DISEASE:
Familial atypical multiple mole melanoma syndrome

NAME:	Familial atypical multiple mole melanoma syndrome
DESCRIPTION:	Familial atypical multiple mole melanoma (FAMMM) syndrome is an inherited genodermatosis characterized by the presence of multiple melanocytic nevi (often >50) and a family history of melanoma as well as, in a subset of patients, an increased risk of developing pancreatic cancer (see this term) and other malignancies.
ORPHACODE:	404560
SYNONYMS:	B-K mole syndrome FAMM-PC syndrome FAMMM syndrome Familial atypical mole syndrome Familial atypical multiple mole melanoma-pancreatic carcinoma syndrome Familial dysplastic nevus syndrome Melanoma-pancreatic cancer syndrome
XREF(S):	Orphanet ICD-10 OMIM OMIM
ANALYTE(S):	CDKN2A
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RELATED CONTENT

Related Genetic Tests

- Hereditary Melanoma Panel (7 genes)
- Hereditary cancer (gene panel)
- Melanoma / Familial Atypical Multiple Mole Melanoma Syndrome (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- cyclin dependent kinase inhibitor 2A

Related Gene Panels

- Hereditary Melanoma Panel (7 genes) - ULG
- Hereditary predisposition to cancer (47 genes) - IPG
- Melanoma and Familial Atypical Multiple Mole Melanoma Syndrome (8 genes) - KUL