Belgian Genetic Tests database

DISEASE:
Mowat-Wilson syndrome due to monosomy 2q22

NAME:	Mowat-Wilson syndrome due to monosomy 2q22
ORPHACODE:	261537
SYNONYMS:	Hirschsprung disease and intellectual disability due to 2q22 microdeletion Hirschsprung disease and intellectual disability due to del(2)(q22) Hirschsprung disease and intellectual disability due to monosomy 2q22 Mowat-Wilson syndrome due to 2q22 microdeletion Mowat-Wilson syndrome due to del(2)q(22)
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	ZEB2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/1076

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