

15) Prader Willi Syndrome (PWS)

- Clinical features in infancy
 - √ Hypotonia
 - √ feeding problems
- Clinical features in childhood
 - √ Hyperphagia
 - ✓ Central obesity
 - √ Hypogonadism
 - ✓ Short stature
 - ✓ Small hand and feet
 - ✓ Mild intellectual disability
 - ✓ Almond-shaped eyes
 - ✓ Behavioral disorder







Mode of inheritance?

- ✓ Sporadic
- ✓ Paternal derived deletion 15q11–13 (80%)
- ✓ Maternal UPD (20%)

It is the most common syndromic form of obesity

- Which medication had FDA approval to be used with PWS?
 - ✓ Growth hormone therapy
- Why they develop short stature, central obesity
 - & hypogonadism?
 - ✓ Secondary to Hypothalamic and pituitary dysfunction

Management Of Prader Willi Syndrome

Xiang Xie

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