ANALYTE:
HPGD

NAME:	15-hydroxyprostaglandin dehydrogenase
SYMBOL:	HPGD
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	15-hydroxyprostaglandin dehydrogenase (NAD(+)) SDR36C1 short chain dehydrogenase/reductase family 36C, member 1
XREF(S):	Orphanet OMIM Reactome SwissProt Ensembl Genatlas HGNC
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Intellectual disability (virtual gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)

Related Diseases

- Cranio-osteoarthropathy
- Isolated congenital digital clubbing
- Pachydermoperiostosis

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Intellectual disability (gene panel)
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

- Skin disorders - UGent
