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Congenital disorder of glycosylation caused by the mutation of ATP6AP1 gene (c.1036G>A) in a Chinese infant : A case report

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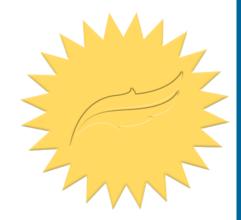
for clarity, consistency, and correctness according to the requirements and guidelines specified by the client.

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