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Clinical and molecular characterization of a boy with intellectual disability, facial dysmorphism, minor digital anomalies and a complex *IL1RAPL1* intragenic rearrangement

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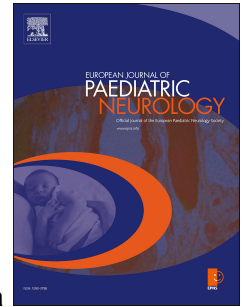
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