

## **Coagulation Factor II (Prothrombin) G20210A**

This mutation is located in the 3' UTR of the Coagulation Factor II propeptide gene near a putative polyadenylation site and occurs in about 2 % of Caucasians. Biochemically, it is associated with raised levels of circulating levels of prothrombin. Clinically, it is associated with deep venous thromboembolism, recurrent miscarriages and portal vein thrombosis in cirrhotic patients.

### References

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