REPORT

TAF1 Variants Are Associated with Dysmorphic Features, Intellectual Disability, and Neurological Manifestations

Jason A. O’Rawe,1,2 Yiyang Wu,1,2 Max J. Dörfel,1 Alan F. Rope,3 P.Y. Billie Au,4 Jillian S. Parboosingh,4 Sungjin Moon,5 Maria Kousi,5 Konstantina Kosma,6,7 Christopher S. Smith,4 Maria Tzetis,6,7 Jane L. Schuette,6,9 Robert B. Hufnagel,10,11 Carlos E. Prada,10,12 Francisco Martinez,13 Carmen Orellana,13 Jonathan Crain,1 Alfonso Caro-Llopis,13 Silvestre Oltra,13 Sandra Monfort,13 Laura T. Jiménez-Barrón,1,14 Jeffrey Swensen,15 Sara Ellingwood,16 Rosemarie Smith,17 Han Fang,1 Sandra Ospina,18 Sander Stegmann,19 Nicolette Den Hollander,20 David Mittelman,21 Gareth Highnam,21 Reid Robison,22,23 Edward Yang,24 Laurence Faivre,25,26 Agathe Roubertie,27,28 Jean-Baptiste Rivière,29 Kristin G. Monaghan,29 Kai Wang,30 Erica E. Davis,8 Nicholas Katsanis,5 Vera M. Kalscheuer,31 Edith H. Wang,32 Kay Metcalfe,33 Tjitske Kleefstra,34 A. Micheil Innes,4 Sophia Kitsiou-Tzeli,6 Monica Rosello,13 Catherine E. Keegan,8,9 and Gholson J. Lyon1,2,22,2, *

We describe an X-linked genetic syndrome associated with mutations in TAF1 and manifesting with global developmental delay, intellectual disability (ID), characteristic facial dysmorphism, generalized hypotonia, and variable neurologic features, all in male individuals. Simultaneous studies using diverse strategies led to the identification of nine families with overlapping clinical presentations and affected by de novo or maternally inherited single-nucleotide changes. Two additional families harboring large duplications involving TAF1 were also found to share phenotypic overlap with the probands harboring single-nucleotide changes, but they also demonstrated a severe neurodegeneration phenotype. Functional analysis with RNA-seq for one of the families suggested that the phenotype is associated with downregulation of a set of genes notably enriched with genes regulated by E-box proteins. In addition, knockdown and mutant studies of this gene in zebrafish have shown a quantifiable, albeit small, effect on a neuronal phenotype. Our results suggest that mutations in TAF1 play a critical role in the development of this X-linked ID syndrome.

Transcription factor II D (TFIID) consists of TATA binding protein (TBP) and 12–14 TBP-associated factors (TAFs). TFIID promotes transcriptional initiation by recognizing promoter DNA and facilitating the nucleation of other general transcription factors for assembly into a functional pre-initiation complex,1–5 and it also functions as a co-activator by interacting with transcriptional activators.1 Subunits of TFIID have been suggested to play a possible role in neurodegenerative diseases and developmental delay when they are disrupted.5–12 Indeed, variants in TAF2 (MIM: 604912) and TBP (MIM: 600075) have been implicated in intellectual disability (ID) and developmental delay with or without corpus callosum hypoplasia.10,11,13 Recent work toward understanding the molecular basis of Cornelia de Lange syndrome (CdLS [MIM: 122470, 300590, 300882, 614701, 610759]) has also suggested

1Stanley Institute for Cognitive Genomics, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY 11724, USA; 2Graduate Program in Genetics, Stony Brook University, Stony Brook, NY 11794, USA; 3Department of Medical Genetics, Northwestern Keller Permanentine, Portland, OR 97227, USA; 4Department of Medical Genetics and Alberta Children’s Hospital Research Institute, Cumming School of Medicine, University of Calgary, Calgary, AB T2J 4L6, Canada; 5Center for Human Disease Modeling, Duke University, Durham, NC 27708, USA; 6Department of Medical Genetics, Medical School, University of Athens, Athens 11527, Greece; 7Research Institute for the Study of Genetic and Malignant Disorders in Childhood, Aghia Sophia, Children’s Hospital, Athens 11527, Greece; 8Department of Pediatrics, Division of Genetics, University of Michigan, Ann Arbor, MI 48109, USA; 9Department of Human Genetics, University of Michigan, Ann Arbor, MI 48109, USA; 10Division of Human Genetics, Cincinnati Children’s Hospital, Cincinnati, OH 45229, USA; 11Unit on Pediatric, Developmental, and Genetic Ophthalmology, Ophthalmic Genetics and Visual Function Branch, National Eye Institute, NIH, Bethesda, MD 20892, USA; 12Centro de Medicina Genómica y Metabolismo, Fundación Cardiovascular de Colombia, Bucaramanga, Santander 681004, Colombia; 13Unidad de Genética, Hospital Universitario y Politécnico La Fe, Valencia 46026, Spain; 14Centro de Ciencias Genomicas, Universidad Nacional Autonoma de Mexico, Cuernavaca, Morelos 62210, Mexico; 15Cine Genomic, Arizona, AZ 85040, USA; 16Division of Genetics, Pediatric Specialty Care, Maine Medical Partners, ME 04102, USA; 17Department of Pediatrics, Division of Genetics, Barbara Bush Children’s Hospital, Maine Medical Center, Portland, ME 04102, USA; 18Fundacion CardioInfantil and Universidad del Rosario, Bogota 681004, Colombia; 19Department of Clinical Genetics, 6202 AZ Maarstricht, the Netherlands; 20Department of Clinical Genetics, Leiden University Medical Center, 2300 RC Leiden, the Netherlands; 21Gene by Gene Ltd., Houston, TX 77008, USA; 22Utah Foundation for Biomedical Research, Salt Lake City, UT 84107, USA; 23Vite Genomics, 150 South 100 West Street, Provo, UT 84601, USA; 24Department of Radiology, Boston Children’s Hospital, Boston, MA 02115, USA; 25Génétique des Anomalies du Développement EA4271, Université Bourgogne Franche-Comté, 21000 Dijon, France; 26Centre de Génétique, Centre de Référence Maladies Rares “Anomalies du Développement et Syndromes Malformatifs,” Centre Hospitalier Universitaire de Dijon, 21000 Dijon, France; 27Institut des Neurosciences de Montpellier, Université de Montpellier, BP 74103, 34950 Montpellier Cedex 5, France; 28Service de Neuropédiatrie, Centre Hospitalier Universitaire Gui de Chauliac, 34000 Montpellier, France; 29GeneDx, Gaithersburg, MD 20877, USA; 30Zilkha Neurogenetic Institute, Department of Psychiatry and Preventive Medicine, University of Southern California, Los Angeles, CA 90089, USA; 31Research Group Development and Disease, Max Planck Institute for Molecular Genetics, 14195 Berlin, Germany; 32Department of Pharmacology, University of Washington, Seattle, WA 98195, USA; 33Clinical Genetics Service, Manchester Centre for Genomic Medicine, Saint Mary’s Hospital, Central Manchester University Hospitals NHS Foundation Trust, Oxford Road, Manchester M13 9WL, UK; 34Department of Human Genetics, Radboud University Medical Center, 6500 HB Nijmegen, the Netherlands

*Correspondence: gholsonjlyon@gmail.com
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that mutations in TAF6 (MIM: 602955), encoding a component of TFII D, play an important role in the pathogenesis of this syndrome.14 CdLS is a phenotypically and genetically heterogeneous syndrome characterized by distinct facial features, hirsutism, developmental delay, ID, and limb abnormalities,15 and mutations in several different genes have been implicated in contributing to this heterogeneous clinical presentation.16–19 Two mutations in TAF6 were implicated in disease pathogenesis in some individuals with phenotypic features of CdLS and were shown, through biochemical investigations, to reduce binding of TAF6 to other core components of TFII D.14

A recent paper nominated TAFI (MIM: 313650) as a candidate gene for ID on the basis of segregation of missense variants in two different pedigrees; however, no clinical information other than ID was provided.20 TAF1 encodes the largest subunit of the TFII D complex and has been ranked 53rd among the top 1,003 constrained human genes in a recent population-scale study,21 suggesting that TAF1 plays a critical role in normal cellular function. Previous work in Drosophila cells has shown that TAF1 depletion increases the magnitude of the initial transcription burst and causes delay in the shutoff of transcription upon removal of the stimulus.22 The authors showed that the magnitude of the transcriptional response to the same signaling event, even at the same promoter, can vary greatly depending on the composition of the TFII D complex in the cell. In addition, and consistent with the notion that TAFI is important in controlling the binding patterns of TFII D to specific promoter regions, this study not only demonstrates severe and progressive disease-related sequence variation. These included whole-genome sequencing, exome sequencing, targeted gene-panel sequencing, and microarray-based strategies (Figures S1–S5 and Tables S1–S4). Sanger sequencing was used for validating sequence variations. Many of the families studied here underwent genotyping via clinical microarrays or gene-specific sequencing for a small number of genomic regions. See the Supplemental Data for more-detailed descriptions of the sequencing and analysis methods used for each family. All 14 affected individuals were found to contain TAF1 sequence variants, the majority of which (11 out of 14) are missense variants. One proband (II-1 in
family 5; 5A in Figure 2) was found to have a variant that influences TAF1 splicing, and two probands (10A and 11A) were found to have a duplication that involves TAF1. As stated above, the two probands with duplications exhibit less overlap with those who harbor single-nucleotide changes (see Table 1) and exhibit severe progressive neurologic impairment. All of the mutations reported here, including the duplications, are de novo or co-segregate with the phenotype in other affected male individuals (see Figures 2 and 3).

Families 1 and 10 were tested for X chromosome inactivation, which showed that female carriers of TAF1 mutations and duplications demonstrate highly skewed inactivation (99:1). The female carriers in families 5 and 11 were not informative for the polymorphic CAG repeat in the human androgen-receptor gene.

All missense variants were found to affect evolutionarily conserved residues (Figure 2B) and were not present at any frequency in public databases such as dbSNP137, 1000 Genomes phase 1, the NHLBI Exome Sequencing Project Exome Variant Server (ESP6500), or ExAC Browser version 0.2, which contains allelic information derived from ~60,000 exome sequences. The TAF1 missense variants were also predicted to be deleterious by a range of prediction software (CADD, SIFT, GERP++, and phyloP) (Figure 2B). The splice-site variant discovered in family 5 was not predicted to be deleterious by the prediction software listed. However, this variant does not change the amino acid content of the predicted protein; instead, it affects TAF1 splicing in both the mother and the proband (Figure S11). In addition, the single-nucleotide variants (SNVs) described here fall within TAF1 regions that are relatively sparsely covered by non-synonymous sequence variations reported in the ExAC Browser (Figure 2C). Indeed, four of these missense variants (c.2419T>C [p.Cys807Arg], c.2926G>C [p.Asp976His], c.3736C>T [p.Arg1246Trp], and c.3708A>G [p.Arg1228Ilefs*16]) were found to be within the two TAF1 regions that are significantly underrepresented by non-synonymous sequence variation reported in the ExAC Browser when restricted to European and Latin populations (Figure 2C). Additional variants found in families 6 and 7 (c.4549A>C [p.Asn1517His] and c.4355G>A [p.Arg1431His]) also fall within a region that is underrepresented by non-synonymous sequence variation, although this last cluster is not statistically significant (p value = 0.29) (see Supplemental Data for further details).

Recent structural work in yeast points to an epigenetic role of the TAF1-TAF7 complex in general TFIID function and/or assembly of the pre-initiation complex (PIC).4,5

Figure 1. Images of the Facial Phenotype from Families 1, 2, and 4–9
Cardinal facial dysmorphologies include prominent supraorbital ridges (seen in 1A, 1B, 2A, 6A, 8A–8C, and 9A), down-slanted palpebral fissures (1A, 1B, 2A, 4A, 5A, 6A, 8A–8C, and 9A), sagging cheeks (1A, 1B, 5A, 8A–8C, and 9A), a long philtrum (1A, 1B, 2A, 4A, 5A, 8A–8C, and 9A), low-set and protruding ears (1A, 1B, 2A, 4A, 5A, 6A, 8A–8C, and 9A), a long face (1A, 1B, 2A, 5A, 6A, 8A–8C, and 9A), a high palate (5A, 6A, 8A–8C, and 9A), a pointed chin (1A, 1B, 2A, 4A, 5A, 6A, and 8A–8C), and anteverted nares (2A, 4A, 5A, 7A, 8A–8C, and 9A).
Human TAF1 is a 1,893 amino acid multifunctional protein that has been reported to possess activities related to DNA promoter binding, histone acetylation, and protein phosphorylation.\textsuperscript{28–31} The histone acetyltransferase (HAT) activity of TAF1 can be blocked by TAF7 binding.\textsuperscript{4,32,35} Studies suggest that phosphorylation of TAF7 at Ser264 causes release from TFIID and thus alleviates its inhibitory effect on TAF1.\textsuperscript{34} Therefore, mutations that disrupt this inhibitory protein-protein interaction could have inhibitor effect on TAF1.\textsuperscript{34} p.Asp976His substitution also has the potential to disrupt TAF1-TAF7 binding. The acidic Asp976 maps to a separate TAF1-TAF7 protein interface within DUF3591 and undergoes intermolecular hydrogen binding in the highly conserved glycine-rich loop of TAF1. This loop interacts extensively with a highly conserved Arg-rich motif in TAF7. The acidic-to-basic amino acid change (p.Asp976His) has the potential to disrupt the architecture of the glycine-rich motif and its ability to effectively bind to TAF7. The p.Arg1246Trp substitution described in this work and the published p.Arg1246Trp substitution described in this work and the published p.Arg1246Trp substitution described in this work and the published p.Arg1246Trp substitution described in this work and the published p.Arg1246Trp substitution described in this work.

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**Figure 2. TAF1 Domains, Variant Scores, and ExAC Sequence Variation Plot**

(A) Pedigree drawings of the nine families who were found to harbor TAF1 SNVs (NCBI Gene ID: 6872 according to the GRCh37.p13 assembly). Black dots indicate maternal carriers.
(B) All nine SNVs are listed and annotated with CADD, SIFT, GERP++, and phyloP scores (which indicate conservation across 99 vertebrate genomes and humans). All of the SNVs are considered to be potentially deleterious by all of the listed annotations, except for c.3708A>G, which is a splice-site variant and as a consequence is not necessarily expected to be categorized as deleterious by any of the listed scores, because it does not affect amino acid composition of the predicted protein.
(C) Known TAF1 domains are shown with respect to their corresponding genic positions. All but non-synonymous variants reported in the ExAC Browser for TAF1 are plotted below as lines; white and gray indicate exon boundaries. Red lines indicate the relative positions of the eight missense variants described in this paper (see Table 2). Numerals link the sequence variants shown on the ExAC plot to their familial origin, and those noted with a star fall within TAF1 of the eight missense variants described in this paper (see Table 2). Numerals link the sequence variants shown on the ExAC plot to their familial origin, and those noted with a star fall within TAF1 of the eight missense variants described in this paper (see Table 2). Numerals link the sequence variants shown on the ExAC plot to their familial origin, and those noted with a star fall within TAF1 of the eight missense variants described in this paper (see Table 2).
each of which consists of a bundle of four alpha helices that form a hydrophobic pocket that can recognize acetylated lysines found on the N-terminal tails of histones. A close examination of the published TAF1 bromodomain structure revealed that the p.Arg1431His substitution in family 7 is a surface residue on one face of the alpha helix. It forms hydrogen bonds with two residues on a nearby helix, most likely playing a supporting role in maintaining the bromodomain fold. Substitution p.Arg1431His could affect the stability of the acetyl-lysine ligand binding site in TAF1 and alter promoter recognition.

As an initial attempt to explore whether a biological signature could be seen on the level of the transcriptome for at least one of the mutations in the above families, we conducted RNA sequencing (RNA-seq) with RNA isolated from the blood of family 1. Blood was collected from the two probands, their parents, and the maternal grandparents. Except for the single blood draws for the grandparents, two blood draws were taken on separate days for each subject. RNA libraries were generated, and the final pooled library was sequenced on a HiSeq 2000 across three lanes (paired-end 100 bp).

### Table 1. Summary of the Clinical Features of TAF1 ID Syndrome

| Features (Human Phenotype Ontology Nos.) | 1A | 1B | 2A | 3A | 4A | 5A | 6A | 7A | 8A | 8B | 8C | 9A | 10A | 11A |
|------------------------------------------|----|----|----|----|----|----|----|----|----|----|----|----|----|----|
| Sex                                      | M  | M  | M  | M  | M  | M  | M  | M  | M  | M  | M  | M  | M  | M  |
| Age (years)                              | 15 | 13 | 5  | 6  | 9  | 3  | 22 | 11 | 9  | 4  | 1  | 3  | 16 | 8  |
| Postnatal growth retardation             | +  | +  | +  | +  | +  | -  | -  | +  | +  | +  | +  | +  | +  | UK |
| (HP: 0008897)                            |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Delayed gross motor development          | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  |
| (HP: 0002194)                            |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Delayed speech and language development  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  |
| (HP: 000750)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Oral-pharyngeal dysphagia                | +  | +  | +  | UK | +  | UK | -  | +  | +  | UK | +  | UK |    |    |
| (HP: 0200136)                            |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Prominent supraorbital ridges             | +  | +  | -  | +  | UK | -  | +  | +  | +  | +  | +  | +  | +  | +  |
| (HP: 0000336)                            |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Downslanted palpebral fissures            | +  | +  | -  | +  | +  | +  | -  | +  | +  | +  | +  | +  | +  | +  |
| (HP: 000494)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Sagging cheeks                            | +  | +  | -  | -  | +  | +  | +  | -  | +  | +  | +  | +  | +  | +  |
|    |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Long philtrum                            | +  | +  | +  | +  | +  | -  | +  | +  | +  | +  | +  | -  |    |    |
| (HP: 000343)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Low-set ears                             | +  | +  | +  | +  | +  | +  | -  | +  | +  | +  | +  | +  | -  | +  |
| (HP: 000369)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Protruding ears                          | +  | +  | +  | +  | +  | +  | -  | +  | +  | +  | +  | +  | -  | +  |
| (HP: 000411)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Long face                                | +  | +  | -  | -  | UK | +  | +  | -  | +  | +  | +  | +  | +  | +  |
| (HP: 000276)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| High palate                              | UK | UK | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  |
|    |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Pointed chin                             | +  | +  | -  | -  | +  | +  | +  | +  | +  | +  | +  | -  |    |    |
| (HP: 000307)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Anteverted nares                          | -  | -  | +  | +  | +  | +  | -  | +  | +  | +  | +  | -  | +  | +  |
| (HP: 000463)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Hearing impairment                       | +  | +  | +  | +  | UK | -  | -  | +  | +  | +  | -  | UK |    |    |
| (HP: 000365)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Chromic otitis media                     | +  | +  | -  | +  | +  | +  | +  | +  | +  | -  | UK |    |    |    |
| (HP: 000389)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Strabismus                               | +  | +  | +  | UK | +  | +  | -  | -  | +  | -  | -  | -  | +  | -  |
| (HP: 000486)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Microcephaly                             | +  | +  | +  | +  | +  | -  | +  | +  | +  | -  | -  | -  | -  | -  |
| (HP: 000252)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Hypoplasia of the corpus callosum        | +  | +  | +  | UK | +  | +  | +  | UK | +  | +  | +  | +  | UK |    |
| (HP: 0002079)                            |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Generalized hypotonia                    | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | -  | +  |
| (HP: 0001290)                            |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Unusual gluteal crease with sacral caudal| +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  | +  |
| remnant and sacral dimple (abnormal sacral|    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| segmentation [HP: 0008468] and prominent |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| protruding coccyx [HP: 0008472]          |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Joint hypermobility                      | +  | +  | -  | +  | UK | -  | -  | +  | +  | +  | -  | UK |    |    |
| (HP: 0001382)                            |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Autistic behaviors                       | +  | +  | -  | +  | UK | +  | +  | +  | +  | +  | -  | +  |    |    |
| (HP: 000729)                             |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Intellectual disability                  | +  | +  | +  | +  | UK | +  | +  | +  | +  | +  | +  | +  | -  | +  |
| (HP: 0001249)                            |    |    |    |    |    |    |    |    |    |    |    |    |    |    |

This table demonstrates clinical features shared by eight or more probands across all affected individuals in the families. See Table S5 for a more comprehensive phenotypic table that includes phenotypic and clinical idiosyncrasies. Abbreviations are as follows: M, male; and UK, unknown. *Probands containing duplications; they are generally less similar to the probands containing SNVs and share fewer common clinical features.
Table 2. Summary of TAF1 Variants across All Affected Individuals in This Study

| Proband | Inheritance | Genetic Background | TAF1 Mutation (hg19) |
|---------|-------------|--------------------|----------------------|
| 1A      | maternal    | European decent    | chrX: g.70621541T>C; p.Ile1337Thr |
| 1B      | maternal    | European decent    | chrX: g.70621541T>C; p.Ile1337Thr |
| 2A      | de novo     | European decent    | chrX: g.70607243T>C; p.Cys807Arg |
| 3A      | de novo     | European decent    | chrX: g.70618477C>T; p.Arg1246Trp |
| 4A      | de novo     | European decent    | chrX: g.70601686T>A; p.Ile505Asn |
| 5A      | maternal    | Ecuadorian         | chrX: g.70618449A>G; r.(3708a>g; 3681_3708del28); p.Arg1228Ilefs*16 |
| 6A      | de novo     | European decent    | chrX: g.70643003A>C; p.Asn1517His |
| 7A      | de novo     | British            | chrX: g.70627912G>A; p.Arg1431His |
| 8A      | maternal    | Colombian           | chrX: g.70602671C>T; p.Pro596Ser |
| 8B      | maternal    | Colombian           | chrX: g.70602671C>T; p.Pro596Ser |
| 9A      | de novo     | Colombian           | chrX: g.70602671C>T; p.Pro596Ser |
| 9A      | maternal    | Spanish             | chrX: g.70612503G>C; p.Asp976His |
| 10A     | maternal    | Albanian            | 0.423 Mb duplication including TAF1 and other genes at Xq13.1(70,370,794–70,794,385); deletion containing KANSL1 and other genes at 17q21.31 (0.63 Mb) |
| 11A     | de novo     | Greek               | 0.42 Mb duplication including TAF1 and other genes: arr Xq13.1(70,287,519–70,711,110)x2 |

49,792,652 (SD = 11,666,119) properly paired reads were generated, and a mean spliced mapping percentage of 85.41% (SD = 7.1%) per sample was observed (Table S6). HISAT39 was used for spliced alignment to UCSC Human Genome Browser hg19, and Stringtie40 was used for quantifying known transcripts. Cuffdiff41 was used for performing differential expression analysis, and CummeRbund42 was used for analyzing, filtering, and visualizing the results from the Cuffdiff differential-expression analysis. Gene-set enrichment analyses were performed with WebGestalt,43 which uses Molecular Signatures Database (MSigDB) for transcription-factor targets,44 Kyoto Encyclopedia of Genes and Genomes (KEGG),45 Gene Ontology (GO),46 and Human Phenotype Ontology (HPO)47 to obtain gene annotations and set-inclusion information. We used various analysis tools in the CummeRbund package42 to evaluate the quality of our RNA-seq data and analysis for family 1 (Figure S10).

We found 213 genes (Table S7) to be differentially expressed between the two affected male probands from family 1 and their unaffected parents and grandparents. Of these 213 genes, 179 were expressed less (downregulated) and 34 were expressed more (upregulated) in the affected individuals. Among those genes that were upregulated in the affected individuals, 24 out of the 34 were expressed with a log2 fold-change value greater than 1, and no genes were found to be upregulated with a log2 fold-change value greater than 2. Among those genes that were downregulated in the affected individuals, 161 out of 179 were expressed with a log2 fold-change value less than −1, and 41 out of 179 were expressed with a log2 fold-change value less than −2.

Among the genes that were upregulated in the affected boys, we found no obvious interpretable biological signal. We present the results from the gene-set enrichment analysis performed on the set of genes that were downregulated in the affected individuals with a log2 fold-change value less than −1, because this set represents the genes that were significantly downregulated and to a potentially biologically relevant level. Transcription-factor-target enrichment analysis revealed a significant enrichment of genes regulated by E-box proteins (CANNTG promoter motifs, Benjamini-Hochberg [BH]-corrected p value of 0.0052). KEGG-pathway enrichment analysis revealed an enrichment of genes involved in Parkinson, Alzheimer, and Huntington diseases and cardiac-muscle contraction (BH-corrected p values of 2.95e−7, 1.54e−6, 2.13e−6, and 2.13e−6, respectively). Enrichments in genes associated with HPO annotations included type 1 muscle-fiber predominance, and laxity of ankles, fingers, and wrists (all four phenotypes were reported with a BH-corrected p value of 0.0481).

It is important to note that the RNA-seq results are preliminary and potentially confounded by age- or sex-specific expression differences between the affected and unaffected groups, and the results here were derived from a single family (because these were the samples that we were able to collect to date). Complete blood counts were not performed on the blood samples used for RNA-seq; thus, our results could also be confounded by secondary differences in mRNA abundances. The RNA-seq and whole-genome sequencing data have been deposited to the Sequence Read Archive (SRA) under BioProject ID PRJNA301337.
To evaluate structural CNS defects, we performed in vivo functional modeling of TAF1 by using transient knockdown in developing zebrafish embryos. The optic tectum is a neuroanatomical structure that occupies the majority of the space within the midbrain, and its size is a proxy for head size (e.g., microcephaly,48 which is one of the most robustly observed phenotypes).

We designed a splice-blocking morpholino (MO) targeting the donor site of exon 9 of the sole zebrafish taf1 ortholog (Ensembl: ENSDART00000051196; Figure S12) and performed stainings for acetylated tubulin to evaluate the structure of the optic tectum. In total, we analyzed 134 control embryos, 133 embryos injected with taf1 MO, 109 embryos injected with taf1 MO and wild-type (WT) TAF1 RNA, and 78 embryos injected with WT TAF1 mRNA by measuring the area of the optic tectum 3 days after fertilization. We found that the relative area of the optic tectum was reduced by approximately 10% in embryos injected with a MO targeting the donor site of exon 9 of D. rerio taf1 (p < 0.0001; Figures 4A and 4B and S12). The effect was specific to the MO knockdown, given that the phenotype could be reliably restored by co-injection of MO and WT human TAF1 mRNA (p < 0.0001). Notably, overexpression of WT human TAF1 mRNA did not result in a phenotype that was significantly different from that of the controls (p = 0.79).

To confirm these findings, we also used CRISPR/Cas9 to disrupt taf1. For the CRISPR experiments, taf1 guide RNA was generated as described.49 In agreement with our observation that the area of the optic tectum was smaller in embryos injected with taf1 MO than in control embryos, F0 embryos with CRISPR-disrupted taf1 showed a smaller area of this structure than did un-injected controls (p < 0.001) (Figures 4C and S12). Unfortunately, the relatively small effects seen with MO knockdown and CRISPR-mediated mosaicism in F0 embryos precluded us from being able to show differences between WT and mutated TAF1 constructs in terms of rescuing the neuroanatomical defect (data not shown).

In conclusion, we have presented evidence showing that mutations involving TAF1 contribute to the phenotype described here. Differences in genetic background and the environment can certainly account for the phenotypic differences among the various males in these families.50–56 Other studies also suggest a functional link between developmental delay and the TFIID multi-protein complex,10–12 although the phenotypic variability and expression of other variants in TAF1 and in other TAFs remains to be determined. We have also provided the results of initial studies...
that suggests a possible regulatory role for at least one variant presented here and have shown that zebrafish knockdown and mutant studies for this gene have demonstrated a quantifiable, albeit small, effect on a neuronal phenotype. In our current study, we did not see a clear mechanistic link between the missense and splice-site variants in some of the probands and the duplications in two of the probands, and the probands with duplications share fewer phenotypic features with the rest of the disease cohort. There are many possible explanations for why a duplication of a gene might mimic the effect of a disruptive missense or splice-site variant, and our current study lacks the necessary experimental evidence to point to any particular scenario with certainty. Further work is needed for teasing apart the contributions of these duplications and their constituent genes to this more complex phenotype. It should also be noted that exome and/or whole-genome sequencing has not yet been performed in family 10 or 11, so other mutations could be contributing to the phenotypes of the probands.

Accession Numbers

The RNA-seq and whole-genome sequencing data reported in this paper have been deposited to the Sequence Read Archive (SRA) under BioProject ID PRJNA301337. The study accession number is SRA: SRP065848, and the sample accession numbers are SRA: SRS1153874, SRS1153892, SRS1153893, SRS1153896, SRS1153897, SRS1153898, SRS1153899, SRS1153900, SRS1153901, SRS1153902.

Supplemental Data

Supplemental Data include Supplemental Material and Methods, a Supplemental Note, 13 figures, 7 tables, and 8 movies and can be found with this article online at http://dx.doi.org/10.1016/j.ajhg.2015.11.005.

Conflicts of Interest

G.J.L. serves on advisory boards for GenePeeks Inc. and Omicia Inc. and is a consultant to Good Start Genetics. D.M. was until recently an employee and shareholder of Gene By Gene Inc. and is currently chief scientific officer of Tute Genomics Inc. K.W. is a board member and shareholder of Tute Genomics Inc., and R.R. is an employee, chief executive officer, and shareholder of Tute Genomics Inc. This study did not involve these companies and did not use products from these companies. The sequencing service by Complete Genomics was provided via a data-analysis grant, but the company was not involved in any subsequent analyses.
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Web Resources

The URLs for data presented herein are as follows:

1000 Genomes, http://www.1000genomes.org/data
ANNOVAR, http://annovar.openbioinformatics.org/en/latest/
dbsNP, http://www.ncbi.nlm.nih.gov/snp
DECIPHER, http://decipher.sanger.ac.uk/
Ensembl, http://www.ensembl.org
Exome Aggregation Consortium (ExAC) Browser, http://exac.broadinstitute.org/
GeneSplicer, https://ccb.jhu.edu/software/genesplicer/
Golden Helix GenomeBrowse, http://www.goldenhelix.com/GenomeBrowse/index.html
HomoloGene, http://www.ncbi.nlm.nih.gov/homologene
Human Gene Mutation Database, http://www.hgmd.org/
Human Splicing Finder, http://www.umd.be/HSF/
MaxEntScan, http://genes.mit.edu/burgelab/maxent/Xmaxentscan-scoreseq.html

NHLBI Exome Sequencing Project (ESP) Exome Variant Server, http://evs.gs.washington.edu/EVS/
OMIM, http://www.ncbi.nlm.nih.gov/omim
PennCNV, http://penncnv.openbioinformatics.org/en/latest/
PolyPhen-2, http://genetics.bwh.harvard.edu/pph2/
SIFT, http://sift.jcvi.org/

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