

Clinical and genetic analysis of multi-system pseudohypoaldosteronism type 1 caused by a novel splice site mutation of the beta subunit gene of epithelial sodium channel (ENaC)

Sicui Hu¹, hongxiu yang¹, cheng Li¹, qiannan jiang¹, lingyan qiao¹, conghui hu¹, and Tang Li¹

¹Qingdao Women and Children's Hospital

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Abstract

This article aims to provide a comprehensive review of the clinical features and genetics basis of multi-system pseudohypoaldosteronism type 1 caused by SCNN1B gene mutations.

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