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DOB 10/02/1953 (71 Yrs FEMALE)

Copy to
Clinical Notes WGT LOSS FOR INVESTIGATION

Requested 28/02/2024
Collected 29/02/2024 12:11
Received 29/02/2024 12:13

Methylene Tetrahydrofolate Reductase Gene Mutation

MTHFR c.665C>T	Not Detected
MTHFR c.1286A>C	Homozygous

Comments

Homozygosity for the c.1286A>C variant is not associated with disease. This finding is of little clinical significance (Genet. Med. 15, 153-6,2013).

MTHFR test information:
Detection of the c.665C>T and c.1286A>C variants of the MTHFR gene is performed by melt curve genotyping PCR using the Tib Molbiol MTHFR C667T and MTHFR A1298C Light mix IVD kits.
Other variants in the MTHFR gene are not tested and therefore will not be identified.

NATA ACCREDITATION NO 2178

Reported on 01-03-2024 15:38