Does your patient have Prader-Willi syndrome?

The presence of **all** the below findings at the age indicated is sufficient to justify investigation for Prader-Willi syndrome.

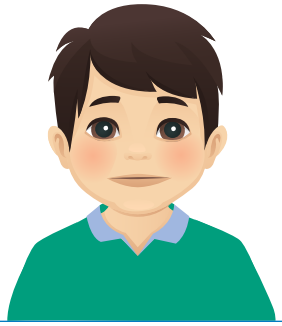
Find out more at: www.ipwso.org

Neonatal period to two years

- Hypotonia with poor appetite and suck in the neonatal period
- Global developmental delay

Age two to six years

- Hypotonia with history of poor suck
- Global developmental delay

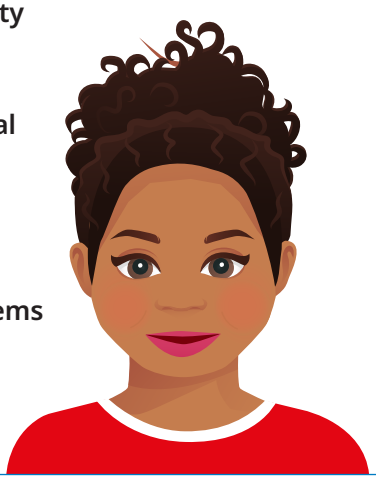


Age six to 12 years

- History of hypotonia with poor suck (hypotonia often persists)
- Global developmental delay
- Excessive eating with central obesity if uncontrolled externally

Age 13 years to adulthood

- Cognitive impairment, usually mild intellectual disability
- Excessive eating and hyperphagia with central obesity if uncontrolled externally
- Hypogonadism and/or typical behaviour problems



How can you diagnose*? • Conduct a DNA methylation test • Discuss with a clinical geneticist

Scan me

We can help and advise, please contact **office@ipwso.org** or visit **www.ipwso.org** for more information

IPWSO
International Prader-Willi Syndrome Organisation

*IPWSO offers free diagnostic testing in some cases

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