Elucidation of the phenotypic spectrum and genetic landscape in primary and secondary microcephaly

Paranchai Boonsawat, MSc1, Pascal Joset, PhD1, Katharina Steindl, MD1, Beatrice Oneda, PhD1, Laura Gogoll, MD1, Silvia Azzarello-Burri, MD1, Frenny Sheth, PhD2, Chaitanya Datar, MD3, Ishwar C. Verma, MD4, Ratna Dua Puri, MD4, Marcella Zollino, MD5, Ruxandra Bachmann-Gagescu, MD1, Dunja Niedrist, MD1, Michael Papik, MSc1, Joana Figueiro-Silva, PhD1, Rahim Masood, MD1, Markus Zweier, PhD1, Dennis Kraemer, MSc1, Sharyn Lincoln, MS6, Lance Rodan, MD6,7, Undiagnosed Diseases Network (UDN), Sandrine Passemard, MD8,9, Séverine Drunat, PhD9, Alain Verloes, MD, PhD9, Anselm H. C. Horn, PhD10, Heinrich Sticht, PhD10, Robert Steinfeld, MD11, Barbara Plecko, MD11,12, Beatrice Latal, MD13, Oskar Jenni, MD13, Reza Asadollahi, MD1 and Anita Rauch, MD1,14,15

Purpose: Microcephaly is a sign of many genetic conditions but has been rarely systematically evaluated. We therefore comprehensively studied the clinical and genetic landscape of an unselected cohort of patients with microcephaly.

Methods: We performed clinical assessment, high-resolution chromosomal microarray analysis, exome sequencing, and functional studies in 62 patients (58% with primary microcephaly [PM], 27% with secondary microcephaly [SM], and 15% of unknown onset).

Results: We found severity of developmental delay/intellectual disability correlating with severity of microcephaly in PM, but not SM. We detected causative variants in 48.4% of patients and found divergent inheritance and variant pattern for PM (mainly recessive and likely gene-disrupting [LGD]) versus SM (all dominant de novo and evenly LGD or missense). While centrosome-related pathways were solely identified in PM, transcriptional regulation was the most frequently affected pathway in both SM and PM.

Unexpectedly, we found causative variants in different mitochondria-related genes accounting for ~5% of patients, which emphasizes their role even in syndromic PM. Additionally, we delineated novel candidate genes involved in centrosome-related pathway (SPAG5, TEDC1), Wnt signaling (VPS26A, ZNRF3), and RNA trafficking (DDX1).

Conclusion: Our findings enable improved evaluation and genetic counseling of PM and SM patients and further elucidate microcephaly pathways.

Keywords: primary microcephaly; secondary microcephaly; MCPH; genetic counseling; mitochondria

INTRODUCTION

Microcephaly is a clinical finding defined as an occipitofrontal head circumference (OFC) of >2 SDs below the mean for age, sex, and ethnicity, which affects approximately 2–3% of the population worldwide. Individuals with microcephaly, especially those with an OFC <−3 SD, can manifest neurological features that require medical attention and a search for the underlying etiology among environmental or, more commonly, genetic factors.2 Microcephaly is classified into primary (PM) if present at birth, and secondary (SM) if developing thereafter.3 Accordingly, PM has been shown frequently to result from early defects in neurogenesis due to abnormal regulation of mitotic division, while SM has been often linked to disruptions of later developmental processes such as myelination and synapse formation owing to abnormal endosome regulation, vesicle membrane transport, or synaptic structural support.4,5 However, neuronal migration, DNA repair, and transcription...
Microcephaly can be nonsyndromic or present as an associated feature in a variety of genetic syndromes. Currently, there are over 900 OMIM phenotype entries and almost 800 genes linked to microcephaly with variable expressivity. Particularly, 18 of these genes constitute a distinct PM subclass, termed autosomal recessive primary microcephaly or microcephaly primary hereditary (MCPH), a form of microcephaly that is relatively consistent and thus far better characterized. On the other hand, SM and non-MCPH PM show considerable heterogeneity; this has not been properly studied so far and hence remained largely elusive.

Previous studies on patients with microcephaly using clinical and radiological information as well as metabolic and targeted genetic testing were able to identify causes in a small fraction of the patients (<20%) (refs. 2,8). Since the advent of next-generation sequencing (NGS), mainly mixed cohorts of neurodevelopmental disorders (NDDs) have been assessed where microcephalic patients accounted for ~15–41% of the cases and on average ~47% of them were identified with a definite cause using exome (ES) or genome sequencing (GS). Until now, there are only two studies that used Mendelome sequencing or ES to evaluate known disease-causing or candidate genes in exclusive microcephaly cohorts. The first study determined a molecular diagnosis for ~29% of the cases (11/38), but did not differentiate between PM and SM. The other study was focused on PM and MCPH from mainly consanguineous families showing the difficulties in their clinical definitions and common overlap with microcephalic primordial dwarfism, and proposed reconsideration of phenotypic boundaries.

Here, we performed a comprehensive genetic study on a cohort of 62 unselected clinically well-characterized patients with syndromic or nonsyndromic microcephaly of different onset using combined high-resolution chromosomal microarray analysis (CMA) and ES. Our approach sheds light on the genetic landscape of PM and SM and delineates their respective clinical and molecular characteristics. In addition to novel clinical and molecular findings in known disease genes, we identified several novel NDD/microcephaly candidate genes.

**MATERIALS AND METHODS**

**Patient recruitment**

Sixty-two unrelated patients, including both syndromic and nonsyndromic, were recruited from 2015 to 2017, clinically assessed in detail, and subjected to defined genetic evaluations (Figure S1). Inclusion criteria consisted of (1) an OFC >2 SDs below the mean at birth or later, based on World Health Organization (WHO) and established growth charts; (2) no clear evidence for an acquired etiology or history of perinatal infection; and (3) without an unequivocal etiological diagnosis after clinical assessment by pediatricians and clinical geneticists (Figure S1). We performed CMA and ES for all patients, and conventional karyotyping for 45 patients including all those who remained undiagnosed after CMA and ES analysis. Genetic testing was performed as part of a research study approved by the ethics commission of the Canton of Zurich or referral centers. Written informed consent for genetic testing, publication of clinical information, and/or photographs were obtained.

**CMA**

CMA for evaluation of rare coding copy-number variants (CNVs) was performed on DNA extracted from peripheral blood using Affymetrix Cytoscan HD or cytogenetic 2.7 M arrays as previously described.

**ES and Sanger sequencing**

ES was performed on DNA extracted from peripheral blood using Agilent SureSelect XT Clinical Research Exome Kit (V5) or Human All Exon (V6) on a HiSeq 2500 System (Illumina, CA, USA) with 125-bp paired-end reads as described elsewhere. ES was done as trios (index patient and parents) in 58 families and duos (index patient and mother due to the lack of paternal DNA) in 4 families. ES coverage for targeted bases and off-target mitochondrial bases, and their distribution among diagnosed and undiagnosed patients, are shown in Fig. 1a. Coding plus flanking intronic (±6 bp) regions as well as 666 previously reported mitochondrial DNA variants in 37 mitochondrial genes from the MITOMAP database were analyzed using the NextGENe Software (SoftGenetics, PA, USA) (Figure S1). A second allele search for all de novo variants in recessive OMIM morbid genes or in high-level candidate genes was performed (Supplementary Materials and Methods). Selected variants from ES were confirmed by Sanger sequencing using an AB3730 capillary sequencer (Applied Biosystems, CA, USA).

**Variant classification**

Rare coding CNVs were classified according to Miller et al. Rare (minor allele frequency [MAF] ≤2%) sequence variants (SVs) affecting genes known to cause Mendelian disorders were classified according to the American College of Medical Genetics and Genomics (ACMG) guidelines. De novo, X-linked maternal, or biallelic variants affecting other genes were classified as suspected candidates, candidates, or high-level candidates according to our defined criteria (Figure S1).

**Functional evaluations of selected variants**

Structural modeling, cell culture, reverse transcriptase polymerase chain reaction (RT-PCR) and quantitative RT-PCR (qRT-PCR), immunoblotting, immunofluorescence, and imaging were performed to evaluate functional consequences of selected variants (Supplementary Materials and Methods).
RESULTS

Cohort characteristics
We enrolled 62 unrelated patients (29 females, 33 males) with microcephaly of unknown etiology from 62 families (Table S1). PM and SM were determined in 36 (58.1%) and 17 (27.4%) patients, respectively (Table 1 and Fig. 1b). In the other 9 (14.5%) patients, the onset of microcephaly could not be determined. The median age at last investigation was 5.4 years (mean: 6.5 years, 0.8–18), for PM 4.5 years (mean: 5.3 years), and for SM 4.3 years (mean: 5.7 years). The majority were of European descent (77.4%) and the remaining were of Middle Eastern/North African (12.9%) or Indian (9.7%) ancestry. Nine (14.5%) patients were born to consanguineous parents. Seven patients had one or more affected siblings. Notably, follow-up OFC measurements showed a pattern of progressive microcephaly in both PM and SM with a statistically significantly higher OFC reduction in PM than in SM patients (p < 0.001, Wilcoxon rank-sum test) (Fig. 1b). However, 61.3% of PM and 70.6% of SM patients did not show a decline in length or height similar to that in OFC, indicating a disproportionate microcephaly in the majority of our patients (Fig. 1b). Apart from microcephaly, varying degrees of different neurological signs were reported, among which abnormal developmental milestones (developmental delay [DD] or ID) and abnormal cerebral magnetic resonance image (MRI) represented the most common associated features (Table 1). Importantly, we observed that the severity of DD/ID was significantly correlated with the severity of microcephaly among our PM patients (Figure S2A, r = −0.43, p = 0.01, Spearman rank correlation with Bonferroni
Table 1 Summary of main clinical features in our cohort of 62 patients

| Main clinical features                  | Number of cases |
|----------------------------------------|-----------------|
| Microcephaly                           | 62/62 (100%)    |
| Primary                                | 36/62 (58.1%)   |
| Secondary                              | 17/62 (27.4%)   |
| Unknown onset                          | 9/62 (14.5%)    |
| DDa                                    | 58/61b (90.3%)  |
| Mild                                   | 13/61 (21.3%)   |
| Mild to moderate                       | 8/61 (13.1%)    |
| Moderate                               | 12/61 (19.7%)   |
| Moderate to severe                     | 7/61 (11.5%)    |
| Severe                                 | 15/61 (24.6%)   |
| Severity not determined                | 3/61 (4.9%)     |
| IDb                                    | 24/28c (85.7%)  |
| Mild                                   | 7/28 (25%)      |
| Mild to moderate                       | 1/28 (3.6%)     |
| Moderate                               | 8/28 (28.6%)    |
| Moderate to severe                     | 5/28 (17.9%)    |
| Severe                                 | 3/28 (10.7%)    |
| Abnormal cerebral MRI                  | 27/43 (62.8%)   |
| Epilepsy/seizures                      | 16/61 (26.2%)   |
| Ataxia or movement disorder            | 15/61 (24.6%)   |
| Behavioral problems                    | 14/61 (23%)     |
| Strabism                               | 6/61 (9.8%)     |
| Hearing problems                       | 6/61 (9.8%)     |
| Short stature                          | 20/61 (32.8%)   |
| Complex congenital heart defect        | 4/61 (6.6%)     |

*a* Developmental delay (DD) and intellectual disability (ID) were classified based on the Diagnostic and Statistical Manual of Mental Disorders (DSM-5).37

b1/62 cases was a fetus.

b2/61 patients were above the age of 5 years at last investigation and 28/32 were evaluated for severity of ID. 3/28 patients had learning disability and 1/28 had normal intelligence.

c6/1 cases was a fetus.

dCerebral magnetic resonance image (MRI) was done for 43 patients.

eCerebral microcephaly.

**Genetic findings**

We identified pathogenic or likely pathogenic (P/LP) causative variants in 48.4% of the patients (Table 2), and variants of uncertain significance (VUS) in another 4.8% of the patients (Fig. 1c and Table S1). Furthermore, we found likely deleterious variants affecting our novel high-level candidate genes in another 8.1%, affecting our novel (suspected) candidate genes in another 17.7%, and we found no (candidate) causative variant in 21% of the patients (Fig. 1c). We did not find a second disease allele for any patient with inherited heterozygous likely gene-disrupting (LGD) variants in established genes known to cause recessive disorders by our alternative methods (Supplementary Materials and Methods). In six (9.7%) patients, we found P/LP inherited heterozygous variants as secondary findings (Supplementary Results).

**P/LP variants**

We identified pathogenic CNVs in six (9.7%) patients (4 deletions, 2 duplications; 5 [assumed] de novo, 1 X-linked recessive inheritance) (Fig. 1c and Table 2). In one of these patients (ID74601) who had a de novo pathogenic ~1.5-Mb duplication, we identified an additional pathogenic de novo sequence variant (SV) c.3555_3556insA, p.(Ala1186Serfs*5) in KA76A (NM_001099412.1), which likely contributes to the severity of his NDD phenotype (Table 2 and S1). Furthermore, we identified possible additional hits which may contribute to the expressivity of microcephaly in another patient (ID70688) who was identified with a pathogenic correction but not among SM patients or the total cohort (Figure S2B–D). In addition, we found a significant correlation between the severity of DD/DD and abnormal cerebral MRI among the total cohort (Figure S2E, F, p < 0.01, Fisher’s exact test with Bonferroni correction).
| No. | Patient ID | Age | Gender | Microcephaly subgroup | Main clinical feature | Cerebral MRI finding | Genetic finding | Disorder | Function/Pathway |
|-----|------------|-----|--------|-----------------------|----------------------|---------------------|------------------|----------|-----------------|
| 1   | 59571      | 10.9 PM | Male   | Moderate global DD, moderate–severe ID, muscular hypotonia, myopia, short stature, scoliosis | Abnormal gyrus, corpus callosum dysgenesis, everted hippocampi | No abnormality in cerebral MRI | CDK5RAP2: c.[4546G>T];[3928G>T], p.(Glu1516*);(Glu1310*), AR (CH) | MCPH3 (MIM 604804) | Cerebro, spine, and microtubule organization |
| 2   | 73869      | 8.8 PM | Female | Mild motor DD, moderate speech delay, mild feeding difficulties, cryptorchidism | Dysmorphic corpus callosum with absence of the splenium, hypoplasia of frontal horns of lateral ventricles | No abnormality in cerebral MRI | CDK5RAP2: c.[558_559del];[4441C>T], p.(Glu186Aspfs*32);(Arg1481*), AR (CH) | MCPH3 (MIM 604804) | Cerebro, spine, and microtubule organization |
| 3   | 74812      | Aborted at 23 GW | | | | Autopsy: Absence of corpus callosum | PLK4: c.[1111C>T];[881T>G], p.(Arg371*);(Ile294Ser), AR (CH) | PLK4-related disorder (MIM: 616171) | Centriole duplication |
| 4   | 77804      | 0.8 PM | Male   | Moderate global DD, movement disorder | Dysmorphic corpus callosum, larger cerebellum and brain stem relative to the supratentorial region | No abnormality in cerebral MRI | PLK4: c.[212C>T];[760_761insAC], p.(Ser71Phe);(Leu254Tyrfs*2), AR (CH) | PLK4-related disorder (MIM: 616171) | Centriole duplication |
| 5   | 68978      | 2.1 PM | Male   | Severe global DD, seizure | Microlissencephaly with simplified cortical pattern | | ASPM: c.[3796G>T];[3796G>T], p.(Glu1266*);(Glu1266*), AR (Homo) | MCPH5 (MIM 608716) | Spindle organization and orientation |
| 6   | 57602      | 3.8 PM | Male   | Severe global DD, spastic hemiparesis, patent ductus arteriosus, vesicoureteral reflux | Dysmorphic corpus callosum, enlarged cerebellum, xanthochromic CSF, and brain stem relative to the supratentorial region | | DHTKD1: c.[2185G>A];[2185G>A], p.(Gly729Arg);(Gly729Arg), AR (Homo) | 2-aminoadipic-2-oxoadipic aciduria (MIM 204750) | Amino acid degradation, mitochondrial biosynthesis |
| 7   | 75595      | 3 PM | Male   | Severe global DD, decreased reflexes, severe failure to thrive, congenital cataract, callosal hypomyelination | Dysmorphic corpus callosum, enlarged cerebellum, xanthochromic CSF, and brain stem relative to the supratentorial region | | ERCC6: c.[850G>T];[850G>T], p.(Glu284*);(Glu284*), AR (Homo) | Cockayne syndrome (MIM 133540) | DNA repair |
| 8   | 75892      | 2 PM | Female | Severe global DD, spastic hemiparesis, patent ductus arteriosus, vesicoureteral reflux | Dysmorphic corpus callosum, enlarged cerebellum, xanthochromic CSF, and brain stem relative to the supratentorial region | | TRAPPC9: c.[3214C>T];[3214C>T], p.(Arg1072*);(Arg1072*), AR (Homo) | Autosomal NF-kappa B signaling | Transcriptional regulation |
| 9   | 53792      | 8.3 PM | Male   | Severe global DD, mild ID, hypotonia, seizures, short stature, uvula bifida, mild truncal adiposity | Dysmorphic corpus callosum, enlarged cerebellum, xanthochromic CSF, and brain stem relative to the supratentorial region | | TRMT10A: c.[379C>T];[379C>T], p.(Arg127*);(Arg127*), AR (Homo) | TRMT10A-related disorder (MIM 616033) | DNA repair |

**Note:** The table provides a summary of main clinical features and genetic findings in patients with P/LP or high-level candidate variants in established disease genes.
| No. | Patient ID | Age | Microcephaly subgroup | Main clinical feature                        | Cerebral MRI finding                                      | Genetic finding                                                                 | Disorder                                                                 | Function/pathway |
|-----|------------|-----|-----------------------|---------------------------------------------|----------------------------------------------------------|-------------------------------------------------------------------------------|---------------------------------------------------------------|------------------|
| 10  | 74601      | 2.3 | PM                    | Severe global DD, cerebral movement disorder, craniosynostosis, delayed visual maturation, strabismus, hyperopia, astigmatism, torticollis, atrial septal defect | Right and left plagiocephalus                              | 1.48-Mb dup in chr2:96732519-98212850, DN                                      | 2q11.2 duplication (Riley et al.38)                            | –                |
|     |            |     |                       |                                             |                                                          | KAT6A: c.3555_3556insA, p.(Ala1186Serfs*5), DN                      | Autosomal dominant mental retardation 32 (MIM 616268)                         | Transcriptional regulation                                      |
| 11  | 74444      | 4.5 | PM                    | Mild global DD, muscular hypotonia, epilepsy, left central cataract, left chorioretinal coloboma, left strabismus and nystagmus | Multiple subependymal heterotopia in lateral ventricles   | 2-Mb del in chr1:145940520-147922681, DN                                | Chromosome 1q21.1 deletion syndrome (MIM 612474)                    | –                |
| 12  | 74579      | 2.2 | PM                    | Mild global DD, generalized hypotonia, bilateral cleft lip and cleft palate, recurrent obstipation | NA                                                       | 7.1-Mb del in chr1:172427631-179864641, DN                                | 1q24q25 deletion (Chatron et al.39)                             | –                |
| 13  | 70547      | 7.1 | PM                    | Moderate speech delay, moderate–severe ID, concentration deficit, EEG abnormalities, mitral valve prolapse, cyanotic episode, contractures, hip dysplasia | Enlarged subarachnoidal space                              | HDAC8: c.522C>A, p.(Tyr174*), DN                                      | Cornelia de Lange syndrome 5 (MIM 300882)                        | Transcriptional regulation                                   |
| 14  | 32410      | 15.3| PM                    | Moderate global DD, moderate ID, bilateral hearing loss, severe progressive optic atrophy, moderate hyperopia, distal spasticity, neurogenic clubfeet, growth deficiency, steroid-resistant focal segmental glomerulosclerosis | Cerebellar hypoplasia, bilateral atrophy of optical nerves | MT-ATP6: m.9185T>C, p.(Leu220Pro), DN                                      | MT-ATP6-related mitochondriopathy                              | ATP synthesis    |
| 15  | 70688      | 1.6 | PM                    | Moderate global DD, high palate, inguinal hernia, bilateral undescended testes, sacral dimple | Abnormal signal intensity in the posterior horns of lateral ventricles, prominent Robin–Virchow area | 736 Kb dup in chrX:53480222-54215972, and 660 Kb dup in chr16:29580020-30240227, XLR | Chromosomes Xp11.22 and 16p11.2 microduplications syndromes (MIM 300705 and 614671) | –                |
| 16  | 75473      | 2.5 | PM                    | Severe global DD, 2 episodes of seizure | Hyoplasia of cerebellar vermis                           | SLC9A6: c.[615_616insT]; [0], p.[Arg206Serfs*58]; [0], XLR                  | Christianson syndrome (MIM 300243)                              | Endosome regulation                                          |
| No. | Patient ID | Age | Microcephaly subgroup | Main clinical feature | Cerebral MRI finding | Genetic finding | Disorder | Function/pathway |
|-----|------------|-----|-----------------------|----------------------|---------------------|-----------------|----------|------------------|
| 17  | 72526      | 2.8 | SM                    | Mild–moderate global DD, brachycephaly, plagiocephaly, atactic gait, seizures, mild thrombopenia, discrete microcytic anemia | Simplified gyri, slight ventricular dilatation, slightly flattened nucleus caudatus and small pituitary gland | DYRK1A: c.665-4del, r.665_951del, p.(Ile222Aspfs*22), DN | Autosomal dominant mental retardation 7 (MIM 614104) | Neuronal proliferation, differentiation, plasticity and death |
| 18  | 59484      | 8   | SM                    | Severe global DD, severe ID, hypotonia, coordination problem, epilepsy, strabismus convergence, uvula bifida, growth deficiency | No abnormality in cerebral MRI | FBXO11: c.1868C>G, p.(Thr623Arg) (Gregoret al.,35) | FBXO11-related disorder (MIM 618089) | Protein ubiquitination |
| 19  | 57570      | 3.5 | SM                    | Severe global DD, muscular hypotonia, growth deficiency, complex pulmonary atresia with ventricular septal defect, immune deficiency (T-cell lymphopenia), left vesicoureteral reflux | N/A | KMT2A: c.8724del, p.(Glu2908Aspfs*21), DN | Wiedemann–Steiner syndrome (MIM 605130) | Transcriptional regulation |
| 20  | 72938      | 2.9 | SM                    | Severe global DD, cyanotic seizures, pain insensitivity, stereotypic movements, inappropriate laughter, tonus dysregulation, cataract, dysphagia | Delayed myelination | NACC1: c.892C>T, p.(Arg298Trp), DN | NACC1-related disorder (MIM 617393) | Transcriptional regulation |
| 21  | 69444      | 0.8 | SM                    | DD, muscular hypotonia | No abnormality in cerebral MRI | POGZ: c.1580A>G, p.(Asp527Gly), DN | White–Sutton syndrome (MIM 618364) | Mitotic progression |
| 22  | 73824      | 0.9 | SM                    | Mild–moderate motor DD, mild speech delay, hypotonia, seizures, pulmonary stenosis | No abnormality in cerebral MRI | PTPN11: c.923A>G, p.(Asn308Ser), DN | Noonan syndrome (MIM 163950) | Cell growth, differentiation, mitotic cycle |
| 23  | 67093      | 5.8 | SM                    | Moderate global DD, muscular hypotonia, behavior abnormality, feeding difficulties, recurrent subfebrile temperature, left iris coloboma | N/A | TLK2: c.968+2T>G, p.T, DN | Autosomal dominant mental retardation 57 (MIM 618050) | DNA repair |
| 24  | 74956      | 2.7 | SM                    | Mild–moderate DD, muscular hypotonia, strabismus (Duane syndrome), short stature, vesicoureteral reflux due to renal tubular ectasia, joint laxity, chronic obstruction | Partial agenesis of the corpus callosum | 509 Kb del in chr17:43703801-44212416, assumed DN (absent in mother) | Koolen–de Vries syndrome (MIM 610443) | – |
| 25  | 76870      | 16.2| Unknown onset         | Moderate–severe global DD, moderate–severe ID, epilepsy, behavior abnormalities, hearing loss, retinal dystrophy, abnormality of dental enamel, recurrent infections | N/A | KARS: c.[1772A>T]; [1772A>T], p.[(Asn591Gln)]; [(Asn591Gln)], AR (Homo) | Autosomal recessive deafness 89 (MIM 613916) | Protein translation |
### Patients with P/LP findings in established disease genes

| No. | Patient ID | Agea | Microcephaly subgroup | Main clinical feature | Cerebral MRI finding | Genetic finding | Disorder | Function/pathway |
|-----|------------|------|-----------------------|----------------------|---------------------|----------------|----------|------------------|
| 26  | 45969      | 15.3 | Unknown onset         | Severe global DD, severe ID, epilepsy, ataxic movement disorder, stereotypy, sleep disturbances, intermittent hyperventilation, impulsivity, scoliosis | N/A                 | STXBP1: c.586C>T, p.(Arg122*), DN | Early infantile epileptic encephalopathy 4 (MIM 612164) | Neurotransmitter release |
| 27  | 55113      | 18   | Unknown onset         | Learning disability (no DD), ADHD, myopia, short stature | Arachnoidal cyst | TRIO: c.4615-2del, p.?, DN | Autosomal dominant mental retardation 44 (MIM 617061) | Neuronal migration |
| 28  | 31773      | 17.8 | Unknown onset         | Mild motor DD, moderate speech delay, mild ID, mild muscular hypotonia, ataxia, growth deficiency | N/A                 | 10.8-Mb del in chr7:87365891-98118059, DN | – | – |
| 29  | 65891      | 12.8 | Unknown onset         | Moderate–severe global DD, moderate ID, axial hypotonia, distal hypertonia (passive), spastic paraparesis, hyperactivity, sleep apnea, short stature, scoliosis, hypoplasia of the pituitary gland | N/A                 | MECP2: c.1138_1144del[0];[0], p.[(Val350Cysfs*27)];[0], XLR | Syndromic X-linked mental retardation 13 (MIM 300055) | Transcriptional regulation |
| 30  | 66916      | 10.5 | Unknown onset         | Fine motor problems, moderate–severe speech delay, moderate ID, dysmetria, short stature | N/A                 | PQBP1: c.459_462del[0];[0], p.[(Arg153Serfs*41)];[0], XLR | Renpenning syndrome (MIM 309500) | Transcriptional regulation |

### Patients with likely deleterious variants in high-level candidate genes

| No. | Patient ID | Agea | Microcephaly subgroup | Main clinical feature | Cerebral MRI finding | Genetic finding | Protein | Function/pathway |
|-----|------------|------|-----------------------|----------------------|---------------------|----------------|---------|------------------|
| 1   | 81652      | 2.3  | PM                    | Mild speech delay, short stature | No abnormality in cerebral MRI | SPAG5: c.[3189C>T];[1223_1224insAC], r.[3189_3198del][=], p.[(Gly1064Glu*3)];[(Lys409Profs*19)], AR (CH) | Sperm associated antigen 5 | Centriole duplication |
| 2   | 68629      | 5.7  | PM                    | Moderate global DD, right plagiocephaly, bilateral hyperopia, short stature, congenital primary hypothyroidism (treated), bilateral cryptorchidism | No abnormality in cerebral MRI | TEDC1 (C14orf80): c.[227-5C>G][1111del], r.[227_267del][=], p.[(Glu76Glyfs*11)];[(Ala371Glnfs*12)], AR (CH) | Tubulin epsilon and delta complex 1 | Centriole stability |
| No. | Patient ID | Age | Microcephaly subgroup | Main clinical feature | Cerebral MRI finding | Genetic finding | Protein | Function/pathway |
|-----|------------|-----|-----------------------|----------------------|---------------------|----------------|---------|------------------|
| 3   | 74091      | 1.8 | PM                    | Severe global DD, spasticity, decreased hearing of both sides, short stature, low weight, vesicoureteral reflux | Atrophy of the white matter, a Dandy-Walker variant and a generalized hypertrophy of the cerebral and cerebellar structures, pontocerebellar hypoplasia | DDX1: c.[1333G>A]; [1333G>A], p.[Val445Ile]; [Val445Ile], AR (Homo) | Asp-Glu-Ala-Asp (DEAD) box helicase 1 | RNA trafficking |
| 4   | 75822      | 4.5 | PM                    | Moderate–severe global DD, hirsutism | Bilateral frontal pachygria, delayed myelination, partial agenesia of posterior corpus callosum | VPS26A: c.[404C>T]; [404C>T], p.[Thr135Ile]; [Thr135Ile], AR (Homo) | VPS26, retromer complex component A | Retrograde transport of proteins from endosomes to the trans-Golgi network; Wnt signaling |
| 5   | 60361      | 4.8 | PM                    | Mild speech delay, hyperopia, left strabismus convergens, coarctation of the aorta, hypoplastic aortic arch, ventricular septal defect, atrial septal defect, bicuspid aortic valve, and persistent left superior vena cava, and ectodermal dysplasia, brachydactyly, mainly distally shortened phalanges, nail hypoplasia, lacrimal duct obstruction, oligodontia, sensitive and dry skin | N/A | ZNRF3: c.311T>C, p.(Leu104Pro), DN | Zinc and ring finger 3 | Negative regulator of the Wnt signaling pathway |

ADHD: attention deficit–hyperactivity disorder, AR: autosomal recessive, ATP: adenosine triphosphate, CH: compound heterozygous, DD: developmental delay, DN: de novo, EEG: electroencephalogram, GW: gestational weeks, Homo: homozygous, ID: intellectual disability, MCPH: microcephaly primary hereditary, MRI: magnetic resonance imaging, N/A: not available, P/LP: pathogenic or likely pathogenic, PM: primary microcephaly, SM: secondary microcephaly, XLR: X-linked recessive.

*Age at last investigation.
Xp11.22 microduplication affecting HUWE1, PHF8, and FAM120C, a CNV known to cause X-linked ID (MIM 300705) without microcephaly (Table 2 and S1). These hits include an additional microcephaly-related 16p11.2 microduplication (MIM 614671) and a hemizygous nonsense unreported variant c.901C>T, p.(Arg301*) in the last exon of ASB11 (NM_008073.2), which has not been yet linked to any disorder, but encodes an E3 ubiquitin protein ligase with an established role in canonical Notch signaling to regulate proper neurogenesis. Therefore, it is possible that these two additional variants may contribute to the manifestation of microcephaly in this patient.

Among the other 56 patients, we identified P/LP SVs affecting 22 different genes in 24 patients (CDK5RAP2 and PLK4 each in two patients), adding up to a total diagnostic yield of 48.4% (Fig. 1c). Among the diagnosed patients (n = 30), only one (~3%) PM patient was born to consanguineous parents. Considering the two microcephaly subgroups, we found comparable diagnostic yields of 44.5% in PM and 47.1% in SM (Fig. 1d). Notably, we observed recessive inheritance in 68.8% and dominant de novo variants in 31.2% of the diagnosed PM patients (n = 16), but dominant de novo variants in all of the diagnosed SM patients (n = 8). In PM, we observed mainly LGD disease alleles (~80%), while in SM, LGD and missense disease alleles were equally detected (Fig. 1d and Table 2). The affected genes in our PM subgroup belong to a variety of pathways including centrosome-associated pathways, regulation of mitotic division, transcriptional regulation, mitochondria-related function, NF-kappa-B signaling, endosome regulation, and DNA repair, whereas the affected genes in the SM subgroup encode proteins playing roles in transcriptional regulation, cell growth and differentiation, protein ubiquitination, mitotic progression, and DNA repair (Table 2).

Of the P/LP SVs, seven (25%) were recurrent variants previously reported, and 21 (75%) were novel. Among the novel variants, we found a de novo noncanonical splice-site variant c.665-4del in DYRK1A (NM_001396.3, NG_009366.1), which was not predicted to have a splice effect, but was demonstrated by us to cause an aberrant splicing at messenger RNA (mRNA) level ( exon 6 deletion, r.665_951del, p. [Ile222Aspfs*22]) (Figure S3). We also found in an aborted fetus (ID74812, Fig. 2e, f) a pathogenic nonsense PLK4 variant c.1111C>T, p.(Arg371*) in trans with a likely pathogenic serine substitution c.881T>G, p.(Ile294Ser). The latter variant, which was absent in an unaffected sibling, is the first to be located in the phosphodegron element of PLK4 and predicted to create an additional phosphorylation site likely leading to a reduced protein level via accelerated autodestruction (Table 2 and S2, Figure S4). Phenotypically, this patient presented with previously unreported organ anomalies found in autopsy (ID74812, Table 2 and S2). In an unrelated child (ID77804) with different PLK4 causative variants, we found a novel MRI finding of a large cerebellum and brain stem relative to the supratentorial region (Table 2 and S2). Other novel clinical findings in our study include uvula bifida in a patient (ID53792) with TRMT10A-related microcephaly, short stature, and impaired glucose metabolism 1 (MIM 616033), and a forgotten concept of smaller pituitary glands in Rett syndrome patients by our similar observation of Noonan syndrome— and a history of perinatal asphyxia, which may contribute as an environmental factor to her microcephaly as an unusual presentation of Noonan syndrome (Table S1).

Importantly, in 1 of the 62 patients we identified a likely pathogenic variant m.9185T>C, p.(Leu220Pro) in the mitochondrial gene MT-ATP6 (NC_012920.1) from ES data (ID32410, Table 2 and S1, Fig. 2a, b). This variant was observed in 59% of the reads (Figure S5A), which was also detected by a targeted panel of mitochondrial disease genes in 82% of urothelial cells (data not shown). In a phenotypically similar patient (ID76870, Table 2 and S1, Fig. 2c, d), we found a homozygous deleterious missense variant c.1772A>T, p. (Asn591Ile) in a nuclear gene KARS (NM_001130089.1) which encodes a mitochondria-related protein. Our structural modeling revealed that the isoleucine substitution likely affects the protein structure and/or stability (Figure S5B). We also identified a likely pathogenic variant in another nuclear gene DHTK1D1 that encodes a mitochondrial protein (Table 2 and S1). Altogether, we identified three patients with likely pathogenic variants in mitochondria-related genes, accounting for 4.8% of the total cohort.

**High-level candidate genes**

We identified likely deleterious variants affecting five different high-level candidate genes in five (8.1%) patients without P/LP variants or VUS in established disease genes (Table 2). Four of them (SPAG5, TELD1, VPS26A, DDX1) were affected by biallelic variants, and one (ZNFR3) by a de novo variant. SPAG5 (sperm-associated antigen 5) encodes a mitochondrial protein and has been shown to be required for regulation of mitotic spindles and recruitment of the known microcephaly gene CDK5RAP2 to the centrosome during mitosis. In a patient (ID81652) with PM, mild speech delay, and short stature, we found an unreported de novo frameshift variant c.1223_1224insAC, p.(Lys409Profs*19) in SPAG5 (NM_006461.3) and, by a second allele search, a maternally inherited synonymous variant c.3189G>G, p. (Gly1063Gly) with extremely low MAF (Fig. 3a and Table S1). Sequencing of mRNA from the patient fibroblast showed a deletion of 11 exonic bp resulting in a predicted premature stop codon (c.3189_3198del, p.[Gly1064Glu*3]) (Fig. 3b). Cycloheximide (CHX) rescue treatment showed that both aberrant alleles were subjected to nonsense-mediated mRNA decay (NMD) with some leakiness of the splicing effect (Fig. 3b). Consistently, qRT-PCR (~75 ± 22%) and immunoblotting (~80 ± 26%) revealed a significantly reduced amount of the wild-type SPAG5 at both mRNA and protein levels.
We also observed a reduced SPAG5 intensity mainly in the centrosomal regions where it normally appears more condensed during prophase to telophase (Fig. 3e). However, morphology of the patient’s fibroblasts during different cell cycle phases appeared with no obvious abnormality in the majority of cells (>95%) (Fig. 3e), with apparently unaffected localization of the SPAG5 interacting partner CDK5RAP2 (Figure S6). Nonetheless, since we observed higher mRNA expression levels of SPAG5 in normal human induced pluripotent stem cell–derived neural progenitor cells (NPCs) compared with fibroblasts and other cell types (Fig. 3f), SPAG5 reduction may only pose deleterious effects on highly proliferative NPCs during embryonic development, which could lead to the clinical manifestations in the patient.

TEDC1 (tubulin epsilon and delta complex 1), previously known as CI4ORF80, has been shown to be required for centriole stability. In a patient (ID68629, Fig. 2g, h and Table 2) with PM, primordial dwarfism, and moderate global DD, we identified a noncanonical splice variant c.227-5C>G (intron 2) in trans with a frameshift variant c.1111del, p. (Ala371Glnfs*12) (last exon) in TEDC1 (NM_001134875.1) (Table 2 and Fig. 3g). Sequencing of mRNA from the patient’s fibroblasts showed a deletion of the first 41 bp of exon 3.
Fig. 3 Functional evaluations of high-level candidate variants in SPAG5 and TEDC1. (a) Determination of the allelic location of the de novo frameshift SPAG5 variant c.1223_1224insAC. A portion of SPAG5 sequence containing the frameshift variant and a nearby single-nucleotide polymorphism (SNP, rs113667723) was analyzed by Sanger sequencing of the patient’s blood DNA, which confirmed that the frameshift SPAG5 variant was located in the paternal allele by a distinct frameshift pattern of three bases around the SNP position. Blue sequence, paternal; pink sequence, maternal; black and underlined, variants. (b) Sanger sequencing of messenger RNA (mRNA) from the patient’s fibroblast (ID81652) showed a reduced amount of an aberrantly spliced transcript (due to the synonymous SPAG5 variant c.3189C>T with splice effect), which lacks the last 11 bp of exon 20, resulting in an out-of-frame mutation and a premature stop codon p.(Gly1064Glufs*11). In the magnified electropherogram of CHX, asterisk indicates rescued frameshift allele (nucleotide C in blue), leaked splice-site variant allele (nucleotide T in red), and rescued aberrantly spliced allele (nucleotide G in black). This means that the frameshift allele and the aberrantly spliced allele were rescued upon CHX treatment. CHX cycloheximide, DMSO dimethyl sulfoxide, WT wild type. (c) Quantitative reverse transcription polymerase chain reaction (qRT-PCR) showed significantly reduced SPAG5 mRNA levels (~75%) in the patient’s fibroblasts (untreated and vehicle DMSO, p<0.05, Welch t test), which were rescued upon treatment with CHX. Experiment was done in a triplicate. (d) Immunoblotting against the C-terminal terminal of SPAG5, detecting the two SPAG5 isoforms (full-length and short) and β-actin on protein extracts showed a significant reduction (~80%) of SPAG5 protein in the patient’s fibroblasts (ID81652) (p<0.05, Welch t test). Note that the short isoform lacks a small portion of N-terminal of which the function has not yet been characterized. Experiment was done in a triplicate. (e) Immunostaining against SPAG5, PCNT, and α-Tubulin shows a reduced SPAG5 intensity mainly in the centrosomal regions where it is more condensed in the control during prophase to telophase. However, morphology of the patient’s fibroblasts appears with no obvious abnormality in the majority of cells (>95%). The nuclei were visualized by DAPI staining (in blue). The scale bar represents 10 μm. (f) RT-PCR showed higher expression levels of SPAG5 in normal human induced pluripotent stem cell–derived neural progenitor cells (NPCs) compared with fibroblasts and other cell types including testis (positive control), heart (negative control), HeLa cell line (highly proliferative control), and NPC-derived neuronal culture at 3 (NC3wks) or 5 (NC5wks) weeks. (g) Sanger sequencing of mRNA from the patient’s fibroblast (ID68629) showed a reduced amount of an aberrantly spliced transcript (due to the noncanonical splice-site TEDC1 variant c.227-5C>G that increases the activity of the cryptic splice acceptor), which lacks the first 40 bp of exon 3, resulting in an out-of-frame mutation and a premature stop codon p.(Glu76Glyfs*11). The levels of the aberrant transcript were rescued upon CHX treatment, indicating that the aberrant transcript was subjected to nonsense-mediated decay (NMD) (see also Figure S7). On the other hand, the sequencing of the other TEDC1 variant c.1111del, which is located in the last exon, did not show a reduced amount of the aberrant transcript. Nevertheless, this variant leads to a frameshift and premature stop codon p. (Ala371Glnfs*12) that removes the last 50 amino acids, likely leading to a deleterious effect on the function of the TEDC1 protein, which remains to be characterized. Bar graphs show the mean ± SEM.

(r.227_267del) predicted to result in a truncated protein (p. [Glu76Glyfs*11]) and CHX rescue treatment confirmed NMD of the aberrantly spliced transcript (Fig. 3g and S7). The other variant was not affected by NMD, but likely results in a C-terminally truncated protein (Fig. 3g and S7).

Our other high-level candidate variants, which were identified in three patients with PM and mild to severe DD, affected ZNRF3 (patient 60361, Fig. 2i, j), a negative regulator of the Wnt signaling; VPS26A, a mediator of Wnt transport; and DDX1 (patient 74091, Fig. 2k, l), a DEAD box RNA helicase, respectively (Table 2). Structural modeling for these missense variants predicts a variety of adverse consequences, including loss of binding affinity to the interacting protein R-spondin for ZNRF3, loss of the ability to form a water-mediated interaction to neighboring residues for VPS26A, and steric clashes with adjacent residues for DDX1, all likely affecting the protein domain stability and therefore probably contributing to the patients’ clinical presentation (Table 2 and Figure S8). Notably, via GeneMatcher, we found an additional patient with unreported biallelic variants (c.133-8T>C, p.[?]; c.839C>T, p. [Thr280Arg]) affecting DDX1. The effects of the splice-site variant remain unknown because of no access to any other sample from this patient. However, our predictions based on the UniProt and PhosphoSitePlus databases suggest that the Thr280Arg change may cause the loss of a phosphorylation site and also interfere with posttranslational modifications of the adjacent residue Lys281, which likely affects the regulation of DDX1 interaction and/or degradation. Moreover, both patients with the recessive DDX1 variants presented with comparable neurological features including severe global DD, spastic quadriaparesis, abnormal sleeping pattern, and abnormal movements/seizures, providing additional support for their pathogenicity. Nonetheless, severe microcephaly was only present in the first patient (ID74091), probably due to the contribution of possible other recessive variants in his multiple large runs of homozygosity (Table S1).

Candidate and suspected candidate genes
We found a total of 22 candidate and 26 suspected candidate genes in our cohort. Of these genes, 9 candidate (RNF113A, CEP350, SIK2, RFX7, C2CD5, KIF23, IRS2, UNC13A, PRTG) and 5 suspected candidate (NMI, LARP4B, SEC14L5, PHB2, RAB40AL) genes were identified in 11 (17.7%) patients without P/LP, VUS, or high-level candidate variants.

DISCUSSION
We have elucidated the phenotypic spectrum and genetic landscape including novel findings in PM and SM by detailed clinical assessment and combined CMA and ES of 62 unselected microcephalic patients.

In our cohort, we confirm previous findings of commonly microcephaly–associated features including DD/ID, abnormal cerebral MRI, seizures, and short stature, but in addition also frequently found movement disorders and behavioral problems. With reference to our total cohort, we corroborate previous studies showing no correlation between the degree of microcephaly and developmental performance, however, when stratifying patients for PM and SM, we unexpectedly show here such a correlation among patients with PM. This implies that prenatal onset of OFC deceleration may pose
stronger adverse effect on the developmental outcome. Nevertheless, our evidence of the correlation between abnormal cerebral MRI and the severity of DD/ID substantiates a previous observation of abnormal brain scans as a better reflection of developmental performance in microcephalic patients.\textsuperscript{20} Interestingly, Shaheen et al.\textsuperscript{7} observed two patterns of head growth in congenital microcephaly with severe and progressive microcephaly (pattern A) in the majority of their patients, and largely stable microcephaly (pattern B) in some patients. However, we observed pattern A, only, in PM, which might be explained by different sets of genes identified or different time points of OFC measurement. This may implicate postnatal functions of the affected genes other than only prenatal roles in proliferation of neural progenitor cells.

Etiologically, we identified P/LP variants in almost half of the cohort (~48%), accounting for a diagnostic yield that is within the higher range achieved by NGS studies on NDDs,\textsuperscript{9–14} but is more than three times that of the previous study evaluating 680 microcephalic children (15%) using non-NGS methods,\textsuperscript{2} further supporting the effectiveness of ES for routine diagnostic testing. In addition, we have identified VUS and candidate variants in ~31% of the patients. Therefore, our diagnostic yield will likely increase over time as further supporting evidence for the affected genes becomes available.

We also highlighted the importance of evaluating relevant noncanonical splice-site variants through our examples of a synonymous exonic variant in \textit{SPAG5}, and a ~4 intronic variant in \textit{DYRK1A}, both of which caused aberrant splicing and subsequent NMD. Therefore, it is crucial to investigate such variants, and to validate those with benign in silico predictions that might be false negative due to the complexity of splicing control.

Previously, inborn errors of metabolism including mitochondrialopathies have been identified in 3% of microcephalic patients.\textsuperscript{2} However, the specific percentage of molecularly diagnosed mitochondrial disorders in microcephalic patients has not been reported so far. Our identification of LP variants in mitochondrial and mitochondria-related nuclear genes in ~5% of the patients highlights the significance of mitochondrial disorders even in PM where mitochondrialopathies may have been underdiagnosed. Notwithstanding, due to the mitochondrial heteroplasmy and highly variable coverage of mitochondrial genes in ES data (Fig. 1a),\textsuperscript{30} a targeted assessment of the mitochondrial DNA should be considered.

Despite the comparable diagnostic yields between PM (~44%) and SM (~47%) in our cohort, we illustrate different predominant modes of inheritance and types of causative variants between them. Our observation of predominantly recessive inheritance and biallelic LGD variants in PM patients suggests that complete protein absence may represent the most common cause of PM, which is in line with the findings in MCPH genes.\textsuperscript{5} On the other hand, dominant de novo LGD or assumed loss-of-function (LoF) missense variants, which we frequently observed in SM patients, suggest haploinsufficiency as a frequent pathomechanism in SM. This difference in inheritance pattern is not explained by a consanguinity bias in diagnosed PM patients, since only 1 of 16 diagnosed PM patients is an offspring of consanguineous parents. Consistent with previous studies,\textsuperscript{3,5} disease-causing genes identified in our cohort also encode proteins of various pathways, among which transcriptional regulation and DNA damage response are the most frequent in both PM and SM. However, centrosome-associated pathways are exclusively implicated in PM with autosomal recessive inheritance, which highlights their crucial function in cell division during neurogenesis.\textsuperscript{4,5} Notably, we observed the progressiveness of microcephaly not only in SM patients, but also in all our PM patients, which implicates postnatal defects in neural maintenance and synaptogenesis in both microcephaly subgroups.

Within the undiagnosed patients, we were able to identify five high-level candidate genes, all in patients with PM. Of these five genes, two (\textit{SPAG5} and \textit{TEDC1}) encode centrosomal proteins, two (\textit{ZNRF3} and \textit{VPS26A}) Wnt signaling-related proteins,\textsuperscript{24,25} and one (\textit{DDX1}) an RNA trafficking protein.\textsuperscript{3} In addition to the known centrosomal functions in regulating neuronal progenitor proliferation,\textsuperscript{26,27} Wnt signaling has been shown to be essential for transition between symmetrical and nonsymmetrical cell division in human neural stem cells\textsuperscript{28} and RNA trafficking to be involved in mRNA translation control of proteins that regulate the balance between maintenance and differentiation of radial glial progenitors and thereby development of the embryonic cortex.\textsuperscript{31} Therefore, compromise in the function of these proteins may in fact lead to defects in neurogenesis and hence primary microcephaly. However, we suggest considering all our candidate genes for NDD in general, due to the variable presentation of microcephaly in non-MCPH patients\textsuperscript{34,35}. This variability has been recently demonstrated for \textit{FBXO11}-related NDD, in which fewer than 25% of the patients presented with microcephaly.\textsuperscript{35}

Clinical variability is often observed in NDDs, even in those with established causative genes, which has been, in some instances, attributed to additional genetic factors.\textsuperscript{36} In our cohort, we were able to identify additional genetic hits or a perinatal event likely contributing to the severity of ID or the presence of microcephaly in three patients. However, individualized explanation for all variable NDD presentations will require a comprehensive understanding of an individual’s genetic as well as epigenetic status.

In conclusion, we showed that microcephaly is highly heterogeneous both phenotypically and genetically. By using a combined high-resolution CNV and ES analyses, we achieved an effective diagnostic yield of ~48% and in addition proposed five novel NDD/microcephaly candidate genes with supporting evidence. We also shed some light on distinct as well as common characteristics of the two microcephaly subclasses PM and SM, which helps with better management of the patients and understanding of the underlying pathways involved in human brain development.
SUPPLEMENTARY INFORMATION
The online version of this article (https://doi.org/10.1038/s41436-019-0464-7) contains supplementary material, which is available to authorized users.

ACKNOWLEDGEMENTS
We sincerely thank the affected individuals and their families for participation and their permission to publish the results. This research was supported by ERA-NET grant “Euromicro” (SNF 31ER30_154238 to AR) and radiz–Rare Disease Initiative Zurich, clinical research priority program, University of Zurich. An additional patient with DDX1 biallelic variants reported in this manuscript was found via collaboration with the Undiagnosed Diseases Network (UDN) supported by the National Institutes of Health (NIH) Common Fund, through the Office of Strategic Coordination/Office of the NIH Director under award number U01HG007690. The content of this manuscript is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health.

DISCLOSURE
The authors declare no conflicts of interest.

Publisher's note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

REFERENCES
1. von der Hagen M. Diagnostic approach to primary microcephaly. Neuropediatrics. 2017;48:133–134.
2. von der Hagen M, Pivarcsi M, Liebe J, et al. Diagnostic approach to microcephaly in childhood: a two-center study and review of the literature. Dev Med Child Neurol. 2014;56:732–741.
3. Woods CG, Parker A. Investigating microcephaly. Arch Dis Child. 2013;98:707–713.
4. Alcantara D, O’Driscoll M. Congenital microcephaly. Am J Med Genet C Semin Med Genet. 2014;166C:124–139.
5. Seltzer LE, Paciorkowski AR. Genetic disorders associated with postnatal microcephaly. Am J Med Genet C Semin Med Genet. 2014;166C:140–155.
6. Jayaraman D, Bae BI, Walsh CA. The genetics of primary microcephaly. Annu Rev Genomics Hum Genet. 2018;19:177–200.
7. Shaheen R, Maddirevula S, Ewida N, et al. Genomic and phenotypic delineation of congenital microcephaly. Genet Med. 2018 Sep 14; https://www.nature.com/articles/s41436-018-0140-3.
8. Dahlgren L, Wilson RD. Prenatally diagnosed microcephaly: a review of etiologies. Fetal Diagn Ther. 2001;16:323–326.
9. Hamdan FF, Srour M, Capo-Chichi JM, et al. De novo mutations in moderate or severe intellectual disability. PLoS Genet. 2014;10: e1004772.
10. Najmabadi H, Hu H, Garshabili M, et al. Deep sequencing reveals 50 novel genes for recessive cognitive disorders. Nature. 2011;478:57–63.
11. Need AC, Shashi V, Hitomi Y, et al. Clinical application of exome sequencing in undiagnosed genetic conditions. J Med Genet. 2012;49:353–361.
12. Gilissen C, Hehir-Kwa JY, Thung DT, et al. Genome sequencing identifies major causes of severe intellectual disability. Nature. 2014;511:344–347.
13. Alazami AM, Patel N, Shamseldin HE, et al. Accelerating novel candidate gene discovery in neurogenetic disorders via whole-exome sequencing of prescreened multiplex consanguineous families. Cell Rep. 2015;10:148–161.
14. Thevenon J, Duffourd Y, Masurel-Paulet A, et al. Diagnostic odyssey in severe neurodevelopmental disorders: toward clinical whole-exome sequencing as a first-line diagnostic test. Clin Genet. 2016;89:700–707.
15. Rump P, Jazayeri O, van Dijk-Bos KK, et al. Whole-exome sequencing is a powerful approach for establishing the etiological diagnosis in patients with intellectual disability and microcephaly. BMC Med Genomics. 2016;9:7.
16. Asadollahi R, Oneda B, Joset P, et al. The clinical significance of small copy number variants in neurodevelopmental disorders. J Med Genet. 2014;51:677–688.
17. Plecko B, Zweier M, Begemann A, et al. Confirmation of mutations in PROSC as a novel cause of vitamin B 6-dependent epilepsy. J Med Genet. 2017;54:809–814.
18. Miller DT, Adam MP, Aradhya S, et al. Consensus statement: chromosomal microarray is a first-tier clinical diagnostic test for individuals with developmental disabilities or congenital anomalies. Am J Hum Genet. 2010;86:749–764.
19. Richards S, Aziz N, Bale S, et al. Standards and guidelines for the interpretation of sequence variants: a joint consensus recommendation of the American College of Medical Genetics and Genomics and the Association for Molecular Pathology. Genet Med. 2015;17:405–424.
20. Sartori da Silva MA, Tee JM, Paridaen J, et al. Essential role for the d-Asb11 cuIS Box domain for proper notch signaling and neural cell fate decisions in vivo. PLoS One. 2010;5:e14023.
21. Murakami JW, Coucheshe E, Haas RH, Press GA, Yeung-Coucheshe R. Cerebellar and cerebral abnormalities in Rett syndrome: a quantitative MR analysis. AJR Am J Roentgenol. 1992;159:177–183.
22. Kodani A, Yu TW, Johnson JR, et al. Centriolar satellites assemble centrosomal microcephaly proteins to recruit CDK2 and promote centriole duplication. eLife. 2015;4:e07519.
23. Breslow DK, Hoengendoor S, Kopp AR, et al. A CRISPR-based screen for Hedgehog signaling provides insights into ciliary function and ciliopathies. Nat Genet. 2018;50:460–471.
24. Hao HK, Xie Y, Zhang Y, et al. ZNRF3 promotes Wnt receptor turnover in an R-spondin-sensitive manner. Nature. 2012;485:195–200.
25. Yang PT, Lawhorne CJ, Stilhankova M, Coudreuil DB, Betist MC, Koroswagen HC. Wnt signaling requires retromer-dependent recycling of MIG-14/Wntless in Wnt-producing cells. Dev Cell. 2008;14:140–147.
26. Sobreira N, Schiettecatte F, Valle D, Hamosh A. GeneMatcher: a matching tool for connecting investigators with an interest in the same gene. Hum Mutat. 2015;36:928–930.
27. Bolduc FC, Shevell MI. Corrected head circumference centiles as a possible predictor of developmental performance in high-risk neonatal intensive care unit survivors. Dev Med Child Neurol. 2005;47:766–770.
28. Baxter PS, Rigby AS, Rotaert MH, Wright I. Acquired microcephaly: causes, patterns, motor and IQ effects, and associated growth changes. Pediatrics. 2009;124:590–595.
29. Custer DA, Vezina LC, Vaught DR, et al. Neurodevelopmental and neuroimaging correlates in nonsyndromic microcephalic children. J Dev Behav Pediatr. 2000;21:12–18.
30. Picardi E, Pesole G. Mitochondrial genomes gleaned from human whole-exome sequencing. Nat Methods. 2012;9:523–524.
31. Verssey JP, Amadei G, Burns SE, Kiebler MA, Kaplan DR, Miller FD. An asymmetrically localized Staufen2-dependent RNA complex regulates maintenance of mammalian neural stem cells. Cell Stem Cell. 2012;11:517–528.
32. Hu WF, Chahour MH, Walsh CA. The diverse genetic landscape of neurodevelopmental disorders. Annu Rev Genomics Hum Genet. 2014;15:195–213.
33. Bengoa-Vergnory N, Gororno-Etxebarria I, Gonzalez-Salazar I, Kypa RM. A switch from canonical to noncanonical Wnt signaling mediates early differentiation of human neural stem cells. Stem Cells. 2014;32:3196–3208.
34. Laugel V, Dalloz C, Durand M, et al. Mutation update for the CSB/ERCC6 and CSA/ERCC8 genes involved in Cockayne syndrome. Hum Mutat. 2010;31:113–126.
35. Gregor A, Sadler LG, Asadollahi R, et al. De novo variants in the F-box protein FBXO11 in 20 individuals with a variable neurodevelopmental disorder. Am J Hum Genet. 2018;103:305–316.
36. Pizzo L, Jensen M, Polyak A, et al. Rare variants in the genetic background modulate cognitive and developmental phenotypes in individuals carrying disease-associated variants. Genet Med. 2018
37. American Psychiatric Association. Diagnostic and statistical manual of mental disorders, fifth edition (DSM-5). American Psychiatric Association Publishing, Arlington, VA. 2013.

38. Riley KN, Catalano LM, Bernat JA, et al. Recurrent deletions and duplications of chromosome 2q11.2 and 2q13 are associated with variable outcomes. Am J Med Genet A. 2015;167A:2664–2673.

39. Chatron N, Haddad V, Andrieux J, et al. Refinement of genotype-phenotype correlation in 18 patients carrying a 1q24q25 deletion. Am J Med Genet A. 2015;167A:1008–1017.

40. Bernardini L, Palka C, Ceccarini C, et al. Complex rearrangement of chromosomes 7q21.13-q22.1 confirms the ectrodactyly-deafness locus and suggests new candidate genes. Am J Med Genet A. 2008;146A:238–244.