## **Ms NICOLE CHESTER**



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Collected: 16-Jun-2016
34 PLATEAU PARADE
BRAY PARK QLD 4500
Lab id: 3433573 UR#:

Date of Birth: 09-Sep-1949

REWIND HEALTH AND WELLNESS UNIT 1/39 WAGAWN STREET TUGUN QLD 4224

## **GENOMIC ASSESSMENTS**

BLOOD SPOT Result Range Units

## MTHFR Gene Mutation, Bloodspot

MethyleneTetraHydroFolate Reductase (MTHFR) Gene Mutation.

MTHFR Gene Mutation (A1298C): Heterozygous for the mutation. MTHFR Gene Mutation (C677T): Heterozygous for the mutation.

## Comment:

The patient has one copy of the MTHFR C677T mutation and one copy of the  $\tt A1298C$  mutation.

This is associated with increased in plasma homocysteine levels.

MethyleneTetraHydroFolate Reductase (MTHFR) is a regulatory enzyme in folate-dependent homocysteine remethylation.

A common polymorphism in the MTHFR gene at position 677 is associated with a thermolabile enzyme with decreased activity. The prevalence of the homozygous mutation ranges from 8-18% in various populations.

Clinically, homozygotes for the mutation have an increased risk of thromboembolism as well as premature vascular disease.

A second mutation (A1298C) has also been described. This mutation is associated with an increased risk of thromboembolism, when only found together with the C677T mutation.

Tests ordered: IMPEI,MTHFRbs