

Phenotype Presentation and Molecular Diagnostic Yield in Non-5q Spinal Muscular Atrophy

Gorka Fernández-Eulate, MD, Julian Theuriet, MD, Christopher J. Record, MD, Giorgia Querin, MD, PhD, Marion Masingue, MD, Sarah Leonard-Louis, MD, Anthony Behin, MD, Nadine Le Forestier, MD, Antoine Pegat, MD, Maud Michaud, MD, Jean-Baptiste Chanson, MD, Aleksandra Nadaj-Pakleza, MD, Celine Tard, MD, PhD, Anne-Laure Bedat-Millet, MD, Guilhem Sole, MD, Marco Spinazzi, MD, Emmanuelle Salort-Campana, MD, Andoni Echaniz-Laguna, MD, PhD, Vianney Poinsignon, PharmD, Philippe Latour, PhD, Mary M. Reilly, MD, Françoise Bouhour, MD, and Tanya Stojkovic, MD

Correspondence
Dr. Stojkovic
tanya.stojkovic@aphp.fr

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Abstract

Background and Objectives

Spinal muscular atrophy (SMA) is mainly caused by homozygous *SMN1* gene deletions on 5q13. Non-5q SMA patients' series are lacking, and the diagnostic yield of next-generation sequencing (NGS) is largely unknown. The aim of this study was to describe the clinical and genetic landscape of non-5q SMA and evaluate the performance of neuropathy gene panels in these disorders.

Methods

Description of patients with non-5q SMA followed in the different neuromuscular reference centers in France as well as in London, United Kingdom. Patients without a genetic diagnosis had undergone at least a neuropathy or large neuromuscular gene panel.

Results

Seventy-one patients from 65 different families were included, mostly sporadic cases (60.6%). At presentation, 21 patients (29.6%) showed exclusive proximal weakness (P-SMA), 35 (49.3%) showed associated distal weakness (PD-SMA), and 15 (21.1%) a scapuloperoneal phenotype (SP-SMA). Thirty-two patients (45.1%) had a genetic diagnosis: *BICD2* (n = 9), *DYNC1H1* (n = 7), *TRPV4* (n = 4), *VCP*, *HSBP1*, *AR* (n = 2), *VRK1*, *DNAJB2*, *MORC2*, *ASAHI*, *HEXB*, and unexpectedly, *COL6A3* (n = 1). The genetic diagnostic yield was lowest in P-SMA (6/21, 28.6%) compared with PD-SMA (16/35, 45.7%) and SP-SMA (10/15, 66.7%). An earlier disease onset and a family history of the disease or consanguinity were independent predictors of a positive genetic diagnosis. Neuropathy gene panels were performed in 59 patients with a 32.2% diagnostic yield (19/59). In 13 additional patients, a genetic diagnosis was achieved through individual gene sequencing or an alternative neuromuscular NGS.

Discussion

Non-5q SMA is genetically heterogeneous, and neuropathy gene panels achieve a molecular diagnosis in one-third of the patients. The diagnostic yield can be increased by sequencing of other neuromuscular and neurometabolic genes. Nevertheless, there is an unmet need to cluster these patients to aid in the identification of new genes.

From the Nord/Est/Ile-de-France Neuromuscular Reference Center (G.F.-E., G.Q., M. Masingue, S.L.-L., A.B., T.S.), Institut de Myologie, Pitié-Salpêtrière Hospital, Paris; Electromyography and Neuromuscular Department (J.T., A.P., F.B.), Hospices Civils de Lyon; Centre for Neuromuscular Diseases (C.J.R., M.M.R.), UCL Queen Square Institute of Neurology, London, United Kingdom; Neurology Department (N.L.F.), Pitié-Salpêtrière Hospital, Paris; Nord/Est/Ile-de-France Neuromuscular Reference Center (M. Michaud), Central Nancy University Hospital; Nord/Est/Ile-de-France Neuromuscular Reference Center (J.-B.C., A.N.-P.), Strasbourg University Hospitals; Nord/Est/Ile-de-France Neuromuscular Reference Center (C.T.), U1172, Lille University Hospital; Nord/Est/Ile-de-France Neuromuscular Reference Center (A.-L.B.-M.), Rouen University Hospital; Neuromuscular Reference Center 'AOC' (G.S.), Bordeaux University Hospitals (Pellegrin Hospital); Neuromuscular Reference Center (M.S.), Angers University Hospital; Neuromuscular and ALS Reference Center (E.S.-C.), La Timone University Hospital, Marseille; French National Center for Rare Neuropathies (A.E.-L.), Neurology Department, Bicêtre University Hospital, INSERM U1195, Paris-Saclay University; Molecular Genetics Lab (V.P.), Bicêtre University Hospital, Le Kremlin Bicêtre; and Center for Biology - East (P.L.), Neurological Hereditary Disorders Unit, Hospices Civils de Lyon, France.

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Glossary

ALS = amyotrophic lateral sclerosis; **CK** = creatine kinase; **CMAP** = compound motor action potentials; **CMT** = Charcot-Marie-Tooth; **DTR** = deep tendon reflexes; **ENMG** = electroneuromyography; **HMN** = hereditary motor neuropathies; **HSP** = hereditary spastic paraplegia; **NGS** = next-generation sequencing; **NHNN** = National Hospital for Neurology and Neurosurgery; **NIV** = noninvasive ventilation; **SMA** = spinal muscular atrophy; **SNAP** = sensory nerve action potentials; **VUS** = variant of unknown significance; **WES** = whole exome sequencing; **WGS** = whole genome sequencing.

Hereditary motor neuropathies (HMN) are a heterogeneous group of diseases characterized by peripheral motor neuron weakness with minimal or absent clinical and neurophysiologic sensory impairment, which differentiates them from Charcot-Marie-Tooth (CMT) disease and the hereditary sensory neuropathies.¹

Distal HMN (dHMN) represent the most common clinical presentation of HMN.^{2,3} The term spinal muscular atrophy (SMA) is mainly used when referring to HMNs with proximal weakness at onset, either isolated (P-SMA) or concomitant with distal (PD-SMA) or scapulo-peroneal (SP-SMA) motor weakness.⁴ A homozygous exon 7 and 8 deletion in the *SMN1* gene represents the main cause of SMA,⁵ ranging from 50-94% of patients depending on the series.^{6,7} In the absence of this deletion (non-5q SMA), several genes have been described, such as *TRPV4*, *DYNC1H1*, *BICD2*, *GLE1*, *UBA1*, *ASAH1*, *VRK1*, *EXOSC3*, *SLC52A3* and *SLC52A2*, *VAPB*, *HEXB*, *SETX*, *LMNA*, and *TFG* with different clinical phenotypes and ages at onset of disease.^{4,6,8-12} These and other genes may overlap with other nosological entities such as dHMN, CMT type 2, amyotrophic lateral sclerosis (ALS), hereditary spastic paraplegia (HSP), or even mitochondrial disease and myopathy.^{4,13} Nevertheless, around 70% or more of non-5q SMA patients do not have a definitive molecular diagnosis.⁶

Unfortunately, although some reviews of non-5q SMA patients have been published,^{4,14} clinical series of patients are lacking and the value of routine neuropathy next-generation sequencing (NGS) gene panels is largely unknown. The aim of this study was to describe the clinical and genetic landscape of a large multicentric series of patients with non-5q SMA and evaluate the performance of neuropathy gene panels in these patients.

Methods

Inclusion and Exclusion Criteria

Between April and May 2021, we retrospectively reassessed the clinical records of adult patients (18 years or older) with an HMN and predominant proximal weakness (SMA) from 10 neuromuscular reference centers in France. Sixty-one patients were included based on the following criteria: (1) Patients had a motor neuropathy or motor neuronopathy on electroneuromyography (ENMG), defined by the presence of large-size (>2 mV in amplitude) simple-shaped motor unit potentials and reduced recruitment. In addition, reductions of compound motor action potential (CMAP) amplitudes and

less than 30% reduction of normal sensory nerve action potential (SNAP) amplitudes (reference values for adult patients: >10 μ V in lower limbs, >8 μ V for ulnar, and >15 μ V for median and radial nerves) were accepted. (2) Patients were negative for the *SMN1* deletion and (3) had a confirmed genetic diagnosis or at least undergone a neuropathy NGS gene panel. Patients with clinical or neurophysiologic disease progression at 6 months were excluded.

Furthermore, clustered data from 10 adult patients followed in the Centre for Neuromuscular Diseases, UCL Queen Square Institute of Neurology, London (United Kingdom) were obtained following the same inclusion and exclusion criteria.

Based on the clinical presentation, patients were classified in 3 groups: (1) Patients with P-SMA had pure proximal motor weakness at first visit, (2) patients with PD-SMA had proximal and distal weakness but pure proximal involvement at disease onset or proximal-predominant weakness at first visit, and (3) patients with SP-SMA had scapulo-peroneal weakness. Demographic (date of birth, sex) and clinical information (age at onset, family history of neuropathy, consanguinity, neurologic, orthopedic, and respiratory manifestations) were collected. Additional data such as ENMG findings, creatine kinase (CK) levels, the need of noninvasive ventilation (NIV), and molecular biology results were collected when available, as well as loss of ambulation as functional outcome.

Molecular Biology Investigations

The molecular biology investigations performed in each patient are shown in eTable 1 (links.lww.com/NXG/A614).

Neuropathy Gene Panels

Fifty-nine neuropathy gene panels were performed in total. Forty-eight were performed at the molecular genetic laboratory of Hospices Civils de Lyon. DNA was extracted from blood samples using Nucleospin Blood L (Macherey-Nagel) on automat Microlab STARlet (Hamilton). Capture was performed with Roche KAPA Hypercap v3.0. Sequencing was performed using Illumina NextSeq 500. Copy number variations were detected using DeCovA_1.6.0. Sequencing depth at 30x was over 99,95%. One hundred two genes are included in this panel (eAppendix 1, links.lww.com/NXG/A613).

Eleven additional neuropathy gene panels were sequenced by the National Hospital for Neurology and Neurosurgery (NHNN) in London (n = 4, 15–32 genes) and the molecular

genetic laboratories from Kremlin-Bicêtre (n = 4, 127 genes), Limoges (n = 2, 124 genes), and Angers (n = 1, 148 genes).

Alternative Neuromuscular Gene Panels and Individual Gene Sequencing

Eight patients underwent an alternative large neuromuscular gene panel: whole genome sequencing (WGS) with a ~180 gene virtual panel performed by the NHNN in London (n = 3), a whole exome sequencing (WES) focused on known neuromuscular genes (n = 1), and large neuromuscular gene panels in Paris (n = 2), Strasbourg (n = 1), and Nancy (n = 1).

In 8 patients, a genetic diagnosis was confirmed through individual sequencing of other clinically suspected neuromuscular or neurometabolic genes.

Whole Exome Sequencing

Furthermore, in 20 patients without a molecular diagnosis after a neuropathy NGS panel, WES was performed at Hospices Civils de Lyon. Exome enrichment was performed with the SeqCap EZ MedExome Target Enrichment Kit (Roche Diagnostics). Sequencing was performed on a NextSeq 500 (Illumina) platform, aligned using Borrows-Weeler Aligner to a hg19 reference genome; variants were called using Genome Analysis Tool Kit Unified genotyper software, and filtering of data were performed based on an in-house pipeline: 1000 genomes frequency <0.01, maximum ExAC occurrences heterozygotes 250, maximum ExAC occurrences homozygotes 2, minimal read depth >7, mosaicism >0.15, coding exons ± 10 .

Variants were classified according to the 2015 American College of Medical Genetics Standards and Guidelines for the interpretation of sequence variants,¹⁵ and patients with pathogenic or likely pathogenic variants were considered as genetically confirmed.

Data Analysis and Statistics

Data analysis was performed using JASP version 0.16.2. Mean and median values, SD, and range of quantitative variables and absolute or relative frequencies of categorical variables were reported. Association was studied through the Student *t* test (continuous dependent variables) and χ^2 test (categorical dependent variables). A multivariate logistic regression was performed to assess the association of relevant clinical variables with a definitive molecular diagnosis. Normality was verified through the Shapiro-Wilk test. Statistical significance was established at $p \leq 0.05$.

Ethics Approval

In accordance with French legislation, the study was registered and is accessible to the French data protection authority (CNIL, Commission nationale de l'informatique et des libertés). This study was approved by the ethics committee of Île-de-France VI on 01/04/2021.

Data Availability

Anonymized data from this study will be shared by request from any qualified investigator.

Results

Clinical Description of Patients and Ancillary Testing

Seventy-one patients from 65 different families were included. Thirty-four patients (47.9%) were female, and most patients had no family history of neuromuscular disease or consanguinity and were therefore classified as sporadic (60.6%). Onset of the disease ranged from 0 to 58 years, and 36 patients (50.7%) had an early onset (younger than 18 years). At presentation (38.2 ± 14.9 years), 21 patients (29.6%) showed exclusive proximal weakness (P-SMA), 35 (49.3%) associated distal weakness (PD-SMA), and 15 patients (21.1%) a scapulo-peroneal phenotype (SP-SMA). Clinical and ancillary data are shown in Table 1. Differences in clinical variables between the P-SMA, PD-SMA, and SP-SMA groups are shown in eTable 2 (links.lww.com/NXG/A615).

On ENMG, all patients had neurogenic motor unit potentials as per inclusion criteria, with 27 (38%) showing reduction of CMAP amplitude and 6 (8.5%) a minimal reduction of SNAPs associated with the peripheral motor neuron involvement.

For 60 patients, follow-up data were available and 5 patients (8.3%) had lost ambulation at a median of 5 years (range 4–54 years) from symptom's onset.

Genetic Diagnosis and Predictive Variables

In 32 patients (45.1%), a genetic diagnosis was achieved. The molecular diagnostic yield was lowest in the P-SMA (6/21, 28.6%) vs the PD-SMA (16/35, 45.7%) and SP-SMA (10/15, 66.7%) clinical groups. The different genes found in each non-5q SMA clinical group can be seen in Figure 1.

Fifty-nine neuropathy NGS gene panels were performed, with a positive genetic diagnosis in 19/59 (32.2%). There was no significant difference in the performance of the neuropathy gene panel from Hospice Civils de Lyon (n = 48) vs other neuropathy gene panels in this series (n = 11) (eTable 3, links.lww.com/NXG/A616). In 8 additional patients, a genetic diagnosis was achieved through individual gene sequencing: *BICD2* (3 patients from a previously reported family with 6 affected individuals),¹⁶ *AR* (n = 2), *HSPB1*, *HEXB*, and *COL6A3*. Reasons for this were genetic testing before routine gene panel sequencing in family members were an index case was found to carry a variant in the gene through exome sequencing (*BICD2*),¹⁶ genetic testing before routine gene panel sequencing and a plausible clinical picture (*HSPB1*), a particular or noncorresponding pattern on muscle MRI (*HEXB* and *COL6A3*), gynecomastia and elevated CK levels (patient with Kennedy disease), and a negative neuropathy gene panel and elevated CK levels (second patient with Kennedy disease). Finally, 5 patients were diagnosed through an alternative neuromuscular NGS: *BICD2* (n = 2), *DYNC1H1*, *VCP*, and *ASAH1*. Furthermore, 20 patients underwent WES, which did not confirm any additional

Table 1 Clinical Features and Ancillary Examinations of Non-5q Spinal Muscular Atrophy (SMA) Patients

| | French (n = 61) | IoN (n = 10) | Total |
|---|--------------------|-----------------|------------------------|
| Female, n (%) | 31 (50.8) | 3 | 34 (47.9) |
| Current age, y, mean ± SD | 48.6 ± 15.1 | 55 ± 15.8 | 49.6 ± 15.2 |
| Age at onset of symptoms <18 y | 31 (50.8) | 5 | 37 (50.7) |
| Family history | | | |
| AD | 13 (21.3) | 5 | 18 (25.4) |
| AR or consanguinity | 10 (16.4) | 0 | 10 (14.1) |
| Sporadic | 38 (62.3) | 5 | 43 (60.6) |
| Age at first consultation, y, mean ± SD | 37 ± 14.4 | 45.5 ± 16.2 | 38.2 ± 14.9 |
| SMA subtype, n (%) | | | |
| Proximal (P-SMA) | 21 (34.4) | 0 | 21 (29.6) |
| Proximo-distal (PD-SMA) | 28 (45.9) | 7 | 35 (49.3) |
| Scapuloperoneal (SP-SMA) | 12 (19.7) | 3 | 15 (21.1) |
| Asymmetric weakness | 6 (9.8) | 5 | 11 (15.5) |
| Facial weakness | 3 (4.9) | 0 | 3 (4.2) |
| Axial weakness | 17 (27.9) | 3 | 20 (28.2) |
| Deep tendon reflexes | | | |
| Brisk | 7 (11.5) | 6 | 13 (18.3) |
| Normal lower limbs | 18 (29.5) | 2 | 20 (28.2) |
| Absent lower limbs | 36 (59) | 2 | 38 (53.5) |
| Spasticity | 0 | 1 | 1 (1.4) |
| Tremor | 14 (23) | 1 | 14 (21.1) |
| Orthopedic | | | |
| Contractures | 10 (17.9) | 2 | 12 (18.2) |
| Arthrogryposis | 5 (8.2) | 0 | 5 (7) |
| Pes cavus/planus | 24 (41.4) | 1 | 25 (36.8) |
| Respiratory insufficiency | 9 (15.8) | 1 | 10 (14.9) |
| Nerve conduction studies^a | | | |
| Reduction of CMAP amplitude | 25 (41) | 2 | 27 (38) |
| Mild reduction of SNAP amplitude | 3 (4.9) | 3 | 6 (8.5) |
| CK levels >300 U/L | 20 (40.1) | 6 (100) | 26 (47.3) ^b |

Abbreviations: CK = creatine kinase; CMAP = compound motor action potentials; CMT = Charcot-Marie-Tooth; IoN = Institute of Neurology UCL; NGS = next-generation sequencing; SMA = spinal muscular atrophy; SNAP = sensory nerve action potential.

The percentage is calculated according to the number of observations. Mild reduction of SNAP amplitude was defined as <30% of expected amplitude. ^a All patients had neurogenic motor unit potentials in EMG as per inclusion criteria.

^b Four patients had CK levels >1000 U/L.

pathogenic or likely pathogenic variants in a known neurologic disease-causing gene.

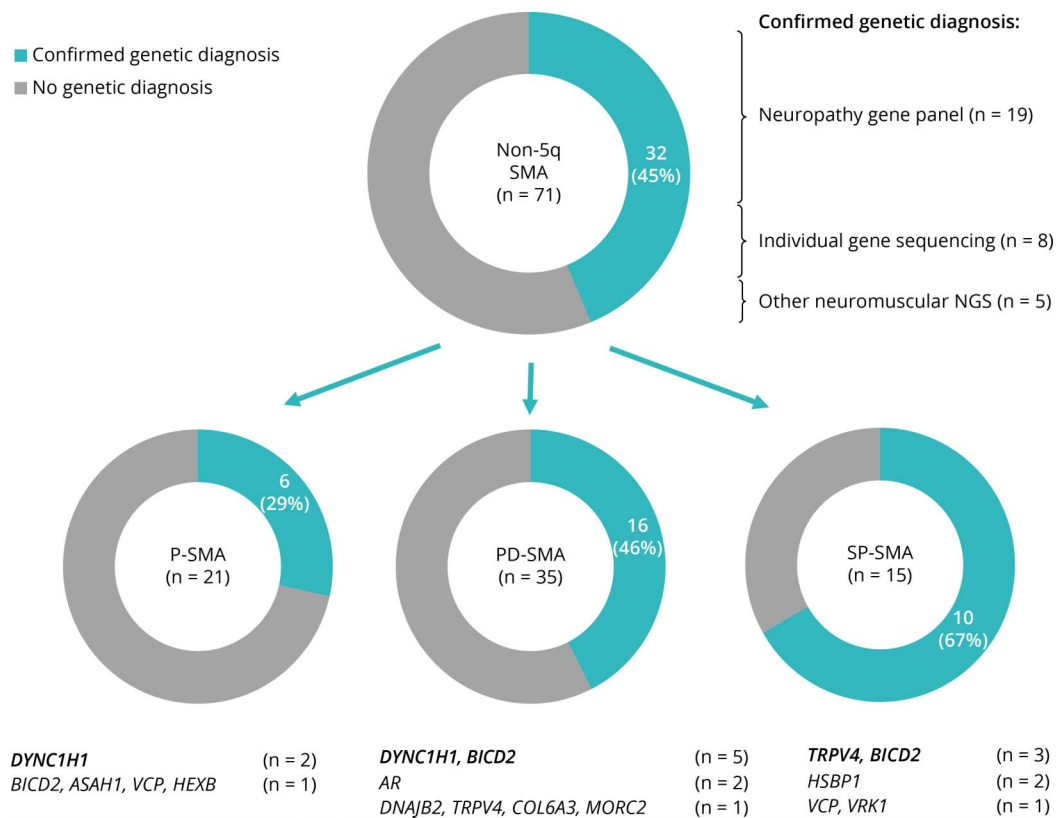
Patients who received a definitive genetic diagnosis were more likely to have an earlier age at onset, a family history of the disease or consanguinity, and contractures and/or arthrogryposis. In a multivariate logistic regression, an earlier onset and a family history of the disease or consanguinity retained statistical significance (Table 2).

Clinical Features of the Most Common Non-5q SMA-Associated Genes

The most common genes in non-5q SMA patients were *BICD2* (n = 9) and *DYNCH1* (n = 7), associated with SMA with lower extremity predominance (SMA-LED), and *TRPV4* (n = 4), implicated in SP-SMA. These 3 genes are dominantly inherited; however, only 13/20 patients had a clear AD family history of the disease, the rest were therefore categorized as sporadic cases. We summarize below the features of these 3 most common genes.

1. *BICD2* (9 patients from 7 different families). Variants: c.1502G>C (n = 3) from the previously reported family,¹⁶ c.2042C>T (n = 2), c.2108C>T, c.2080C>T, c.1617C>G, and c.380A>G. The latter 2 had never been reported: The c.1617C>G variant segregated correctly within the family and predicted as likely pathogenic in silico; the c.380A>G variant was predicted as a likely pathogenic in silico (because it is absent from population databases, well conserved, and situated in a mutational hotspot, with a clinical and muscle MRI phenotype compatible with *BICD2* SMA-LED).¹⁷ However, given that SMA-LED is not a disease with a single genetic etiology and that segregation was not performed, we retain this variant as a variant of unknown significance (VUS) for now. Globally, although all types of phenotypical presentations were possible between and within families [PD-SMA (n = 5), SP-SMA (n = 3) or P-SMA (n = 1)], *BICD2* patients frequently showed some degree of distal weakness on presentation (7/9) and a disease onset younger than 10 years (8/9). Normal or brisk deep tendon reflexes (DTR) (6/9) with 1 patient presenting some degree of spasticity, *scapula alata* (4/9), and axial weakness (3/9) were relatively frequent. Two patients had mild respiratory insufficiency, 1 needing overnight NIV. No patient had lost ambulation at follow-up. All patients had a motor neuropathy on ENMG, one with mild sensory involvement.
2. *DYNCH1* (7 patients from 7 families). Variants: c.2327C>T (n = 2), c.752G>A, c.1792C>T, c.752G>T, c.1427T>C, c.596A>C. The latter 2 variants have never been reported and were predicted as likely pathogenic in silico. Segregation for the c.596A>C confirmed a de novo origin; however, segregation for the c.1427T>C variant could not be performed because the patient was adopted. Clinically, *DYNCH1* patients presented most frequently a PD-SMA (5/7). Disease onset was always in younger

Figure 1 Distribution of Patients With a Positive Genetic Diagnosis According to the Spinal Muscular Atrophy (SMA) Subtype



Thirty-two patients with a genetic diagnosis achieved through neuropathy gene panel (n = 19: *DYNC1H1* x6, *BICD2* x4, *TRPV4* x4, *HSPB1*, *VCP*, *VRK1*, *DNAJB2*, *MORC2*), individual gene sequencing (n = 8: *BICD2* x3, *AR* x2, *HSPB1*, *HEXB*, and *COL6A3*), and other neuromuscular NGS (n = 5: *BICD2* x2, *DYNC1H1*, *VCP*, and *ASAH1*). Neg: negative; NGS: next-generation sequencing; P-SMA: proximal SMA; PD-SMA: proximo-distal SMA; SP-SMA: scapulo-peroneal SMA.

than 5 years, 3/7 patients had neonatal arthrogryposis, and 3/7 learning difficulties since childhood. DTRs were mostly abolished (6/7). Only 1 patient had moderate respiratory insufficiency in the context of obesity. Of the 4 patients who were followed up, none had lost ambulation. All patients had a pure motor neuropathy on ENMG.

- TRPV4 (4 patients from 3 families). Variants: c.557G>A (n = 2) and c.694C>T (n = 2), all previously reported. These patients showed predominantly a SP-SMA (3/4), although 1 patient presented with a PD-SMA without *scapula alata*. Although onset of the first progressive symptom was in adult age (18–53 years), 2 patients had a personal history of arthrogryposis and one of scoliosis and *pes cavus*. DTRs were always abolished. Three patients had some degree of respiratory insufficiency, none requiring ventilation. Three of these patients also had vocal cord paralysis with hypophonia. One patient showed a minimal reduction of normal SNAP on ENMG. All but 1 patient remain ambulant, the latter losing ambulation at age 5 years (before the onset of weakness) because of arthrogryposis.

Other Non-5q SMA-Associated Genes

Other variants in genes implicated in HMNs were found. In 2 SP-SMA patients with lower-limb distal involvement, *scapula*

alata, and a disease onset in adult life, *HSPB1* variants were found (AD transmission). Two patients (SP-SMA and P-SMA) carried variants in the *VCP* and one (SP-SMA) in the *VRK1* gene (both AD), all 3 with an adult disease onset. A previously reported patient with an adult-onset PD-SMA carried a variant in the *MORC2* gene.¹⁸ One patient with an infantile-onset PD-SMA carried a homozygous pathogenic variant in the *DNAJB2* gene (*AR*). The specific genetic variants can be seen in eTable 1 (links.lww.com/NXG/A614).

It is important that 2 patients with a PD-SMA phenotype without clinical nor ENMG sensory involvement and an onset of symptoms in their late 20s carried CAG repeat expansions in the *AR* gene (AD transmission). None of these patients had bulbar symptoms, and only one had very high CK levels (2000U/L) and gynecomastia.

Furthermore, 2 patients carried biallelic variants (*AR* transmission) in genes implicated in lysosomal storage disorders (LSD). One patient carries 2 new compound heterozygous pathogenic variants in the *HEXB* gene (c.881A>G and c.1082+2T>C), compatible with a diagnosis of Sandhoff disease. This patient presented with a pure P-SMA, an onset of symptoms at 11 years, and no other neurologic manifestation of the disease (cerebellar, extrapyramidal, cognitive, or

Table 2 Predictors of a Positive Genetic Diagnosis

| | Positive genetic diagnosis (n = 32), n (%) | Negative genetic diagnosis (n = 39), n (%) | p Value ^a | OR (95% CI), ^b p Value |
|-------------------------------|---|---|----------------------|-----------------------------------|
| Early onset (0–17 y) | 23 (71.9) | 13 (33.3) | 0.001 | 0.5 (0.14 to 1.3) 0.016 |
| Positive family history (yes) | 18 (56.3) | 10 (25.6) | 0.009 | 3.9 (0.13 to 2.6) 0.030 |
| Contractures (yes) | 10 (31.3)* | 2 (5.1) | 0.008 | 3.7 (–0.5 to 3.1) 0.144 |

Abbreviations: CI = confidence interval; OR = odds ratio; SP-SMA = scapulo-peroneal SMA.

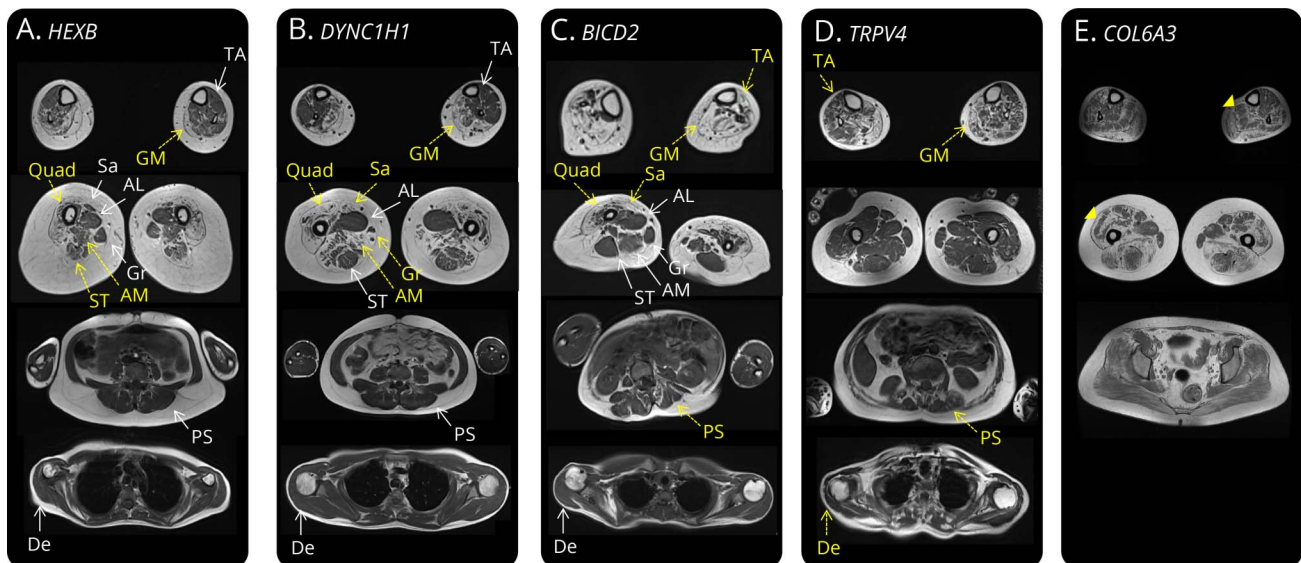
Variables significantly associated with a positive genetic diagnosis in a univariate χ^2 test (a) were tested in a multivariate logistic regression model (b). The multivariate model was statistically significant ($p = 0.001$), and the performance diagnostic showed an area under the curve (AUC) of 0.801.

* Five patients had arthrogryposis.

psychiatric); DTRs were normal, and there was no upper limb tremor. ENMG was compatible with a motor neuronopathy. Importantly, the patient was found to carry a VUS in the *DYNC1H1* gene, but the muscle MRI pattern did not correspond to the one previously described in *DYNC1H1*-associated SMA (Figure 2),¹⁹ so a leukocyte hexosaminidase analysis was performed, finding deficient A and B enzymatic activities. A second patient which we have previously reported²⁰ carries compound heterozygous variants in the *ASAH1* gene (c.77C>G and c.125+1G>A) and has a deficient leukocyte ceramidase activity. This patient presented with a pure P-SMA, an onset of symptoms at 19 years and again no other neurologic manifestation of the disease (epilepsy). The patient had scoliosis and club feet, upper

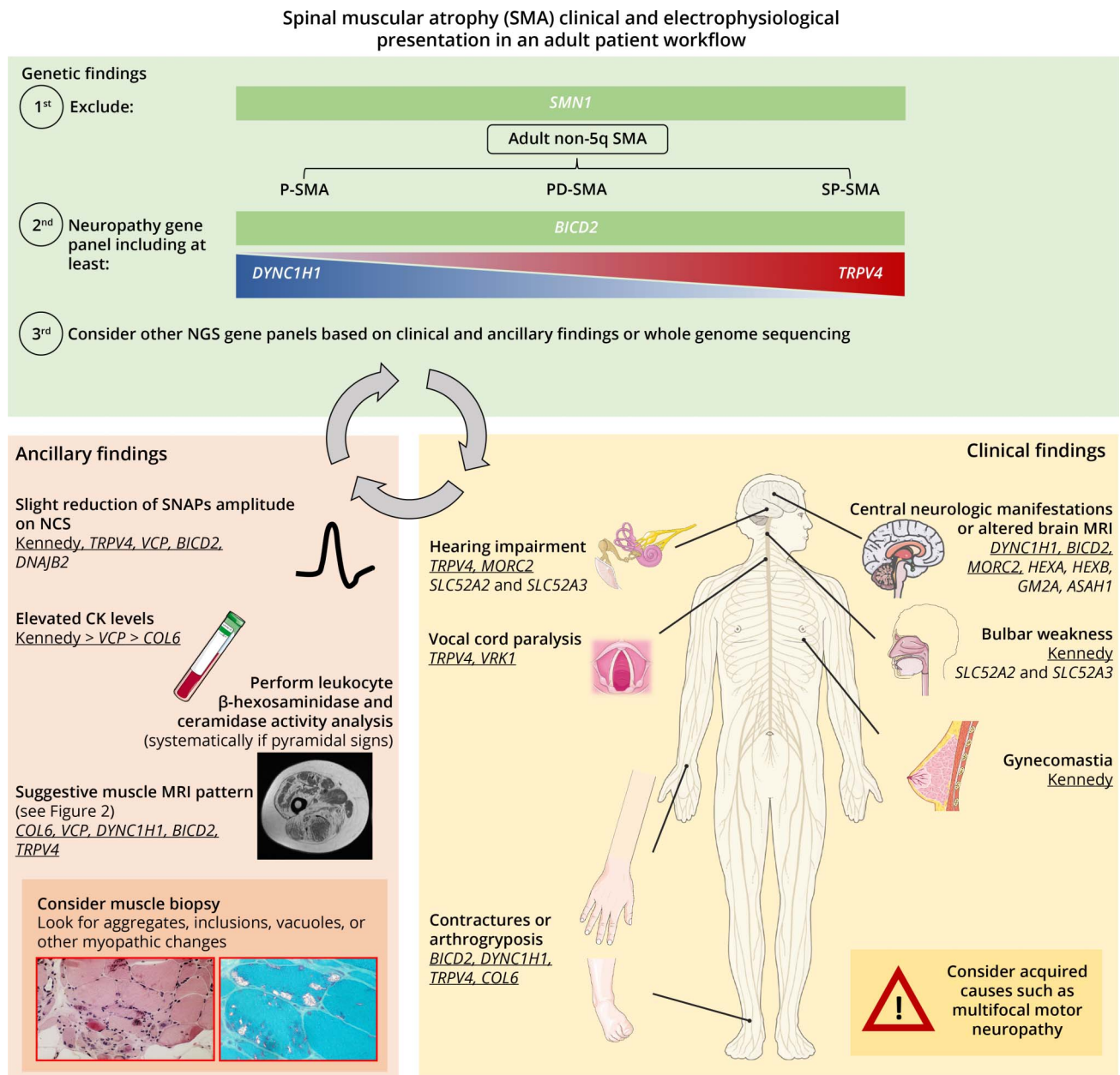
limb tremor, and retained DTRs. ENMG was compatible with a motor neuronopathy.

Finally, in 1 patient with a PD-SMA, retained DTRs, contractures in all 4 limbs since early childhood, and a neurogenic ENMG and muscle biopsy with no evidence of myopathy, who had no genetic diagnosis (after a 102 gene neuropathy panel, *SMN1* deletion, and hexosaminidase analysis); a characteristic “sandwich” sign on muscle MRI, typical of collagenopathies, was observed (Figure 2).²¹ Direct sequencing of the *COL6A3* gene found a previously unreported likely pathogenic homozygous c.5867A>G variant as well as a previously described heterozygous missense variant (c.7447A>G), which probably acts as a modulator of

Figure 2 Differing MRI Muscle Involvement in Non-5q Spinal Muscular Atrophy (SMA) Patients

T1-weighted MRI sequences of 5 patients with variants in the 3 most frequent non-5q SMA genes (*DYNC1H1*, *BICD2*, and *TRPV4*) as well as in Sandhoff disease (*HEXB*) and *COL6A3*. Muscles with yellow arrows had marked fat replacement; muscles with white arrows were completely or partially spared. (A) Quadriceps, semitendinosus, and adductor magnus muscle degeneration in a patient with Sandhoff disease presenting as P-SMA. (B) The previously described quadriceps muscle degeneration with adductor longus and semitendinosus sparing in a patient with *DYNC1H1* PD-SMA.¹⁹ In this case, sartorius and gracilis muscle fat replacement is also observed. (C) Quadriceps and sartorius muscle degeneration with more important lower limb involvement in a patient with *BICD2* PD-SMA. Notice some degree of paraspinal muscle atrophy and important scoliosis. (D) Tibialis anterior and gastrocnemius medialis involvement with sparing of muscles of the thigh and paraspinal and deltoid muscle degeneration in a patient with *TRPV4* SP-SMA. (E) Finally, the characteristic muscle fat replacement in a “sandwich pattern” of soleus and quadriceps muscles (yellow arrowheads) in a patient with *COL6A3* mutations and a pure SMA phenotype. AL = adductor longus; AM = adductor magnus; De = deltoid; GM = gastrocnemius medialis; Gr = gracilis; PS = paraspinal; Quad = quadriceps; Sa = sartorius; ST = semitendinosus; TA = tibialis anterior.

Figure 3 Flow Diagram With the Recommended Diagnostic Approach in Non-5q Spinal Muscular Atrophy (SMA) Patients



All 3 different sections (genetic, clinical, and ancillary findings) are critical for a correct final diagnosis. Findings in each section should be confronted to findings in the other 2 sections before establishing a definitive diagnosis. Genes underscored show an autosomal dominant (AD) transmission while the rest show an autosomal recessive (AR) transmission, with the exception of the *COL6* genes (*COL6A1, COL6A2, COL6A3*), which may be both AD and AR. The Figure was partly generated using Servier Medical Art, provided by Servier, licensed under a Creative Commons Attribution 3.0 unported license. CK = creatine kinase; NCS = nerve conduction studies; NGS = next-generation sequencing; SNAP = sensory nerve action potential.

the disease.²² A sibling, who presents a similar clinical phenotype and a predominant neurogenic ENMG and muscle biopsy, carried the same variants. Their mother was heterozygous for the c.5867A>G variant, and the father was deceased.

Discussion

We have described the clinical and genetic spectrum of non-5q SMA, whether P-SMA, PD-SMA or SP-SMA, in a large

series of patients without the *SMN1* deletion. Furthermore, this study provides evidence that the diagnostic yield of neuropathy NGS gene panels in non-5q SMA is still low (almost one-third of patients were diagnosed) but differs across the different clinical presentations.

This study first confirms that variants in 3 genes, *BICD2*, *DYNC1H1*, and *TRPV4*, account for almost two-thirds of genetically confirmed non-5q SMA patients. Nevertheless, each gene shows a distinctive clinical presentation, and while

both *BICD2* and *DYNC1H1* patients had an early disease onset and some degree of distal weakness, *BICD2* patients had more frequently *scapula alata* and preserved or brisk DTRs, while *DYNC1H1* patients had more frequently orthopedic manifestations and learning difficulties. *TRPV4* patients present most frequently with SP-SMA and even though orthopedic manifestations since childhood are frequent, onset of motor weakness occurs in adult life.

Variants in 8 additional genes were found and account for a significant genetic heterogeneity in non-5q SMA, with multiple possible pathophysiologic mechanisms. These genes can overlap with other nosological entities such as CMT and dHMN (*HSBP1*, *DNAJB2*, *MORC2*),^{2,23} Kennedy disease (*AR*),²⁴ myopathies (*VCP*, *COL6A3*), ALS (*VCP*),^{25,26} LSDs (*ASAHI* and *HEXB*),²⁷ and other developmental disorders (*VRK1*, *MORC2*).²⁸⁻³¹

Non-5q SMA is also a clinically heterogeneous disorder, with different ages at onset of symptoms, patterns of muscle weakness, and associated signs such as abnormal DTRs or orthopedic manifestations. It is important that these clinical features affect the performance of genetic studies. While an earlier onset and a family history of the disease or consanguinity are independent predictors of a positive genetic diagnosis, contractures and/or arthrogyrosis as well as SP-SMA phenotype also seem to increase the likelihood of a definitive genetic diagnosis. A previous genetic-focused study testing the diagnostic yield of a panel of 62 non-5q SMA genes vs a broad 479 neuromuscular gene panel⁶ reported a diagnostic yield of 13% for the non-5q SMA panel vs a 33% for the broad neuromuscular panel; the latter comparable with the performance of neuropathy gene panels in our study. Therefore, neuropathy gene panels (coupled to in-depth phenotyping) seem an adequate strategy in the genetic diagnosis of non-5q SMA patients.

With the increase throughput capability of NGS technologies, comes an increase in the number of VUS, which require an important amount of time and expertise to interpret the variants and avoid misclassification. An example of this is the patient with Sandhoff disease, who was previously found to carry a VUS in the *DYNC1H1* gene. In this case, careful analysis of the muscle MRI prompted the search for other alternative diagnosis. This further shows that ancillary testing, and in particular muscle MRI, is crucial to increase the proportion of correctly diagnosed non-5q SMA patients in the advent of NGS techniques. Muscle MRI can not only help discriminate between neuropathy and myopathy³² but also show particular patterns of muscle fat replacement described in *DYNC1H1* and *BICD2* patients^{17,19} or characteristic MRI signs, like the “sandwich” sign in collagenopathies.²¹

Nevertheless, a significant proportion of non-5q SMA patients do not have a definitive genetic diagnosis. Strategies that may increase the number of genetically diagnosed

patients can be extracted from the results of this study. First, Kennedy disease should be more frequently searched for, even in the absence of other features such as gynecomastia or sensory involvement, which are not systematic.³³ Second, the performance of NGS panels in non-5q SMA could be increased if other neurometabolic disease-associated genes were to be included. This is especially critical because some motor neuron mimics such as *SLC52A2* and *SLC52A3* genes show a dramatic clinical response to riboflavin³⁴ and that therapeutic trials are ongoing in LSDs such as Sandhoff disease (NCT04221451). Third, the frontier between neuropathies and myopathies is increasingly porous. A well-known example of this is the *VCP* gene, with varied neurologic presentations (myopathy, neuropathy, or ALS) for most genetic variants.³⁵ Furthermore, almost half of the patients showed moderate elevations of CK levels, a feature which is fairly common in diseases involving motor nerve damage.³⁶ Unexpectedly, 1 patient with a pure SMA (based on repeated ENMG and muscle biopsy findings) but contractures and a muscle MRI compatible with a collagenopathy was found to carry likely pathogenic variants in the *COL6A3* gene. Indeed, collagen VI animal models show alterations in myelination and axonal growth,³⁷ and in humans, a previous *COL6A3* patient with myopathic and neuropathic changes on ENMG has been described.²² However, this is a report of a patient with pure SMA probably because of mutations in the *COL6A3* gene. Finally, WGS, and particularly long-read WGS, may be a more suitable approach than WES in genetically unconfirmed non-5q SMA to search for intronic variants as well as structural variants (SV) because these may account for significant part of the missing heritability in non-5q SMA, as is the case in other neuromuscular diseases.³⁸⁻⁴⁰ In Figure 3, the reader can find our recommended diagnostic workflow for an adult patient with a non-5q SMA.

In summary, non-5q SMA is a clinically and genetically heterogeneous disorder, and neuropathy NGS gene panels achieve a genetic diagnosis in about one-third of patients followed in specialized neuromuscular reference centers, with an earlier onset and a family history of the disease being an independent predictor of a positive genetic diagnosis. Two-thirds of these patients carry pathogenic variants in 3 genes (*BICD2*, *DYNC1H1*, and *TRPV4*); however, non-5q SMA may also be caused by other overlapping genes, whose identification is laborious if not included in the NGS gene panel. Unfortunately, more than half of the patients do not have a precise genetic diagnosis despite gene panels and WES. Thus, there is an unmet need to cluster these rare patients in view of the identification of new common genes through higher throughput NGS techniques.

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Appendix Authors

| Name | Location | Contribution |
|-------------------------------------|--|--|
| Gorka Fernández-Eulate, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Institut de Myologie, Pitié-Salpêtrière Hospital, Paris, France | Drafting/revision of the manuscript for content, including medical writing for content; major role in the acquisition of data; study concept or design; analysis or interpretation of data |
| Julian Theuriet, MD | Electromyography and Neuromuscular Department, Hospices Civils de Lyon, France | Major role in the acquisition of data |
| Christopher J. Record, MD | Centre for Neuromuscular Diseases, UCL Queen Square Institute of Neurology, London, United Kingdom | Major role in the acquisition of data |
| Giorgia Querin, MD, PhD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Institut de Myologie, Pitié-Salpêtrière Hospital, Paris, France | Analysis or interpretation of data |
| Marion Masingue, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Institut de Myologie, Pitié-Salpêtrière Hospital, Paris, France | Drafting/revision of the manuscript for content, including medical writing for content |
| Sarah Leonard-Louis, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Institut de Myologie, Pitié-Salpêtrière Hospital, Paris, France | Major role in the acquisition of data |
| Anthony Behin, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Institut de Myologie, Pitié-Salpêtrière Hospital, Paris, France | Major role in the acquisition of data |
| Nadine Le Forestier, MD | Neurology department, Pitié-Salpêtrière Hospital, Paris, France | Major role in the acquisition of data |
| Antoine Pegat, MD | Electromyography and Neuromuscular Department, Hospices Civils de Lyon, France | Major role in the acquisition of data |
| Maud Michaud, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Central Nancy University Hospital, France | Major role in the acquisition of data |
| Jean-Baptiste Chanson, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Strasbourg University Hospitals, France | Major role in the acquisition of data |
| Aleksandra Nadaj-Pakleza, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Strasbourg University Hospitals, France | Major role in the acquisition of data |

Appendix (continued)

| Name | Location | Contribution |
|---------------------------------------|--|---|
| Celine Tard, MD, PhD | Nord/Est/Ile-de-France Neuromuscular Reference Center, U1172, Lille University Hospital, France | Major role in the acquisition of data |
| Anne-Laure Bedat-Millet, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Rouen University Hospital, France | Major role in the acquisition of data |
| Guilhem Sole, MD | Neuromuscular Reference Center 'AOC', Bordeaux University Hospitals (Pellegrin Hospital), France | Major role in the acquisition of data |
| Marco Spinazzi, MD | Neuromuscular Reference Center, Angers University Hospital, France | Major role in the acquisition of data |
| Emmanuelle Salort-Campana, MD | Neuromuscular and ALS Reference Center, La Timone University Hospital, Marseille, France | Major role in the acquisition of data |
| Andoni Echaniz-Laguna, MD, PhD | French National Center for Rare Neuropathies, Neurology department, Bicêtre University Hospital, INSERM U1195, Paris-Saclay University, Le Kremlin Bicêtre, France | Drafting/revision of the manuscript for content, including medical writing for content |
| Vianney Poinignon, PharmD | Molecular genetics lab, Bicêtre University Hospital, Le Kremlin Bicêtre, France | Major role in the acquisition of data |
| Philippe Latour, PhD | Center for Biology - East, Neurological hereditary disorders unit, Hospices Civils de Lyon, France | Drafting/revision of the manuscript for content, including medical writing for content; major role in the acquisition of data |
| Mary M. Reilly, MD | Centre for Neuromuscular Diseases, UCL Queen Square Institute of Neurology, London, United Kingdom | Drafting/revision of the manuscript for content, including medical writing for content |
| Francoise Bouhour, MD | Electromyography and Neuromuscular Department, Hospices Civils de Lyon, France | Drafting/revision of the manuscript for content, including medical writing for content |
| Tanya Stojkovic, MD | Nord/Est/Ile-de-France Neuromuscular Reference Center, Institut de Myologie, Pitié-Salpêtrière Hospital, Paris, France | Drafting/revision of the manuscript for content, including medical writing for content; study concept or design |

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Phenotype Presentation and Molecular Diagnostic Yield in Non-5q Spinal Muscular Atrophy

Gorka Fernández-Eulate, Julian Theuriet, Christopher J. Record, et al.

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