
DISEASE:
Floating-Harbor syndrome

NAME:	Floating-Harbor syndrome
DESCRIPTION:	A multiple congenital anomalies/dysmorphic syndrome-intellectual disability that is characterized by facial dysmorphism, short stature with delayed bone age, and expressive language delay.
ORPHACODE:	2044
XREF(S):	<u>Orphanet</u> <u>OMIM</u> <u>MeSH</u> <u>ICD-10</u>
ANALYTE(S):	<u>SRCAP</u>
CREATED:	13 May 2019 - 01:02
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Source URL: <http://gentest.healthdata.be/disease/810>

RELATED CONTENT

Related Genetic Tests

- [Floating Harbor](#)
- [Short Stature \(gene panel\)](#)
- [Short stature/ Growth retardation/ \(gene panel\)](#)

Related Laboratories

- [Centre de Génétique-Institut de Pathologie et de Génétique \(IPG\)](#)
- [Centrum Medische Genetica - UZ Antwerpen](#)

Related Analytes

- [Snf2 related CREBBP activator protein](#)

Related Gene Panels

- [Growth retardation/short stature \(genepanel\) - UZA](#)
- [Short Stature \(46 genes\) - IPG](#)

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