

2895650160 bfr11

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2895650160 BFR11: Unlocking Your BFR11 Potential

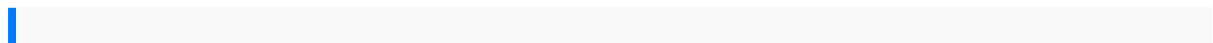
Q: What is 2895650160 BFR11? A: 2895650160 BFR11 is a unique identifier assigned to the BFR11 gene, which provides instructions for creating a protein involved in regulating blood clotting.

Q: What is the function of BFR11? A: BFR11 plays a critical role in the coagulation cascade, a complex process that prevents excessive bleeding. It inhibits the activity of activated Factor IX (FIXa), a key protein involved in the formation of blood clots.

Q: Why is 2895650160 BFR11 important? A: Mutations in the BFR11 gene can lead to bleeding disorders such as hemophilia B, a rare condition characterized by prolonged bleeding due to a deficiency in FIX. Understanding the genetics of BFR11 helps in diagnosis, treatment planning, and carrier screening.

Q: How is 2895650160 BFR11 tested? A: Genetic testing can identify mutations in the BFR11 gene. This testing typically involves collecting a blood sample and analyzing the DNA for variations in the BFR11 sequence.

Q: What are the implications of a positive test result? A: A positive test result for a mutation in 2895650160 BFR11 indicates that an individual is either affected with or a carrier for hemophilia B. The severity of the bleeding disorder depends on the specific mutation and its impact on BFR11 function. Genetic counseling and medical management are recommended to optimize care and prevent complications.



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