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# 46,XX gonadal dysgenesis

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. [46,XX gonadal dysgenesis](#). ORPHA:243*

46,XX gonadal dysgenesis (46,XX GD) is a primary ovarian defect leading to premature ovarian failure (POF; see this term) in otherwise normal 46,XX females as a result of failure of the gonads to develop or due to resistance to gonadotrophin stimulation.