
DISEASE:
Turcot syndrome with polyposis

NAME:	Turcot syndrome with polyposis
DESCRIPTION:	Turcot syndrome with polyposis or Turcot syndrome type 2 is a form of familial adenomatous polyposis, characterized by the concurrence of thousands of colonic adenomatous polyposis or colorectal cancer (CRC) and a primary central nervous system tumor (principally medulloblastoma). It is also associated with pigmented ocular fundus lesions.
ORPHACODE:	99818
XREF(S):	<u>OMIM</u>
CREATED:	03 Aug 2018 - 10:51
CHANGED:	03 Aug 2018 - 10:51

Source URL: <http://gentest.healthdata.be/disease/31>

RELATED CONTENT

Related Genetic Tests

- [Hereditary Polyposis Panel \(11 genes\) - ULG](#)

Related Laboratories

- [Centre de Génétique Humaine - CHU Sart-Tilman](#)

Related Gene Panels

- [Hereditary Polyposis Panel \(11 genes\) - ULG](#)

Source URL: <http://gentest.healthdata.be/disease/31>