

Hemophilia A F8 Gene, Intron 22 Inversion Known
Mutation, Whole Blood

Patient ID SA00141426	Patient Name TESTINGRNA, VALIDATION	Birth Date 1999-09-09	Gender M	Age 21
Order Number SA00141426	Client Order Number SA00141426	Ordering Physician CLIENT,CLIENT	Report Notes	
Account Information C7028846 DLMP Rochester		Collected 12 Jan 2021 07:00		

HA F8 Intron 22 Inversion KM, B

F822B Interpretation

1 MCR

These results are consistent with a diagnosis of hemophilia A. This mutation has been identified in patients with severe hemophilia A. Since we have documented the presence of a mutation within the F8 gene in an affected family member, carrier testing for at risk individuals is possible.

A genetic consultation may be of benefit.

MCC Negative: Using a panel of polymorphic markers to compare maternal and fetal specimens, there was no evidence of maternal cell contamination in the fetal specimen.

ADDITIONAL INFORMATION

Genomic DNA is digested with Ksp22 I restriction enzyme, ligated with T4 DNA ligase, and amplified by PCR with primers specific for the F8 intron 22 inversion mutations.

HA F8 Intron 22 Inversion KM, B

MCR

Positive

Reference Value
Not applicable

An intron 22 inversion was detected within the F8 gene.

HA F8 Int22 KM Reason for Referral

MCR

A F8 gene mutation was previously identified in an affected family member. Test for the presence of the familial mutation in the F8 gene.

HA F8 Int22 KM Reviewed By

MCR

BENJAMIN BAILEY

Received: 13 Jan 2021 11:33

Reported: 13 Jan 2021 15:38

Laboratory Notes

- 1 This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

Performing Site Legend

Code	Laboratory	Address	Lab Director	CLIA Certificate
MCR	Mayo Clinic Laboratories - Rochester Main Campus	200 First Street SW, Rochester, MN 55905	William G. Morice M.D. Ph.D	24D0404292