DISEASE:
Acrofacial dysostosis, Weyers type

NAME:	Acrofacial dysostosis, Weyers type
DESCRIPTION:	A rare ectodermal dysplasia syndrome with bone abnormalities characterized by onychodystrophy; anomalies of the lower jaw, oral vestibule and dentition; post-axialpolydactyly; moderately restricted growth with short limbs; and normal intelligence. Although it closely resembles Ellis-van Creveld syndrome (see this term), an allelic disorder and another type of ciliopathy, WAD is usually a milder disease without the presence of heart abnormalities and is inherited in an autosomal dominant manner.
ORPHACODE:	952
SYNONYMS:	Curry-Hall syndrome Weyers acrodental dysostosis Weyers acrofacial dysostosis
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	EVC EVC2
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