

# Prenatal Diagnosis of Pfeiffer Syndrome Patient with FGFR2 C.940-1G>C Variant: A Case Report

This article was published in the following Dove Press journal:  
*The Application of Clinical Genetics*

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**Background:** Pfeiffer syndrome (PS) is an autosomal dominant disorder caused by mutations in fibroblast growth factor receptor FGFR1 and FGFR2 genes, occurring in approximately 1:100,000 live births. PS has a wide range of clinical expression and severity, so early prenatal diagnosis is difficult and genetic counseling is desirable. We describe a PS newborn with her ultrasound and molecular studies.

**Case Report:** We describe a female term newborn with cloverleaf-shaped skull, facial hypoplasia, low ears, exophthalmos and wide, broad and deviated thumbs and hallux. The patient was diagnosed by ultrasound at 29 WGA and referred to a tertiary care hospital for her follow-up. Molecular test revealed a heterozygous pathogenic variant in intron 8 of the FGFR2 gene (FGFR2: c.940-1G>C). It was a de-novo mutation. At 17 days of life, craniostylosis correction and a Lefort-III frontomaxillary advancement were performed.

**Conclusion:** Pfeiffer syndrome is a devastating genetic disorder. Prenatal diagnosis according PS morphological features in prenatal ultrasound allows timely genetic counseling, early referral to third-level centers, and close follow-up in the prenatal and postnatal stages.

**Keywords:** Pfeiffer syndrome, acrocephalosyndactylia, craniostylosis, fibroblast growth factor receptor 2

## Introduction

Pfeiffer syndrome (PS) was firstly described by Rudolph Pfeiffer as an acrocephalosyndactylia: bicoronal craniostylosis, middle facial hypoplasia, digital anomalies of hands and feet, presenting in some cases, partial cutaneous syndactylia.<sup>1</sup> PS prevalence is one of 100.000 live births, without sex preference.<sup>2</sup> PS has an autosomal dominant inheritance pattern, generated by two types of mutations affecting the 1 or 2 gene of the fibroblast growth factor receptor (FGFR1 or FGFR2).<sup>2</sup> There are three PS phenotypes to determine neurodevelopment and survival prognosis.<sup>3</sup> In type 1, the patient presents mild physical manifestations without neurodevelopmental impairment. Types 2 and 3 are more frequent and severe. Type 1 is generally associated with FGFR1 and Type 2 and 3 with FGFR2. They could develop craniostylosis and hypertelorism with external proptosis and cloverleaf skull in case of type 2.<sup>4,5</sup>

The diagnosis in PS patients is usually made after the birth. It is difficult a comprehensive management that could worsen the quality of life of the patients and their family. Prenatal diagnosis in utero is essential for an early approach in a tertiary care center. We describe a Type 2 PS patient diagnosed in prenatal stage and molecularly confirmed after birth through a new generation sequencing panel in a high-complexity hospital in Colombia.

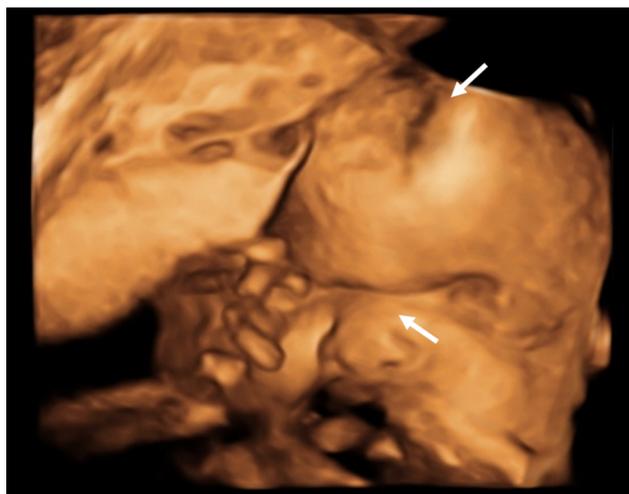
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## Case Report

A 32-year old pregnant woman with no relevant past medical history underwent a routine ultrasound scan showing a foetus of 29 weeks of gestational age (WGA). Findings included frontal prominence and a cloverleaf skull suggestive of craniosynostosis and hypoplastic nasal bone (Figure 1). In this ultrasound scan, feet and hands had no abnormalities. The woman was considered high-risk pregnancy and was referred to a tertiary care center for her weekly follow-up and the caesarean was scheduled at 38 WGA. After delivery, the female newborn had APGAR scores of 8, 9 and 10 at minutes 1, 5 and 10, respectively. Birth weight was 2862 grams, height was 48 cm and head circumference was 33 centimeters. The physical exam showed a cloverleaf skull, hypertelorism, facial hypoplasia, low ears, exophthalmos, sacral dimple, and wide, short and deviated thumbs and hallux (Figure 2).

The 3D CT head scan confirmed the cloverleaf skull, complete bilateral frontoparietal craniosynostosis, incomplete craniosynostosis of the sagittal and bilateral lambdoid suture, orbital alteration with bilateral proptosis secondary to hypertelorism and trans ependymal edema (Figure 2). The neonatal cranial sonography showed dysgenesis of corpus callosum with tentorium cerebelli high implantation. She also had persistent arterial ductus with bidirectional shunt without hemodynamic instability. There were no renal and ophthalmic malformations.

A new generation sequencing panel study was performed. It studies 241 genes potentially responsible for one or more Mendelian diseases related to hereditary osteodysplasia, collagen diseases and short stature. We found a heterozygous



**Figure 1** 3D-ultrasound at 29 WGA. The foetus has frontal prominence and a cloverleaf skull suggestive of craniosynostosis and hypoplastic nasal bone (white arrows).

pathogenic variant in intron 8 of the FGFR2 gene (FGFR2: c.940-1G>C). The finding was confirmed using an ABI 3500 sequencer (Applied Biosystems, Thermo Fisher Scientific, Waltham, MA, USA) by Sanger sequencing. Bioinformatic analysis of the variant was performed; FATHMM classified the variant as “damaging,” while MutationTaster classified it as “causing disease”. Currently, it is classified as “pathogenic” according to the American College of Medical Genetics and Genomics (ACMG) guidelines (PVS1, PM2, PP3).<sup>6</sup> It was a de-novo mutation as the parents were unaffected. At 17 days old, a craniosynostosis correction and a Lefort-III frontomaxillary advancement were performed. In the postoperative period, she had hydrocephaly and ventricular dilatation requiring a ventriculoperitoneal catheter (VPC). The VPC was discontinued after 6 days and the patient was discharged at 24 days-old with a multidisciplinary outpatient follow-up.

At the time of the writing this report, the patient was 12 months and was diagnosed with psychomotor development retardation. A polysomnogram diagnosed sleep apnea-hypopnea syndrome. A nasolaryngoscopy showed multiple turbinoseptal synechiae and right choana atresia.

This case report was approved by the institutional IRB to publish the case details. Written informed consent was provided by the patient’s legal guardian to have the case details and any accompanying images published.

## Discussion

Prenatal diagnosis of Pfeiffer syndrome is challenging in prenatal care. PS low incidence and the wide variability of morphological findings, which can also be related to other conditions, makes it difficult to suspect this syndrome early pregnancy stages.<sup>7</sup> Pfeiffer syndrome is a clinically and autosomal dominant disorder caused by mutations in FGFR1 and FGFR2.<sup>2</sup> Only 17 cases of diagnosis in the prenatal stage have been reported in the world.<sup>2</sup> We present the clinical findings of a patient with SP diagnosed in-utero due to a variant in FGFR2 gene.<sup>8-13</sup>

Although most cases are diagnosed in their neonatal stage, prenatal diagnosis is possible. Three-dimensional obstetric ultrasound is the first-line diagnostic tool for suspected PS, being useful to verify suture closure in the third trimester of pregnancy for the most severe cases<sup>14</sup> although the diagnosis with lower gestational ages has been reported.<sup>15</sup> Adding to craniosynostosis, principal ultrasound prenatal findings suggestive to PS are thumbs deformations and big toes.<sup>7</sup> Even when the cloverleaf skull is the most common finding in PS, and in our patient, it was the most important finding to perform PS diagnosis, the absence of



**Figure 2** Physical findings at birth. (A and B). Female newborn with cloverleaf skull, hypertelorism, facial hypoplasia, low ears, exophthalmos, (C) short and deviated hallux and (D) thumbs.

this characteristic does not exclude the diagnosis, as well as its presence does not directly certify this syndrome, because other diseases can also present.<sup>7</sup> The presence of wide fingers, sacral appendix, vertebral fusions and coronal clefts in the prenatal ultrasound should lead the clinician to perform prenatal molecular tests to rule out PS.<sup>2,7</sup> Almost all cases of PS are a consequence of new mutations because the parents are unaffected. Nevertheless, there is a percentage where the abnormal gene can be inherited from either parent. In the cases were FGFR2 mutations could be inherited from the father, Cell-free DNA extracted from maternal plasma can provide valuable molecular information and identify the genetic diagnosis in utero.<sup>16</sup>

Our patient had a Pfeiffer syndrome type 2. Her cloverleaf skull it is a consequence of premature fusion of all sutures, except the metopic and squamous suture. This condition produces the frontal prominence of both middle fossae. The patient also presented hydrocephalus and dysgenesis of the corpus callosum. Our patient also had hypertelorism and severe ocular proptosis. This condition can lead to

endophthalmitis and rupture of the eyeball, so periodic evaluation by the ophthalmologist is necessary. The upper airway is also compromised in PS type 2 patients given the facial hypoplasia that characterizes them. Although the patient had a wide nasal bridge, she had no obstruction of the upper airway.

The karyotype in all PS cases reported is normal so the cytogenetic study is not recommended.<sup>2</sup> In PS, 94% of mutations occur in exon 8 or 10 of the FGFR2 gene, which is also associated with other syndromic craniosynostosis, often sporadic, such as Apert syndrome, Crouzon among others;<sup>17</sup> In our patient, the variant detected in intron 8 of the FGFR2 gene, in a heterozygous state, is an intronic variant that alters the splicing of RNA whose alteration contributes to the presentation of the disease.

An early and accurate diagnosis is essential for multidisciplinary management. Prenatal diagnosis allows for rapid genetic counseling, early referral to third-level centers, and close follow-up in the prenatal and postnatal stages. Therefore, the implementation of new generation

sequencing panels that include genes associated with genetic syndromes related to craniosynostosis is a useful strategy to improve the diagnosis. In addition to this, clinical suspicion is essential to detect more cases in the prenatal stage, as well as early referral to tertiary care centers for a multidisciplinary evaluation is necessary to improve the patient's living conditions.

## Conclusion

Pfeiffer syndrome type 2 is a genetic disorder caused by mutations in the FGFR2 gene, which encodes the human fibroblast growth factor type 2 receptor that consequently generates a wide range of abnormalities, mainly craniosynostosis. Recognizing morphological features in prenatal ultrasound allow a timely and comprehensive management, improving the quality of life of the PS patients and their family.

## Disclosure

The authors report no conflicts of interest in this work.

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