



# The genomic and clinical landscape of fetal akinesia

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**Purpose:** Fetal akinesia has multiple clinical subtypes with over 160 gene associations, but the genetic etiology is not yet completely understood.

**Methods:** In this study, 51 patients from 47 unrelated families were analyzed using next-generation sequencing (NGS) techniques aiming to decipher the genomic landscape of fetal akinesia (FA).

**Results:** We have identified likely pathogenic gene variants in 37 cases and report 41 novel variants. Additionally, we report putative pathogenic variants in eight cases including nine novel variants. Our work identified 14 novel disease–gene associations for fetal akinesia: *ADSSL1*, *ASAH1*, *ASPM*, *ATP2B3*, *EARS2*, *FBLN1*, *PRG4*, *PRICKLE1*, *ROR2*, *SETBP1*, *SCN5A*, *SCN8A*, and *ZEB2*. Furthermore, a sibling pair harbored a homozygous copy-number variant in *TNNT1*, an ultrarare congenital myopathy gene that has been linked to arthrogryposis via Gene Ontology analysis.

**Conclusion:** Our analysis indicates that genetic defects leading to primary skeletal muscle diseases might have been underdiagnosed, especially pathogenic variants in *RYR1*. We discuss three novel putative fetal akinesia genes: *GCN1*, *IQSEC3* and *RYR3*. Of those, *IQSEC3*, and *RYR3* had been proposed as neuromuscular disease–associated genes recently, and our findings endorse them as FA candidate genes. By combining NGS with deep clinical phenotyping, we achieved a 73% success rate of solved cases.

*Genetics in Medicine* (2020) 22:511–523; <https://doi.org/10.1038/s41436-019-0680-1>

**Keywords:** fetal akinesia; arthrogryposis; myopathy; exome; copy-number variation

## INTRODUCTION

Fetal akinesia (FA) describes a clinical syndromic entity characterized by reduced or absent fetal movements leading to multiple phenotypic abnormalities. These abnormalities may include intrauterine growth restriction (IUGR), cranio-facial dysmorphic features, limb pterygia, pulmonary hypoplasia, and arthrogryposis. These multiple contractures are commonly known as arthrogryposis multiplex congenita (AMC) or, when associated with pterygia, multiple pterygium

syndrome. It was initially described as the Pena–Shokeir phenotype, and is further subdivided into 20 clinical subtypes that are currently linked to at least 37 different genetically defined entities.<sup>1,2</sup> “Arthrogryposis” has been commonly used as a descriptive term for congenital contractures of the joints in at least two different body parts, in contrast to the most common isolated congenital contracture, the congenital clubfoot or pes equinovarus.<sup>3</sup> And arthrogryposis may occur isolated or as part of a broad spectrum of syndromic

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Submitted 29 June 2019; accepted: 1 October 2019  
Published online: 4 November 2019

conditions. The phenotypical spectrum includes both the clinical entities of AMC and the fetal akinesia deformation sequence (FADS). Therefore, we propose to use fetal akinesia (FA) as an overarching term covering the entire phenotypical spectrum from a mild AMC phenotype with a reasonable quality of life for the patient to a severe FADS phenotype with a prenatally lethal outcome.

Even though there are 166 genes linked to an arthrogryptic or fetal akinetic phenotype (Supplementary Table S4), currently many patients with FA have no causal genetic diagnosis. Therefore counseling of affected patients and their families with regard to prognoses and recurrence risks remains challenging.<sup>2</sup> Furthermore, this lack of knowledge is a major obstacle for the development of molecular therapies for these patients. There is an unmet clinical need to decipher the genetic basis and defective pathways in AMC and FADS.

Here, we present novel genomic insights gained from implementing next-generation sequencing (NGS) analysis in a cohort of patients presenting with FA, with a phenotypic severity ranging from mild musculoskeletal defects to prenatally lethal phenotypes.

## MATERIALS AND METHODS

### Inclusion criteria

Approval for the research performed in this publication has been granted by the ethics board of the University of Cologne (sign 19–1269). Written consent for publication from all included patients and/or legal guardians has been acquired and archived. The patients included in this cohort were recruited primarily with regard to the presence of multiple joint contractures manifested at or even before birth as the key diagnostic criterion for the fetal akinesia spectrum. Patients were referred to the authors when no causative pathogenic variant had been determined by candidate gene sequencing, array comparative genomic hybridization (array CGH), or karyogram. Exclusion criteria were a nongenetic etiology such as maternal antibodies or a prenatal infection. Prenatal fetal abnormalities, dysmorphic facial features, neurological defects, cognitive disabilities, limb deformations, and dysfunction involving other organs besides the musculoskeletal system were clinically phenotyped in depth if possible. Wherever accessible, the clinical findings from follow-up examinations were included in the clinical description of the patient. Clinical findings were recorded using a clinical core data form for the relevant clinical data supplied by the primary specialized health-care provider of the patient or by the fetal pathologist and the parents in cases of prenatally lethal FA. Four of the novel variants (cases 2, 11, 12, 14; see Tables 1 and 2) were previously published by the authors.<sup>4–6</sup>

### Next-generation sequencing

DNA used for NGS was extracted using a standard extraction protocol (details in Supplementary Materials file 1). The Mendeliome panel was run initially with Illumina TruSight One covering 4813 genes and later the expanded TruSight

One base panel with 6709 genes used by the authors in cooperation with Illumina during the development of TSOOne V2.0.<sup>7,8</sup> Exome sequencing (ES) was performed initially using the NimbleGen SeqCap EZ Human Genome Library v2.0 and later the Agilent SureSelect V6 panel, which was chosen due to better overall performance and target coverage.<sup>9</sup> The average coverage of all currently known FA genes has been analyzed for the different NGS kits used in this study (Supplementary Figs. S9–13). If single ES did not lead to the diagnosis, a trio exome (9 trios) was performed provided parental consent was available.

The data analysis of the generated raw NGS data was described previously<sup>7,10</sup> using up-to-date versions of the algorithms and programs implemented within the Varbank pipeline (<https://varbank.ccg.uni-koeln.de/varbank2/>).

The NGS data were filtered with the aid of the Varbank pipeline to successively remove low-quality variant artifacts, deep intronic variants, variants that were predicted to be benign and variants in genes without a matching disease association that was already documented in public control databases. The remaining variants were then manually curated by gene function, known disease association, and severity of the predicted effect on the protein product to generate a list of potential candidates. Detailed information about the NGS data analysis and filtering strategy is provided in the “Methods” section of Supplementary Materials file 1 and Supplementary Fig. 14.

### Dideoxy sequencing validation and segregation analysis

The validation of the variants determined by NGS was performed by dideoxy sequencing and segregation analysis was performed whenever parental DNA was available (Supplementary Materials file 2 for detailed pedigrees).

### Variant classification

The remaining candidate variants were graded using both the American College of Medical Genetics and Genomics (ACMG) classification system<sup>11</sup> as well as the proposed European Society of Human Genetics (ESHG) classification system (<https://www.eshg.org/index.php?id=949>). All variants achieving a pathogenic or likely pathogenic ACMG score were considered to be definitely solved; all ACMG variants of uncertain significance (VUS) achieving an ESHG rating of C–D (mildly pathogenic or susceptibility variant) were considered to be potentially solved with a variant of interest.

### Systems biology analysis

The protein–protein interactions between previously published FA genes (Supplementary Table S4) and our new candidates (Supplementary Tables S1 and S2) have been visualized using Cytoscape StringApp,<sup>12</sup> which uses STRINGdb data.

An over-representation analysis (ORA) has been performed for Gene Ontology (GO) biological process (BP), molecular function (MF), and cellular component (CC) classes via R-clusterProfiler library.<sup>13</sup> We also used the Spearman rank

**Table 1** Clinical overview of the cohort

ID/ gender/ age	Prenatal abnormalities	Joint contractures	Dysmorphic features	Neurological abnormalities	Other features
01/m/1	Polyhydramnios, IUGR	Hips, knees	Macrocephaly	Hydrocephalus externus, combined hearing dysfunction, floppy infant syndrome	Pulmonary hypoplasia, dysphagia, gastroesophageal reflux, cryptorchidism, RI, tracheostomy and ventilation
02/f/0	RFM	Elbows, wrists, fingers, knees, ankles	Microcephaly, hypertelorism, low-set ears, left axillary pterygium	Microcephaly, simplified gyration	Pulmonary hypoplasia
03/m/0	Polyhydramnios	Knees, ankles, fingers	Hypertelorism	Atrophic leg musculature	Neck edema
04/f/0	RFM, hydrops fetalis, IUGR	Elbows, fingers, ankles	Hypertelorism, micrognathia, low-set ears, pterygium at knees, elbows, and axillae	Lissencephaly	—
05/f/0	RFM, hydrops fetalis	Elbows, wrists, knees, ankles	Micrognathia	—	—
06/m/5	IUGR	Wrists, fingers, hips, knees, ankles	—	Dilated ventricles, corpus callosum dysgenesis, epilepsy, mixed type polyneuropathy, muscular hypotonia, no motor development	PDA, scoliosis, RI, tracheostomy
07/f/1	IUGR, polyhydramnios, RFM	Hips, ankles, wrists, fingers	Microstomia, micrognathia	—	RI, NIV, gastroesophageal reflux, scoliosis, bilateral hip dysplasia, cholestasis and cholelithiasis, ASD
08/m/1	Polyhydramnios, hydrops fetalis	Wrists, hips, knees, ankles	Hypertelorism, micrognathia, low-set ears	Muscular hypotonia	Pulmonary hypoplasia, RI, hypertirchosis
09/f/1	—	Hips, knees, ankles, elbows, fingers	Microcephaly	Delayed motor development, muscular hypertonia	Dysphagia, hip joint luxation
10/f/1	—	Wrists, fingers, ankles	—	Delayed motor development, delayed speech development	Diaphragm paralysis requiring ventilation, dysphagia, unilateral enophthalmus, camptodactyly, arachnodactyly
11/f/1	Polyhydramnios, RFM	Elbows, fingers, ankles	Whistling face, pterygium colli, hypertelorism, micrognathia, high arched palate	Severe intellectual disability, no gaze fixation, corpus callosum hypoplasia, muscular hypertonia	Scoliosis, joint luxation, dysphagia, episodic tachypnea, apnea, bronchial paroxysms, temperature dysregulation, growth retardation, camptodactyly, rectus diastasis, severe perspiration
12/f/1	Polyhydramnios, RFM	Elbows, hands, hips, ankles	Microstomia, retrogenia, high arched palate, low-set ears	—	Respiratory insufficiency, NIV
13/m/1	—	Elbows, wrists, fingers, ankles	—	Delayed motor development, distal muscle weakness	Dilated renal pelvis, gastroschisis, bilateral hydronephrosis
14/m/1	—	Fingers, ankles	High arched palate	Sensory axonal neuropathy, defective proprioception, muscular hypotonia	Perinatal respiratory dysfunction, dysphagia, scoliosis, rigid spine, distal laxity of extremities, hip dysplasia
15/m/16	RFM	Fingers, ankles	High arched palate, low-set ears, camptodactyly	—	Scoliosis, respiratory dysfunction, NIV
16/f/0	Polyhydramnios, RFM, hydrops fetalis	Elbows, knees, fingers, toes	Low-set ears	—	Thorax hypoplasia, hydrothorax
17/f/0	Polyhydramnios, RFM, hydrops fetalis	Elbows, knees, fingers, talus verticalis	Low-set ears	—	Thorax hypoplasia
18/m/1	Polyhydramnios, RFM	Elbows, wrists, fingers, hips, knees, ankles, jaw joint, shoulders	High arched palate	—	RI, scoliosis
19/m/0	Hydrops fetalis, IUGR, RFM	Elbows, knees, shoulders	Cleft palate, low-set ears, pterygium colli, micrognathia	Microcephaly	Malrotated heart
20/m/0	Hydrops fetalis	Knees, ankles	Cleft palate	Microcephaly	Bilateral hydrothorax, hydronephrosis
21/f/3	RFM	Complete akinesia	Cleft palate	Intellectual disability	Dysphagia, RI requiring tracheostomy
22/f/0	Hydrops fetalis, IUGR, RFM	Elbows, shoulders, hips, knees, thumbs	High arched palate	—	Pulmonary hypoplasia, hydrothorax
23/f/0	—	Shoulder, elbows, hips, knees, ankles	Macrocephaly, hypertelorism, protrusio bulbi, cleft palate, microretrognathia, pterygia	—	Bilateral camptodactyly, pulmonary hypoplasia, cardiac hypertrophy, streak gonads
24/f/1	—	Knees, ankles	Mild dyscrania, high arched palate	Areflexia, muscular hypotonia	—
25/m/4	—	Hips, knees, elbows, fingers, ankles	—	Ophthalmoplegia, intracranial hemorrhage, delayed motor development, centronuclear myopathy, reduced DTR	RI, reduced spontaneous movement, scoliosis
26/m/3	Polyhydramnios, RFM	Neck, shoulders, elbows, wrists, fingers, hips, knees, ankles, toes	High arched palate	Microcephaly, ptosis	RI, tracheostomy and ventilation, scoliosis, right pulmonary hypoplasia, hypothyreosis, umbilical and inguinal hernias
27/f/5	—	Multiple joint contractures	Micrognathia	Severe intellectual disability	Esophageal herniation with reflux, tracheomalacia
28/f/1	IUGR	Knees, wrists, fingers, ankles	Hypertelorism, micrognathia, low-set ears, high arched palate, anteverted nostrils, upslping palpebral fissures, epicanthus	Cavum septi pellucidi, seizures, subdural hemorrhage, muscular hypotonia, reduced DTR	Irregular breathing activity, camptodactyly, missing interphalangeal creases
29/f/0	—	Shoulders, elbows, hips, knees, ankles	—	Cerebral atrophy, cerebellar hypoplasia, corpus callosum agenesis	—
30/f/0	—	Knees, elbows, shoulders	Hypertelorism, neck edema, protrusio bulbi, micrognathia, bilateral clinodactyly of the fifth digit	Hypoplastic frontal brain with aprosencephaly, neuronal migration defect, missing fissura sylvii, neuroblast layer between cortex and subventricular zone, microcephaly	Two accessory spleens

**Table 1** continued

ID/ gender/ age	Prenatal abnormalities	Joint contractures	Dysmorphic features	Neurological abnormalities	Other features
31/m/0	RFM	Shoulders, elbows, wrists, knees, ankles	Hypertelorism, telecanthus, high arched palate, low-set ears, pterygia at knees, elbows, and axillae, micrognathia, hyperelorism, cleft palate, pterygium coli	Macrocephaly	Syndactyly right hand 3th–5th digit, incomplete syndactyly right foot 2nd–4th digit, pectus excavatum
32/m/0	IUGR	Shoulders, elbows, wrists, hips, knees, ankles	Micrognathia, hyperelorism, low-set ears	Microcephaly, porencephaly, lissencephaly	—
33/m/0	—	Wrists, elbows, knees, ankles	Retrognathia, low-set ears	—	—
34/m/0	—	Elbows, knees, wrists, hips, shoulders, right pes planus, left pes equinovarus	Hypertelorism, epicanthus, micrognathia, high arched palate, low-set ears	—	Pulmonary hypoplasia
35/m/2	Polyhydramnios	Knees, elbows, fingers, ankles	Micrognathia, nuchal pterygium, cubital pterygium, low-set ears, high arched palate, neck hygroma	Corpus callosum hypoplasia, subcortical cerebral atrophy, cerebellar vermis hypoplasia, hydrocephalus internus, muscle atrophy	Cardiomegaly, cardiac insufficiency, atroseptal defect, pulmonary hypoplasia, RI, kyphoscoliosis, thorax hypoplasia, cryptorchidism, arachnodactyly, bilateral nephrocalcinosis
36/m/2	RFM, oligohydramnios	Elbows, knees, ankles	High arched palate	Tetraparesis, hydrocephalus, corpus callosum dysgenesis, cerebral atrophy, intellectual disability, muscular hypotonia, left torticollis	Scoliosis, CO-1 vertebral dysgenesis, cryptorchidism
37/m/2	—	Wrists, ankles	Caput quadratum, low-set ears	Corpus callosum agenesis, cerebrotendinous atrophy, developmental delay, bilateral loss of hearing	PFO, hypospadias, hydronephrosis, hypothyroidism
38/m/0	—	Multiple joint contractures	—	Lissencephaly	—
39/m/0	Polyhydramnios, RFM	Hips, knees, ankles, elbows, wrists, fingers	Micrognathia, trismus	—	RI, scoliosis, rigid spine, deformed chest
40/f/3	Polyhydramnios	Fingers, hips, knees, ankles	Clinodactyly right fifth digit, brachymesophalangia	—	Reduced skeletal calcination, RI, NIV, PFO, kyphoscoliosis, proximal dystonia, dysphagia, gastroesophageal reflux
41/f/3	RFM	Hips, knees	—	Muscular hypotonia, spontaneous myoclonic episodes, fasciculations, reduced DTR	RI, NV
42/f/0	Fetal vascular thrombopathy	Fingers, ankles	Bilateral popliteal pterygia	—	—
43/f/1	Polyhydramnios, RFM	Wrists, fingers	—	Axial muscle weakness, atrophied shoulder musculature, elevated diaphragm	—
44/f/28	RFM	Elbows, hips, knees, wrists, fingers, ankles	Cleft palate, dolichocephaly, high arched palate	Muscular hypotonia, absent DTR	Growth retardation, scoliosis, rigid spine, pectus excavatum
45/m/0	—	Fingers, wrists, ankles	Bilateral elbow pterygia	—	Scoliosis, VSD, intestinal malrotation
46/m/1	RFM	Hips, knees, elbows, wrists, fingers, ankles	—	Reduced DTR, distal muscle weakness, delayed motor development, microcephaly	Bilateral hip dysplasia, hyperlordosis
47/m/1	—	Knees, ankles	—	Hydrocephaly, porencephaly, reduced corpus callosum, ventriculomegaly, atrophic leg musculature	Cryptorchidism, ventricular septal defect, perinatal respiratory dysfunction
48/f/0	Amnion band constrictions, polyhydramnios, RFM	Elbows, wrists, knees, ankles	Hypertelorism, low-set ears	—	—
49/f/9	—	Hip, knees, ankles	Pterygium coli, bilateral clinodactyly	Muscle atrophy, scapula alata, Marcus-Gunn syndrome	Rigid spine, scoliosis, flat thorax, growth retardation
50/m/13	—	Knees, ankles	Cleft palate, ptosis, microstomia, hypomimia	Increased muscle tonus especially in upper limbs	Short stature, bilateral inguinal herniation
51/f/1	IUGR, RFM	Shoulders, elbows, wrists, fingers, hips, ankles	Ptosis, cleft palate, pterygium coli, deformed thorax	Encephalomalacia, periventricular cystic defects, pontine and cerebellar hypoplasia, generalized epileptic seizures, muscular hypotonia	Perinatal respiratory dysfunction, gastroesophageal reflux

Clinical details of the fetal akinesia cohort sorted by patient ID, gender, and age in years at inclusion in this study (age 0 denotes induced abortions, spontaneous miscarriages, stillbirths, and children dying <1 month after birth), prenatal abnormalities, location of joint contractures, dysmorphic features, neurological abnormalities, and other features. Cases 2, 11, 12, and 14 were previously published by the authors.<sup>4–6</sup> ASD atrial septal defect, DTR deep tendon reflexes, IUGR intrauterine growth restriction, NIV noninvasive ventilation, PDA persisting ductus arteriosus, PFO persisting foramen ovale, RFM reduced fetal movements, RI respiratory insufficiency, VSD ventricular septal defect.

**Table 2 List of primary candidate pathogenic variants**

ID	Affected gene	Reference sequence ID	DNA change	Predicted AA change	Consanguinity/zygosity	ACMG	ESHG	Supporting evidence for pathogenicity
01	ACTA1	NM_001100.3	c.[739G>A];[=]	p.(G247R)	De novo heterozygous	Known pathogenic variant	Known pathogenic variant	ClinVar variation ID: 381639; known FA gene PMID: 21984750
02	ASCC1	NM_001198799.3	c.[710+1C>T]; [710+1C>T]	Essential splice site	Yes/ homozygous	Pathogenic	Highly pathogenic (A)	Known FA gene PMID: 26924529
03	CHRD	NM_000751.2	c.[452G>C];[452G>C]	p.(C151S)	Yes/ homozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 18252226
04	CHRG	NM_005199.4	c.[710_711delinsAA]; [(710_711delinsAA)]	p.(I237K)	homozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 27245440
05	CHRG	NM_005199.4	c.[710_711delinsAA]; [(710_711delinsAA)]	p.(I237K)	homozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 27245440
06	CNTNAP1	NM_003632.2	c.[69C>G];[69C>G]	p.(Y23*)	Yes/ homozygous	Pathogenic	Highly pathogenic (A)	Known FA gene PMID: 28374019
07	CNTNAP1	NM_003632.2	c.[1906G>A];[=]	p.(V636M)	De novo heterozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 28374019
08	GBE1	NM_000158.3	c.[1693C>T]; [1693C>T]	p.(R565W)	Yes/ homozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 27546458
09	GLDN	NM_181789.2	c.[1178G>A]; [1428C>A]	p.[(R393K)]; [(F476L)]	Compound heterozygous	VUS/likely pathogenic	Mildly pathogenic (C)/pathogenic (B)	Known FA gene PMID: 27616481, absent from controls; aa highly conserved; functional domain affected; variants cosegregates; CADD 17
10	LGII	NM_139284.2	c.[504G>C]; [1031T>A]	p.[(W168C)]; [(L344Q)]	Compound heterozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 28318499
11	NALCN	NM_052867.2	c.[950T>G];[=]	p.(F317C)	De novo heterozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 27214504
12	NALCN	NM_052867.2	c.[1783G>T];[=]	p.(V595F)	De novo heterozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 27214504
13	NALCN	NM_052867.2	c.[191A>G];[=]	p.(Y64C)	heterozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 27214504
14	PIEZO2	NM_022068.2	c.[1384C>T]; [1384C>T]	p.(R462*)	homozygous	Pathogenic	Highly pathogenic (B)	Known FA gene PMID: 27974811
15	PIEZO2	NM_022068.2	c.[8057G>A];[=]	p.(R2686H)	Heterozygous	Known pathogenic variant	Known pathogenic variant	ClinVar variation ID: 137629; known FA gene PMID: 24726473
16	RAPSN	NM_005055.4	NC_000011.9:g.(? -47459239)_(47460480-?)del	—	homozygous CNV deletion	Pathogenic	Highly pathogenic (A)	Known FA gene PMID: 28495245
17	RAPSN	NM_005055.4	NC_000011.9:g.(? -47459239)_(47460480-?)del	—	homozygous CNV deletion	Pathogenic	Highly pathogenic (A)	Known FA gene PMID: 28495245
18	RAPSN	NM_005055.4	c.[272G>T];[794C>T]	p.[(R91L)]; [(A265V)]	Compound heterozygous	Pathogenic/likely pathogenic	Highly pathogenic (A)/pathogenic (B)	Known FA gene PMID: 28495245
19	RYR1	NM_001042723.1	c.[2167G>A]; [14647 -15_14649del]	p.[(G723R)]; essential splice site	Two heterozygous variants	Likely pathogenic/ pathogenic	Pathogenic (B)/ highly pathogenic (A)	Known FA gene PMID: 26932181
20	RYR1	NM_001042723.1	c.[2167G>A]; [14647 -15_14649del]	p.[(G723R)]; essential splice site	Two heterozygous variants	Likely pathogenic/ pathogenic	Pathogenic (B)/ highly pathogenic (A)	Known FA gene PMID: 26932181
21	RYR1	NM_001042723.1	c.[2500_2501dupCG]; [8024C>A]	p.[(P836Gfs*49)]; [(I2675K)]	Compound heterozygous	Pathogenic/likely pathogenic	Highly pathogenic (A)/pathogenic (B)	Known FA gene PMID: 26932181
22	RYR1	NM_001042723.1	c.[5618delA]; [10018G>A]	p.[(E1873Gfs*57)]; [(V3340M)]	Two heterozygous variants	Pathogenic/likely pathogenic	Highly pathogenic (A)/pathogenic (B)	Known FA gene PMID: 26932181
23	RYR1	NM_001042723.1	c.[1835C>A]; [1835C>A]	p.(A612D)	Yes/ homozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 26932181
24	RYR1	NM_001042723.1	c.[4405C>T]; [13983G>A]	p.[(R1469W)]; essential splice site	Compound heterozygous	Likely pathogenic	Pathogenic (B)	Known FA gene PMID: 26932181
25	RYR1	NM_001042723.1	c.[7298T>C]; [9579C>G]	p.[(L2433P)]; [(C3193W)]	Compound heterozygous	Likely pathogenic/ pathogenic	Pathogenic (B)/ highly pathogenic (A)	Known FA gene PMID: 26932181

**Table 2** continued

ID	Affected gene	Reference sequence ID	DNA change	Predicted AA change	Consanguinity/zygosity	ACMG	ESHG	Supporting evidence for pathogenicity
26	SCN4A	NM_000334.4	c.[2018T>C]i[=]	p.(L673P)	De novo heterozygous	Likely pathogenic VUS	Pathogenic (B)	Known FA gene PMID: 27587986
27	UNC50	NM_014044.5	c.[287C>G]i[=]	p.(T96S)	Yes/ heterozygous	VUS	Mildly pathogenic (C)	Absent from controls; aa highly conserved; CADD 13; known FA gene PMID: 29016857
28	ADSSL1	NM_152328.3	c.[741delC]; [741delC]	p.(K248Rfs*23)	Homozygous	Pathogenic	Highly pathogenic (A)	Distal myopathy associated disease gene PMID: 28268051
29	ASAH1	NM_004315.4	c.[136G>A]i[539G>T]	p.(D46N);(G180V)	Two heterozygous variants	Likely pathogenic	Pathogenic (B)	SMA and Farber disease-associated disease gene PMID: 22703880; fits phenotype
30	ASPM	NM_018136.4	c.[2863C>T]i[3082+1G>C]	p.[(Q955*)]; essential splice site p.(S66L)	Compound heterozygous	Pathogenic/likely pathogenic	Highly pathogenic (A)/mildly pathogenic (C)	Microcephaly associated disease gene PMID: 28674240
31	ATP2B3	NM_001001344.2	c.[197C>T]	p.(S66L)	Hemizygous	Likely pathogenic	Pathogenic (B)	Spinocerebellar ataxia associated disease gene PMID: 25953895
32	EARS2	NM_001083614.1	c.[814G>A]; [1277_1279dupCTC]	p.[(A272T)]; [(T426_R427insP)]	Two heterozygous variants	Likely pathogenic	Pathogenic (B)	Leukoencephalopathy-associated disease gene PMID: 27206875; fits phenotype
33	FBLN1	NM_006486.2	c.[1991G>A]i[=]	p.(R664Q)	Heterozygous	VUS	Susceptibility variant/VOI (D)	Synpolydactyly associated disease gene PMID: 11836357; AF <0.001; aa highly conserved; CADD 28
34	PRG4	NM_001127708.1	c.[3446G>A]; [3446G>A]	p.(G1149D)	Homozygous	VUS	Susceptibility variant/VOI (D)	CACP-associated disease gene PMID: 23290693; AF <0.01; aa conserved; functional domain affected; CADD 23
35	PRICKLE1	NM_153026.2	c.[1682A>T]i[=]	p.(Y561F)	Heterozygous	VUS	Susceptibility variant/VOI (D)	Corpus callosum agenesis and polymicrogyria-associated disease gene PMID: 26272662; fits phenotype; absent from controls; aa conserved; animal model fits; CADD 13
36	ROR2	NM_004560.3	c.[808A>G]; [1675G>A]	p.[(I270V)]; [(G559S)]	Compound heterozygous	Likely pathogenic	Pathogenic (B)	Robinow syndrome-associated disease gene PMID: 26284319; fits phenotype
37	SETBP1	NM_015559.2	c.[2612T>C]i[=]	p.(R871T)	Yes/de novo heterozygous	Known pathogenic variant	Known pathogenic variant	ClinVar variation ID: 1031; Schinzel-Giedion syndrome-associated disease gene PMID: 20436468
38	SCN5A	NM_001099404.1	c.[5213C>T]i[=]	p.(S1738F)	Heterozygous	Likely pathogenic	Pathogenic (B)	Brugada syndrome-associated disease gene PMID: 28341781
39	SCN8A	NM_014191.3	c.[719T>C]i[=]	p.(I240T)	De novo heterozygous	Likely pathogenic	Pathogenic (B)	Epileptic encephalopathy-associated disease gene PMID: 28676440
40	TNNT1	NM_003283.5	NC_000019.9; g.(?_55652166)_(55660604?)del	—	Homozygous CNV deletion	Pathogenic	Highly pathogenic (A)	Nemaline myopathy-associated disease gene PMID: 15665378
41	TNNT1	NM_003283.5	NC_000019.9; g.(?_55652166)_(55660604?)del	—	Homozygous CNV deletion	Pathogenic	Highly pathogenic (A)	Nemaline myopathy-associated disease gene PMID: 15665378
42	ZEB2	NM_014795.3	c.[444T>G]i[=]	p.(F148L)	Heterozygous	VUS	Susceptibility variant/VOI (D)	Mowat-Wilson syndrome-associated disease gene PMID: 24715670; absent from controls; aa highly conserved; animal model fits; CADD 15
43	GCM1	NM_006836.1	c.[3581C>A]i[=]	p.(A1194E)	De novo heterozygous	VUS	Susceptibility variant/VOI (D)	Absent from controls; aa conserved; CADD 22
44	IQSEC3	NM_001170738.1	c.[1058T>C]; [3546G>A]	p.[(L353P)]; cryptic splice site p.(D4605G)	Compound heterozygous	VUS	Susceptibility variant/VOI (D)	First variant absent from controls; second variant AF <0.001; aa highly conserved; cryptic splice site activation; CADD 26; 14
45	RYR3	NM_001036.3	c.[13814A>G]i[=]	p.(D4605G)	Heterozygous	VUS	Susceptibility variant/VOI (D)	AF <0.0001; aa highly conserved; animal model fits; CADD 32

List of candidate variants sorted by patient ID, affected gene, RefSeq ID, DNA change, AA change, consanguinity, and zygosity. Additionally, ACMG and ESHG scores as well as supporting evidence of pathogenicity for VUS are given. If for compound heterozygous variants the ACMG and ESHG scores are identical, then the score is listed only once. Those variants that have already been independently published as pathogenic are listed with the corresponding ClinVar ID. Novel disease genes are marked with a blue background. Please see Supplementary Table 1 for additional details. Terms used in evidence of pathogenicity are as follows: conserved—conserved in mammals; highly conserved—conserved in teleosts, amphibians, avians, mammals; very highly conserved—conserved in fungi, nematodes, insects, teleosts, avians, mammals; absent from controls—was not found in the control databases ExAC, gnomAD, 1000 Genomes as well as GME Variome for patients of Middle Eastern ancestry. The variant nomenclature was confirmed via [www.mutalyzer.nl](http://www.mutalyzer.nl) to conform with Human Genome Variation Society (HGVS) standards.

AA amino acid; ACMG American College of Medical Genetics and Genomics; AF allele frequency according to the control databases ExAC, gnomAD, 1000 Genomes as well as GME Variome for patients of Middle Eastern ancestry; CADD Combined Annotation Dependent Depletion score;<sup>39</sup> CNV copy-number variant; ESHG European Society of Human Genetics; FA fetal akinesia; PMID PubMed ID; SMA spinal muscular atrophy; VOI variant of interest in the ESHG classification system; VUS variant of uncertain significance in the ACMG classification system.

order correlation to identify statistical significant differences between categorical classifications for phenotypical distribution, etiological classification, NGS success, and molecular function distribution (further details in Supplementary Materials file 1, “Methods” section).

## RESULTS

### Fetal akinesia phenotype classification

We recruited 51 patients from 47 families, including four affected sibling pairs, who were diagnosed with FA. Of 51 individuals in the cohort, 26 were female and 25 were male. In 21 cases the pregnancy was either terminated prematurely, the child was stillborn, or the child died shortly after birth. The age range of the other patients are 0–28 years (median age 1 year).

The cohort can be grouped into five categories based on their clinical phenotype (Table 1 and Supplementary Table S5). We propose to expand the clinical classification first introduced by Hall,<sup>3</sup> and recently critically reviewed,<sup>14</sup> as in our opinion the viability of the phenotype is an important feature. Category I consists of ten patients with a phenotype limited primarily to limb contractures. Category II contains patients with limb contractures and other systemic malformations and is split into IIa, indicating a viable phenotype with 14 patients, and IIb, indicating the seven patients who died either prenatally or within their first year of life. Similarly, category III contains patients with limb contractures and neurological abnormalities, and again IIIa contains the 12 patients with viable phenotypes whereas IIIb contains the eight lethal phenotypes (Fig. 1a, b).

### Molecular findings

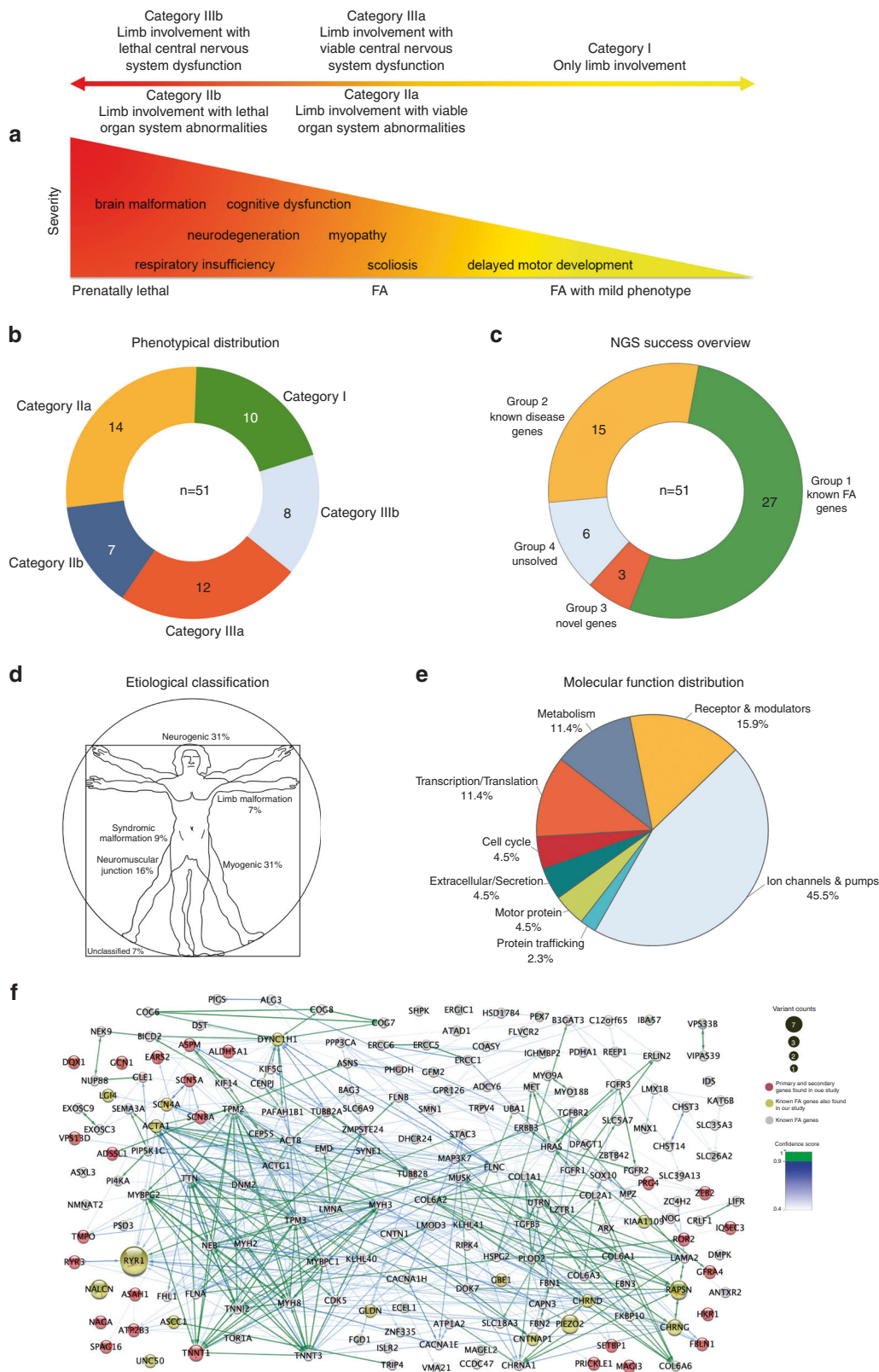
The pathogenic variants in 45 of 51 cases can be divided into the following three categories (Fig. 1c):

In 27 cases we found the underlying pathogenic variants in known FA-associated genes, including a sib pair with a homozygous deletion in *RAPSN* (cases 16 and 17; Supplementary Fig. S15b). Two of the variants in *ACTA1* (case 1) and *PIEZO2* (case 15) respectively were already known according to ClinVar (ClinVar IDs 381639 and 137629). The remaining 31 variants are to our knowledge novel pathogenic variants.

In a further 15 cases, the underlying pathogenic variants are in known disease genes that until now had not been linked to FA. These novel FA candidate genes containing 18 pathogenic variants are as follows: *ADSSL1*, *ASAH1*, *ASPM*, *ATP2B3*, *EARS2*, *FBLN1*, *PRG4*, *PRICKLE1*, *ROR2*, *SETBP1*, *SCN5A*, *SCN8A*, and *ZEB2*, as well *TNNT1* with a homozygous copy-number variant (CNV) deletion in a sibling pair (cases 40 and 41; Supplementary Fig. S15a). The pathogenic variant in *SETBP1* (case 37) is a known pathogenic variant (ClinVar ID 1031) for Schinzel–Giedion syndrome (OMIM 269150), which fits the patient's phenotype. A homozygous variant in *PRG4* (case 34) and the homozygous CNV deletion in *TNNT1* mark the first time these genes have been linked to FA beyond in silico predictions.<sup>15</sup>

Furthermore, in three cases we have found likely pathogenic variants in putative novel FA candidate genes without clear and convincing prior FA-disease link until now: *GCNI*, *IQSEC3*, and *RYR3*. A de novo heterozygous variant in *GCNI* and compound heterozygous variants in *VPS13D* were found in case 43. While *GCNI* appears to play an important role in translational regulation during stress response, the scientific literature provides no clear disease association beyond a tenuous link to schizophrenia.<sup>16,17</sup> *VPS13D*, on the other hand, is involved in mitochondrial regulation and has been linked to ataxia and intrauterine fetal death.<sup>6,18</sup> However, the variant in *GCNI* is a confirmed de novo variant that is absent from controls with a Combined Annotation Dependent Depletion (CADD) score of 22, the gene has a Residual Variation Intolerance Score (RVIS) of  $-3.77$ , is expressed in skeletal muscle and cerebellum brain tissue according to GTEx, and according to the Gene Network 2.0<sup>19</sup> is coregulated with the FA gene *DYNCH1H1* ( $p$  value  $6.7 \times 10^{-42}$ ), whereas one of the compound heterozygous variants in *VPS13D* has a rather high allele frequency. Therefore, we consider the variant in *GCNI* as the primary candidate while the compound heterozygous variants in *VPS13D* might be modifiers or contribute to an oligogenic phenotype. The *VPS13D* variants are therefore listed as secondary variants. In case 44 we found compound heterozygous variants in *IQSEC3*, a gene known to play a role in the regulation of synapse formation<sup>20</sup> and recently linked to a neurodevelopmental phenotype in two cases.<sup>21</sup> One of the two variants causes an amino acid change, is absent from controls, and reaches a CADD score of 26.5, while the other one is predicted to trigger a cryptic splice site activation. According to GTEx, *IQSEC3* is almost exclusively expressed in the brain, and particularly in the cerebellum. It has a RVIS score of  $-0.84$  and is also coregulated with another novel FA candidate gene, *ATP2B3* ( $p$  value  $8.8 \times 10^{-21}$ ), according to the Gene Network 2.0. Therefore, we conclude that given its function in neuronal development *IQSEC3* is a likely candidate as a novel FA gene. A highly conserved heterozygous variant in *RYR3* has been found in case 45 with a high CADD score of 32. *RYR3* is a  $\text{Ca}^{2+}$  ion channel for excitation–contraction coupling with a RVIS score of  $-5.87$  and closely related to the known FA gene *RYR1*. A stringApp analysis of the interactome showed that it interacts closely with the FA genes *RYR1* and *STAC3* (see Supplementary Fig. S25). It has recently been linked to nemaline myopathy<sup>22</sup> as well as a phenotype with global developmental delay.<sup>23</sup> Given its expression in both brain and muscle tissue and the murine animal knockout model with a severe neuromuscular phenotype it is a promising novel FA disease gene candidate.<sup>24</sup>

This translates to a 73% (37/51) success rate of solved cases and in total 88% (45/51) of possibly solved cases including potentially pathogenic variants linked to the observed phenotype. Detailed overviews of the clinical features of each patient and of the proposed causative pathogenic variants can be found in Tables 1 and 2, respectively.



**Disease association of candidate genes**

With regard to known disease associations and anatomical-physiological locations of the primary defect, the disease candidate genes were grouped (Fig. 1d) as follows.

Eleven genes are known to cause neurogenic disease phenotypes: *ASAH1*, *ASPM*, *ATP2B3*, *CNTNAP1*, *EARS2*, *GLDN*, *LGI4*, *NALCN*, *PRICKLE1*, *SCN8A*, and *ZEB2*. Another seven genes are linked to myogenic phenotypes:



**Fig. 1 Fetal akinesia (FA) cohort overview.** (a) Phenotypical spectrum of the cohort ranging from prenatally lethal FA to a mild arthrogrypotic phenotype. We introduce a modified version of the classification first proposed by Hall.<sup>3</sup> Category I consists of patients with a phenotype limited primarily to limb contractures. Category II contains patients with limb contractures and other systemic malformations, and is split into IIa, indicating a viable phenotype, and IIb, indicating patients who died either prenatally or within their first year of life. Similarly, category III contains patients with limb contractures and neurological abnormalities, and again IIIa contains the viable phenotypes whereas IIIb contains the lethal phenotypes. (b) The cohort has been sorted into the five phenotypical categories based on the modified Hall classification described above. (c) The cohort has been sorted into the following categories with regard to the success of next-generation sequencing (NGS) in identifying potential pathogenic variants: group 1—identified variants or copy-number variants (CNVs) in known AMC/FADS genes; group 2—identified variants in known disease genes; group 3—identified putative variants in genes previously not linked to a disease; group 4—currently unsolved cases. (d) Etiological distribution grouped into primarily neurogenic or myogenic pathomechanisms, pathomechanisms involving the neuromuscular junction, limb malformation and syndromic malformation, as well as unclassified in three cases where no prior disease association is known. (e) Breakdown of the molecular function of the genes carrying pathogenic variants. (f) This network map shows the gene interactions generated from STRINGdb. Gray nodes represent the previously published FA-related genes that are not present in our study; yellow nodes represent the previously published FA genes that are also found in our study; red nodes represent the new primary and secondary candidate genes that have been found in our study. Size differences illustrate the variant counts for a particular gene in this study. Blue lines between nodes represent the combined confidence score of interaction between 0.4 and 0.9 (medium confidence–high confidence). Green lines represent the combined confidence score above 0.9 (highest confidence).

*ACTA1*, *ADSSL1*, *GBE1*, *RYR1*, *SCN4A*, *SCN5A*, and *TNNT1*. The third group includes four genes that are associated with dysfunction of the neuromuscular synaptic junction: *CHRND*, *CHRNG*, *RAPSN*, and *UNC50*. Additionally, *ASCC1*, *PIEZO2*, and *SETBP1* are linked to complex developmental syndromic malformations including the nervous system. Another set of genes are known to cause limb malformations—this group includes *FBLN*, *PRG4*, and *ROR2*. For detailed information regarding these disease associations please refer to the PubMed ID (PMID) links provided in Supplementary Table S1. Recently, the first disease associations have been reported for *IQSEC3*<sup>21</sup> and *RYR3*.<sup>22,23,25</sup> *GCNI* had been associated with schizophrenia in genome-wide association studies (GWAS).<sup>16</sup> All primary pathogenic variant candidates are listed in Table 2, and Supplementary Table S1 contains additional information for the primary pathogenic variant candidates. In addition to the candidate genes listed above, secondary variants besides the primary candidates survived our filtering strategy in several cases: *ALDH5A1*, *DQX1*, *DYNC1H1*, *GFRA4*, *HKR1*, *KIAA1109*, *MAGI3*, *NAGA*, *PIEZO2*, *SPAG16*, *TMPO*, and *VPS13D* (Supplementary Table S2). They might function as disease phenotype modifiers or form part of an oligogenic phenotype in the case of likely pathogenic variants in *ALDH5A1*, *KIAA1109*, or *PIEZO2*.

### Gene functions

The potential candidate genes discovered in our cohort cover a broad spectrum of molecular functions based on the gene function descriptors of OMIM (Fig. 1e) as expected in the case of phenotypes such as AMC and FADS with their heterogeneous genetic etiology. The most common group among our candidates are the ion channel or ion pump genes such as *ATP2B3*, *CHRNG*, *CHRND*, *NALCN*, *PIEZO2*, *RYR1*, *SCN4A*, *SCN5A*, *SCN8A*. Functionally related are the receptor protein genes such as *ROR2*, ion channel modulators like *RAPSN*, as well as myelination modulators like *CNTNAP1* and *LG14*. Another major group contains genes involved in the regulation of transcription and translation such as *ASCC1*, *GCNI*, *EARS2*, *PRICKLE1*, *SETBP1*, and *ZEB2* as well as

*UNC50*, a gene involved in protein trafficking. Motor protein genes like *ACTA1*, *DYNC1H1*, or *TNNT1* are also potential disease candidates for some of our patients, while the extracellular matrix structure gene *FBLN1* and the proteoglycan gene *PRG4* contain other potentially pathogenic variants. In other cases, pathogenic variants are found in genes involved in cell cycle regulation (*ASPM*) or cell development (*GLDN*). The last major group of disease candidate genes contains several genes involved in cytosolic, mitochondrial, or lysosomal metabolic function such as *ADSSL1*, *ASAHI*, *GBE1*, and *IQSEC3*. Figure 1f illustrates the protein–protein interaction network based on STRINGdb and contains all proteins linked to an FA phenotype as well as the primary and secondary candidates found in this study.

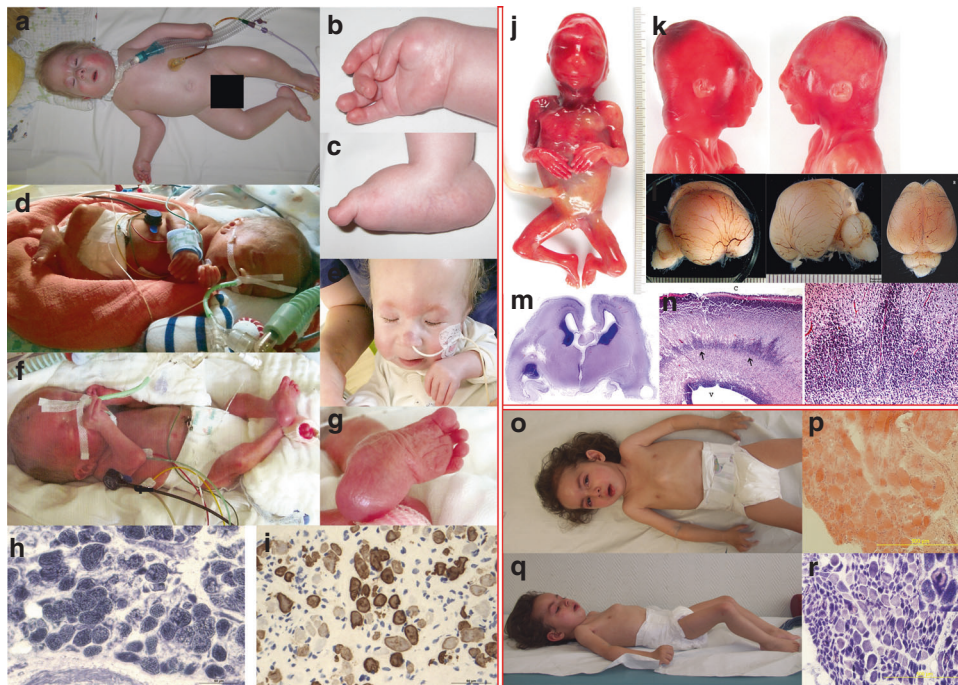
### Correlation analysis

A correlation analysis between the phenotype of the patients, the etiology or the pathomechanism of the disease-associated genes, and their molecular function (based on the data collated in Supplementary Table S5) revealed (Supplementary Fig. S22) that the more severe phenotypes (IIb and IIIb) could all be solved whereas the milder phenotypes (I, IIa, and IIIa) contain several cases without any promising candidate genes that could be discovered via NGS. While the molecular function of genes where the pathomechanism leads to primary muscle diseases is concentrated on motor proteins, ion channels, and cell metabolism proteins, the genes with a neurogenic pathomechanism appear more diverse in their molecular function.

### DISCUSSION

The aim of this study was to determine the underlying genetic cause for FA by NGS, a genetically extremely heterogeneous syndromic disease entity. This heterogeneity comprises hundreds of genes containing causative pathological variants (Supplementary Table S4).

The results from this study suggest a major role for pathogenic variants in ion channel genes and genes coding for ion channel modulators in the pathogenesis of FA (Fig. 1e).



**Fig. 2 Clinical presentation of cases 26, 30, and 40.** (a) Fetal akinesia (FA) patient 26 with a de novo *SCN4A* variant with respiratory support at 24 months. (b) Close-up view of the right hand of patient 26 with an ulnar deviation of the hand at 24 months. (c) Close-up view of the right foot of patient 26 with characteristic joint contractures at 24 months. (d) Patient 26 at 2 months with prominent joint contractures. (e) Close-up view of the facial features of patient 26 at 12 months. (f) Patient 26 shortly after birth with prominent joint contractures. (g) Congenital contractures of the left foot observed shortly after the birth of patient 26. (h) NADH stain of muscle tissue biopsy of patient 26 showing darkly stained type 1 fibers, myopathic presentation. (i) Slow myosin immunostain of type 1 muscle fibers (dark) of patient 26 showing an increased variation of fiber size. No fiber grouping is observable, myopathic presentation. (j) Patient 30: aborted fetus in the 18th gestational week with microcephaly and arthrogryposis due to compound heterozygous *ASPM* pathogenic variants. (k) Lateral view left/right of the microcephalic head of patient 30. (l) Lateral view left/right and axial view of the fetal brain of patient 30 with hypoplastic forebrain and lack of gyration. (m) Histological overview of a fetal brain tissue slice of patient 30. (n) Increased magnification of fetal brain tissue of patient 30 with reduced size of the cortical plate (C) and neuroblast stretch between cortex and lateral ventricle (V) resembling subcortical band heterotopia. (o) Patient 40 with prominent pectus carinatum, facial features, and multiple joint contractures including jaw contractures at the age of 2 due to a homozygous copy-number variant (CNV) deletion in *TNNT1*. (p) Hematoxylin and eosin (H&E) stain of muscle tissue biopsy of patient 40 with several hypertrophic and numerous small atrophic fibers and pathological caliber variance, myopathic presentation. (q) Lateral view of patient 40 at the age of 2 years with flexed hips due to contractures. (r) NADH stain of muscle tissue biopsy of patient 40 with trilaminar fibers, in the upper right corner, is an enhanced picture detail showing a trilaminar fiber.

*RYR1* alone contained the causative pathogenic variants in five unrelated cases and one sibling pair of a total of 37 solved cases. Our study proves the importance of pathogenic variants in *RYR1* as a potential cause for FA in a large cohort together with an earlier description by Alkhunaizi.<sup>26</sup>

Some of our findings are particularly noteworthy. *TNNT1* was first linked to human diseases by Johnston *et al.*,<sup>27</sup> who proved its link to a subtype of nemaline myopathy (OMIM 605355) occurring in old order Amish families. The first case reported outside the old order Amish population by van der Pol *et al.*<sup>28</sup> shares remarkably similar clinical symptoms such as pectus carinatum, kyphoscoliosis, multiple joint contractures, respiratory insufficiency, and dysphagia with our patients (40 and 41) from one afflicted family of Eastern European (Romania/the former Yugoslavia) descent (Fig. 2). The reported results of the muscle biopsy generally mirror our findings with highly divergent muscle fiber calibers and increased endo- and perimysial connective tissue. Our findings differ in that we see trilaminar fibers (Fig. 2r) that have not yet been reported in *TNNT1* cases whereas no

classical nemaline rods were present in the examined tissue samples.<sup>29</sup> To our knowledge, this is the first time that a pathogenic gene variant has been discovered for myopathy with trilaminar fibers.<sup>30</sup> While in all previous cases the causative pathogenic variant was homozygous or there were compound heterozygous small mutations, here we report a homozygous CNV deletion in the two affected siblings as a likely cause for the phenotype, as reported very recently in another case with different myopathic phenotype.<sup>31</sup>

*ADSSL1* has only recently been discovered as a myopathy gene.<sup>32,33</sup> Here we report the first case of a pathogenic homozygous frameshift *ADSSL1* variant in a patient of Turkish descent (patient 28). Previously, *ADSSL1* pathogenic variants (OMIM 617030) have been reported in the Korean population. Unlike the eight Korean patients, our unrelated patient presented with congenital joint contractures and a more severe neurological phenotype, while all previously described patients with recessive *ADSSL1* pathogenic variants suffered from adolescent-onset myopathy and did not show any contractures. A possible explanation for the earlier onset

and more severe phenotype could be the fact that all previously described mutations resulted in amino acid exchanges except for one compound heterozygous frameshift variant. This frameshift variant truncates the sequence of the protein after a substrate binding domain, unlike the frameshift variant we present here. Another factor could be the presence of a frameshift variant in *ALDH5A1*, which might modify the phenotype of the patient as a multilocus phenotype.<sup>34</sup>

Our study emphasizes that neuronal migration defects may lead to fetal akinesia. We present the histopathological examination of the brain of a fetus (patient 30) that suffered from an *ASPM* nonsense variant at—to our knowledge—the earliest developmental stage published (Fig. 2). The neuronal migration was delayed and resulted in a neuroblast layer in the subventricular cortical zone still present at gestational week 18 and consequently a hypoplastic cortex—in particular in the frontal cortex (Fig. 2).<sup>35</sup> The results of Passemard *et al.*<sup>36</sup> agree with our findings that *ASPM* defects in humans primarily affect cortical development, which in turn leads to reduced intrauterine movement and consequently an FA phenotype.

With a success rate of about 88%, we could determine the underlying genetic defect for most of the patients in the cohort presented here. The initially employed targeted panel approach using the Mendeliome proved unsatisfactory because only 41% (21 cases) of all cases could be solved this way. For the remaining initially unsolved 30 cases, the ES analysis had a success rate of 53% (16 cases). This includes the four cases that could only be solved by systematic CNV analysis, which significantly increased the power of pathogenic variant detection in ES data sets.

Both of the currently employed ES kits provided a satisfactory average coverage of 20× or above and allowed the analysis of all 166 known FA genes, whereas the initially used Mendeliome enrichment kit did not sufficiently cover 45 known FADS disease genes. Even the improved version of the Mendeliome enrichment kit failed to cover 20 genes.

Another issue of recent interest due to the increasing availability of ES data is multilocus or oligogenic pathogenic variations and their combined effect on the phenotype of individual patients. The work of Posey *et al.*<sup>34</sup> and Karaca *et al.*<sup>37</sup> indicated that pathogenic multilocus variants are a valid alternative to classical phenotypic expansion of a known disease gene when it comes to explaining divergent phenotypes in consanguineous families. Here, we demonstrate the validity of this hypothesis also in nonconsanguineous cases. The potential multilocus candidates that survived the filtering process are provided in Supplementary Table S2.

The genes containing presumed pathogenic variants in our study vary in both molecular function and disease phenotype association, but most can be broadly grouped into five groups. The two major groups contain myogenic and neurogenic genes, while the three smaller groups consist of genes linked to neuromuscular junction dysfunction, limb malformations, and syndromic malformations involving multiple organ systems (Fig. 1d, Supplementary Table S5).

A total of 192 genes were included in the systems biology approach. One hundred fifty of those were previously published with an FA association but not found in our study; 16 were found in both gene lists and 26 were found only in our study (Supplementary Tables S1, S2, S4). The STRINGdb analysis (Fig. 1f) performed for this study showed that most of the novel genes proposed to be involved in the development of an FA phenotype are interacting directly with genes known to cause AMC/FADS. This supports the idea that pathogenic variants in these novel genes adversely affect the same pathways as pathogenic variants in known FA disease genes. An example of this would be *SCN8A*, which interacts with *SCN4A* and *CNTNAP1*, or *ASPM*, which interacts with *KIF14*, *CENPJ*, and *CEP55*. Moreover, even within the group of the known FA disease genes, there are multiple outliers with no direct interaction with any other known FA disease gene, indicating that there might be gaps in our knowledge regarding the protein networks and metabolic pathways involved in the FA phenotypical spectrum. This of course means that even if no direct interaction with a known disease gene can be found via this method, a novel gene carrying a pathogenic variant should not be excluded based solely on this lack of interaction partners, as it might belong to the outlier group with no direct interaction with the large majority of known FA disease genes.

The Gene Ontology enrichment analysis (Supplementary Figs. S16–21) showed that our novel candidate genes reveal a similar enrichment pattern as the known FA genes used as a comparison group with regard to the biological process affected, the molecular function of the protein, and cellular compartment localization, favouring ion channels, primary muscle/myogenic and excitation-coupling genes/pathways. The notable outcome was that the over-representation analysis (ORA) performed with known FA genes had a higher statistical significance compared with the ORA performed with our candidates for all GO classes (known FA genes GO:BP ORA  $p > 2 \times 10^{-6}$ ; all candidates GO:BP ORA  $p > 0.01$ ; known FA genes GO:CC ORA  $p > 0.001$ ; all candidates GO:CC ORA  $p > 0.025$ ). This was to be expected given the difference in the number of genes used in the analysis. It was also supported by the fact that the enriched genes are often common to both gene lists (Supplementary Figs. S16–21).

Overall, our findings support the usage of exome sequencing and stringent variant filtering combined with thorough clinical assessment of patients together with refiltering as an essential tool to uncover the genetic etiology of complex syndromic diseases. Applying a selected panel of disease genes—the Mendeliome—can certainly be helpful in identifying novel pathogenic variants in known disease genes as we solved 41% of all cases this way. However, by analyzing not only a particular set of enriched genes as in a gene panel but examining the entirety of the patients' exome with a superior target coverage previously unnoticed pathogenic variants even in known disease genes can be detected. In addition, the effect of multiple pathogenic variants on an oligogenic phenotype

might become evident, and CNVs with pathogenic effect are observable, as we have shown here (Supplementary Fig. S15).

The methodical analysis presented in this publication and subsequent efforts will hopefully enable clinicians to determine the underlying genetic cause of many FA cases that until now have been labeled as sporadic and allow for improved genetic counseling and prenatal diagnosis. This makes ES a tool that is most suited to uncover potential issues for couples planning further pregnancies, in particular given the recent developments of rapid exome sequencing and ultrarapid genome sequencing.<sup>38</sup>

### SUPPLEMENTARY INFORMATION

The online version of this article (<https://doi.org/10.1038/s41436-019-0680-1>) contains supplementary material, which is available to authorized users.

### ACKNOWLEDGEMENTS

We thank the patients and their families contributing to this clinical study. Furthermore, we thank Brunhilde Wirth of the Institute of Human Genetics of the University of Cologne as well as Peter Herkenrath from the Institute of Pediatrics of the University Hospital Cologne, Ulrike Schara from the University Children's Hospital Essen, Mehmet Saracoglu from the Akdeniz University Hospital Antalya, Tacy Dudding-Byth of the NSW Genetics of Learning Disability Service, and also Birgit Kampschulte, Axel Weber, and Markus Waitz from the Department of Pediatrics and Neonatology at the university hospital of Justus Liebig University Gießen for their contributions toward patient acquisition for this study as well as the collection of their patients' clinical data. We thank Monia Al-Areeqi from the Center for Molecular Medicine Cologne (CMMC) for her technical assistance as well. We furthermore thank the Regional Computing Center of the University of Cologne (RRZK) for providing computing time on the Deutsche Forschungsgemeinschaft (DFG)-funded high-performance computing (HPC) system CHEOPS as well as technical support. The research published in this paper was funded by the DFG Emmy Noether Grant (CI 218/1-1) to S.C.

### DISCLOSURE

The authors declare no conflicts of interest.

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