

Heterozygous Variants in *KMT2E* Cause a Spectrum of Neurodevelopmental Disorders and Epilepsy

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We delineate a *KMT2E*-related neurodevelopmental disorder on the basis of 38 individuals in 36 families. This study includes 31 distinct heterozygous variants in *KMT2E* (28 ascertained from Matchmaker Exchange and three previously reported), and four individuals with chromosome 7q22.2-22.23 microdeletions encompassing *KMT2E* (one previously reported). Almost all variants occurred *de novo*, and most were truncating. Most affected individuals with protein-truncating variants presented with mild intellectual disability. One-quarter of individuals met criteria for autism. Additional common features include macrocephaly, hypotonia, functional gastrointestinal abnormalities, and a subtle facial gestalt. Epilepsy was present in about one-fifth of individuals with truncating variants and was responsive to treatment with anti-epileptic medications in almost all. More than 70% of the individuals were male, and expressivity was variable by sex; epilepsy was more common in females and autism more common in males. The four individuals with microdeletions encompassing *KMT2E* generally presented similarly to those with truncating variants, but the degree of developmental delay was greater. The group of four individuals with missense variants in *KMT2E* presented with the most severe developmental delays. Epilepsy was present in all individuals with missense variants, often manifesting as treatment-resistant infantile epileptic encephalopathy. Microcephaly was also common in this group. Haploinsufficiency versus gain-of-function or dominant-negative effects specific to these missense variants in *KMT2E* might explain this divergence in phenotype, but requires independent validation. Disruptive variants in *KMT2E* are an under-recognized cause of neurodevelopmental abnormalities.

KMT2E (GenBank: NM_182931.2, MIM: 608444) encodes a member of the lysine N-methyltransferase 2 (KMT2) family. This family of enzymes plays a vital role in regulating post-translational histone methylation of histone 3 on

lysine 4 (H3K4).¹ Proper H3K4 methylation is required to maintain open chromatin states for regulation of transcription. There are at least eight known monogenic disorders that impair regulation of H3K4 methylation and that

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present with neurodevelopmental syndromes^{2–8} (Table S1). In addition to these Mendelian disorders, dysregulated H3K4 methylation is believed to play a role in the pathogenesis of schizophrenia and autism.⁹ Truncating variants in *KMT2E* have previously been reported in three unrelated males in a large sequencing study of non-syndromic autism, but phenotypic data were limited.^{10–12} In this report, we present 35 additional individuals with heterozygous variants in *KMT2E* in an effort to define a *KMT2E*-related neurodevelopmental disorder.

New cases were ascertained from GeneMatcher through the Matchmaker Exchange Network and MyGene2 between September 2016 and August 2018.^{13,14} Microdeletions were detected by chromosomal microarrays in some individuals, whereas all other individuals were found to have variants in *KMT2E* via exome or genome sequencing. Written consent for publication of photographs was provided from the individuals' parents or legal guardians. Additional phenotype data and genetic findings for individuals are summarized in Table S2.

KMT2E is constrained for protein-truncating variation in the general population. The Genome Aggregation Database (gnomAD) is a large-scale reference database with high-quality, jointly processed exome or genome data from more than 140,000 individuals.¹⁵ Constraint analysis performed on the gnomAD dataset shows that *KMT2E* is a candidate haploinsufficient gene. *KMT2E* is very depleted (presumably as a result of negative selection) for protein-truncating variants; there is a probability of loss-of-function intolerance (pLI) score of 1.0 and an observed/expected ratio of 0.01 (showing 1% [0–0.06 95% CI] of expected loss-of-function variation in gnomAD).

We reviewed the 28 loss-of-function variants present in gnomAD v2.1 (Table S3). The majority of these variants are not expected to result in protein truncation for a variety of reasons, including annotation artifacts (n = 8), sequence errors at a simple repeat (n = 5), somatic mosaicism (n = 1), and a splice-site rescue (n = 1). Four variants are part of a complex variant found in one individual; this complex variant, when resolved, is not expected to result

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in truncation. Four variants found in eight individuals in gnomAD are in the last exon; two are expected to result in truncation of the last exon, and two will result in protein extension. Of note, the two protein-extension variants are located close to the variant in individual 28 (c.5453_5460delTGGCCCTG [p.Val1818Alafs*48]). The inheritance of this variant is unknown because the father is not available for testing, although it is not present in his mother, so this remains a variant of uncertain significance.

After review, we found that five variants in gnomAD appear to result in protein truncation. These are found in three males and two females between the ages of 30 and 70. All five are absent from the control-only subset in gnomAD (although it should be noted that gnomAD does not contain cohorts recruited for severe, pediatric-onset disease; rather, it contains cohorts recruited for common adult-onset diseases such as cardiovascular disease and type II diabetes). By reviewing the data subsets, we found two variants that appear to be from neurologic cohorts and three from non-neurologic and non-cancer cohorts. Overall, very few variants present in this large general population reference database are likely to result in protein truncation of *KMT2E*.

We identified 38 individuals with *KMT2E* variants in association with a neurodevelopmental phenotype. Including the three previously reported cases,^{10–12} 34 individuals from 32 families were found to have single-nucleotide or indel variants in *KMT2E*, and four additional individuals had copy-number variants encompassing *KMT2E* (Figure 1, Table 1, Table S2). The *KMT2E* variants arose *de novo* in 26 individuals in our cohort. The variant was maternally inherited in one previously reported individual (maternal phenotype unknown).¹² Inheritance of the variant was unknown in four families where neither parent was available for testing. In one family, a variant was found in three affected male siblings. The variant was not found in their mother. The father was not available for testing but was reported to have an intellectual disability. 30 variants were protein-truncating variants: 24 were indels, four were nonsense variants, and two were variants at essential splice sites (Figure 1A). Only one variant was seen in two independent families (c.1776_1780delAAAGA [p.Lys593Argfs*17]); it was found in a male (individual 9) and a female (individual 10). 23 of these variants are predicted to produce transcripts that would be subject to nonsense-mediated decay. Five of the protein-truncating variants fall in the terminal exon of the gene, potentially escaping nonsense-mediated decay; three (found in individuals 26, 27, and 28) of these five variants extend the open reading frame. The last exon in individuals 26 and 27 in our cohort has a frameshift variant that alters the last 244–259 amino acids of *KMT2E*, whereas individual 28 has an alteration in the last 48 amino acids. We evaluated the impact of this on protein structure. Wild-type *KMT2E* has a very disordered C terminus, but these upstream frameshifts result in increased

stability and the formation of a predicted homeodomain (Figure S1). CADD scores are summarized in Table 1.

Four of the individuals had *de novo* missense variants, three of which occurred at highly conserved positions and/or regions of the gene (Figure 1B). Pro1376 is not well conserved, and serine is present in some mammalian species. None of the *KMT2E* variants are reported in public databases (gnomAD, Exome Variant Server, or 1000 Genomes),^{15–17} although another missense change is seen at Pro1376 in gnomAD (p.Pro1376Leu, allele frequency 0.015%).

To understand the biophysical consequence of *KMT2E* protein sequence changes, we used structural-prediction programs (HMMER,¹⁸ PHYRE2,¹⁹ InterProScan,²⁰ and NetPhos²¹) that evaluate the presence of protein domains and major secondary structure elements (e.g., helices, strands, loops, disorder, post-translational modification sites, etc.). A large protein of 1,858 amino acids, *KMT2E* has two N-terminal domains: a SET enzymatic domain (aa 282–445), which is predicted to be inactive, and a Zn-finger PHD domain (aa 120–165), and most of the protein has few scattered helices and strands, as well as a disordered C terminus. There was no clustering of the missense variants; one is in the SET domain, one is in the PHD domain, and two are not in identified domains. *KMT2E* is not significantly constrained for missense variation in the general population (Z score +1.42, observed/expected ratio of 0.87 [0.82–0.92 95% CI] for missense variation in gnomAD). All four missense variants might significantly change local structure by introducing rotamers (c.418G>A [p.Val104Ile]),²² or by changing the charge and hydrophobicity of local sequences (c.850T>C [p.Tyr284His], c.2720A>T [p.Asp907Val], and c.4126C>T [p.Pro1376Ser]). Additionally, p.Tyr284His abolishes and p.Pro1376Ser creates potential phosphorylation sites. Changing rotamers, electrical charge, and hydrophobicity might alter *KMT2E* binding properties.

For the four individuals with chromosome microdeletions encompassing *KMT2E*, all deletions occurred *de novo*. Deletion sizes range from 0.052 to 3.2 Mb. The 0.052 Mb deletion in individual 30 involves only *KMT2E*, whereas the other three deletions include additional genes.²³ Figure 1C illustrates the genes included in these deletions. Median maternal and paternal age across the cohort was 30 and 36 years, respectively. There were phenotypic differences between individuals with protein-truncating, missense, and copy-number variants, as summarized below.

For the 30 individuals with protein-truncating variants in *KMT2E*, 22 were male and eight were female (Figure 2). Age at most recent evaluation ranged from 19 months to 24 years. Prenatal and neonatal courses were largely uncomplicated for most individuals with protein-truncating variants. One individual was born prematurely at 35 weeks. Several individuals had neonatal jaundice, one had hypoglycemia, one had sinus tachycardia, and two had neonatal feeding difficulties. Individual 10 developed respiratory arrest at 14 h of life and had a hypoxic-

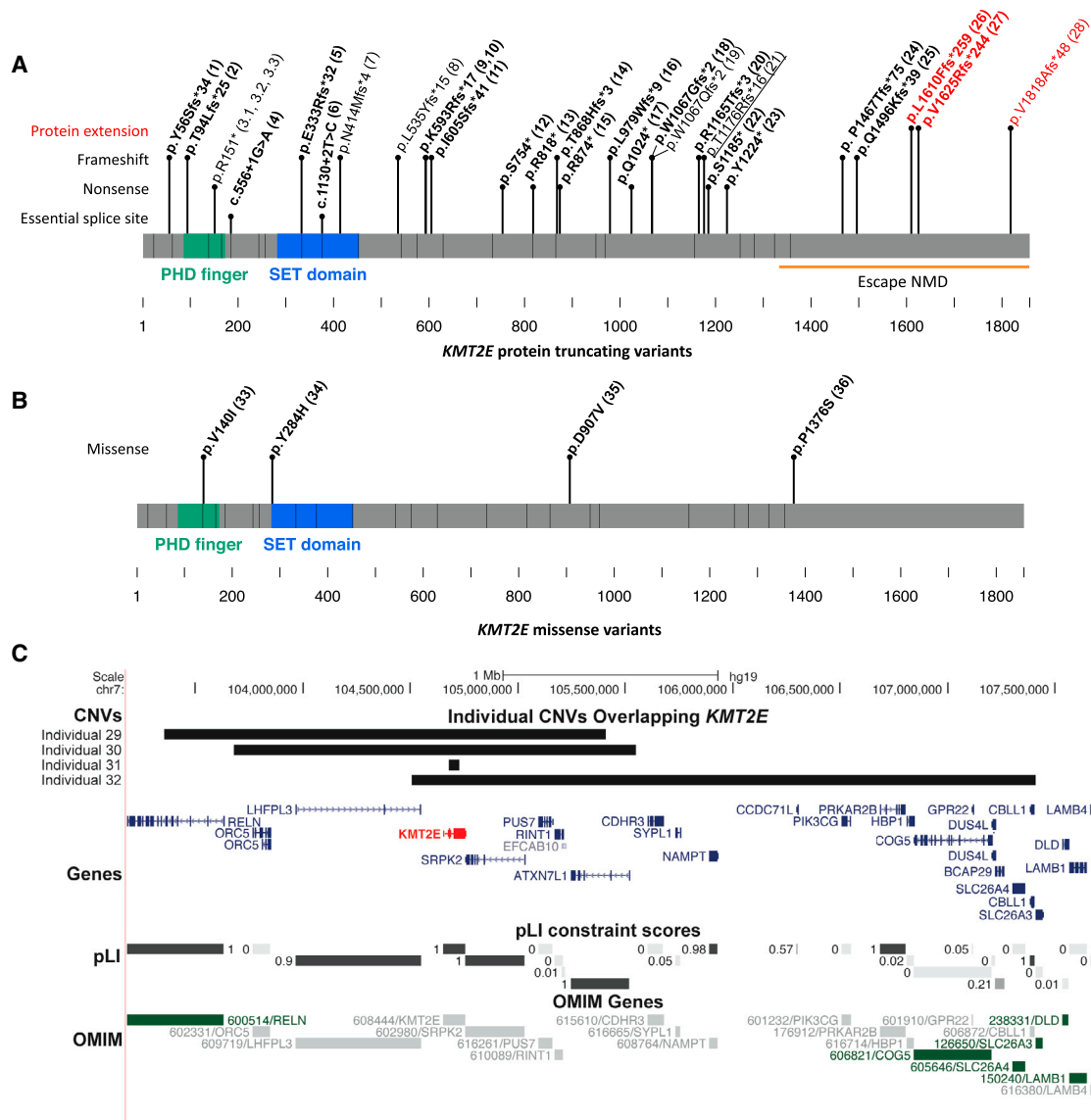


Figure 1. KMT2E Variants in 38 Individuals

(A) 28 protein-truncating variants in *KMT2E* identified in 30 individuals. Variants in bold are *de novo* in the proband, whereas the underlined variant was inherited. In some cases, both parents are unavailable and the *de novo* status is unknown (non-bold). Variants in the last exon are predicted to escape nonsense-mediated decay (individuals 24–28), whereas the last three variants (red) also result in protein extension (individuals 26–28).

(B) *De novo* missense variants in *KMT2E* in individuals 33–36.

(C) *De novo* deletions overlapping *KMT2E* were identified in individuals 29–32. All OMIM gene-disease associations (green) are for recessive disease.

ischemic injury with typical sequelae seen on neuroimaging. She has spastic quadriplegia and epilepsy, and she is not included in the analysis below because her acquired injury significantly influences her phenotype and is most likely not representative of the disorder itself (although it cannot be excluded that the genetic disorder predisposed her to the injury).

Of the remaining 29 individuals in this group (i.e., excluding individual 10), 24 had early developmental delay documented. For three individuals without documented developmental delay, these are cases previously reported from autism studies where only limited clinical

information is available.^{10–12} The mean age of independent walking in this group was 20 months (range 12 to 48 months, Figure 3). All individuals are currently able to walk independently. 12 of the 29 individuals have hypotonia. Individual 15 had normal initial motor development but developed progressive spastic diplegia at 14 months of age. Neuroimaging in this individual demonstrated cerebral white-matter abnormalities.

The mean age of acquired first word in this group was 20 months (range 12 to 48 months, Figure 3). Although this information is not available for all individuals, 14 (out of 17) individuals are verbal, but seven are noted to

Table 1. Summary of *KMT2E* Variants Found in 38 Individuals with Neurodevelopmental Phenotypes

Individual	Sex, Age	Variant, GenBank: NM_182931.2	Consequence	Inheritance	CADD	ID	Autism	Delay	Epilepsy	Macrocephaly ^a
1 ¹¹	male, 11 y	c.167delA, (p.Tyr56Serfs*34)	frameshift, expect NMD	<i>de novo</i>	30	mild	yes	NA	no	no
2	female, 12 y	c.280delA, (p.Thr94Leufs*25)	frameshift, expect NMD	<i>de novo</i>	33	moderate	no	yes	no	yes
3.1	male, 9 y, 6 m	c.450dupT, (p.Arg151*)	nonsense, expect NMD	unknown	34	NA	yes	yes	NA	no
3.2	male, 7 y	c.450dupT, (p.Arg151*)	nonsense, expect NMD	unknown	34	NA	yes	yes	NA	no
3.3	male, 6 y	c.450dupT, (p.Arg151*)	nonsense, expect NMD	unknown	34	NA	yes	yes	NA	no
4	male, 5 y, 9 m	c.556+1G>A	essential splice site, expect NMD	<i>de novo</i>	34	NA	no	yes	yes	no
5	male, 12 y, 2 m	c.997delG, (p.Glu333Argfs*32)	frameshift, expect NMD	<i>de novo</i>	33	NA	no	yes	no	yes
6	male, 3 y, 1 m	c.1130+2T>C	essential splice site, expect NMD	<i>de novo</i>	33	yes	no	yes	no	yes
7	female, 21 y	c.1239delC (p.Asn414Metfs*4)	frameshift, expect NMD	unknown	34	moderate	no	yes	yes	yes
8	female, 8 y	c.1603delC (p.Leu535Tyrfs*15)	frameshift, expect NMD	unknown	25	NA	no	yes	NA	relative
9	male, 11 y, 4 m	c.1776_1780delAAAGA, (p.Lys593Argfs*17)	frameshift, expect NMD	<i>de novo</i>	34	yes	no	yes	no	yes
10	female, 3 y, 6 m	c.1776_1780delAAAGA, (p.Lys593Argfs*17)	frameshift, expect NMD	<i>de novo</i>	34	yes	no	yes	yes	no
11	female, 1 y, 10 m	c.1812delG, (p.Ile605Serfs*41)	frameshift, expect NMD	<i>de novo</i>	26	NA	NA	yes	no	no
12	male, 3 y, 7 m	c.2261delC, (p.Ser754*)	nonsense, expect NMD	<i>de novo</i>	34	low-normal	no	yes	no	no
13	male, 4 y, 3 m	c.2452C>T, (p.Arg818*)	nonsense, expect NMD	<i>de novo</i>	37	mild	no	yes	no	no
14	male, 8 y	c.2602_2605delACTA, (p.Thr868Hisfs*3)	frameshift, expect NMD	<i>de novo</i>	35	NA	yes	yes	no	no
15	male, 1 y, 7 m	c.2620C>T, (p.Arg874*)	nonsense, expect NMD	<i>de novo</i>	39	NA	no	yes	no	no
16	female, 3 y, 6 m	c.2936delT, (p.Leu979Trpfs*9)	frameshift, expect NMD	<i>de novo</i>	23	NA	no	yes	no	yes
17	male, 4 y, 8 m	c.3070C>T, (p.Gln1024*)	nonsense, expect NMD	<i>de novo</i>	38	NA	no	no	no	yes
18 ¹⁰	male, 12 y	c.3198delC, (p.Trp1067Glyfs*2)	frameshift, expect NMD	<i>de novo</i>	35	mild	yes	NA	no	yes
19	female, 6 y, 5 m	c.3198_3234del, (p.Trp1067Glnfs*2)	frameshift, expect NMD	unknown	35	mild	no	yes	no	yes
20	male, 5 y, 10 m	c.3494_3495delGA, (p.Arg1165Thrfs*3)	frameshift, expect NMD	<i>de novo</i>	34	NA	no	yes	no	yes
21 ¹²	male, NA	c.3527_3530delCAGA, (p.Thr1176Argfs*16)	frameshift, expect NMD	maternally inherited	20	NA	yes	NA	no	NA
22	female, 9 y	c.3554C>G, (p.Ser1185*)	nonsense, expect NMD	<i>de novo</i>	35	mild	no	yes	yes	no
23	male, 6 y	c.3672_3673delTA, (p.Tyr1224*)	frameshift, expect NMD	<i>de novo</i>	24	NA	no	yes	no	yes
24	male, 5 y	c.4397_4398ins19, (p.Pro1467Thrfs*75)	frameshift, last exon, escape NMD	<i>de novo</i>	NA	mild	no	yes	no	yes
25	male, 12 y, 10 m	c.4485_4486delTC, (p.Gln1496Lysfs*39)	Frameshift, last exon, escape NMD	<i>de novo</i>	24	mild	NA	yes	no	no
26	male, 6 y, 7 m	c.4829dupT, (p.Leu1610Phefs*259)	frameshift, protein extension	<i>de novo</i>	34	low-normal	NA	NA	no	yes

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Table 1. Continued

Individual	Sex, Age	Variant, GenBank: NM_182931.2	Consequence	Inheritance	CADD	ID	Autism	Delay	Epilepsy	Macrocephaly ^a
27	male, 8 y, 8 m	c.4872dupC, (p.Val1625Argfs*244)	frameshift, protein extension	<i>de novo</i>	24	yes	no	yes	no	yes
28	male, 24 y	c.5453_5460delTGGCCCTG, (p.Val1818Alafs*48)	frameshift, protein extension	unknown	35	moderate	no	yes	no	relative
29	female, 12 y, 11 m	7:103354482-105407628x1, 2.05 Mb	microdeletion	<i>de novo</i>	NA	moderate	yes	yes	no	yes
30	female, 18 y	7:104678742-104730547x1, 0.052 Mb	microdeletion	<i>de novo</i>	NA	moderate	no	yes	yes	no
31	male, 22 y	7:103679146-105547471x1, 1.87 Mb	microdeletion	<i>de novo</i>	NA	mild/moderate	no	yes	yes	no
32 ^{2,6}	male, 7 y	7:10409959-107002808x1, 2.9 Mb	microdeletion	<i>de novo</i>	NA	mild	no	yes	yes	yes
33	male, 16 y, 3 m	c.418G>A (p.Val140Ile)	missense	<i>de novo</i>	25	NA	yes	yes	yes	NA
34	male, 2 y, 5 m	c.850T>C (p.Tyr284His)	missense	<i>de novo</i>	24	severe	NA	yes	yes	no
35	female, 2 y, 11 m	c.2720A>T (p.Asp907Val)	missense	<i>de novo</i>	24	severe	no	yes	yes	microcephaly
36	female, 36 y	c.4126C>T (p.Pro1376Ser)	missense	<i>de novo</i>	11	mild	no	yes	yes	microcephaly

NA = not available; NMD = nonsense-mediated decay.

^aMacrocephaly is defined here as a head circumference >2 standard deviations (SD) above mean for age and microcephaly as >-2 SD below mean for age. Relative macrocephaly is defined here as a head circumference 1 SD above the SD of the height.

speak poorly or to have articulation problems. Three of the individuals were reported to have speech regression. Intelligence quotient (IQ) data were available for only seven out of the 29 individuals: the mean IQ was 74 (range 62–98). Seven of the individuals have been diagnosed with autism. One additional individual was diagnosed with a sensory integration disorder, and another diagnosed with difficulty in social interaction but did not meet the criteria for autism. At least two of the individuals have been diagnosed with attention-deficit/hyperactivity disorder (ADHD). Additional behavioral concerns were reported in 11 of the individuals; these included stereotypies, skin-picking behavior, self-injurious behavior, aggression, and anxiety.

14 of the 30 individuals had macrocephaly, defined by a head circumference equal to two or more standard deviations above the mean, or in the 95th percentile or greater. An additional two individuals have relative macrocephaly, defined here as head circumference one standard deviation higher than the standard deviation for the height. Individual 6 also had a *de novo* pathogenic *PTEN* (GenBank: NM_000314.6, MIM: 601728) c.493G>A (p.Gly165Arg) variant, which can also account for his macrocephaly. Other growth parameters were variable for individuals in this group, but most were in the normal range for height and weight.

Excluding individual 10, who had a hypoxic-ischemic injury, only four of the individuals in this group (4, 7, 8, and 22) had epilepsy (two or more unprovoked seizures); an additional individual (9) had a history of just one seizure at eight years of age. There was no consistent seizure semiology or epilepsy syndrome described across the individuals. Only one of the four individuals with epilepsy (7) had treatment-resistant epilepsy. 19 of the individuals had undergone at least one brain MRI. MRI findings were normal or non-specific, and there were no consistent abnormalities (Table S2). Noted abnormalities included thinning or partial agenesis of the corpus callosum (individuals 5, 12, and 15); various cysts including pineal, epidermoid, arachnoid, and ependymal (individuals 6, 7, 9, and 19, respectively); increased white-matter signal (individuals 8 and 17); hyperintense signal in the basal ganglia (individual 10); decreased volume (individuals 5, 10, 12, and 15); delayed myelination (individual 19); small areas of heterotopia (individual 20); and Chiari I malformation (individual 14).

Many of the individuals were reported to have gastrointestinal symptoms, including reflux, vomiting, or bowel motility issues; these are issues commonly seen in individuals with hypotonia. All individuals tested had normal hearing. There were no significant ophthalmological findings. There were no other recurrent health complications noted in this group. When we compared individuals with truncating variants in the terminal exon of *KMT2E* to those with earlier-truncating variants, there were no clear phenotypic differences, although the number of individuals available for comparison is small.

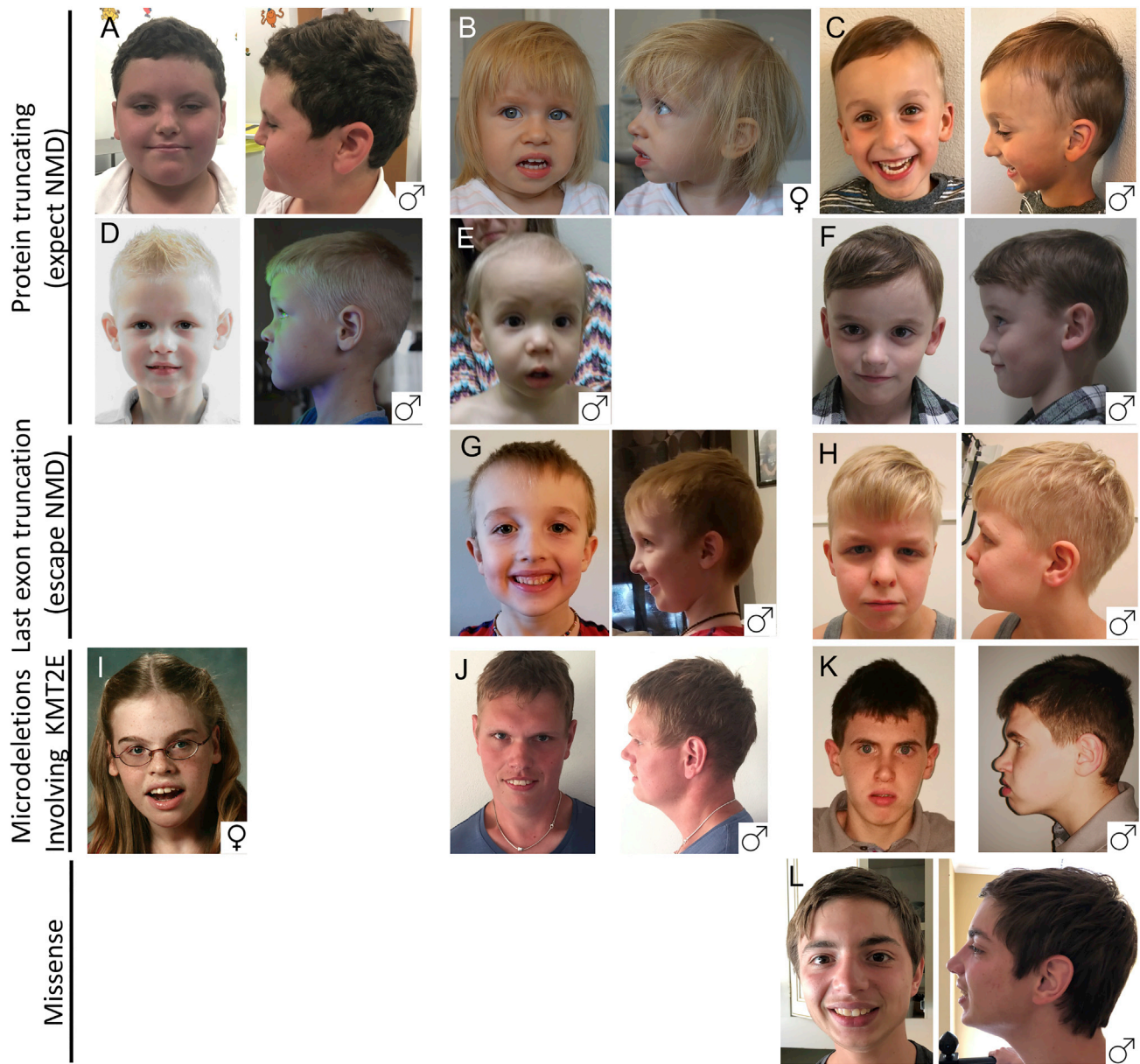


Figure 2. Photos of Individuals with *KMT2E* Variants

Each individual is noted with the corresponding number used throughout the manuscript. Included on the bottom right of each cluster is the individual's sex.

- (A) Individual 9, 11 years old
- (B) Individual 11, 1 year, 10 months old
- (C) Individual 12, 4.5 years old
- (D) Individual 13, 6 years old
- (E) Individual 15, 1 year, 7 months old
- (F) Individual 20, 6 years old
- (G) Individual 24, 5 years old
- (H) Individual 25, 12 years old
- (I) Individual 30, 18 years old
- (J) Individual 31, 22 years old
- (K) Individual 32, 7 years old
- (L) Individual 33, 16 years old

Consistent facial features include dolichocephaly, large foreheads, and deep-set eyes, often with down-slanting palpebral fissures, peri-orbital fullness, prominent cheeks, and prominent nasolacrimal folds.

It is notable that 22 out of the 30 individuals with protein-truncating variants were male. It is possible that decreased penetrance or variable expressivity of the condi-

tion in females means that fewer female individuals with *de novo* protein-truncating variants come to diagnostic attention. Additionally, the expressivity of certain aspects

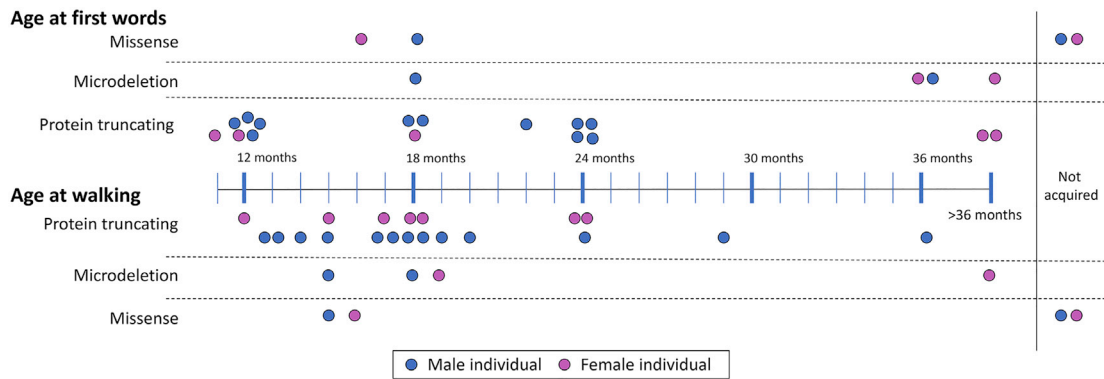


Figure 3. Developmental Milestones in Individuals with Variants in *KMT2E*

Most children with protein-truncating variants acquire first words and walking by 24 months of age, though a minority are more significantly delayed. Only individual 12, who experienced a cardiac arrest and injury, did not acquire these skills. A majority of individuals with a microdeletion had significant delay in speech development but walked at a similar time to individuals with protein-truncating variants. Of those with missense variants, those with severe infantile epilepsy had significant delays.

of the phenotype is variable between males and females (Table 2). Although the rates of intellectual disability and macrocephaly were similar, interestingly, epilepsy was seen in 43% of females but in only 5% of males ($p = 0.047$, Fisher's exact test), whereas autism was seen in 35% of males and in none of the females ($p = 0.14$, Fisher's exact test) with protein-truncating variants in *KMT2E*. These sex-related differences in phenotype parallel differences in the epidemiology of autism and epilepsy: autism is four times more common in males than in females,²⁴ whereas polygenic idiopathic generalized epilepsies are more common in females.²⁵

Of the four individuals with *de novo* 7q22.2-22.3 chromosome deletions including *KMT2E*, two were male and two were female (Figure 2). The age at most recent evaluation ranged from 7 to 22 years. Clinically, individuals with deletions presented similarly to those with truncating variants. Although the sample size is small, there appear to be more severe developmental delays in this group. The average age of first words was 34.5 months (range 18 to 48 months, Figure 3). Only two of the four individuals are verbal. Walking was delayed in all; age of walking ranged from 15 to 42 months. Three of the four individuals in this group have epilepsy (30, 31, and 32). Two of the four individuals in this group have macrocephaly (29 and 32).

Individual 32 has been previously reported.²⁶ He presented with global developmental delay, overgrowth, macrocephaly, delayed bone age, and treatment refractory generalized epilepsy. MRI of the brain demonstrated reduction of cerebral white matter, corpus callosum hypoplasia, right cerebellar hypoplasia, and an enlarged cisterna magna. Brain imaging was also performed in individuals 30 and 31. The MRI of individual 31 demonstrated global cerebral atrophy, and the MRI of individual 30 demonstrated a possible focal cortical dysplasia.

Of the four individuals with *de novo* missense variants in *KMT2E*, two were male and two were female (Figure 2).

The age at most recent evaluation ranged from 29 months to 36 years. All four of the individuals with missense variants had epilepsy. Individual 33 had five generalized tonic-clonic seizures, starting at the age of 15 years. Individuals 34, 35, and 36 all presented with infantile epileptic encephalopathy. Individual 34 developed seizures at 6 months of age, and individuals 35 and 36 both developed seizures in the neonatal period. Reported seizure semiologies include generalized tonic-clonic, tonic, atonic, and myoclonic seizures and epileptic spasms. The initial EEG in individual 35 showed burst suppression and subsequently evolved into hypsarrhythmia. The EEG in individual 36 also showed hypsarrhythmia. The EEG in individual 34 demonstrated background disorganization and multifocal and generalized epileptiform discharges. All three individuals have treatment-resistant epilepsy. Individual 34 was started on the ketogenic diet at 14 months of age, but this diet did not improve seizure control.

In our cohort, individuals with missense variants also had more severe developmental delays than did the individuals with truncating variants. Only two of the four individuals can walk independently, and none of the individuals were verbal at most recent follow-up (Figure 3). Two of the four individuals in this category have microcephaly, and the other two are normocephalic. Three of these individuals have had a brain MRI: one individual had delayed myelination, one had cerebral atrophy, and one had an incidental abnormality in the right cerebral peduncle.

Comparison of the facial features of eleven of the individuals in our cohort suggests some commonalities, including macrocephaly, dolichocephaly, high forehead, deep-set eyes, periorbital fullness, prominent cheeks, and prominent nasolabial folds (Figures 2 and 4). Utilizing Face2Gene (FDNA, Boston, MA) facial recognition software, we created a composite image from frontal photographs of these 11 individuals (excluding individual 30,

Table 2. Summarized Phenotypes by Variant Type

Subset	#	Intellectual Disability	Autism	Epilepsy	Macrocephaly	Microcephaly
Protein-Truncating Variants						
Total	30	81% (13/16)	26% (7/27)	15% (4/26)	55% (16/29)	0% (0/29)
Male	22	82% (9/11)	35% (7/20)	5% (1/19)	52% (11/21)	0% (0/21)
Female	8	80% (4/5)	0% (0/7)	43% (3/7)	63% (5/8)	0% (0/8)
Microdeletion						
Total	4	100% (4/4)	25% (1/4)	75% (3/4)	50% (2/4)	0% (0/4)
Missense						
Total	4	100% (3/3)	33% (1/3)	100% (4/4)	0% (0/3)	66% (2/3)

who wore glasses in the photograph) to represent the common facial gestalt.

KMT2E encodes a histone methyltransferase protein, a transcriptional regulator reported to play key roles in diverse biological processes, including cell-cycle progression, maintenance of genomic stability, adult hematopoiesis, and spermatogenesis. The gene is highly expressed in the brain, particularly during fetal development.¹¹ *KMT2E* appears to be distinct from other members of the KMT2 family. Most KMT2 proteins contain an enzymatically active SET domain that possesses methyltransferase function.^{9,27} Although the *KMT2E* protein contains a SET domain, its sequence and location within the protein are different from those of other members of the KMT2 family, and studies suggest that it might lack intrinsic methyltransferase activity.²⁸ However, the SET domain is still highly conserved in *KMT2E*, and it has been proposed that *KMT2E* might have an indirect effect on H3K4 methylation, possibly through transcriptional regulation of additional histone-modifying enzymes. Most members of the KMT2 family contain multiple PHD finger domains that function as H3K4 methylation readers. In contrast, *KMT2E* contains a single PHD finger domain. PHD fingers typically bind to specific epigenetic histone marks in order to recruit transcription factors and nucleosome-associated complexes to chromatin. Finally, whereas most members of the KMT2 family function as global activators of open chromatin, *KMT2E* is believed to be a repressor, although the precise mechanisms involved in *KMT2E* regulation of gene transcription have not yet been elucidated.²⁹

Of the individuals in our cohort, those who have protein-truncating *KMT2E* variants present with syndromic intellectual disability. Most individuals are functioning in the low normal to mild range of intellectual disability. Seven of the male individuals (including three of the previously reported individuals^{10–12}) have also been formally diagnosed with autism. There appears to be a subtle common facial gestalt among the individuals whose images were available for review. Additional features, albeit not obligate or specific, include macrocephaly, hypotonia, and GI dysmotility. Neuroimaging is normal or non-spe-

cific. Epilepsy was not common among individuals with protein-truncating variants. There were no significant phenotypic differences between individuals with truncating variants in the terminal exon of the gene and those with earlier-truncating variants, suggesting a probable common pathophysiology of haploinsufficiency.

Whereas, in our cohort, only approximately 14% of the individuals with protein-truncating variants have epilepsy, all of the individuals we report as having missense variants have epilepsy. This association met statistical significance ($p = 0.0026$, Fisher's exact test). Three of the individuals with missense variants fall in the category of an infantile-onset epileptic encephalopathy. In addition, these individuals have more severe developmental delays, and two have microcephaly. We hypothesize that the phenotype of epileptic encephalopathy could be variant specific and might relate to an alternate mechanism such as a gain-of-function or dominant-negative effect. Recently, distinct developmental disorder phenotypes have been identified to result from PTVs and missense variants in the same gene.^{30,31} Additional cases and further functional studies are required to clarify this.

Overall, the individuals with chromosome 7q22.2-22.3 microdeletions encompassing *KMT2E* presented similarly to those with truncating variants, further supporting haploinsufficiency as the disease mechanism. Although the sample size was small, these individuals appeared to have more severe developmental delays than did those individuals with truncating variants, which is likely explained by the influence of additional genes included in their deletions. The 7q22.2-22.3 region contains multiple additional genes involved in the regulation of the cell cycle; such genes include *SRPK2* (MIM: 602980), *RINT1* (MIM: 610089), and *LHFPL3* (MIM: 609719).²⁶ In particular, *SRPK2* and *LHFPL3* show depletion of loss-of-function variation from expectation in the gnomAD database (pLI of 1.0 and 0.9, respectively) and are expressed in the central nervous system. *SRPK2* encodes a cell-cycle-regulated protein kinase that phosphorylates serine and arginine domain-containing proteins and modulates pre-mRNA splicing in neurons.³² *LHFPL3* is a transmembrane protein, but little is known about its function to date.

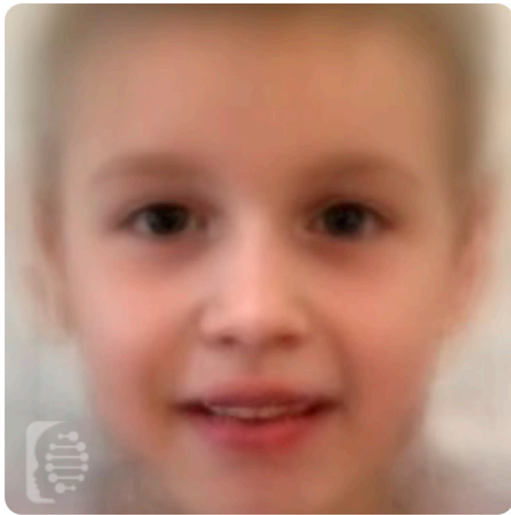


Figure 4. Composite Photo from Face2Gene
Individuals in Figure 2 were used in this analysis, excluding individual 30, who is wearing glasses.

Several *Kmt2e* (*Mll5*)-deficiency mouse models have been created and characterized.^{29,33–36} These mice present with growth restriction and increased mortality, as well as impaired hematopoiesis. A neurological phenotype in these mice has not been reported. Both homozygous and heterozygous loss of *Kmt2e* in mice results in DNA damage and elevated levels of reactive oxygen species (ROS).³⁶ The cellular effects were effectively reversed by supplementation with the glutathione precursor N-acetylcysteine (NAC).³⁶ This has interesting therapeutic implications for humans because NAC supplementation has been used in the treatment of glutathione depletion in acetaminophen overdose as well as rare inborn errors of metabolism associated with increased free-radical damage. Further studies are required if we are to establish whether humans haploinsufficient for *KMT2E* are also vulnerable to increased ROS and whether there might be a benefit in treating with NAC or other antioxidants. This evaluation could include clinically measuring urine F2 isoprostanes and blood glutathione levels.³⁷

In this report, we define a *KMT2E*-related neurodevelopmental disorder, which adds to the growing list of *KMT2* gene family disorders. Most individuals with protein-truncating variants appear to present with generally mild developmental delay and intellectual disability. Autism is also relatively common. Additional common, but not obligate, features include relative macrocephaly, hypotonia, and functional gastrointestinal disturbances. There appears to be a subtle facial gestalt. Epilepsy was not common among individuals with protein-truncating variants. We suspect haploinsufficiency as the disease mechanism. The similar phenotype seen in individuals with microdeletions of this region is consistent with this hypothesis. In contrast, individuals with missense variants all presented with epilepsy, including infantile-onset epileptic encephalopathy, and more severe developmental delays. Variant-

specific alterations in *KMT2E* function, possibly even gain-of-function alterations, might explain this divergence in phenotype. Further studies are required if we are to further understand the correlation between genotype and phenotype. There is no established therapy for *KMT2E*-related disorders, although based on animal data, there might be a role for NAC or other antioxidant treatments.

Supplemental Data

Supplemental Data can be found online at <https://doi.org/10.1016/j.ajhg.2019.03.021>.

Consortia

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here are those of the authors and not necessarily those of the Wellcome or the Department of Health.

Declaration of Interests

A.T., C.R., H.M.M., I.M.W., K.M., R.H., and R.P. are employees of GeneDx, a wholly-owned subsidiary of OPKO Health. F.K. and L.L.P.R. are employees of Mendelics Genomics Analysis.

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Web Resources

CADD, <https://cadd.gs.washington.edu/>

ClinVar, <https://www.ncbi.nlm.nih.gov/clinvar>

DECIPHER, <https://decipher.sanger.ac.uk/>

GenBank, <https://www.ncbi.nlm.nih.gov/genbank/>

GeneMatcher, <https://genematcher.org/>

Genome Aggregation Database (gnomAD), <https://gnomad.broadinstitute.org>

HMIMER, <http://hmmer.org/>

InterProScan, <https://www.ebi.ac.uk/interpro/search/sequence-search>

MyGene2, NHGRI/NHLBI University of Washington-Center for Mendelian Genomics (UW-CMG), Seattle, WA, <https://www.mygene2.org/MyGene2/>

NetPhos 3.1, <http://www.cbs.dtu.dk/services/NetPhos/>

Online Mendelian Inheritance in Man (OMIM), <https://omim.org/>
Phyre2, <http://www.sbg.bio.ic.ac.uk/~phyre2/html/page.cgi?id=index>

UCSC Human Genome Browser, <http://www.genome.ucsc.edu>

Variant Validator, <https://variantvalidator.org/variantvalidator/>

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Supplemental Data

Heterozygous Variants in *KMT2E* Cause a Spectrum of Neurodevelopmental Disorders and Epilepsy

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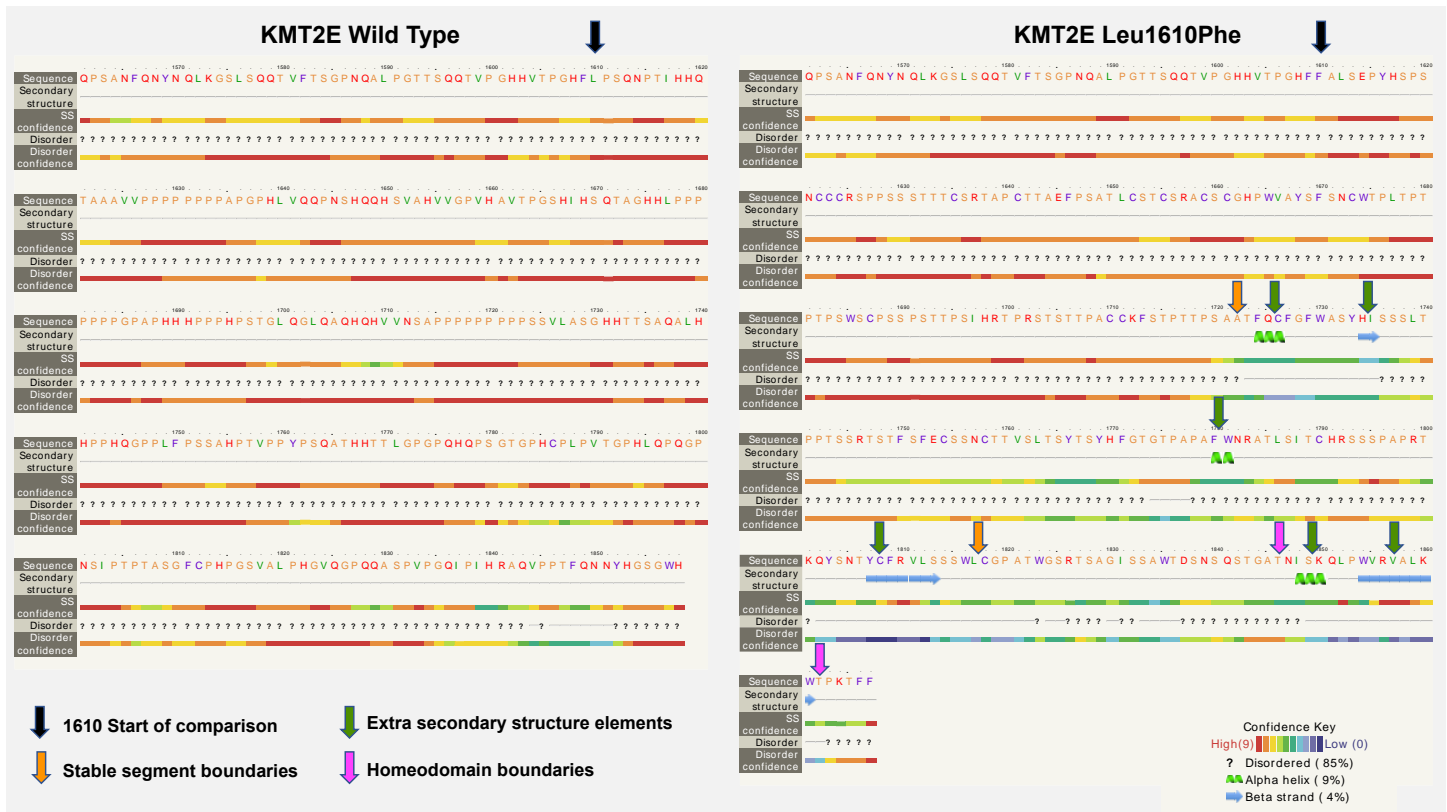


Figure S1. Frameshifting variants acquire stabilizing secondary structure elements. PHYRE¹⁹ secondary structure report for the C-terminal fragments of KMT2E wild type (left panel, amino acids 1560-1858) and KMT2E frameshifting variant Leu1610Phe (right panel, amino acids 1560-1867). Black arrows indicate the start of comparison at position 1610, after which the two variants are completely different due to a frameshift variant. The orange arrows indicate the boundaries of the stabilizing segment present in p.Leu1610Phefs*259 and p.Val1625Argfs*244 but not in wild type. The green arrows indicate the helices and strands not present in wild type. The pink arrows indicate the boundaries of the C-terminal predicted homeodomain present in p.Leu1610Phefs*259 and p.Val1625Argfs*244 but not in wild type. The confidence key (bottom right) indicates the reliability of secondary structure prediction (helix, strand, disorder) at each position.