

Investigation of *H19/RsaI* Polymorphism in Children With Low Birth Weight in Pernambuco, Brazil

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Background: *H19* is a strong candidate gene for influencing birth weight variation and is exclusively imprinted maternally. In an attempt to understand the relationship of this gene polymorphism with low birth weight children, we investigated association of *H19/RsaI* polymorphism with low birth weight and normal birth weight in children and their mothers.

Objectives: The aim of our study was to establish the association between *H19* gene polymorphism and LW in children born in Pernambuco, state of Brazil.

Patients and Methods: It were selected 89 children, 40 low birth weight (LW) and 49 normal birth weight (NW) and 71 mothers (40 mothers of newborns NW and 31 mothers of newborns LW) attended at Dom Malan Hospital, Petrolina, Pernambuco - Brazil. Peripheral blood samples were collected from patients and genomic DNA was extracted and detected by electrophoresis agarose gel, stained by Blue Green Loading Dye. DNA PCR amplification was done using the primers *H1* (*sense*) and *H3* (*antisense*). PCR products were digested with *RsaI* and electrophoresed on agarose gel stained by ethidium bromide. Statistical analyses were performed using the program BioEstat version 5.0.

Results: The *RsaI* polymorphism in the *H19* gene showed that genotype frequencies did not differ statistically between low birth weight (AA = 12.5%, AB = 45%, BB = 42.5%) and control (AA = 8.6% AB = 36.73%, BB = 55.10% groups) and the allele frequencies were not significantly different ($P = 0.2897$). We also did not observe any association between maternal *H19* allele polymorphism and low birth weight newborns ($P = 0.7799$) or normal birth weight children ($P = 0.8976$).

Conclusions: The small size of sample may be the explanation for these results; future studies with more patients are needed to confirm the effect of *H19/RsaI* polymorphism on birth weight of LW newborns.

Keywords: Birth Weight; Infant Low Birth Weight; Genotype; Alleles; Polymorphism Genetic

1. Background

Neonatal (0 to 28 days of life) mortality represents about 70% of infant mortality in Brazil, an indicator of living conditions and health of the population. Birth weight < 2,500 g is pointed out as the most influential factor in determining neonatal morbidity and mortality. The fetal growth is regulated by genetic, environmental, hormonal, nutritional, and placental factors (1).

IGF2 and *H19* genes are of special interest, since in addition to their reciprocal imprinting patterns, they are closely linked on chromosome 11 in humans and are strong candidate genes for influencing birth weight variation (2). The human *H19* is an untranslated gene that lies within 200 kb downstream of the paternally expressed allele (3). The maternally expressed *H19* gene itself does not encode a protein, but the RNA has growth potentially

suppressing functions (4) through inhibiting translation of *IGF2* RNA (5). *IGF2* appears to be an important growth factor and low expression of its gene might be associated with intrauterine fetal life damaging resulting in low birth weight (LW) of newborn and may predispose the individual to chronic diseases in post-natal life such as obesity and hypertension (6). Studies have shown an association between birth weight with polymorphisms of *IGF2* and *H19* genes (7).

2. Objectives

The aim of our study was to establish the association between *H19* gene polymorphism and LW in children born in Pernambuco, state of Brazil.

Table 1. Allele and Genotype Frequencies of the Polymorphism in the Gene *H19/RsaI* in Newborns With LW and NW and Mothers of Newborns With LW and NW^{a, b}

Frequencies	Children NW	LW	Mothers NW	LW
Genotypes				
AA	4 (8.16)	5 (12.5)	2 (5)	6 (19.35)
AB	18 (36.73)	18 (45)	17 (42.5)	12 (38.71)
BB	27 (55.10)	17 (42.5)	21 (52.5)	13 (41.94)
Alleles				
A	26 (26.53)	28 (35)	21 (26.25)	24 (38.71)
B	72 (73.47)	52 (65)	59 (73.75)	38 (61.29)

^a Abbreviations: HW, Hard-Weinberg equilibrium; LW, Low weight; NW, normal weight.

^b Data are presented as No. (%).

Table 2. P value and OR (95% CI) Calculation for Different Types of Allels and Genotypes^a

	Genotypes				Alleles			
	Children NW × LW	Mothers NW × LW	Children NW × Mothers NW	Children LW × Mothers LW	Children NW × LW	Mothers NW × LW	Children NW × Mothers NW	Children LW × Mothers LW
P Value	0.4873	0.1689	0.7705	0.7169	0.2897	0.1613	0.8976	0.7799
OR (95% CI)					0.67 (0.35-1.27)	0.56 (0.28-1.15)	1.01 (0.52-1.98)	0.85 (0.43-1.69)

^a Abbreviations: HW, Hard-Weinberg equilibrium; LW, Low weight; NW, normal weight.

3. Patients and Methods

3.1. DNA Extraction

We selected 89 children, 40 LW and 49 normal birth weight (NW) and 71 mothers (40 mothers of NW newborns and 31 mothers of LW newborns) who attended Hospital Dom Malan, Petrolina, PE-Brazil. Genomic DNA was extracted from peripheral blood. Ethical approval for this study was obtained from the Ethics Committee of the Faculty of Medicine/UFPE. Written consent was obtained from parents of the subjects.

3.2. DNA PCR and Genotyping

DNA PCR amplification was based on Petry and Ong (8) with *H1* (sense) and *H3* (anti-sense) primers. The PCR products were digested with *RsaI* 4 hour at 37°C and electrophoresed on 1.5% agarose gel and stained by ethidium bromide.

3.3. Statistical Analysis

Statistical analyses were performed using the program BioEstat version 5.0. Genotype and allele frequencies in groups were compared by I^2 test with significance set for a P value < 0.05. Hardy-Weinberg test was used to verify if the genotypes of the control group NW and LW were in equilibrium.

4. Results

Allele and genotype frequencies *H19/RsaI* polymor-

phism are shown in Table 1. We observed in this study that the frequencies of AA, AB and BB genotypes did not differ significantly between LW and NW children ($P = 0.4873$) as well as the allele frequencies ($P = 0.2897$; OR = 0.6). Also no association was observed between maternal *H19* allele polymorphism and LW newborns ($P = 0.7799$; OR = 0.85) or NW newborns ($P = 0.8976$; OR = 1.01). Genotype frequencies in both groups did not differ significantly. In the present study, we observed a higher frequency of allele B for both groups in relation to gene polymorphism *H19/RsaI*. It was also shown that the value of genotypic frequency of allele B of the gene *H19* was higher than the A allele for both NW children (55.10%) and mothers of NW children (52.5%).

5. Discussion

It was believed that the relationship of *IGF2* and *H19* could influence birth weight. SNP located in *H19* gene that may alter mRNA structure could influence the imprint of *IGF2*, which is a major fetal growth factor (9). However our results did not reveal an association of *H19/RsaI* polymorphism with LW children, outcomes that are consistent with those described by Araujo (10). Study of polymorphism of the *IGF2* gene in this same population found no association of this polymorphism with LW children (11).

Despite the few studies on the association of *H19* gene polymorphism with LW children, the findings detect association of the polymorphism with the condition of children born with LW. SNPs in the 5' region of the *H19* (rs2067051, rs2251375, and rs4929984) gene were asso-

ciated with birth weight (2). Petry and Ong found that birth weight of offspring showed association with *H19* 2992C >T SNP genotype of mothers (8). The small size of sample may be the explanation for these results; future studies with more patients are needed to confirm the effect of *H19/RsaI* polymorphism on birth weight of LW newborns.

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Authors' Contributions

Paula Maia: Concept/design, acquisition of data, drafting of the manuscript. Paulo Souza: drafting of the manuscript, critical revision of the manuscript. Hildson Dornelas Angelo: drafting of the manuscript, critical revision of the manuscript. Igor Santos: critical revision of the manuscript. Danyelly Martins: acquisition of data, data analysis. Jose Lima Filho: critical revision of the manuscript and approval of the article. Maria Mascena Maia: drafting of the manuscript, critical revision and approval of the article.

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