

JP (aged 15yr 9m) was referred to the paediatric endocrine team with a diagnosis of thyrotoxicosis (T3 41.9, TSH <0.1)

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During the consultation it was noted that JP's pubertal progression was markedly delayed. Further investigations were undertaken, the results of which demonstrated primary ovarian failure (LH 16.9ui/L FSH 71.5iu/L) Genetics report a few weeks later confirmed diagnosis of Turner Syndrome.

This presentation will examine the care of JP since referral to the paediatric endocrine team 10 months ago. Following a very brief overview of the investigations, diagnosis and treatments given, it will then open up for discussion on the role of the endocrine specialist nurse during this turbulent time for the family.