Primary Ciliary Dyskinesia Diagnostic Challenges: Understanding the Clinical Phenotype of the Puerto Rican 
RSPH4A Founder Mutation

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Abstract: Primary ciliary dyskinesia (PCD) is a rare, heterogeneous ciliopathy resulting in chronic oto-sino-pulmonary disease, bronchiectasis, newborn respiratory distress, and laterality defects. PCD diagnosis can be achieved by following diagnostic algorithms that include electron microscopy, genetics, and ancillary testing. Genetic mutations in more than 45 genes, including RSPH4A, can lead to PCD. RSPH4A mutations located on chromosome six, affect radial spokes and results in central complex apparatus abnormalities. The RSPH4A [c.921 + 3_6delAAGT] founder mutation was described as one cause of PCD without laterality defects in Puerto Rico. Additionally, there are further diagnostic challenges present in the Puerto Rican population to diagnose PCD. We describe the demographics, clinical features, and RSPH4A genetic variants in 13 patients with clinical PCD affecting 11 Puerto Ricans from unrelated families.

Keywords: RSPH4A; primary ciliary dyskinesia; founder mutation; cilia; Puerto Rico

1. Introduction

Primary ciliary dyskinesia (PCD) is a genetically heterogeneous, autosomal recessive disorder characterized by motile cilia dysfunction with a prevalence of approximately 1:15,000 individuals in the United States [1]. The actual prevalence of PCD in Puerto Rico is unknown. Genetic mutations in more than 45 human genes can lead to PCD, affecting the function of around 200 ciliary structural proteins [2]. These abnormalities result phenotypically in neonatal respiratory distress in approximately 80% despite a full-term gestation, oto-sino-pulmonary disease since birth, infertility, and organ laterality defects in 40–50% of cases [3]. Diagnostic algorithms for suspected PCD patients recommend evaluation of individuals with at least 2 of the 4 key clinical features including unexplained neonatal respiratory distress in term infants, year-round daily cough and/or daily nasal congestion beginning before 6 months of age, and laterality defects [4,5].

PCD diagnosis has become a challenge for physicians since it is easily misdiagnosed with similar genetic syndromes such as cystic fibrosis and other chronic suppurative lung diseases [6]. Previous studies have shown that approximately 1 of 3 patients have a normal ciliary structure on electron microscopy, which increases the difficulty in confirming the PCD diagnosis [7]. As a key diagnostic test, electron microscopy evaluates the 9 + 2 ciliary normal configuration and assesses for ciliary ultrastructure defects. Updated guidelines incorporate genetic testing as part of the algorithm for PCD diagnosis [4]. Additional PCD mutations have been discovered as genetic testing has become more accessible to healthcare providers and patients.
Biallelic pathogenic variants in the RSPH4A gene, located on chromosome 6, affect ciliary radial spokes which alter the configuration of the central complex apparatus [8]. Mutations in RSPH4A affect the normal planar beating pattern of the cilia into a circular ciliary motion [9]. The abnormal ciliary function impairs effective mucociliary clearance in PCD patients [10]. PCD mutations may interfere with the placement of the left-right organ, which is established during embryogenesis [11]. Normally, the embryonic node cilium lacks a central pair of microtubules following a 9 + 0 axonemal structure, which specifies the left-right axis [12]. Therefore, PCD mutations that affect radial spokes, like RSPH4A, will not present laterality defects like dextrocardia or complete situs inversus [13]. RSPH4A [c.921 + 3_6delAAGT] founder mutation was described as a PCD pathogenic variant without laterality defects. The splice site mutation [c.921 + 3_6delAAGT] in RSPH4A has been described as a common cause of PCD in the Hispanic population, specifically Puerto Rico [14]. We aim to describe our experience with clinical features, pulmonary function test, chest imaging, genetic profile, and electron microscopy findings diagnosing native Puerto Ricans with RSPH4A mutations. A better understanding of the clinical-genetic correlation of RSPH4A genetic variants will help to expose the diagnostic challenges for PCD patients without laterality defects in Puerto Rico.

2. Materials and Methods

A descriptive retrospective chart review of 31 Puerto Rican subjects was completed. Patients were followed up at a single private outpatient clinic in Puerto Rico with expertise in PCD from September 2018 to September 2020. Detailed histories, physical examinations, pulmonary function tests, imaging studies, and nasal ciliary biopsies were completed. Criteria for PCD evaluation were achieved by documenting the presence of 2 of the 4 key clinical features described on the suggested diagnostic algorithm as previously published [5]. Measurement of nasal nitric oxide (nNO) was not available in our institution. The sweat test was completed to rule out cystic fibrosis. Genetic testing was evaluated using commercially available sequence analysis (Invitae Corporation, San Francisco, CA), using buccal swabs or saliva samples. Gene panel testing evaluated for deletion/duplication on 35 PCD-related genes. The cystic fibrosis transmembrane conductance regulator (CFTR) gene was also evaluated as part of the initial patient PCD evaluation. Genetic testing included a sequence of exons and introns for the RSPH4A gene (Transcript: NM_001010892.2). Nasal biopsies were submitted for pathology review at Focus Pathology Medical Laboratories in Chester, New York. A total of thirteen (n = 13) PCD patients with the presence of RSPH4A mutations were included in the analysis. High-Resolution chest CT (HRCT) scans were reviewed, if available. Pulmonary function tests were completed in accordance with the American Thoracic Society guidelines [15]. Airflow limitation was determined using Z-score values of −1.64 as the lower limit of normal. The Global Lung Function Initiative-2012 (GLI) was used as a reference equation. Descriptive statistics presented in percentages and means. The Institutional Review Boards approved data collection and analysis for the Protection of Human Subjects from University of Puerto Rico, Medical Sciences Campus in San Juan, Puerto Rico.

3. Results

3.1. Demographics and Clinical Characteristics

Thirteen patients had clinical PCD phenotype, at least 2 of the key clinical features for PCD evaluation, presence of RSPH4A genetic variants, and normal or abnormal electron microscopy. Ages ranged from 1 year to 51 years old and 62%, (8/13) were females. All patients were of Hispanic ethnicity (100%, 13/13). PCD related symptoms in our cohort included: Year-round wet cough (100%, 13/13), neonatal respiratory distress (69%, 9/13), bronchiectasis (89%, 8/9), Year-round daily nasal congestion (100%, 13/13), chronic secretory otitis media (30%, 4/13), and hearing loss (69%, 9/13). In our cohort, only two females homozygous for RSPH4A [c.921 + 3_6delAAGT] were sexually active and self-reported to
be diagnosed with infertility. Laterality defects were present in 0%, (0/13) of the subjects with RSPHA4 mutations (Table 1 and Table A1).

Table 1. Demographics and clinical characteristics.

| Characteristics                                               | Value (N = 13) |
|---------------------------------------------------------------|----------------|
| Gender (F/M), N (%)                                           | 8 (62)         |
| Age, median ± SD, (years)                                    | 13 ± 15.2      |
| Ethnicity, N (%), Hispanics, Puerto Ricans                    | 13 (100)       |
| PCD-related symptoms/sign N (%)                               | -              |
| Year-round wet cough                                         | 13 (100)       |
| Year-round daily nasal congestion                             | 13 (100)       |
| Neonatal respiratory distress                                 | 9 (69)         |
| Hearing loss                                                  | 9 (69)         |
| Bronchiectasis                                                | 8/9 (89)       |
| Chronic secretory otitis media                                | 4 (30)         |
| Infertility                                                   | 2 (15)         |
| Laterality defects                                            | 0 (0)          |

F: Female; M: Male, SD: Standard Deviation

3.2. Electron Microscopy Findings

Ciliary biopsies were abnormal in 100%, (11/11). Abnormal findings on electron microscopy were related to central apparatus defects and abnormal microtubule configuration, including loss of central and outer doublets, presence of central solitary microtubule, and extra gain of central singlets, triplets, and quadruplets (Figure 1).

![Figure 1. Ciliary biopsy findings of RSPH4A subjects. A total of 11 samples were submitted for electron microscopy, representative images of abnormal findings are shown. (a) Normal ciliary configuration [9 + 2]; (b) Abnormal peripheral microtubule configuration, [7 + 2]; (c) Single microtubule outside the 9 outer doublet rings [9 + 1]; (d) Extra central microtubule, [9 + 3]; (e) Electron dense material in central area surrounded by eight outer doublets [8 + 0]; (f) Single microtubule in position of the central pair [9 + 1]. (g, h) Abnormal microtubule organization and configuration, with displacement of peripheral doublets [9 + 0]; (i) Supernumerary central microtubules, [9 + 5]. (j) Central displacement of a peripheral doublet [8 + 4].](image-url)
3.3. RSPH4A Genetic Sequencing

RSPH4A genetic variants were identified in all 13 subjects from 11 unrelated families. Biallelic pathogenic variants in RSPH4A gene were noted in 62%, (8/13) subjects of the cohort (Table A1). Rare RSPH4A genetic variants classified as a variant of unknown significance (VUS) were present in 23%, (3/13) of the cohort. Most common RSPH4A pathogenic variants were the RSPH4A [c.921 + 3_6delAAGT]. A single heterozygous copy of the pathogenic genetic variant RSPH4A [c.1103T > G] (p. Val368Gly) was present in one patient. Two subjects from unrelated families had one heterozygous copy of the RSPH4A [c.902A > C (p.Gln301Pro)], classified as a VUS. These two rare genetic variants have been previously described on population databases: RSPH4A [c.1103T > G] (p.Val368Gly); (rs747419302, ExAC 0.001%) and RSPH4A [c.902A > C (p.Gln301Pro)] (rs755128358, ExAC 0.009%). Both genetic variants have been reported in the literature on individuals with RSPH4A-related disease [www.lovd.nl/RSPH4A].

3.4. Imaging

HRCT were available for review in a total of 9 of 10 patients. Homozygous RSPH4A [c.921 + 3_6delAAGT] patients displayed bronchiectasis on 100% (8/8) of those above age seven. Case 9 and 10 with compound heterozygous genetic variants for RSPH4A displayed bronchiectasis on HRCT. The most common findings on HRCT were: cylindrical and varicose bilateral bronchiectasis, bibasilar parenchymal scarring tissue, centrilobular nodules, and tree-in-bud opacities. Representative views of the HRCT of both pediatric and adult PCD patients who are homozygous, compound heterozygous, or VUS for RSPH4A are presented in Figure 2.
Figure 2. High-Resolution chest CT scans (HRCT) of PCD patients with positive RSPH4A genetic variants. (a). 15-year-old male homozygous for RSPH4A [c.921 + 3_921 + 6del (Intronic)] founder mutation shows a tree-in-bud pattern at left lower lobe (LLL), lingula, right lower lobe (RLL), right middle lobe (RML), right upper lobe (RUL). HRCT showed bilateral ground-glass appearance but no bronchiectasis. (b). 51-year-old male, homozygous for RSPH4A [c.921 + 3_921 + 6del (Intronic)] founder mutation who also had a heterozygous Variant of Unknown Significance (VUS) in DNAH5 [c.1250C > G (p.Thr417Ser)]. HRCT showed bibasilar bronchiectasis and parenchymal scarring tissue more evident at the LLL. (c). 13-year-old male with two pathogenic heterozygous genetic variants in RSPH4A [c.921 + 3_921 + 6del (Intronic)] and RSPH4A [c.1103T > G (p.Val368Gly)] and one heterozygous VUS in DNAH8 [c.9839A > T (p.Gln3280Leu)]. HRCT showed cylindrical and varicose bilateral bronchiectasis, multiple centrilobular nodules, and tree-in-bud opacities. (d). A 33-year-old female with one heterozygous RSPH4A [c.902A > C (p.Gln301Pro)] VUS who showed extensive bilateral bronchiectasis, reticular-nodular infiltrates with tree-in-bud changes at RML and RLL. Case 4 also had an heterozygous VUS on DNAAF3 [c.1672G > T (p.Glu558*)] and DNAH11 [c.11678C > T (p.Thr3893Met)]. All four cases had central apparatus abnormalities on the cilia ultrastructure examined by electron microscopy.

3.5. Pulmonary Function Profile

Pulmonary function test was reviewed on subjects above age 5. Airflow limitation was noted abnormal in a total of 78%, (7/9). Obstructive and restrictive ventilatory defects were present in 55%, (5/9) and 22%, (2/9) of patients, respectively as defined by spirometry classification. Post bronchodilator challenge was positive (increase in [Forced Expiratory Volume in 1 s], FEV1 > 12%) in 40%, (2/5) of those patients with baseline airway flow obstruction. Residual volume (RV), RV/TLC, and specific airway resistance (sRaw) were noted elevated in 100%, (5/5) of patients.
4. Discussion

Diagnostic challenges in rare lung disorders like PCD could be numerous depending on the prevalence, clinical phenotype, and genetic variability of the population to be studied. In Puerto Rico, there is a lack of information about rare lung diseases, including PCD. The actual prevalence of PCD on the island is unknown. Considering a prevalence of PCD of 1:15,000 and the actual population of Puerto Rico to be around 2.8 million individuals, we can estimate a population of 187 patients living with PCD on the island. This estimation doesn’t count the presence of a founder genetic mutation on the RSPH4A gene previously described [14], which may contribute to an increased amount of patients living in Puerto Rico with PCD. The fact that in 24 months our clinic was able to diagnose a total of 13 patients with RSPH4A, shows that PCP in Puerto Rico has been underdiagnosed. We understand that genetic testing and electron microscopy are limitations for the Puerto Rican population to follow the established diagnostic algorithms as recommended by the PCD foundation in the United States. The lack of nNO test as a screening and diagnostic PCD tool for Puerto Rico and the Caribbean are limitations in some challenging cases. The genetic variability of the Puerto Rican population introduces additional challenges as commercial genetic testing panels with 35 genes may not be able to unmask rare PCD genetic variants as pathogenic to confirm the diagnosis when more than 45 PCD genes have been described. For instance, VUS genetic variants may need to be analyzed taking into consideration the clinical phenotype of PCD patients in Puerto Rico. To date a total of 36 cases are reported on LOVD3 (Leiden Open Variation Database), (www.lovd.nl/RSPH4A) for RSPH4A mutations. A total of 17 unique cases are reported for the RSPH4A [c.921 + 3_6delAAGT] mutation with Puerto Rican heritage. We add a total of new 13 RSPH4A cases and the other 11 RSPH4A [c.921 + 3_6delAAGT] unique mutations from native Puerto Ricans.

In this article, we describe 13 patients, from 11 unrelated families, who were evaluated for PCD and presented RSPH4A mutations on genetic testing. Although 8 of 13 (62%) patients were positive for the presence of homozygous genetic mutations on RSPH4A [c.921 + 3_6delAAGT], 5 of 13 patients showed uncertain results, classified as VUS. While we know that RSPH4A [c.921 + 3_6delAAGT] genetic mutation has been detected on Hispanics living in the United States with Puerto Rican ancestry, our study confirms this observation in native Puerto Ricans with clinical PCD symptoms. Interestingly, patients with biallelic pathogenic variants in RSPH4A [c.921 + 3_6delAAGT] mutation showed most of the clinical PCD features including neonatal respiratory distress at birth, bronchiectasis, and sinus diseases as the main phenotype. However, chronic secretory otitis media was presented only in 33%, (2/6) of individuals with biallelic homozygous RSPH4A [c.921 + 3_6delAAGT] mutation. As compared with other genetic variants published [14,16], we saw fewer chronic secretory otitis media cases. Central complex defects on electron microscopy have been associated with severe otologic features in children with PCD [17]. Our results showed that central complex defects were present in 100% of the cohort. However, the severity of otologic features was not as prevalent as expected. Laterality defects were absent in our cohort as previously documented on mutations that affect the central complex apparatus. We know that RSPH4A mutations are not necessarily a requirement for abnormal left-right axis orientation [12].

The genetic diversity of Puerto Rican ancestry may explain the range and combination of VUS genetic variants found. Three patients (Cases 4, 7 and 8) were “purely” homozygous for the RSPH4A [c.921 + 3_6delAAGT] mutation. All other patients had a combination of pathogenic RSPH4A with other PCD related genetic heterozygous VUS. The interactions of VUS with pathogenic PCD variants are unknown. Additional studies are needed to explore the role of digenic inheritance in PCD as seen in other ciliopathies [18]. On the other hand, we show in Table 1A, four cases classified as compound heterozygous genetic variants with clinical criteria for PCD. Initially, on Case 9, PCD genetic tests revealed one pathogenic variant at the RSPH4A [c.921 + 3_6delAAGT] mutation. Additional one likely pathogenic variant was noted: RSPH4A [c.1103T > G (p.Val368Gly)] plus a VUS
on DNAH8 [c.9839A > T (p.Gln3280Leu)]. Familial PCD genetic studies on Case 7 showed that RSPH4A [c.1103T > G (p.Val368Gly)] variant was maternally inherited. No paternal RSPH4A or DNAH8 variants were detected. As a result of the family analysis, genetic reclassification was made for a VUS in RSPH4A [c.1103T > G (p.Val368Gly)] PCD genetic variant as pathogenic. It is important to fully understand the interaction of multiple heterozygous variants in the clinical phenotype of patients with PCD. Analysis of familial studies plays an important role in the reclassification of VUS PCD genetic variants to get a point score threshold for pathogenicity.

Another atypical patient is presented in Case 10, which exhibited two of the main features: daily wet cough and year-round daily nasal congestion but lack laterality defects or recurrent secretory otitis media. Electron microscopy findings showed central apparatus defects and HRCT revealed extensive bilateral bronchiectasis, reticular-nodular infiltrates with tree-in-bud changes at RML and RLL (Figure 2c). Pulmonary function test demonstrated severe airflow obstruction with an FEV1: 24%. Genetic testing resulted in three VUS genetic variants including RSPH4A, DNAAF3, and DNAH11. Although none of these genetic VUS had been associated with PCD pathogenic phenotype; the medical history, clinical findings, electron microscopy, and the severity of her pulmonary disease evidenced by pulmonary function test and imaging, highly suggest PCD. Ancillary testing like nNO, if available, will help to confirm challenges in PCD cases like this. Two rare RSPH4A variants were detected in our cohort. First, the RSPH4A [c.1103T>G], which replace a valine with glycine at the codon 368 of the RSPH4A protein. This missense mutation found in Case 9 promotes physicochemical differences between both amino acids. Although the effect of this missense mutation is unclear, this variant is currently classified as pathogenic and seen in individuals with PCD phenotype. The second variant found on Case 10 is RSPH4A [c.902A > C], which replace glutamine with proline at the codon 301. This missense mutation was not previously reported on individuals with RSPH4A-related diseases and is currently classified as VUS. Additional testing and evaluation including nNO and electron microscopy will be needed to confirm the diagnosis of PCD.

5. Conclusion

In Puerto Rico, PCD diagnosis remains under-recognized by the general population and healthcare providers. To understand the presence of a PCD founder mutation on the island, it is important to address the medical history of recurrent pulmonary infections, bronchiectasis, and sinus disease during the initial encounter. Although asthma may be a concomitant disease in PCD, a detailed history of PCD symptoms is vital in those with severe persistent asthma not responding to standard care. Referral to knowledgeable medical doctors with PCD expertise is imperative for early patient diagnosis that will prevent or delay additional PCD comorbidities later in life. Awareness about PCD clinical features on the island must be a priority for early recognition of potential PCD cases. Collaboration among neonatologists, otorhinolaryngologists, pediatricians, infertility clinics, pulmonologists, and radiologists, among other providers, is essential to reveal the actual prevalence of PCD on the island. The presence of a founder mutation in RSPH4A warrants the development of a multidisciplinary center for PCD for the evaluation of suspected cases in Puerto Rico. The development of a PCD center is urgently needed to explore new founder mutations in the genetic pool of individuals living in the Caribbean and Latin-America using state-of-the-art diagnostic tools following ATS and PCD foundation diagnosis guidelines. Collaboration with PCD Centers in the United States is important considering that, to date, we do not have the capacity for lung transplantation in Puerto Rico and lung transplantation in Latin-America is limited.
Author Contributions: All authors were involved in the design of the study and editing of this article as follow: Conceptualization, W.D.J.-R.; methodology, W.D.J.-R. and R.A.M.; data analysis and curation, W.D.J.-R., and D.R.-D.J.; writing—original draft preparation, W.D.J.-R.; writing—review and editing, W.D.J.-R., D.R.-D.J., and R.A.M.; supervision, R.A.M. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded by the Hispanic Center of Excellence, University of Puerto Rico School of Medicine, Grant Number: D34HP24463, U.S. Department of Health and Human Services, Health Resources and Services Administration, Bureau of Health Workforce.

Institutional Review Board Statement: The study was conducted according to the guidelines of the University of Puerto Rico, Medical Sciences Campus, and approved by the Institutional Review Board of the University of Puerto Rico, Medical Science Campus, Department of Pediatrics protocol code B1730120 on October 13, 2020.

Informed Consent Statement: Patient consent was waived by IRB due to the methodology used in this article: a retrospective chart review.

Data Availability Statement: All data analyzed during this study are included in this published article.

Conflicts of Interest: The authors declare no conflicts of interest. The funders had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript, or in the decision to publish the results.

Abbreviations

PCD: Primary ciliary dyskinesia
nNO: nasal nitric oxide
HRCT: High-resolution chest computer tomography
CFTR: Cystic fibrosis transmembrane conductance regulator
GLI: Global lung function initiative-2012
LLL: Left lower lobe
RLL: Right lower lobe
RML: Right middle lobe
RUL: Right upper lobe
VUS: Variant of unknown significance
FEV1: Forced Expiratory Volume in 1 second
RV: Residual Volume
TLC: Total Lung Capacity
sRAW: Specific airway resistance
RDS: Respiratory Distress Syndrome
| Case | Sex | Age (y) | Laterality | Ciliary Defects | Ethnicity | Chronology | Neonatal RDS | Bronchiectasis on HRCT | Chronic Rhinorrhea | Chronic Secretory Otitis | Hearing Loss | FEV1 (%) | PCD | Affected Base | Amino-Acid Change | Zygosity | Classification |
|------|-----|---------|------------|----------------|-----------|------------|-------------|----------------------|------------------|------------------------|--------------|-----------|-----|----------------|-----------------|----------|----------------|
| 1    | M   | 1       |            |                | Hispanic  |            |             |                      |                  |                        |              |           |     | RSPH4A         | c.921 + Intronic  | Homo      | Pathogenic     |
|      |     |         |            |                |           |            |             |                      |                  |                        |              |           |     | CCDC15         | 3_921 + Exons 9-11 | Hete      | VUS            |
|      |     |         |            |                |           |            |             |                      |                  |                        |              |           |     | DNAH5          | 6del c.921 +         | Hete      | VUS            |
|      |     |         |            |                |           |            |             |                      |                  |                        |              |           |     | RSPH4A         | c.921 + Intronic  | Homo      | Pathogenic     |
|      |     |         |            |                |           |            |             |                      |                  |                        |              |           |     | DNAH11         | 3_921 + Exons 9-11 | Hete      | Pathogenic     |
|      |     |         |            |                |           |            |             |                      |                  |                        |              |           |     | CCDC11         | 6del c.921 +         | Hete      | Pathogenic     |
|      |     |         |            |                |           |            |             |                      |                  |                        |              |           |     | RSPH4A         | c.921 + Intronic  | Homo      | Pathogenic     |
|      |     |         |            |                |           |            |             |                      |                  |                        |              |           |     | DNAH11         | 6del c.921 +         | Hete      | VUS            |
|      |     |         |            |                |           |            |             |                      |                  |                        |              |           |     | GAS8           | c.921 + Intronic  | Homo      | VUS            |
| 4    | M   | 15      |            |                | Hispanic  |            |             |                      |                  |                        |              |           |     | RSPH4A         | c.921 + Intronic  | Homo      | Pathogenic     |

**Table A1:** Demographics and PCD Genetic Variants in 13 PCD patients from Puerto Cas"
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
| 5 | M | 51 | 0 | CA | Hispanic | Y | Y | Y | Y | N | Y | 62 | RSPH4A | c.921 | + | Intrinsic | Homo | Pathogenic |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   | DNAH5 | 3_921 | + | p.Thr417Ser | Hete | VUS |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   | 6del | c.1250C > G |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
| 6 | F | 35 | 0 | CA | Hispanic | Y | Y | Y | Y | N | Y | 36 | RSPH4A | c.921 | + | Intrinsic | Homo | Pathogenic |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   | DNAAF2 | 3_921 | + | p.Val20Leu | Hete | VUS |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   | 6del | c.58G > C |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
| 7 | F | 26 | 0 | CA | Hispanic | Y | Y | - | Y | N | - | 53 | RSPH4A | c.921 | + | Intrinsic | Homo | Pathogenic |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   | 3_921 | + |   |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   | 6del |   |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
| 8 | F | 19 | 0 | CA | Hispanic | Y | N | Y | Y | N | Y | 59 | RSPH4A | c.921 | + | Intrinsic | Homo | Pathogenic |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   | 3_921 | + |   |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   | 6del |   |

**Compound Heterozygous Genetic Variants**

|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| 9 | M | 13 | 0 | CA | Hispanic | Y | Y | Y | Y | Y | Y | 60 | RSPH4A | c.921 | + | Intrinsic | Homo | Pathogenic |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   | DNAH5 | 3_921 | + | p.Val368Gly | Hete | VUS |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   | 6del | p.Gln3280Leu |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
| 10 | F | 33 | 0 | CA | Hispanic | Y | N | Y | Y | N | - | 24 | RSPH4A | c.902A > C | + | Intrinsic | Homo | VUS |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   | DNAAF3 | 1672G > A | + | p.Glu558* | Hete | VUS |
|   |   |   |   |   |   |   |   |   |   |   |   |   |   | DNAH11 | 11678C > T | + | p.Thr3893Me | Hete | VUS |
|   | F/M | Age | Gender | Hispanic | 50- | 50- | 50- | 50- | 50- | 50- | 50- | Pathogenicity |
|---|-----|-----|--------|----------|-----|-----|-----|-----|-----|-----|-----|----------------|
| 11 | F   | 5   | 0      | -        | Hispanic | Y   | N   | N   | Y   | Y   | Y   | - RSPH4A c.921 + Intronic Hete Pathogenic |
|    |     |     |        |          |          | 3_921 | + | p.Glu257Asp Hete VUS |
|    |     |     |        |          |          | del | 6del | p.Leu3216Val Hete VUS |
|    |     |     |        |          |          | c.971T > G | c.9646C > A | p.Met1096Ile Hete VUS |
|    |     |     |        |          |          | c.3288G > T | |
| 12 | M   | 2   | 0      | CA       | Hispanic | Y   | Y   | -   | Y   | N   | -   | - RSPH4A c.921 + Intronic Hete Pathogenic |
|    |     |     |        |          |          | 3_921 | + | p.Ser62Arg Hete VUS |
|    |     |     |        |          |          | del | 6del | p.Ser2963Tyr Hete VUS |
|    |     |     |        |          |          | c.186C > G | c.12709G > T | p.Val4237Phe Hete VUS |
|    |     |     |        |          |          | c.8888C > A | |

**Heterozygous Genetic Variants.**

|   | F   | 3   | 0      | -        | Hispanic | Y   | N   | -   | Y   | N   | -   | - RSPH4A c.902A > C p.Gln301Pro Hete VUS |

F/M: Female/Male, R*: Rare Cilia present, not diagnostic, CA: Central Apparatus defects, OM: Otitis Media, F#: Infertile Female, HRCT: High-Resolution Chest CT, FEV1: Forced Expiratory Volume in 1 s. RDS: Respiratory Distress Syndrome. Pathogenic*: Reclassified as pathogenic after parental genetic testing.
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