

Genetic Disorder Prediction Report

Patient Age: 8

Gender: Female

Predicted Disorder: Genetic Disorder

Subclass: Leigh syndrome

Date: 27-01-2026

Description:

Leigh syndrome is a rare inherited neurological disorder that primarily affects infants and children. It is characterized by progressive brain damage.

Common symptoms include developmental delay, muscle weakness, breathing difficulties, and neurological abnormalities.

Diagnosis involves clinical evaluation, genetic testing, and imaging. There is currently no cure, and treatment focuses on managing symptoms.

Regular follow-up with neurologists and genetic specialists is recommended.