Case Report: Hepatic Adenomatosis in a Patient With Prader–Willi Syndrome

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Prader–Willi syndrome (PWS) is a genetic disorder caused by the lack of expression of genes on the paternally inherited chromosome region 15q11.2-q13. It is a multisystem disorder that is characterized by severe hypotonia with poor suck and feeding difficulties in early infancy, followed in early childhood by excessive eating and gradual development of morbid obesity. The incidence of type 2 diabetes mellitus is high, particularly in obese patients. Non-alcoholic fatty liver disease has also been reported in some patients with PWS. Liver adenomatosis is a benign vascular lesion of the liver, defined by the presence of >10 adenomas, in the otherwise healthy liver parenchyma. We report the first case of a patient with PWS with severe obesity, type 2 diabetes mellitus, and non-alcoholic fatty liver who also developed liver adenomatosis, review the pediatric literature on liver adenomatosis, and discuss the potential underlying mechanisms.

Keywords: hepatic adenomatosis, Prader Willi syndrome, liver adenoma, oral contraception pills, Glycogen Storage Disease

INTRODUCTION

Prader–Willi syndrome (PWS) is a complex multisystem disorder caused by lack of expression of genes on the paternally inherited chromosome region 15q11.2-q13. In the neonatal period, there is severe hypotonia with poor suck and feeding difficulties followed in infancy or early childhood by excessive eating and gradual development of morbid obesity. The syndrome is considered the most common genetic cause of obesity, occurring in 1:10,000–1:30,000 live births (1). Obesity and its related complications are the most common causes of morbidity and mortality in PWS. The mechanisms underlying the obesity include alterations in hypothalamic pathways that regulate satiety thus resulting in hyperphagia, disruption in hormones regulating appetite and satiety, and reduced energy expenditure (1).

Severe obesity is a strong risk factor for the development of type 2 diabetes mellitus (T2DM) in patients with PWS. T2DM in PWS occurs mostly in adults, but it has also been reported in patients under the age of 18 years (1). The prevalence of metabolic syndrome in obese patients with PWS...
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seems to be similar to other obese patients (2). Patients with PWS are thought to be at a lower risk of developing non-alcoholic fatty liver disease (NAFLD) because of a higher insulin sensitivity as well as insulinopenia (3, 4) with the staging of the NAFLD depending on body composition (3, 4).

Liver adenomatosis is a benign vascular lesion of the liver, defined by the presence of >10 adenomas, in an otherwise healthy liver parenchyma (5). A variety of associations have been reported with liver adenomatosis including glycogen storage disease types 1 and 1V, transfusion-induced hemosiderosis, Fanconi’s anemia, Hurler’s disease, severe combined immunodeficiency, familial adenomatous polyposis, and galactosemia (6).

We report the finding of liver adenomatosis in a case of PWS who is known to have severe obesity, T2DM, and NAFLD. We review all the previous reported cases of liver adenomatosis in pediatric patients and discuss the possible underlying mechanisms. Despite several cases reported in the literature, liver adenomatosis remains a poorly understood disease of unknown etiology.

CASE PRESENTATION

We report a 17-year-old obese female patient with PWS who was born in Egypt and then moved to Qatar at the age of 9 years. She was born at term to non-consanguineous parents with a birth weight of 3.5 kg. The pregnancy was uneventful. In the newborn period, there were feeding difficulties, and on day 12 of life, she had a seizure with frothing, cyanosis, and jaw clenching. She was admitted for 10 days to the neonatal intensive care unit (NICU). There was a delay in achievement of motor milestones, with crawling starting at the age of 1 year 8 months and walking at 2 years. There was also a history of speech delay, and the patient started speaking at the age of 5 years. She started to gain weight from the age of 2 years; however, at the age of 12 years, around puberty, hyperphagia and excessive weight gain were noticed. Her parents had difficulty limiting her food intake, as she had a persistent feeling of hunger to the extent of vomiting after eating large quantities. She initially sought endocrine care at the age of 15 years for an obesity workup.

Ultrasound of the liver was done to rule out fatty liver, and this showed multiple lesions in the liver that were thought to be areas of focal fatty sparing. MRI of the liver was done that showed an enlarged liver with diffuse fat infiltration and multiple numerous focal lesions suggestive of multiple adenomas (adenomatosis; Figure 1). A follow-up ultrasound in 2021 showed that the liver remained markedly echogenic and heterogeneous. The largest measured adenoma was 13 mm superficially in the left lobe (previously up to 11 mm) and generally was of the order of around 6 mm in diameter. Echocardiography showed normal cardiac function and anatomy. The patient undergoes an MRI of the abdomen every 6 months for the liver Adenomatosis. To date, all MRI scans do not show any significant interval changes in the number or size of the adenomas.

During the coronavirus disease 2019 (COVID-19) quarantine, at the age of 16, she gained weight rapidly, with 17 kg in 1 year. The weight upon presentation was 124.1 kg (99.54 centile), height was 138.8 cm (0.01 centile -3.73 Z-Score), and body mass index (BMI) was 64.4 kg/m². The patient also had symptoms suggestive of sleep apnea with mouth breathing, snoring nights, and occasional daytime sleepiness. A sleep study revealed obstructive sleep apnea; hence, she was started on non-invasive ventilation. She had menarche at the age of 13, with irregular menses since then with prolonged heavy menses. In light of the long-term risks of endometrial hyperplasia and carcinoma due to her secondary amenorrhea, the oncology team recommended a minimum of four withdrawal bleeds per year to reduce the risk. The patient was prescribed medroxyprogesterone; she received a short course, 5 days, once every 3 months. She has received three courses since 2019 (15 days total in 2 years).

When she was 17 years of age, blood glucose values were high (250–300 mg/dl) during multiple clinic follow-ups with polyuria, polydipsia, and nocturia. Her blood gas did not show acidosis. HbA1c was 8.1% with high serum insulin and C-peptide levels along with acanthosis nigricans on the neck, axilla, and inguinal region suggestive of insulin resistance. The patient was started on both long-acting and short-acting insulin for glycemic control. Table 1 shows the laboratory test results.

In terms of management, metformin was started initially, but a response was not observed. Liraglutide, a glucagon-like peptide 1 receptor (GLP-1) agonist, was tried for a month to help with

![Figure 1](https://example.com/figure1.png)

**FIGURE 1** | MRI liver findings. (A) Contrast enhanced liver MRI image showing multiple small liver lesions showing early arterial enhancement. (B) Lesions showing retention of contrast after 6 minute delay. (C) Lesions showing retention of contrast after 31 minute delay.
TABLE 1 | Results of laboratory tests done.

| Blood test          | Blood test result | Reference range |
|---------------------|-------------------|-----------------|
| Insulin level       | 618 pmol/L        | 20–571          |
| C-peptide           | 14.9 ng/ml        | (0.9–9.4)       |
| HbA1c               | 8.1%              | <6%             |
| ALT                 | 118 IU/L          | 9–22            |
| AST                 | 95 IU/L          | 15–28           |
| ALP                 | 141               | 48–96           |

ALT, Alanine Aminotransferase; AST, Aspartate Aminotransferase; ALP, Alkaline Phosphatase; HbA1c, Haemoglobin A1c.

weight reduction and reduce insulin resistance. However, she developed a local skin allergy in the form of itchiness on the injection sites, so it was stopped. Her insulin regimen included long-acting insulin (Glargine) 20 units daily and rapid-acting insulin (noveorapid) of 3–4 units three times daily. Despite this, her obesity and T2DM were difficult to manage. A sleeve gastrectomy was done in October 2021, considering the extreme obesity, associated comorbidities, and the relatively poor response to medical therapy for her weight. Her 2-month post-sleeve gastrectomy weight is 113.2 kg (compared to 121 kg). Her blood glucose readings are within range, and she is off all antihyperglycemic medications.

Family History
She is the youngest of three siblings; attended school for children with special needs. Father has T2DM.

Genetic Testing
Methylation studies in 2019, when she was 15 years old, revealed an absence of the paternal allele at 15q11-q13 due to abnormal methylation, thus suggesting a defect in imprinting as the underlying molecular mechanism of the PWS.

Radiology
MRI of the liver was done that showed an enlarged liver with diffuse fat infiltration and multiple numerous focal lesions suggestive of multiple adenomas (adenomatosis; Figure 1). A follow-up ultrasound in 2021 showed that the liver remained markedly echogenic and heterogeneous. The largest measured adenoma was 13 mm superficially in the left lobe (previously up to 11 mm) and generally was of the order of around 6 mm in diameter. Echocardiography showed normal cardiac function and anatomy.

DISCUSSION
Hepatic adenomas are benign vascular lesions of the liver that are usually solitary, associated with young women taking oral contraceptives. However, some patients have multiple adenoma, which is called liver adenomatosis. This was first described as a distinct entity in 1985 by Flejou et al. (7), who defined it as the presence of multiple adenomas (>10) that are not associated with steroid medication or glycogen storage disease. Currently, liver adenomatosis is defined as the presence of more than 10 adenomas in an otherwise normal parenchyma (8). The first clinical presentation due to liver adenomatosis is usually abdominal pain (9). Intraperitoneal bleeding, intratumoral hemorrhage, or necrosis producing acute pain are also reported (10). Only one report refers to malignant transformation of adenomas (11). This shows that even if the disease is benign, the risk of hemorrhage remains a concern.

We reviewed all the previous cases of liver adenomatosis in patients under the age of 18 years. Table 2 summarizes the age of presentation, gender, clinical features of the patients, the use of oral contraceptives, and associated complications. No previous patients with PWS have been reported with liver adenomatosis. The mechanism behind the development of multiple hepatic adenomas is not well established, especially in the pediatric age group. Liver adenomatosis has been reported in association with metabolic conditions, vascular anomalies, with the use of oral contraceptives, and with inactivating mutations in the HNF1A gene.

With regard to metabolic disease, there is a strong association reported between liver adenomatosis and glycogen storage disease with 50%–80% of children with type I or III glycogen storage disease developing multiple hepatic adenomas (19, 20). The impairment in glycogenesis and the accumulation of glycosgen deposits within the hepatocyte lead to hepatocyte hyperplasia, resulting in multiple adenoma formation. NAFLD is also reported to be associated with liver adenomatosis but mostly in adults (20). The increase in the intracellular lipid content could lead to a hyperplastic reaction with changes in oxidative and inflammatory pathways (21). An alternative mechanism of liver adenomatosis due to NAFLD suggests that the fatty tissue may generate continuous local estrogen through the increased activity of the enzyme aromatase, thus leading to the accelerated rate of hepatocyte growth and possible tumor formation (22). Multiple adenomas have been reported in other metabolic diseases such as diabetes, metabolic syndrome, and obesity in the adult population (8, 23).

The vascular hypothesis of liver adenomatosis is based on the association between reported cases of liver adenomatosis and hepatic vascular abnormalities, assuming that irregular vascular flow can result in the development of liver adenomatosis (Table 2). For instance, liver adenomatosis was reported in a 13-year-old male patient by Kawakatsu et al. (13) who had a spontaneous intrahepatic porto-hepatic venous shunt.

The association of liver cell adenomatosis and oral contraceptive or androgenic steroid use is still a point of controversy. Chiche et al. (8) reported that oral contraceptive therapy is not as rarely associated with this liver disease as initially suggested by Flejou et al. (7); 46% of their female patients were on oral contraceptives (9, 15). Adenoma regression was recognized after discontinuing hormonal contraceptives in multiple studies, which suggest that oral contraceptives play a role in the evolution of liver adenomatosis. In our review of the pediatric cases, we did not find a correlation with the use of oral contraceptives and liver adenomatosis, as only 3 out of 13 reported cases used oral contraceptives (Table 2).
In some patients with liver adenomatosis, there is a genetic background to the etiology. This involves biallelic inactivating mutations in the transcription factor, HNF1A, in the hepatic adenomas by the occurrence of two molecular events: either a germline and a somatic HNF1A mutation or two independent somatic events (25). This is usually associated with the Maturity Onset Diabetes of the Young (MODY) in the family history as heterozygous HNF1A mutations are a cause of MODY3.

Our patient with PWS has several risk factors for the development of liver adenomatosis. These include severe obesity (BMI of 64.4 mg/m²), T2DM, NAFLD, and possibly the use of oral contraceptives. Medroxyprogesterone was used for a short period of time, so we do not think that this is a significant risk factor, although we cannot completely rule this out. There was no family history of MODY in this patient. It is difficult to know which of these risk factors is directly linked to the liver adenomatosis but it is likely that the combination of the risk factors is involved. This case highlights that patients with PWS, who have risk factors such as obesity, T2DM, and NAFLD, may develop liver adenomatosis. Physicians should have a low threshold for making this possible diagnosis.

### DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/supplementary material. Further inquiries can be directed to the corresponding author.

### AUTHOR CONTRIBUTIONS

HD and AS collected patient information, recruited the patient, analyzed and interpreted the data, and drafted the article. KH designed the study, obtained funding, and reviewed and edited the article. BH analyzed the data and reviewed and edited the article. All authors contributed to the article and approved the submitted version.

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REFERENCES

1. Muscogiuri G, Barrea L, Faggiano F, Maiorino MI, Parrillo M, Pugliese G, et al. Obesity in Prader-Willi Syndrome: Pathophysiological Mechanisms, Nutritional and Pharmacological Approaches. J Endocrinol Invest (2021) 44(5):607-17. doi: 10.1007/s40618-021-02613-x

2. Brambilla P, Crinò A, Bedogni G, Bosio L, Cappa M, Corrias A, et al. Metabolic Syndrome in Children With Prader-Willi Syndrome: The Effect of Obesity. Nutr Metab Cardiovasc Dis (2011) 21(4):269–76. doi: 10.1016/j.numecd.2009.10.004

3. Talebizadeh Z, Butler MG. Insulin Resistance and Obesity-Related Factors in Prader-Willi Syndrome: Comparison With Obese Subjects. Clin Genet (2005) 67(3):230–9. doi: 10.1111/j.1399-0004.2004.00392.x

4. Haqq AM, Muchibauer MJ, Newgard CB, Gribow S, Freemark M. The Metabolic Phenotype of Prader-Willi Syndrome (PWS) in Childhood: Heightened Insulin Sensitivity Relative to Body Mass Index. J Clin Endocrinol Metab (2011) 96(1):E225–32. doi: 10.1210/jc.2010-1733

5. Barbier L, Nault JC, Dujardin F, Scotto B, Besson M, de Muret A, et al. Natural History of Liver Adenomatosis: A Long-Term Observational Study. J Hepatol (2019) 71(6):1184–92. doi: 10.1016/j.jhep.2019.08.004

6. Davenport M, Pakarinen M. Preface - Pediatric Hepatobiliary Surgery. Pediatr Cardiol (2000) 23(5):558. doi: 10.1007/s00246-000-01574-9

7. Flejou JF, Barge J, Menu Y, Degott C, Bismuth H, Potet F, et al. Liver Adenomatosis. An Entity Distinct From Liver Adenoma? Gastroenterology (1985) 99(5):1132–8. doi: 10.1016/S0016-5085(85)80220-3

8. Chiche L, Dao T, Salameh A, Galais MP, Bovard N, Schmutz G, et al. Liver Adenomatosis: Reappraisal, Diagnosis, and Surgical Management: Eight New Cases and Review of the Literature. Ann Surg (2000) 231(1):74–81. doi: 10.1097/00000658-200001000-00011

9. Ribeiro A, Burgart LJ, Nagorney DM, Gores GJ. Management of Liver Adenomatosis: Results With a Conservative Surgical Approach. Liver Transpl (1998) 4(5):388–98. doi: 10.1002/liv.400040505

10. Lui AF, Hiaratzka LF, Hirose FM. Multiple Adenomas of the Liver. Cancer (1980) 45(5):1001–4. doi: 10.1002/1097-0142(19800301)45:5<1001::aid-cncr2820450528>3.0.co;2-f

11. Leese T, Farges O, Bismuth H. Liver Cell Adenomas. A 12-Year Surgical Experience From a Specialist Hepato-Biliary Unit. Ann Surg (1988) 208(5):558–64. doi: 10.1097/00000658-198811000-00003

12. Chen KT, Bocian JJ. Multiple Hepatic Adenomas. Archives of Pathology & Laboratory Medicine (1983) 107(5):274–5.

13. Kawakatsu M, Vilgrain V, Belghiti J, Flejou JF, Nahum H. Association of Multiple Liver Cell Adenomas With Spontaneous Intrahepatic Porto-portal Shunt. Abdom Imaging (1994) 19(5):438–40. doi: 10.1007/BF00206934

14. Gokhale R, Whittington PF. Hepatic Adenomatosis In an Adolescent. J Pediatr Gastroenterol Nutr (1996) 23:482–6.

15. Babaoglu K, Binnetoglu FK, Aydoğan A, Altun G, Gürbüz Y, Inan N, et al. Hepatic Adenomatosis in a 7-Year-Old Child Treated Earlier With a Fontan Procedure. Pediatr Cardiol (2010) 31(6):861–4. doi: 10.1007/s00246-010-9685-x