Prader-Willi Syndrome is:
■ A non-hereditary birth defect resulting from a disorder of chromosome 15
■ A serious, life-long, and life-threatening medical condition
■ Occurs in 1:12,000 to 1:15,000 births; both sexes, all races
■ Characterized by:
  - Hypotonia (low tone)
  - Hypogonadism (underdeveloped sex organs)
  - Hyperphagia (uncontrollable hunger)
  - Cognitive impairment
  - Challenging behaviors
■ One of the most common conditions seen in genetic clinics
■ The most common genetic cause of life-threatening obesity

A major medical concern is morbid obesity, however with early diagnosis and early intervention, many children can maintain a healthy weight.

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We hope you find these materials helpful and that you consider a donation to PWSA | USA to assist in developing more good work(s) like this. Please see our website, www.pwsausa.org.

Quality of Life Issues
General health is usually good in individuals with PWS. If weight is controlled, life expectancy may be normal, and the individual’s health and functioning can be maximized. The constant need for food restriction and behavior management may be stressful for family members. PWSA | USA can provide information and support.

Adolescents and adults with PWS can function well in group and supported living programs, if the necessary diet control and structured environment are provided.

To date, no medication or surgical intervention has been found that would eliminate the need for strict dieting and supervision around food. Bariatric procedures do not address the central lack of satiety and put the individual at risk for complications.

Studies show improvement in linear growth, fat mass, motor strength, respiratory drive, and bone density with the use of growth hormone in PWS. Precautions need to be taken prior to starting treatment including polysomnography, checking adrenal gland function, and following IGf1 levels.

Cause and Diagnosis of PWS
■ PWS occurs from three main genetic errors. Approximately 70% of cases have a non-inherited deletion in the paternally contributed chromosome 15. Approximately 25% have maternal uniparental disomy (UPD) - two maternal chromosome 15s and no paternal chromosome 15. Also, 2-5% have an error in the “imprinting” process that renders the paternal contribution nonfunctional; rarely, these imprinting defects may be inherited.

■ Diagnostic testing: Individuals who have a number of the clinical findings should be referred for genetic testing. DNA methylation analysis confirms diagnosis of PWS. FISH and DNA techniques can identify the specific genetic cause and associated recurrence risk. Patients who had negative or inconclusive tests with older techniques should be retested.

■ Recurrence risk: Recurrence is significant only for rare cases with imprinting mutations, translocations, or inversions. All families should receive genetic counseling.

Medical Overview
A Diagnosis and Reference Guide on Prader-Willi Syndrome for Physicians and Other Health Professionals
Life Threatening Medical Concerns

- Anesthesia, medication reactions. Unusual reactions to standard dosages of medications and anesthetic agents may occur because of metabolic differences and obesity seen in PWS. A narrow airway may be present. Use extreme caution in giving medications that may cause sedation; prolonged and exaggerated responses have been reported. Several genes for GABA receptor subunits are located in the PWS chromosome region and are missing in patients with the deletion. This decrease in GABA receptors in PWS could alter the response to GABA receptor agonist sedative agents (propofol, benzodiazepines).

- High pain threshold. Lack of typical pain signals is common and may mask the presence of infection or injury. Someone with PWS may have difficulty locating pain or not complain of pain until infection is severe. Parent/caregiver reports of subtle changes in condition or behavior should be investigated for medical cause.

- Respiratory concerns. Risk may be increased for respiratory difficulties. Obesity, hypotonia, weak chest muscles, and sleep apnea are among possible complicating factors. Sleep studies for medical cause.

- Lack of vomiting. Vomiting rarely occurs. Emetesis may be ineffective, and repeated doses may cause toxicity. This characteristic is of particular concern in light of hyperphagia and the possible ingestion of uncooked, spoiled, or otherwise unhealthful food items. The presence of vomiting may signal a life-threatening illness.

- Body temperature abnormalities. Idiopathic hyper- and hypothermia have been reported. Hyperthermia may occur during minor illness and in procedures requiring anesthesia. Fever may be absent despite serious infection.

- Severe gastric illness. Abdominal distension or bloating, pain and vomiting may be signs of life-threatening gastric inflammation or necrosis, more common in PWS than in the general population. Rather than localized pain, there may be a general feeling of unwellness. If an individual with PWS has these symptoms, close observation is needed. A CAT scan of the abdomen and/or endoscopy may be necessary to determine degree of the problem and possible need for emergency surgery.

- Central adrenal insufficiency. Studies suggest an increased incidence of CAI in individuals with PWS. Measurement of cortisol levels during a significant illness and supplementation of cortisol may be indicated.

- Skin lesions and bruises. Skin picking is common in PWS, causing open sores. In some situations, skin and rectal picking can be severe. Individuals with PWS also tend to bruise easily. Appearance of such wounds and bruises may wrongly lead to suspicion of physical abuse.

- Hyperphagia (excessive appetite). Insatiable appetite may lead to life-threatening weight gain, which can be very rapid and occur even on a low-calorie diet. Individuals with PWS must be supervised at all times in all settings where food is accessible. Those who have normal weight have achieved this because of strict external control of their diet and food intake. Water intoxication has occurred in relation to use of certain medications with anti-diuretic effects, as well as from excess fluid intake alone, producing lower electrolytes.

- Obesity-related problems include hypoventilation, hypertension, right-sided heart failure, stasis ulcers, cellulites, and skin problems in fat folds.

Potential Characteristics

Any infant with hypotonia should be tested for PWS. The following common characteristics raise suspicion of a diagnosis of PWS.

- Decreased fetal movement, infantile lethargy, weak cry
- Feeding problems and poor weight gain in infancy
- Insatiable appetite may include soft tooth enamel, thick sticky saliva, poor oral hygiene, teeth grinding, and infrequently rumination. Special toothbrushes can improve hygiene. Products to increase saliva flow are helpful.
- Speech articulation defects and dyspraxia
- Strabismus — esotropia is common; requires early intervention, possible surgery
- Scoliosis — may occur unusually early; may be difficult to detect without X-ray, kyphosis is also common in teens and adults

Resources for Health Care Providers

“Growth Hormone and Prader-Willi Syndrome, 2nd Edition” and the book “Management of Prader-Willi Syndrome” are available from PWSA | USA at www.pwsusa.org, as are many other publications for professionals and parents. Medical crisis support for professionals and parents is available at (941) 312-0400.

In the Event of Death

Reporting of Deaths

The Prader-Willi Syndrome Association | USA has created a research database of reported deaths of individuals with PWS. In the event of death of someone with PWS, please contact PWSA | USA: (941) 312-0400.

Organ Donation for Research

When a child or adult with PWS dies, the family may wish to consider donation of organs for research. Prompt action is essential for tissue preservation. Families are advised to contact the Brain and Tissue Bank directly: 1-800-847-1539 (Maryland).