DISEASE:
Menkes disease

NAME:	Menkes disease
DESCRIPTION:	A rare congenital disorder of copper metabolism with severe multisystemic manifestations that are primarily characterized by progressive neurodegeneration and marked connective tissue anomalies. A pathognomonic feature is the typical sparse, abnormal steely hair.
ORPHACODE:	565
SYNONYMS:	MD Menkes kinky hair disease Menkes syndrome
XREF(S):	Orphanet MedDRA OMIM ICD-10
ANALYTE(S):	ATP7A
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RELATED CONTENT

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- Neuropathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

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