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A Case of Congenital Hereditary Metabolic Disease-urea Cycle Metabolic Disorder

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ABSTRACT

Congenital metabolic diseases are caused by the lack or abnormality of a certain enzyme or its cofactors, which can cause accumulation or lack of specific metabolites. The best outcome for children with inborn errors of metabolism (IEM) depends on recognizing the symptoms and signs of metabolic diseases, promptly assessing them, and transferring them to a medical center familiar with the evaluation and treatment of related diseases. Delayed diagnosis may lead to Acute metabolic decompensation, progressive nerve damage, or death.

EXTERNAL LINK

<https://www.creative-proteomics.com/services/urea-cycle-metabolism-analysis-service.htm>

ATTACHMENTS

A Case of Congenital
Hereditary Metabolic
Disease-urea Cycle
Metabolic Disorder.docx

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