## The association of cytochrome *4F2 rs210622* variation with type II diabetes and its complications

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## Abstract

**Background:** 20-Hydroxyeicosatetraenoic acid (20-HETE) is one of the arachidonic acid metabolites, the omega hydroxylated metabolite, produced by the CYP4F2 enzyme. Moreover, the *CYP 4F2* gene is described as the major gene involved in the synthesis of 20-HETE in humans. The genetic variances in the *CYP 4F2* gene can affect on CYP4F2 enzymatic activity and 20HETE production. Furthermore, pathological diseases such as ischemic cerebrovascular disorders, cardiac ischemia-reperfusion damage, renal diseases, hypertension, and diabetes mellitus have all been linked to higher levels of 20-HETE.

**Aim:** This study will find out the association between the major genetic variant in *CYP4F2* and type II diabetes and its complications among Jordanian patients.

**Methods:** Ninety genomic DNA samples of healthy controls and 90 samples from DM2 patients were genotyped for *CYP4F2* genetic variants. The DNA samples were amplified using polymerase chain reaction (PCR). These products were sequenced using Applied Biosystems Model (ABI3730x1).

**Results:** The frequency of *CYP4F2 rs2108622 C>T* genetic variant was found among healthy subjects and diabetic patients. It is found that there was no significant association between these polymorphisms and the incidence of CVD, CKD, and DL (P>0.05). Also, the baseline triglyceride, low-density lipoprotein, high-density lipoprotein, creatinine, and HbA1c% among Jordanian diabetic and healthy controls subjects were not significantly associated with the *CYP4F2 rs2108622* variant. On the other hand, there was a significant association between the *CYP4F2 rs2108622* genetic variant and the retinopathy among the patients. In addition, the frequency of co-dominant and dominant genotyping models showed statistical differences among all studied groups.

**Conclusion:** The findings of this study indicated that the CYP4F2 SNP of rs2108622 is dominant, and the individual will be two times more likely to develop diabetes. Furthermore, the rs2108622 CYP4F2 SNP was found to be associated with retinopathy in DM2 patients.

Keywords: CYP4F2 gene, Jordanians, genetic variants.