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RELATED CONTENT

Related Genetic Tests

- Ichthyosis (gene panel)
- Periodic Fever (88 genes)

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- adaptor related protein complex 1 subunit sigma 3
- interleukin 36 receptor antagonist

Related Gene Panels

- Ichthyosis and erythroderma (98 genes) - KUL
- Periodic Fever (88 genes) - ULB