TITLE: Novel variants identified in *CKAP2L* in two siblings with Filippi Syndrome

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RUNNING TITLE: Novel variants in Filippi Syndrome

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ABSTRACT

Pathogenic variants in *CKAP2L* have previously been reported in Filippi Syndrome (FS), a rare autosomal recessive, craniodigital syndrome characterized by microcephaly, syndactyly, short stature, intellectual disability, and dysmorphic facial features. To date, fewer than ten patients with pathogenic variants in *CKAP2L* associated with FS have been reported. All of the previously reported probands have presumed loss-of-function variants (frameshift, canonical splice site, starting methionine) and all but one have been homozygous for a pathogenic variant. Here we describe two brothers who presented with microcephaly, micrognathia, syndactyly, dysmorphic features, and intellectual disability. Whole exome sequencing of the family identified a missense variant, c.2066G>A;p.(Arg689His), in *trans* with a frameshift variant, c.1169_1173del;p.(Ile390LysfsTer4), in *CKAP2L*. To our knowledge, these are the first patients with FS to be reported with a missense variant in *CKAP2L* and only the second family to be reported with two variants in *trans*.

CASE PRESENTATION

The male proband was first seen at 14-years-old presenting with microcephaly, dysmorphic facial features including a broad forehead, micrognathia, and thin lips, abnormal hair growth patterns including broad eyebrows and long eyelashes, developmental delay, serrated incisors, 2-3 toe syndactyly, and a seizure disorder. Seizures began at age three. EEGs showed an abundance of epileptiform activity, sometimes occupying 90% of the record. He was prescribed Lamictal and has not had a seizure in the past 6 years. His most recent head circumference measurement at age 17 was 50.2 cm, a -3.8 standard deviation, and height was in the third percentile. In addition, he had small testes secondary to congenital cryptorchidism, which was repaired via orchiopexy at age 1. He was born at 41 weeks after an uncomplicated pregnancy weighing 3260g. Case two is the five-year-old brother of the proband, who is similarly affected with intellectual disability, dysmorphic facial features, microcephaly, small testes, syndactyly of his 2nd and 3rd toes, and congenital cryptorchidism. Given his brother's seizure disorder, an EEG analysis was performed which found abnormal EEG activity putting him at a higher seizure risk. However, seizures have never been documented, and he is not on an anticonvulsant. No further EEGs have been performed. He was born at 40 weeks after an uncomplicated pregnancy weighing 3266g. He has undergone a right inguinal orchiopexy, right inguinal hernia repair, and frenotomy. His most recent head circumference measurement at age eight years four months was 48.1 cm, a -3.5 standard deviation, and height was in the sixth percentile. Both siblings' cases of toe syndactyly were surgically repaired. The constellation of features in both patients are consistent with the common features shared amongst patients with Filippi syndrome (see Supplemental Figure 1 and Table 1). A third male sibling and both parents were asymptomatic and did not share any features with the two affected siblings. No consanguinity was reported in the family.

TECHNICAL ANALYSIS

An Affymetrix CytoScan HD chromosomal microarray with a resolution of 30 kb for deletions and 60 kb for duplications was performed on the proband's younger brother by the Mayo Medical Laboratory. Data was analyzed and reported using human genome build 37.1 (hg19) and did not reveal significant copy number changes. Whole Exome sequencing was performed by the Sanford Medical Genetics Laboratory on genomic DNA isolated from both siblings, mother, and father. The unaffected sibling was not tested. Exome capture was performed using the SureSelect Clinical Research Exome kit V1 (Agilent Technologies) and sequencing was performed on an Illumina HiSeq2500 both according to manufacturers' protocols. The sequence reads were converted to FASTQ format. Alignment, variant calling, and analysis were performing using Codified software utilizing BWA to align to the reference human genome (hg38) and platypus to call variants. Data was interpreted and classified according to ACMG guidelines (Richards et al. 2015). Both affected siblings were found to be compound heterozygous for a paternally inherited frameshift variant, c.1169_1173del;p.(Ile390LysfsTer4), and a maternally inherited missense variant, c.2066G>A;p.(Arg689His), in the *CKAP2L* gene (Table 2). Both variants and *trans* phase in the siblings were confirmed by targeted Sanger sequencing analysis of all family members.

VARIANT INTERPRETATION

CKAP2L contains nine exons that encode a 745 amino acid protein. The C-terminal 319 amino acids encode for a cytoskeleton-associated protein-2 domain that is homologous to CKAP2, a microtubule-associated protein involved in mitotic progression (Seki and Fang 2007). To date, eight other frameshift

variants in *CKAP2L* have been reported in the literature in affected individuals (Hussain et al. 2014; Capecci et al. 2018; Sabir et al. 2019; Karakaya et al. 2021). All are suspected of leading to premature termination (PT) and nonsense mediated decay (NMD) and are found in the N-terminal half of the protein that does not encode for the CKAP2 domain, with a majority found in exon 4. The c.1169_1173del;p.(Ile390LysfsTer4) variant is consistent with these previously reported variants and is also predicted to result in NMD. It is therefore also classified as a pathogenic variant.

In contrast, the c.2066G>A;p.(Arg689His) variant is located in the last exon of the gene. It is present in the gnomAD (Genome Aggregation Database) population database only in the European (non-Finnish) population (8/113738 alleles). The arginine at position 689 is highly conserved in vertebrate species from humans to zebrafish. Prediction algorithms are discordant as to the pathogenicity of this variant (SIFT: Deleterious; PolyPhen-2: Probably Damaging; Align-GVGD: Class C0) and there are currently no studies about the effect of this variant on protein function. Therefore, the c.2066G>A;p.(Arg689His) variant is classified as a variant of uncertain significance (VUS).

DISCUSSION

FS was first described in 1985 in three Sardinian siblings with syndactyly, microcephaly, shared facial abnormalities, and growth and developmental delays (Filippi 1985). Since the original description, fewer than 40 cases have been reported. Additional features such as cleft palate, hypertrichosis, coarse hair, seizures, cerebellar atrophy, precocious puberty, and multiple dental and skeletal abnormalities have been reported in patients suggesting a variable clinical presentation (Orrico and Hayek 1997; Walpole et al. 1999; Schorderet et al. 2002; Sharif and Donnai 2004; Battaglia et al. 2008; Sandhu et al. 2013; Sabir et al. 2019). However, it is important to note that many of these patients either did not have molecular testing or are consanguineous and therefore these additional features may be caused by variants in other genes resulting in a blended phenotype. The molecular identification of variants in *CKAP2L* provides a definitive diagnosis.

At the cellular level, loss of *CKAP2L* protein causes a defect in spindle organization and subsequent disruption of chromosomal segregation (Hussain et al. 2014). However, studies on the mouse homolog gene *radmis* also suggest a gain-of-function mechanism. Yumoto et al. showed that overexpression of *CKAP2L in vitro* also results in defects in spindle formation and mitotic arrest through hyper-stabilization of microtubules (Yumoto et al. 2013). The role of *CKAP2L* in neural progenitor cell division as well as a potential role in regulating apoptosis, purportedly leads to the phenotypic microcephaly and syndactyly of FS, respectively (Hussain et al. 2014).

In summary, we present here two siblings with novel *CKAP2L* variants associated with FS. This is only the second family reported with two different variants in *trans*. Although the frameshift variant is consistent with previously reported pathogenic variants, to our knowledge this is the first missense variant in *CKAP2L* reported to be associated with FS. It is possible that the c.2066G>A;p.(Arg689His) variant leads to a more stable protein resulting in the gain-of-function phenotype observed in the mouse studies. However, it is also possible that there is a yet undetected causative variant in *cis* with the missense variant. While one of the siblings had a negative SNP microarray result, the possibility of a genomic deletion too small to be detected by array yet also too large to be detected by exome sequencing cannot be ruled out. Additionally, the presence of a deep intronic variant leading to abnormal mRNA splicing in *cis* with the c.2066G>A;p.(Arg689His) also cannot be ruled out. Additional studies are

warranted to determine what affect the c.2066G>A;p.(Arg689His) VUS has on cell spindle organization and chromosomal segregation.

ADDITIONAL INFORMATION

Data Deposition and Access

The c.1169_1173del;p.(Ile390LysfsTer4) and c.2066G>A;p.(Arg689His) variants were submitted to ClinVar and can be found under accession numbers VCV001292042.1 and VCV001040806.1 respectfully.

Ethics Statement

This study was approved by the Sanford Research institutional review board and all participants provided written informed consent for genetic testing. The family provided written consent for publication of identifiable images.

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Author Contributions

R.P. and M.L. wrote the manuscript. L.D.K. and J.W. reviewed the manuscript. M.L. analyzed and interpreted the sequencing data. L.D.K. oversaw patient care.

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Competing Interest Statement

The authors have declared no competing interest.

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Table 1.

	Filippi Syndrome Clinical Features^a	Proband	Sibling	Comments
Growth	Low Weight	Y	Y	
	Intrauterine Growth Retardation	N	N	
	Postnatal Growth Retardation	Y	Y	
Head & Neck	Microcephaly	Y	Y	
- Face	Broad Forehead	Y	Y	
	Hairy Forehead	N	N	
	Short Philtrum	N	N	
- Eyes	Poor Vision	N	N	
	Retrobulbar Venous Varix	N	N	
	Proptosis	N	N	
	Optic Atrophy	N	N	
- Nose	Prominent columella	N	N	
	Broad Nasal Bridge	N	N	
	Hypoplastic Alae Nasi	N	N	
- Mouth	Straight Mouth	N	N	
	Thin Lips	Y	Y	
	Small Teeth	N	N	
	Abnormally Shaped Teeth	Y	Y	
	Hypodontia (Rare)	N	N	
	Serrated Incisors (Rare)	Y	Y	
Cardiovascular	Ventricular Septal Defect	N	N	Echocardiography performed to verify absence following negative auscultation.
Genitourinary	Cryptorchidism	Y	Y	
	Ambiguous Genitalia			
Skeletal	Bilateral Cutaneous Syndactyly of Second, Third, and Fourth Fingers	N	N	
- Hands	Fifth-finger Clinodactyly	Y	Y	
- Feet	Bilateral Cutaneous Syndactyly of Second and Third toes	Y	Y	2nd and 3rd - Surgically repaired in both
	Hypoplasia of Fifth Toes			
Hair	Abnormal Hair Growth Pattern	Y	Y	
	Sparse Hair (Rare)	N	N	
	Hypertrichosis (Rare)	N	N	
Neurologic	Developmental Delay	Y	Y	

	Seizures	Y	N	Sibling had abnormal EEG and is therefore high seizure risk, but has not had seizures.
	Dystonic Movements	N	N	
	Dystonic Tongue Protrusion	N	N	
	Cerebellar Atrophy	Unknown	Unknown	
	Arachnoidal Cyst	Unknown	Unknown	
	Diffuse Enlargement of Subarachnoid Spaces and Lateral Ventricles (Rare)	Unknown	Unknown	
	Megacisterna Magna (Rare)	Unknown	Unknown	
	MRI abnormalities	Unknown	Unknown	
^a The list of clinical features is based on the OMIM clinical synopsis (#272440)				

Table 2. Genomic Findings

Gene	Genomic Location	HGVS cDNA	HGVS Protein	Zygoty	Parent of Origin	Variant Interpretation
<i>CKAP2L</i>	GRCh37: Chr2:113513775_113513782 GRCh38: Chr2:112756198-112756205	NM_152515.5 c.1169_1173del	p.Ile390LysfsTer4	Heterozygous	paternal	Pathogenic
<i>CKAP2L</i>	GRCh37: Chr2:113496572 GRCh38: Chr2:112738995	NM_152515.5 c.2066G>A	p.Arg689His	Heterozygous	maternal	Variant of Uncertain Significance

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