

**Name of Journal:** *World Journal of Clinical Cases*

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**Manuscript Type:** CASE REPORT

**Autism with dysphasia accompanied by mental retardation caused by FOXP1 exon deletion: A case report**

Autism with dysphasia accompanied by mental retardation

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What is foxp1-related ID syndrome? ▾

How many patients have been reported with FOXP1? ▾

Is fosp1 a spectrum disorder? ▾

## FOXP1

Protein



Forkhead box protein P1 is a protein that in humans is encoded by the FOXP1 gene. FOXP1 is necessary for the proper development of the brain, heart, and lung in mammals. It is a member of the large FOX family of transcription factors.

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