Prader-Willi Syndrome: Clinical Aspects

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Prader-Willi Syndrome (PWS) is a complex multisystem genetic disorder that shows great variability, with changing clinical features during a patient’s life. The syndrome is due to the loss of expression of several genes encoded on the proximal long arm of chromosome 15 (15q11.2–q13). The complex phenotype is most probably caused by a hypothalamic dysfunction that is responsible for hormonal dysfunctions and for absence of the sense of satiety. For this reason a Prader-Willi (PW) child develops hyperphagia during the initial stage of infancy that can lead to obesity and its complications. During infancy many PW child display a range of behavioural problems that become more noticeable in adolescence and adulthood and interfere mostly with quality of life. Early diagnosis of PWS is important for effective long-term management, and a precocious multidisciplinary approach is fundamental to improve quality of life, prevent complications, and prolong life expectancy.

1. Introduction

PWS (OMIM number 176270) is a complex multisystem genetic disorder originally described in 1956 by three Swiss doctors, Prader et al. [1]. PWS was the first recognized disorder related to genomic imprinting in humans [2] and is caused by the lack of expression of paternally inherited imprinted genes on chromosome 15q11–q13. In several studied populations prevalence has been estimated to be 1/15,000–1/25,000.

The syndrome shows great variability, with changing clinical features during a patient’s life. A newborn might suffer from severe hypotonia with feeding problems and global developmental delay. During infancy these characteristics impede the acquisition of gross motor and language milestones. A PW child develops hyperphagia during the initial stage of infancy that can lead to precocious obesity if left uncontrolled. This is most probably caused by a hypothalamic dysfunction, which impedes the sense of satiety. This hypothalamic dysfunction is also responsible for growth-hormone (GH) and thyroid-stimulating hormone (TSH) deficiencies, central adrenal insufficiency, and hypogonadism. During infancy, the PW child shows a characteristic problematic behavioral pattern, which has been reported to worsen with age. Patients sometimes present psychosis.

Early diagnosis of PWS is important for effective long-term management. In fact, the multidimensional problems of patients with PWS cannot be treated with a single intervention. A precocious multidisciplinary approach is fundamental to improve quality of life, prevent complications, and prolong life expectancy.

2. Diagnosis

PWS diagnosis is based on specific clinical features, and it is confirmed by genetic testing.

2.1. Clinical Features. Although diagnostic molecular testing for PWS is currently available, the clinical identification of patients remains a challenge as many features of PWS are nonspecific while others evolve over time or can be subtle [3].
2.2. Genetic Findings. PWS, together with Angelman syndrome (AS), represents perhaps the best example of genomic imprinting in humans. Genomic imprinting is an epigenetic process by which the male and the female germ lines confer specific marks (imprints) onto certain gene regions. Probably <1% of our genes are imprinted with an expression pattern determined by the parent of origin [6–8].

The PWS region is found in a 5-6 Mb genomic region on the proximal long arm of chromosome 15 (15q11.2–q13; Figure 1). The complex phenotype is due to the loss of expression of several paternally genes on chromosome 15q11.2–q13 [9, 10].

Three main molecular mechanisms result in PWS.

1. **Paternal microdeletion** is responsible for 75–80% of cases.
2. **Maternal uniparental disomy (UPD)** is responsible for 20–25% of cases.
3. **Imprinting defect (ID)** is responsible for 1–3% of cases.
4. **Other defects** is such as balanced and unbalanced translocations, which, together with ID, are responsible for the majority of familial cases.

2.2.1. Genotype-Phenotype Correlation. Genotype-phenotype correlation is not possible, because no features are known to occur exclusively in individuals with one of the genetic classes. However, some studies identify significant statistical differences between the two largest genetic subtypes (deletion and UPD). For example, postterm delivery is more common in UPD patients. They are less likely to show hypopigmentation [11, 12], the typical characteristic facial appearance [11, 13], or possess jigsaw-puzzle skills [14]. In most studies, patients with UPD have a somewhat higher verbal IQ and milder behavioral problems [15–17]. However, psychosis [18] and autism spectrum disorder [19, 20] occur with significantly greater frequency among those with UPD. Individuals with the slightly larger, type-1 deletions (BP1–BP3) show poorer adaptive behavior, and lower intellectual ability and academic achievement than those with type-2 deletions (BP2-BP3) deletions [21, 22]. Two other studies found fewer clinically significant differences between individuals with these two deletion types [23, 24].

2.2.2. Genetic Counseling. Knowing the specific genetic etiology in individuals with PWS is essential for the appropriate genetic counseling of affected families. The risk of PWS recurrence in families with affected children is usually less than 1%, except for inherited mutation in the imprinting center (up to 50%), and translocation inherited mutation with break point in the 15q11.2–q13 region (up to 25%).

2.2.3. PW-Like Patients. The term “PW-like” is used to indicate a patient with clinical features that are very similar to PW, but without the confirmation of a classical genetic subtype that can cause the syndrome.

The literature reports many PW-like cases, but also many reports of the association between these patients and some particular generic anomalies or syndromes.

The most frequently reported are the associations between PW-like and

(i) Fragile X syndrome [25, 26],
(ii) Klinefelter syndrome [27, 28],
(iii) interstitial deletion in 6q [29],
(iv) subtelomeric deletion 1p36 [30, 31].

2.3. Genetic Tests. Various types of genetic test can be used for PWS diagnosis and to characterize the different subtypes (Figure 2) [32].

Methylation-specific-multiplex ligation-dependent probe amplification analysis (MS-MLPA) combines both DNA methylation analysis and dosing analysis across the PWS region and can be considered the gold standard for PWS diagnosis in 99% of cases. This test shows the absence of the paternal allele using a methylation analysis to measure the amplitude of deletion (type 1 or type 2) and exclude suspect UPD or imprinting defects.

The karyotype is useful in association with MS-MLPA to point out balanced translocation whose break point might be in the critical PWS region (15q11–q13).

Microsatellite analysis is necessary only if MS-MLPA does not show deletions in the critical PWS region, but just paternal allele absence at methylation analysis. It shows UPD (both alleles belonging to the mother) and also specifies if there is a heterodisomy (two mother alleles that are different from each other) or isodisomy (the same maternal allele in a double copy). If this analysis shows a biparental pattern, there is indication of mutation or microdeletions of the imprinting center.

3. Clinical Presentation

PWS is a multigenic pathology that shows great clinical variability that shows great clinical variability. Features change from patient to patient and even during the lifetime of the individual patient.

We have designated 4 age classes that can help the physician to outline ideal lifelong PW patient followup taking
into consideration all the problems these patients might have to face.

3.1. From Birth to 3 Years. Pregnancy is generally normal, but some mothers may report decreased fetal activity, and newborns are often found in the breech position at time of delivery. Premature delivery may occur, and newborns that are adequate for gestational age frequently have low weight and length at birth.

The majority of newborns with PWS present marked neonatal axial hypotonia (babies are described as “floppy”); this is associated with lethargy, decreased movement, weak crying, and poor reflexes, including poor sucking, often resulting in failure to thrive [33, 34].

The baby can present dysmorphic characteristics, such as narrow bifrontal diameter, dolichocephaly, almond-shaped eyes, downturned angles of the mouth with abundant and thick saliva, and small hands and feet. These are less pronounced at birth but can become more evident with age.

Newborns generally present the clinical signs of hypogonadism. In males, the penis may be small; more characteristic is a hypoplastic scrotum that is small, poorly rugated, and poorly pigmented. Unilateral or bilateral cryptorchidism is present in 80–90% of males [35, 36]. In females, genital hypoplasia is often overlooked; however, the clitoris and labia, especially the labia minora, are generally small from birth.

Hypotonia and hypogonadism are the first manifestations of a primitive hypothalamic alteration, which many studies indicate to be at the base of PWS. This central deficiency leads to many manifestations, in particular a pituitary hormonal deficit (GH, TSH, central adrenal insufficiency [37]), satiety alteration, sleep disturbances, and a tendency for dysthermoregulation.

Although hypotonia slowly evolves over time, gross motor and language milestones are delayed. Early milestones are reached on average at double the normal age (e.g., sitting at 12 months, walking at 24 months, and saying words at 2 years) [38].

3.2. From 3 to 10 Years. During preschool age, PW patients develop a food obsession; children become overweight as a consequence of an insatiable appetite and compulsive eating, which can lead to morbid obesity in adolescence and adulthood if not kept under control.

PWS eating habits are complex and multifactorial. They are thought to be associated with abnormalities in the hypothalamic circuitry or the peripheral satiety signals [39, 40].

PWS individuals show differences in various gut hormones, including high levels of obestatin (an anorexigenic hormone) in infancy, with markedly elevated levels of ghrelin (an orexigenic hormone) in childhood and adulthood. The structural brain abnormalities present in these individuals might also contribute to appetite aberrations [41]. Functional MRI studies indicate that these individuals assign a high reward value to food with increased activation of the limbic and paralimbic areas of the brain that drive eating.
behavior, even after meal, showing that the brain influencing appetite in this syndrome.

In 2011, McAllister et al. [42] reviewed eating behavior in PWS. They concluded that a genetic abnormality might be the cause of fetal malnourishment or fetal starvation, leading to neonatal failure to thrive and also causing feeding problems. Ghrelin may be involved in the instigation of the binging and hyperphagic stage, and later development of atypical reward circuitry in response to food might be the result of altered pathways generated in the early binging stages in childhood, combined with insensitivity to satiation cues, such as leptin levels. Examining reward pathways by neural imaging in response to food in younger children, for example, may elucidate the development of the hyperresponsive circuitry [42].

Growth restriction is a frequently observed sequel of PWS; approximately 90% of affected individuals are short in stature, probably resulting from GH deficiency [37].

During the first 6 years of life, children with PWS often do not achieve normal levels of cognitive, motor, and language development. Indeed, according to one study, these individuals have a below-average IQ of about 70 [43–45]. A review of cognitive ability among 575 affected individuals confirms this, showing that just 5% of patients had a normal IQ (i.e., >85). Borderline mental retardation was observed in 28% of patients, while 34%, 27%, and 5%, respectively, were mildly, moderately, or severely mentally retarded [45].

During this period, many PW sufferers display a range of behavioral problems that include both excessive appetite and lack of food selectivity. There is also a high incidence of stubbornness, verbal perseverance, skin picking, and temper tantrums.

Despite hypogonadism generally causing incomplete, delayed, and sometimes disordered pubertal development, in approximately 15–20% of patients of both sexes, premature adrenarche or precocious puberty often occurs [35, 46].

3.3. From 10 to 18 Years. During adolescence, behavioral problems and hyperphagia become more noticeable.

In experimental settings, PWS individuals have been seen to consume around three-to-six times more than the normal caloric intake at a given meal [47, 48]. Overeating can lead to stomach rupture [49], and PWS individuals have also been known to steal and hoard food. These individuals have also been reported to eat inappropriate food, such as uncooked chicken, or even to eat nonfood items.

Central obesity results from the combination of uncontrolled food intake, a low metabolic rate, and a decreased activity level (resulting in a decreased total caloric requirement). Obesity-related complications appear, such as cardiorespiratory insufficiency, obstructive sleep apnea, thrombophlebitis, and chronic leg edema, and are the major causes of morbidity and mortality [50].

The severity of behavioral problems increases with age and body mass index and can then diminish in older adults. Psychosis is evident by young adulthood in at least 5–10% of individuals [51–53]. Behavioral and psychiatric problems interfere mostly with quality of life in adolescence and adulthood.

Hypogonadism causes incomplete, delayed, and sometimes disordered pubertal development. Primary amenorrhea or oligomenorrhea are present in females. Infertility is the rule in both sexes although a few instances of reproduction in females have been reported and presented [54, 55]. Although hypogonadism in PWS has long been believed to be entirely hypothalamic, resulting in low gonadotropins and subsequent low gonadal hormones, recent studies have suggested a combination of hypothalamic and primary gonadal deficiencies [56–58].

3.4. The Adult. The quality of life of adults with PWS is largely conditioned by the degree of obesity, the presence of its complications, and behavioral problems.
4. Management

Patient management has to be tailored, due to the clinical variability between patients, but also to variability in the same patient throughout this or her lifetime. Therapeutic decisions and clinical followup need to consider each problem that a patient might have to face. A multidisciplinary team, consisting of neonatologists, pediatricians, endocrinologists, orthopedic surgeons, psychologists, psychiatrists, physiotherapists and urologists, has to deal with all of the patient’s medical and psychological problems.

4.1. Nutrition. The maintenance of adequate and appropriate nutrition is fundamental to the treatment of people with PWS at every age.

A correct approach should consider the two distinct nutritional stages of every PW patient:

(i) stage 1: poor feeding and hypotonia, which can often cause a failure to thrive;
(ii) stage 2: “hyperphagia leading to obesity” [5, 40, 59].

4.1.1. Feeding Support. During the first two months of life, most PWS infants are unable to suck an adequate quantity of milk from the breast or the feeding bottle, and they have to be fed by gavage. The use of a gastrostomy tube (generally using a button-style device) can be avoided in most cases, but if, after considering the risks and benefits of both approaches, a decision to use a gastrostomy tube is made, the device should be promptly removed when no longer needed.

Milk requirement is that of other infants of the same age and weight. Children should not be given simple sugars for a “sweet taste.”

Infants may require feeding support for several months. Caloric needs may sometimes be somewhat reduced in infants with PWS, who typically do not spontaneously demand feeding. The infant’s diet must therefore be adjusted to maintain appropriate weight gain as determined by frequent weight checks. Increased caloric density can be helpful.

4.1.2. Dietary Control. Incorrect eating is one of the most serious disorders affecting the lives of children and adults with PWS. Hyperphagia is a serious chronic problem for children with PWS, together with their families, and it can severely limit independence in adult life due to the risk of life-threatening obesity.

Controlling food-related behavior is complex, aiming to limit the child’s access to food, reduce exposure that can cause the child to think about food, and promote a daily routine that helps obtain good weight control. Relatives and friends also have to understand that “sneaking” food to the child with PWS is not a demonstration of affection as, on the contrary, it undermines the child’s nutritional regimen and sense of wellbeing.

When hyperphagia occurs the caloric intake needs to be reduced. The diet should have a balanced distribution and be rich in fiber, and the caloric intake should be about 75–80% that of a healthy child of the same age.

During adulthood, the caloric intake should be below 1000–1200 kcal/day to maintain stable weight structure, or between 800 and 1000 kcal/day to lose weight. Dietary restriction should anyway be balanced, and complex carbohydrates should be preferred.

Pharmacological treatment, including available anorexigenic agents, has not been of benefit in treating hyperphagia although there are some published placebocontrolled studies [40, 60]. The potential benefits of newer agents, such as endocannabinoid antagonists, are still under examination in PWS. Recent concerns regarding psychiatric side effects need careful monitoring in these patients.

Restrictive bariatric surgery, such as gastric banding or bypass, has not been seen to reduce hyperphagia or achieve long-term weight reduction and are associated with unacceptable morbidity and mortality [61, 62]. Whereas some of the reports using bilipancreatic diversion reported successful weight loss, there have been frequent complications from the resulting intestinal malabsorption. Bariatric surgery should only be considered when the patient’s excessive weight becomes life threatening. Strict diet control and postsurgical followup are mandatory.

4.2. Motor Program. The motor program has to consider the various necessities during a patient’s life; in newborns the goal is to improve axial hypotonia, while in childhood and adulthood the patient’s physical, socials and metabolic condition needs to be focused.

4.2.1. Physiotherapy. The newborn is characterized by a variable grade of hypotonia that influences many aspects of normal development (delayed gross motor milestones) and growth (feeding difficulties).

Even when hypotonia evolves slowly, we suggest the prompt introduction of personal training programs. These programs must be supervised by physiotherapists and maintained by parents. They help counteract the hypotonic PWS infant’s difficulties in overcoming gravity during early life. This is a particularly sensitive period for motor development and skills acquisition and might have consequences regarding cognitive and social development.

4.2.2. Physical Activity. Physical activity and sports are a fundamental therapy for PWS patients.

During childhood, physical activity

(i) improves physical functions,
(ii) promotes socialization,
(iii) helps improve caloric expenditure, together with diet,
(iv) is one of the best ways to limit access to food.

We suggest regular daily physical activity of around 30 minutes. Any kind of sport is possible, and parents should consider the abilities and tastes of their children when choosing.

Although hypotonia improves with age, it persists into adulthood, together with reduced muscle mass. Regular exercise is therefore an important part of everyday life.
In a recent study, Vismara et al. provided an effective and simple home-based training program representing a continuum of the rehabilitation process outside the hospital, which is a crucial issue in chronic conditions. In fact, following six months of daily activity patients were seen to improve their physical function [63].

4.3. Endocrinological Aspects. As already stated, the hypothalamic dysfunction at the base of PWS contributes to the development of multiple endocrine disorders, such as adrenal insufficiency, GH deficiency, LH/FS disorder, and thyroid dysfunction.

4.3.1. Adrenal Insufficiency. Several reports show a mortality rate in PWS estimated at 3% yearly [64, 65]. Disturbances in the hypothalamus-hypophysis-adrenal (HHA) axis are thought to be responsible for these events or, at least, to represent concurrent factors consistent with an inadequate or late response during infection or relevant dehydration episodes. This hypothesis is supported by pathological findings. Adrenal atrophy has been documented autopsically in a number of such cases [66], and small volumes of the hypothalamic paraventricular nuclei with decreased cell number were demonstrated in PWS adults [67]. Although these aspects have been described in PWS, the genetic basis of putative central adrenal insufficiency (CAI) is far from being unraveled, as the molecular mechanisms leading to PWS phenotype are still largely unknown.

From a functional viewpoint, some studies reveal a differing prevalence for this problem. De Lind van Wijngaarden et al. using the metyrapone test found that the hypophyseal response to adrenalin inhibition was insufficient in 15 out of 25 (60%) PWS patients [68]. Two recent reports on 41 and 57 PWS patients found no cases of CAI by employing a low-dose (LDTST) and a standard ACTH stimulation test (250 μg), respectively [69, 70]. The Study Group for Genetic Obesity of the Italian Society of Pediatric Endocrinology and Diabetology (SIEDP/ESPED) designed a study that confirms that clinically relevant CAI in pediatric PWS patients is rare (14.3% to LDTST), with one third of them (4.8%) also having suboptimal response to a second test. The authors suggest [71].

According to the literature, the administration of glucocorticoids during episodes of moderate/severe stress is recommended (hydrocortisone at 30–70 mg/m²/day divided into 3 doses Continuous replacement should be limited to cases with clinical signs of adrenal insufficiency, as for other nonsyndromic forms of CAI.

4.3.2. Growth Hormone (GH) Deficiency. There are many data indicating reduced GH secretion in PWS patients. Low peak GH response to stimulation tests, decreased spontaneous GH secretion, and low serum IGF-1 levels have been documented in at least 15 studies involving about 300 affected children [37]. Clinical features of the condition also support the presence of GHD in PWS. Both PWS and GHD are characterized by short stature, obesity with extra fat deposits over the abdomen, abnormal body composition with reduced muscle mass and decreased bone density, and, in some patients, retarded bone age [4, 72]. Patients with PWS are therefore GH deficient although the degree of GH deficiency may vary from mild to severe.

Guidelines for GH therapy in PWS children have been drawn up by the Italian Drug Agency (AIFA) note number 39 and do not depend on the presence of GH deficiency.

In accordance with the literature [50], we suggest beginning GH therapy during the first year of life after performing the following:

(i) polysomnography and ENT evaluation,
(ii) fasting glucose and OGTT (glucose 1, 75 g/Kg, to maximum 75 g),
(iii) blood sample for: IGF1, fT4, TSH,
(iv) spine X-ray,
(v) cardiologic evaluation with echocardiography.

The recommended dosage is 0, 01-0, and 03 mg/Kg/day, adjusted on IGF1 levels, which should not exceed 2SDS. It is preferable to start with a dose corresponding to 1/3 of the minimal dosage. GH therapy should not be started in the presence of obstructive sleep apnea syndrome (OSAS), adenotonsillar hypertrophy, severe obesity, glucidic intolerance, and/or unstable scoliosis. Interruption of therapy should be evaluated if scoliosis, glucose intolerance, or OSAS deteriorate.

When the individual stops growing in height, evaluation of the GH secretory pattern is required through ITT, or rather using the GnRH-arginine test that has greater tolerance. For adolescents in therapy, evaluation of the GH secretory pattern must be preceded by a therapeutic washout of at least 1-2 months. In patients with GH deficit (lower than 3 μg/L) a therapeutic pattern not involving weight is recommended, starting with a standard dose of 0.2 mg/day, which can be modified based on IGF1 levels.

During GH therapy, it is fundamental to carry out periodic monitoring of

(i) polysomnography and ENT evaluation at 6 weeks–3 months–6 months after the start, and whenever necessary,
(ii) fasting glucose and OGTT, IGF1, fT4, TSH: 1 month after the start, and whenever necessary,
(iii) orthopedic evaluation and spine X-ray whenever necessary,
(iv) cardiovascular function evaluation every 6 months, or whenever necessary.

In children with PWS, the aims of GH treatment are to improve growth during childhood, adult height, and body composition. There is much evidence that this therapy has multiple beneficial effects on growth and body composition. In particular, it decreases fat mass and increases muscle mass, and it may have a beneficial effect on weight gain, and possibly appetite, in individuals with PWS [37, 73]. Infants with PWS treated with GH therapy show improved head circumference, height, BMI, body composition (with
improvement of lean muscle mass and delay of fat tissue accumulation), body proportions, acquisition of gross motor skills, language acquisition, and cognitive scores. Several studies have documented the safety and efficacy of GH treatment in adults with PWS on body composition and quality of life [74, 75].

4.3.3. Disorder of Gonadotropins. Hypothalamic dysfunction can cause LHRH-LH/FSH axis disorders. In general, the patient shows hypogonadism from birth, but in some PW children precocious adrenarche or early puberty can occur.

(a) Hypogonadism: hypogonadism is a consistent feature in both males and females with PWS, and hypogenitalism is present, even at birth. There is increasing evidence to implicate both central and peripheral origins for hypogonadism, at least in males [76].

At birth, both sexes can have clinical signs of hypogonadism: females show genital hypoplasia (clitoris and labia minora), while males have cryptorchidism, scrotal hypoplasia, and small testicular volume. Cryptorchidism is present in over 80% of boys from birth [35, 36]. At present there is no consensus in the literature as to the best treatment to use. Our experience suggests human chorionic gonadotropins (HCG) therapy from 6 to 12 months with a dosage of 250 U, while from 1 to 5 years a dosage of 500 U should be used twice a week for 6 weeks.

Before and during therapy regular clinical evaluation, testosterone dosage and testicular ultrasound are necessary. When therapy does not have positive results, surgical correction needs to be considered before 2 years of life, or as soon as possible when the child is older. During adolescence patients can show delayed or incomplete pubertal development, and it is therefore necessary to consider hormonal treatment for the induction, promotion, or maintenance of puberty. Before any therapy is carried out, the following need to be considered:

(i) dosage of LH, FSH, estradiol (in females), and testosterone (in males) both basal and after GnRH
(ii) pelvic ultrasound in females,
(iii) testicular ultrasound,
(iv) dual-energy X-ray photon absorptiometry (DEXA),
(v) evaluation of thrombophilic status in females.

For the induction and maintenance of puberty there is no consensus regarding best therapy. We suggest testosterone in males (25 mg once a month, which after 3 months can be increased to 50 mg once a month if well tolerated), and estrogen (incremental dosage up to complete development) and progesterone in females (in general after around 24 months from start of estrogen).

Patients with PWS have low bone mineral density (BMD) and are at risk of osteoporosis related to sex-steroid and GH deficiencies, and low muscular activity with elevated biochemical markers of bone turnover [77–80]. Reduced BMD in PWS is associated with high risk of fracture in the long bones, as well as the small bones, of the hands and feet, with some patients suffering multiple fractures [81]. These findings support the need for hormone therapy, particularly sex-steroid replacement, during adolescence, and maintenance during adulthood. Estrogen, and androgen status should be monitored yearly during adolescence and adulthood and BMD assessed as indicated by DEXA.

Infertility is the rule in both sexes although a few instances of reproduction in females have been reported [54, 55].

(b) Precocious adrenarche and early puberty: Isolated premature pubarche has been reported in 14% and precocious puberty in 4% of males and females [35, 82, 83]. There is no consensus as to the management of either of these conditions. Some investigators have suggested the use of hydrocortisone in premature pubarche to decrease adrenal androgens when associated advancement of bone age is present; treatment with GnRH agonists must be reserved for selected patients [35].

In some patients with advanced bone age and hyperandrogenism we propose cyproterone acetate therapy. In these patients we generally obtain a slowdown of both bone age and androgen values. This drug is off-label, and parental consent is therefore necessary.

4.4. Psychological/Psychiatric Support. A characteristic behavioral profile with temper tantrums, stubbornness, controlling and manipulative behavior, compulsivity, and difficulty with change in routine becomes evident in early childhood in 70–90% of individuals with PWS. Behavioral and psychiatric problems most interfere with the quality of life in adolescence and adulthood.

Interventions concerning behavioral problems must be coordinated by specialists (psychologists, psychiatrists, and doctors), primary care providers, parents, and other family members.

Psychological support during infancy is fundamental for parents and children alike. Early logopedic therapy should be carried out to prevent language disorders. Due to their low cognitive level, PW children should receive learning support during school time.

In adolescence and adulthood some patients develop psychosis that requires pharmacological treatment [53, 86, 87]. In our experience, it is advisable to start treatment with psychotropic drugs at low doses, due to the possible hyperresponsiveness or paradoxical effects induced by commonly
used. The drugs used are benzodiazepines, classical antipsy-
chotics and atypical antipsychotics, mood stabilizers, and
selective serotonin reuptake inhibitors. Molecules lacking
orexigenic action, or with a lower capacity to induce
increased appetite, such as risperidone and fluoxetine, are
preferable. Topiramate might help combat the skin-picking
phenomenon [60].

4.5.1. Casa Sora. The nonprofit (ONLUS) foundation “Casa
Sora Per Voi” [88] is the first Italian project to offer PW
children, adolescents, adults, and their families the possibility
of formative stays. This project stems from the need of a
controlled environment, where patients can learn to cope
with their daily lives, socialize, and experience nature and
animals.

Various specialists are involved, such as dieticians and
psychologists, who plan numerous activities. Families, indi-
viduals, and healthcare professionals meet during structured
daily programs for adults and children; the children are
involved in programmed psychological activities, while their
parents and affiliated professionals attend lectures and group
discussions.

The house is in an isolated hilly location near lake
Iseo and is completely surrounded by greenery and uncon-
taminated nature. Daily activities include outdoor physical
exercise, group experiences in contact with nature in a
serene and stimulating environment, and psychospecific
recreational and artistic tasks.

4.5. Other Problematics

4.5.1. Orthopedic Problems. Scoliosis is a frequent feature
observed in children with PWS (30%–70%) and may be
explained partly by hypotonia and obesity [81, 89–91].
Regular clinical assessment is required at each visit, and
periodic spinal X-rays are useful, whether or not the
patient is receiving GH. Surgical treatment is indicated in
severe early-onset scoliosis-kyphosis, and in adolescents near
skeletal maturity. Due to the possibility of complications,
surgical treatment requires a multidisciplinary team with
expertise in the management of scoliosis and PWS.

4.5.2. Orthodontic Problems. Abnormal enamel and frequent
caries have been previously reported, but in a recent survey
PWS patients presented with a more favorable oral health
status than those in previous studies [92]. This status is wors-
ened by poor salivary production, which requires education
for both parents and children. Education for regular daily
drinking and products designed to increase saliva flow might
help prevent dental complications. Orthodontic treatment is
often needed.

4.6. Medical Alerts

4.6.1. Anesthesiological Risk. PWS patients have a higher
anesthesiological risk characterized by an exaggerated
response to hypnotic drugs, ventilation difficulties due to
dysmorphic facial conformation, hypoxia, breathing prob-
lems, and thermoregulation control. These phenomena are
more frequent and severe in obese patients that can have
obstructive apnea and right ventricular hypertrophy and
failure due to pulmonary hypertension. These conditions
require preventive cardiology and pulmonary evaluation.
Moreover, following surgery patients should be sent to
an intensive care unit, and some characteristics must be
considered:

(i) high pain thresholds,
(ii) thermoregulation disorders. Although there is no
indication for a predisposition to malignant hyper-
thermia in PWS, the use of depolarizing neuromus-
cular-blocking drugs (e.g., succinylcholine) should be
avoided unless absolutely necessary,
(iii) dense salivation could compromise airway patency,
especially during extubation,
(iv) hypotonia reduces cough reflex efficiency in clearing
airways.

4.6.2. Acute Gastric Distension. In the literature there are at
least 8 cases of acute gastric distension, and at least 3 cases of
death following gastric rupture in PWS patients. The basis
of this serious and potentially mortal complication is the
presence of hyperphagia, a high pain threshold, the inability
to vomit, and delayed gastric emptying. It is very important
to keep PWS patients under strict regular control in the
presence of large quantities of available food (banquets,
parties, supermarkets, etc.). Furthermore, in the presence of
abdominal pain and/or vomiting, an abdominal X-ray has to
be carried out to exclude gastric perforation.

5. Follow Up

The PW patients need a lot of specialists controls:

(i) axiological evaluation: length/height, weight, head
circumference, BMI (Figure 3),
(ii) nutritionist/dietician evaluation,
(iii) blood sample: oral glucose tolerance test (OGTT),
HbA1c, total cholesterol and HDL, triglyceride level,
uricemia, thyroid function (TSH, fT4), IGF1,
(iv) evaluation of pubertal development: dosage of LH,
FSH, estradiol (in females) and testosterone (in
males) both basal and after GnRH,
(v) bone age (left hand X-ray),
(vi) evaluation of bone density: blood sample for calcium
(Ca), phosphor (P), and magnesium (Mg) levels,
protidemia, PTH, vitamin D3, and to carry out a
vertebro femoral DEXA,
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<th>Evaluation</th>
<th>From birth to 3 years</th>
<th>From 3 to 10 years</th>
<th>From 10 to 18 years</th>
<th>Adult</th>
</tr>
</thead>
<tbody>
<tr>
<td>Axiological evaluation</td>
<td>3-4 months</td>
<td>6 months</td>
<td>6–12 months</td>
<td></td>
</tr>
<tr>
<td>Nutritionist/dietician evaluation</td>
<td>6–12 months</td>
<td>6–12 months</td>
<td>6–12 months</td>
<td>6–12 months</td>
</tr>
<tr>
<td>Blood sample (OGTT, HbA1c, total cholesterol</td>
<td>6 month</td>
<td>6–12 months</td>
<td>6–12 months</td>
<td>1 year</td>
</tr>
<tr>
<td>and HDL, triglyceride level, uricemia, TSH,</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>fT4, IGF1)</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Evaluation of pubertal development (dosage of</td>
<td></td>
<td></td>
<td></td>
<td>According to clinical evaluation</td>
</tr>
<tr>
<td>LH, FSH, estradiol (in females) and testosterone</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(in males) both basal and after GnRH)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bone age (left hand X-ray)</td>
<td></td>
<td>GH therapy: 1 year</td>
<td>According to clinical evaluation</td>
<td></td>
</tr>
<tr>
<td>Evaluation of bone density</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Blood sample for Ca, P and Mg, protidemia,</td>
<td>1-2 years</td>
<td>1-2 years</td>
<td>1-2 years</td>
<td></td>
</tr>
<tr>
<td>PTH, vitamin D3</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DEXA</td>
<td>1-2 years</td>
<td>1-2 years</td>
<td>1-2 years</td>
<td></td>
</tr>
<tr>
<td>Polysomnography</td>
<td>No GH therapy: 1 year</td>
<td>1 year</td>
<td>1 year</td>
<td>In case of problem</td>
</tr>
<tr>
<td></td>
<td>GH therapy: 3–6 months</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ENT with fibroscopy</td>
<td>No GH therapy: 6–12 months</td>
<td>In case of problem</td>
<td>In case of problem</td>
<td></td>
</tr>
<tr>
<td></td>
<td>GH therapy: 3–6 months</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Orthopedic evaluation</td>
<td>1 year (consider the necessity of spine X-ray)</td>
<td>6 months (consider the necessity of spine X-ray)</td>
<td>6 months (consider the necessity of spine X-ray)</td>
<td>In case of problem</td>
</tr>
<tr>
<td>Orthoptic/ophthalmology evaluation</td>
<td>1 year</td>
<td>In case of problem</td>
<td>In case of problem</td>
<td></td>
</tr>
<tr>
<td>Dental evaluation</td>
<td>Annual after 2 years</td>
<td>6 months</td>
<td>6 months</td>
<td>In case of problem</td>
</tr>
<tr>
<td>Dermatological evaluation</td>
<td></td>
<td></td>
<td></td>
<td>In case of skin picking</td>
</tr>
<tr>
<td>Echocardiography and cardiological evaluation</td>
<td></td>
<td></td>
<td></td>
<td>2 years</td>
</tr>
<tr>
<td>Neuropsychiatric and physiatric evaluation</td>
<td>3 months</td>
<td>1 year</td>
<td>1 year</td>
<td>1-2 years</td>
</tr>
</tbody>
</table>

In consideration of the numerous problems that can take place during the PW patient’s life, we have identified various age classes in which we define the best followup, that is resumed in Table 2.

6. Conclusion

Prader-Willi syndrome is a complex multisystem disorder. Patients can be affected by various problems; therefore precocious diagnosis is fundamental to guarantee optimal assistance. Each patient should undergo personally tailored treatment from birth. Therapeutic decisions and clinical followup need to consider all of these possible problems. A multidisciplinary team is required, made up of specialists such as neonatologists, geneticist, pediatricians, endocrinologists, orthopedic surgeons, psychologists, psychiatrists, physiotherapists, and urologists to deal with the numerous medical and psychological problems a PWS patient has to face. Only in this way we can improve quality of life, prevent complications, and prolong life expectancy in patients with PWS.

References


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