

**DISEASE:****Familial adenomatous polyposis due to 5q22.2 microdeletion**

<b>NAME:</b>	Familial adenomatous polyposis due to 5q22.2 microdeletion
<b>ORPHACODE:</b>	261584
<b>SYNONYMS:</b>	Colorectal adenomatous polyposis due to monosomy 5q22.2 FAP due to monosomy 5q22.2 Familial adenomatous polyposis due to del(5)(q22.2) Familial adenomatous polyposis due to monosomy 5q22.2 Familial polyposis coli due to monosomy 5q22.2
<b>XREF(S):</b>	<u>Orphanet</u> <u>ICD-10</u>
<b>ANALYTE(S):</b>	<u>APC</u>
<b>CREATED:</b>	13 May 2019 - 01:02
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## RELATED CONTENT

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### Related Genetic Tests

- Hereditary Polyposis Panel (11 genes) - ULG
- Hereditary cancer (gene panel)

### Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

### Related Analytes

- APC regulator of WNT signaling pathway

### Related Gene Panels

- Hereditary Polyposis Panel (11 genes) - ULG
- Hereditary predisposition to cancer (47 genes) - IPG

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