NKX6-2

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Phenotypic Variance & Individual Differences

NKX6-2-related disorder (SPAX8) is an exceptionally rare condition characterised by a **massive variance in patient phenotypes**. Because the genetic mutations and their physical expressions differ drastically from person to person, no two patients will have the exact same clinical journey. The symptoms, progression, milestone achievements, and therapeutic needs outlined in these FAQs represent a broad spectrum of possibilities and **cannot be relied upon** to predict the specific outcome, timeline, or medical path of any individual child or patient.

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