



Immunopathological manifestations in Kabuki syndrome: a registry study of 177 individuals

A full list of authors and affiliations appears at the end of the paper.

Purpose: Kabuki syndrome (KS) (OMIM 147920 and 300867) is a rare genetic disorder characterized by specific facial features, intellectual disability, and various malformations. Immunopathological manifestations seem prevalent and increase the morbimortality. To assess the frequency and severity of the manifestations, we measured the prevalence of immunopathological manifestations as well as genotype–phenotype correlations in KS individuals from a registry.

Methods: Data were for 177 KS individuals with *KDM6A* or *KMT2D* pathogenic variants. Questionnaires to clinicians were used to assess the presence of immunodeficiency and autoimmune diseases both on a clinical and biological basis.

Results: Overall, 44.1% (78/177) and 58.2% (46/79) of KS individuals exhibited infection susceptibility and hypogammaglobulinemia, respectively; 13.6% (24/177) had autoimmune disease (AID; 25.6% [11/43] in adults), 5.6% (10/177) with ≥ 2 AID

manifestations. The most frequent AID manifestations were immune thrombocytopenic purpura (7.3% [13/177]) and autoimmune hemolytic anemia (4.0% [7/177]). Among nonhematological manifestations, vitiligo was frequent. Immune thrombocytopenic purpura was frequent with missense versus other types of variants ($p = 0.027$).

Conclusion: The high prevalence of immunopathological manifestations in KS demonstrates the importance of systematic screening and efficient preventive management of these treatable and sometimes life-threatening conditions.

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INTRODUCTION

Kabuki syndrome (KS, OMIM 147920 and 300867) is a rare genetic disorder with an estimated prevalence of 1/32,000 people. KS is characterized by specific facial features, intellectual disability, skeletal anomalies, short stature, and dermatoglyphic anomalies, including persistence of fetal fingertip pads.¹ KS is due to pathogenic variants in lysine-specific methyltransferase 2D (*KMT2D*, OMIM 602113)² or lysine-specific demethylase 6A (*KDM6A*, OMIM 300128, an X-linked gene that escape partially to X inactivation).³ Those genes are part of the complex of proteins associated with Set1 (COMPASS), which is involved in chromatin remodeling by modifying epigenetic marks on histones.⁴ For 56% to 75% of KS individuals, the major gene involved is *KMT2D*.⁵ *KDM6A* is mutated in about 5% of KS individuals.⁶

Recurrent infections, mainly otitis media, abnormal immunoglobulin secretion,⁷ decreased memory cells, and poor vaccine response⁸ have been reported in isolated or small series of patients with KS but few authors have emphasized the potential severity of immune complication such as hypogammaglobulinemia and granulomatous–lymphocytic interstitial lung disease (GLILD).⁹ Autoimmune diseases

(AIDs have also been reported; the most frequent manifestations include immune thrombocytopenic purpura¹⁰ with or without concurrent hemolytic anemia or autoimmune neutropenia,¹¹ followed by vitiligo,¹² autoimmune thyroiditis,¹³ diabetes type 1,¹⁴ Crohn disease,¹⁵ and membrane glomerulonephritis type 3.¹⁶ To our knowledge those immune manifestations have never been quantified by epidemiological tools. Actual guidelines for monitoring immune defects of KS are based on single-patient data or small series experience.

By taking advantage of a French collaborative survey on KS, we report a series of 177 individuals with KS due to *KMT2D* or *KDM6A* pathogenic variants; describe the immune phenotype, emphasizing differences by sex, age, gene, and type of pathogenic variant; and propose recommendations for immunologic survey in KS.

MATERIALS AND METHODS

Participants

We collected data for 177 individuals from a national registry (PHRC AOM-07-090 and additional individuals coordinated by Montpellier University Hospital; ClinicalTrials.gov identifier: NCT01314534). All patients gave their written informed

Correspondence: David Genevieve (d-genevieve@chu-montpellier.fr)

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consent before inclusion. Ethical approval was obtained from CPP2007-12-01, CPP Ile de France 2. All patients had a disease-causing variant in *KMT2D* or *KDM6A*.

This study relies in part on prospective data from the registry and retrospective data collected by a questionnaire addressed to 30 different French genetic centers that followed KS individuals from this registry, in coordination with the hematologist from the corresponding center. For subgroup analysis, individuals were divided into three age groups. From an endocrinological standpoint supporting hormonal changes during puberty, children were analyzed at <9 and 10 years old for females and males. Teenagers were analyzed at 9–17 and 10–18 years old for females and males, respectively, and adults at >17 and >18 years old for females and males, respectively.

Immunodeficiency

To assess immunodeficiency, data were extracted from medical records and the PHRC database. After data extraction, the clinician was asked by questionnaire if the individual had any recurrent or severe infections, especially ear, nose, and throat (ENT), pulmonary, or gastrointestinal infections. Recurrent infections are defined as three or more infections (sinusitis, otitis, bronchitis, pneumonia) in one year. Severe infections include those with failure to respond to oral antibiotics and/or the need for intravenous antibiotics or hospitalization. Susceptibility to infection was considered if the individual matched one of the criteria mentioned above. Biological values for complete blood count and quantitative serum immunoglobulin tests extracted from records were collected. A decrease in immunoglobulin levels of ≥ 2 standard deviations according to age was considered hypogammaglobulinemia. In cases of abnormal levels, we asked for results regarding immunoglobulin subclass, postimmunization antibodies, complement levels, and lymphocyte subset counts. Common variable immunodeficiency disorders (CVIDs) were defined by the combination of recurrent, severe, or unusual infections and hypogammaglobulinemia: a marked decreased (at least 2 standard deviations below the mean for age) in serum IgG and IgA.¹⁷

Autoimmunity

For autoimmune diseases, medical history of immune thrombocytopenic purpura (ITP), autoimmune hemolytic anemia (AIHA), thyroiditis, diabetes, vitiligo, etc. were specifically assessed. ITP was defined according to international criteria.¹⁸ AIHA was characterized by hemoglobin level <11 g/dL, a positive direct antiglobulin test, and at least one biological hemolysis criteria (reticulocyte count $>120 \times 10^9/L$, haptoglobin level <10 mg/dL, or total bilirubin level >1 mg/dL). Vitiligo was considered if the individual had a depigmentation disorder of the skin without other etiology secondary to infection or treatment, for example. The diagnosis of autoimmune thyroiditis was based on increased level of antithyroid peroxidase (TPO) antibody (≥ 30 kU/l) or thyroglobulin (Tg).¹³

Variant screening and validation

Genomic DNA was isolated from peripheral EDTA-anticoagulated whole blood by using QIAamp DNA Blood Kits (Qiagen, Valencia, CA, USA). The entire coding sequence and intronic boundaries of *KMT2D* (NG_027827.1) and *KDM6A* (NG_016260.1) were screened for variants by Sanger sequencing of *KMT2D* according to routine diagnostic protocols or by next-generation sequencing (NGS) on a custom capture-based panel including *KMT2D* and *KDM6A*, then confirmed by Sanger sequencing. Obtained sequences were aligned against the hg19 (genome build: GRCh37) genome reference sequences from the National Center for Biotechnology Information (NCBI). Molecular results were analyzed by using the transcript NM_003482.3 for *KMT2D* and NM_021140.3 for *KDM6A*, Alamut Visual (Interactive Biosoftware, Rouen, France) and Ensembl website (<http://grch37.ensembl.org/>). Part of the genetic variants described in this registry were published previously.¹⁹ We consider a missense variant as likely pathogenic or pathogenic (Supplementary Table 3) when it arises de novo and we classified it as likely pathogenic or pathogenic on the VarSome website (<https://varsome.com/>).²⁰

Statistical analysis

Descriptive statistics were reported: quantitative data are expressed with mean \pm SD and range, and qualitative data by number (%). Statistical association between variables was tested by chi-square or Fisher's exact test otherwise (full data in Table 1). In our sample, females were older than males. Therefore, when necessary, we tested the confounding effect of these variables by multivariate logistic regression. Results of logistic regression are presented when they differ from results of univariate analysis.

RESULTS

Our series of 177 KS individuals included 89 males. Full data are available in Supplementary Table 2; 68 individuals were children (<9 or <10 years old) at the time of their last medical appointment; 66 were teenagers (<17 or <18 years old), and 43 were adults. The mean (SD) age at the time of the last medical appointment was 11.7 ± 10 (range 0–66) years. Five individuals died, three from immunopathological complications: acute bronchitis infection for two patients, hypovolemic shock consecutive to gastroenteritis and acute hemolysis at age 24 years due to chronic Evans syndrome for another. We identified *KMT2D* and *KDM6A* pathogenic variants in 162 and 15 KS individuals, respectively. For the type of pathogenic variant in *KMT2D*, we identified 67 nonsense, 50 frameshift, 35 missense, and 10 splice variants. For *KDM6A*, we identified 7 splice, 6 nonsense, 1 frameshift and 1 missense variants.

The prevalence of immunopathological manifestations in this registry is shown in Fig. 1, and the prevalence among subgroups is listed in Table 1. Overall, 78/177 (44.1%) individuals showed susceptibility to infections (Fig. 1). The most frequent infection was repeated ENT infection (47/177). Infections of the lower respiratory tract (6/177) and urinary tract (6/177) were less

Table 1 Comparison of immune phenotype by age, sex, gene, variant type

	Age			Sex		Gene			Variant type		
	Children	Teens	Adults	Female	Male	KDM6A	KMT2D	Missense	Others		
	N = 68	N = 66	N = 43	N = 88	N = 89	N = 15	N = 162	N = 35	N = 142	p	p
Infection susceptibility											
No	42 (62)	35 (53)	22 (51)	47 (53)	52 (58)	10 (67)	89 (55)	19 (54)	80 (56)	0.381	0.977
Yes	26 (38)	31 (47)	21 (49)	41 (47)	37 (42)	5 (33)	73 (45)	16 (44)	62 (44)		
Hypogammaglobulinemia											
No	14 (21)	13 (20)	6 (14)	15 (17)	18 (20)	2 (13)	31 (19)	7 (20)	26 (18)	1 ^a	0.650
Yes	12 (18)	19 (29)	15 (35)	23 (26)	23 (26)	2 (13)	44 (27)	13 (36)	33 (23)		
Autoimmune disease											
No	64 (94)	57 (86)	32 (74)	75 (85)	78 (88)	14 (93)	139 (86)	28 (80)	125 (88)	0.639	0.333
Yes	4 (6)	9 (14)	11 (26)	13 (15)	11 (12)	1 (7)	23 (14)	7 (19)	17 (12)		
Immune thrombocytopenic purpura											
No	67 (99)	61 (92)	36 (84)	81 (92)	83 (93)	15 (100)	149 (92)	29 (83)	135 (95)	0.757	0.024 ^a
Yes	1 (1)	5 (8)	7 (16)	7 (8)	6 (7)	0 (0)	13 (8)	6 (17)	7 (5)		
Autoimmune hemolytic anemia											
No	67 (99)	63 (95)	40 (93)	83 (94)	87 (98)	15 (100)	155 (96)	33 (94)	137 (96)	0.278 ^a	0.604 ^a
Yes	1 (1)	3 (5)	3 (7)	5 (6)	2 (2)	0 (0)	7 (4)	2 (6)	5 (4)		
Thyroiditis											
No	68 (100)	66 (100)	41 (95)	86 (98)	89 (100)	15 (100)	160 (99)	35 (100)	140 (99)	0.46 ^a	1 ^a
Yes	0 (0)	0 (0)	2 (5)	2 (2)	0 (0)	0 (0)	2 (1)	0 (0)	2 (1)		
Vitiligo											
No	66 (97)	62 (94)	40 (93)	85 (97)	83 (93)	14 (93)	154 (95)	34 (97)	134 (94)	0.496 ^a	0.691 ^a
Yes	2 (3)	4 (6)	3 (7)	3 (3)	6 (7)	1 (7)	8 (5)	1 (3)	8 (6)		
Number of autoimmune diseases co-occurring in an individual											
0	64 (94)	57 (86)	32 (74)	75 (85)	78 (88)	14 (93)	139 (86)	28 (80)	125 (88)	0.931 ^a	0.315 ^a
1	3 (4)	5 (8)	6 (14)	7 (8)	7 (8)	1 (7)	13 (8)	5 (14)	9 (6)	1 ^a	
2	0 (0)	3 (5)	4 (9)	4 (5)	3 (3)	0 (0)	7 (4)	1 (3)	6 (4)		
3	1 (1)	1 (2)	1 (2)	2 (2)	1 (1)	0 (0)	3 (2)	1 (3)	2 (1)		

Values are N (column percentage).

^aP values by Fisher's exact test; otherwise, by chi-square test.

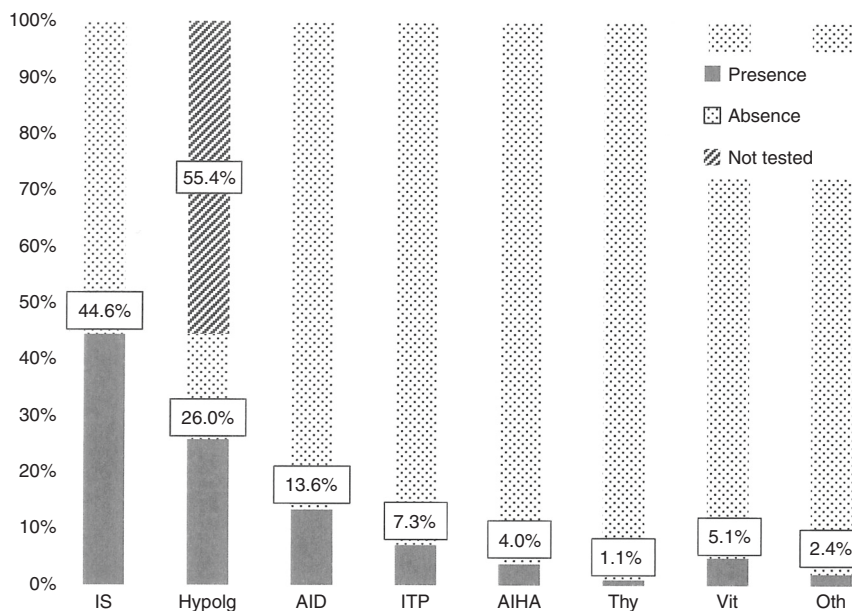


Fig. 1 Prevalence of immunodeficiency and autoimmune disorders in Kabuki syndrome (KS). Presence denotes presence of one of the following items: AID autoimmune disease, AIHA autoimmune hemolytic anemia, HypoIg hypogammaglobulinemia, IS infection susceptibility, ITP immune thrombocytopenic purpura, Oth other, Thy thyroiditis, Vit vitiligo. Absence denotes absence of the event. Not reported data were not available.

frequent. Bones, skin, gastrointestinal tract, cerebral, and ophthalmologic infections were occasionally reported. Two individuals had lung granulomas associated with hypogammaglobulinemia, for granulomatous-lymphocytic lung disease.

Repeated complete blood count was normal in all individuals from this registry. Quantitative serum immunoglobulin test was performed in 79/177 (44.6%) patients; 46/79 (58.2%) individuals had defects in secretion of IgA, IgM, or IgG (Fig. 1). Combined hypogammaglobulinemia was observed in 28/79 (35.4%) individuals. CVID was observed in half (39/79, 49.4%), whereas 22/79 (27.8%) individuals were without hypogammaglobulinemia and infection susceptibility (Supplementary Table 2). Nonadequate protective postimmunization antibodies of antitetanic titers were found in 6/11 individuals with available data. We could not collect lymphocyte subset counts.

In total, 24/177 (13.6%) individuals had AID (Fig. 1); 10/177 (5.6%) had ≥ 2 autoimmune manifestations. ITP was the most frequent autoimmune complication (7.3%, 13/177); 12/13 (92%) individuals had the chronic form lasting more than 12 months. The second hematologic autoimmune manifestation was AIHA, including 7/177 (4.0%) individuals (Table 1); 6/7 individuals with AIHA had simultaneous or dissociated ITP (Evans syndrome). In total, 9/177 (5.1%) individuals had vitiligo (Fig. 1). Two adult females had autoimmune thyroiditis (1.1%). Finally, one individual had diabetes mellitus type 1, one had inflammatory bowel disease, one had antiphospholipid syndrome, and one had juvenile idiopathic arthritis (oligoarthritic form).

In total, 11/39 (28%) individuals with CVID had an AID as compared with only 2/22 (9%) without CVID. The rate of AID was significantly increased for individuals with CVID (p

< 0.1 , Fisher's exact test), whereas association between HypoIg and AID was not significant ($p = 0.3$).

The prevalence of immunopathological manifestations was similar on subgroup analysis by age, gene (*KMT2D* or *KDM6A*), variant subtype, and sex ($p > 0.05$; Table 1). AID manifestations increased with age ($p = 0.013$, Table 1). Fewer than 4/68 (5.9%) children had AID manifestations as compared with 11/43 (25.6%) adults (Supplementary Fig. 2).

Genotype-phenotype correlations showed missense variants frequent in individuals with ITP ($p = 0.027$, Table 1). Also, autoimmune manifestations less frequently affected individuals with *KDM6A* than *KMT2D* variants, although not significantly (6.7% vs. 14.2%, $p = 0.697$; Table 1). The position of the pathogenic variants on the coding sequence was not associated with immune phenotype.

DISCUSSION

In this nationwide collaborative study, almost half of the 177 KS individuals analyzed had features of immunodeficiency and 13% had an AID. Prevalence of AID manifestations increased with age, with one quarter of adults showing at least one disease. Several authors had previously reported on CVID or autoimmunity in KS, but no large registry was studied. We focused on immunopathological manifestations in KS and described their prevalence, detailed features, and any genotype-phenotype correlations. We found that immunopathological manifestations were frequent and also seemed severe in KS.

Our findings may be understood by the loss of function of *KMT2D* and *KDM6A*. Indeed, *KMT2D* and *KDM6A* encode for proteins acting in a same biological complex called COMPASS. This complex allows for open-state chromatin by

adding an activator trimethylation on the lysine 4 of histone 3 and removing the inhibitor trimethylation on the lysine 27.²¹ This situation translates epigenetically into a positive regulation of targeted gene transcription.²² KS individuals feature defective B-lymphocyte terminal differentiation, which probably explains the hypogammaglobulinemia. The literature describes significantly reduced number of total memory (CD27⁺) and class-switched memory B cells (IgM⁻) in Kabuki patients type 1 or without molecular study, compared with controls.²³ The COMPASS complex helps in the transcription of the AID complex gene.²⁴ This complex guides somatic hypermutation and class switch recombination, which is essential to B-cell differentiation, diversity, and efficiency.²⁴ Pathogenic variants in *KMT2D* and *KDM6A* may lead to a dysfunction in this process and could explain the immune deficits observed in people with KS. The COMPASS complex is also involved in the epigenetic regulation of *FOXP3*,²⁵ a key gene during the differentiation of naïve CD4⁺ T cells into T-regulatory cells, which contribute to the maintenance of peripheral tolerance.²⁶ Pathogenic variants in *KMT2D* or *KDM6A* may lead to *FOXP3* dysfunction and therefore a breakdown of T-cell tolerance, which could be the mechanism for the autoimmune phenotype observed in KS.

The strengths of our study are the large and multicentric setting and the retrospective collection of data that relied on genetics and hematoimmunological rare disease networks in our country as well as the genotype–phenotype study. The wide distribution of age in our sample disallows conducting a reliable longitudinal analysis regarding autoimmune diseases. Several biological data regarding immunodeficiency were lacking. Some centers had different follow-up management; therefore, our missing data are mainly due to the intercenter management strategies. However, when data were available, data for 77.2% individuals were concordant for results for susceptibility to infection and hypogammaglobulinemia and thus we could correctly assess CVID in individuals (49.4% [39/79] with CVID and 27.8% [22/79] without hypogammaglobulinemia and infection susceptibility).²⁷ In this series, only 79/177 (44.6%) individuals had a quantitative serum immunoglobulin test performed and registered in their medical record. This low percentage demonstrates that hypogammaglobulinemia is not always tracked by clinicians in France. Therefore, immunoglobulin supplementation is not implemented or not early enough. This situation increases the risk of intercurrent infectious disease, leading to avoidable hospitalizations in these fragile individuals and fosters rare but critical conditions such as GLILD. However, most of the clinical manifestations secondary to immune impairment were mild to moderate, but for the five deaths in the registry, three were from immunological features and highlight the real need for adequate monitoring of these features in KS individuals.

We identify a high prevalence of immune manifestations in KS: 25.6% of our adult cohort had autoimmune manifestations as compared with 7.6–9.4% in the general adult population.²⁸ The most frequent manifestations were isolated

or combined autoimmune cytopenia, ITP, AIHA, or Evans syndrome). Prevalence of ITP usually decreases during childhood in the general population,²⁹ but we observed the opposite in this series ($p = 0.011$; Table 1 and Supplementary Fig. 2). Also, 20% of general children with ITP have a chronic course lasting more than 12 months,³⁰ whereas 92% of our KS sample had a chronic form, with second-line treatment.

One individual died from an acute hemolysis as a result of Evans syndrome. This severe situation is described in individuals with AIHA in the general population,³¹ more often in adults. Therefore, autoimmunity in KS should be carefully monitored and especially from pediatric to adult care.

Vitiligo was ten times more frequent in our series than in the general population.³² The sex ratio was 2 (6 males to 3 females) in our sample but 0.84 in the general population.³³ We do not have an explanation for this discrepancy. The difference might not be significant because of the small number of individuals in this series.

We found thyroiditis in 2 (1.1%) adult females. In the literature, thyroiditis is usually described in adult females (94.5%).³⁴ The prevalence of thyroiditis in the general population is 0.008%³⁴ versus 1.1% in our registry. Here again, this data may be uncertain because of the small sample size.

On univariate analysis, women were more affected, but not significantly, by autoimmunity than were men, as observed in the general population.³⁴

The rate of any autoimmune disease in KS individuals with CVID was 28% versus 9% without immune manifestations. This observation is related to the fact that CVID is frequently associated with AID in the literature,³⁵ which led us to hypothesize that KS individuals with CVID should be supervised even more carefully for autoimmune manifestations. A larger sample of individuals is necessary to demonstrate an increased risk of AID in KS individuals with CVID.

For genotype–phenotype correlations, ITP was more frequent with missense than truncating variant carriage. However, we observed no difference of the distribution of missense pathogenic variants associated to ITP compared with other missense pathogenic variants at the exonic and protein level respectively (Supplementary Table 3 and Supplementary Fig. 3). Immune deficiency or autoimmune manifestations did not differ by type of gene (*KMT2D* or *KDM6A*). Autoimmune manifestations were fewer with *KDM6A* than *KMT2D* (6.7% vs. 14.2%), which is due to the younger median age of *KDM6A* than *KMT2D* individuals (8.1 vs. 13.7).

Patients with KS should have an assessment by an immunologist as soon as there is any doubt of dysimmunity, to avoid chronic diseases such as bronchiectasis or GLILD. The overall management of those patients should include clinical specific assessment, with a special attention to repeated ENT, bronchial, or digestive infection; lymphoproliferation; and symptoms of autoimmune diseases, mainly

cytopenia or thyroiditis. We recommend checking by complete blood cell counts, IgG IgA IgM levels, T, B and NK lymphocytes phenotyping all patients at diagnosis. This blood work should be repeated it in case of recurrent or unusual infections. The immunization schedule should follow standard national guidelines. Measurement of T3, T4, and thyroid-stimulating hormone (TSH) levels is also indicated in case of clinical hypothyroidism. Patients with KS should have an assessment by an immunologist as soon as there is any suspicion of immunopathological manifestations, to avoid chronic diseases such as bronchiectasis or GLILD and to discuss specific treatments.

Prompt and adapted curative anti-infectious treatments, prophylactic antibiotics, and immunoglobulin substitution are to be considered on a case-by-case basis, based on guidelines available for patients with primary immunodeficiency.³⁶

SUPPLEMENTARY INFORMATION

The online version of this article (<https://doi.org/10.1038/s41436-019-0623-x>) contains supplementary material, which is available to authorized users.

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DISCLOSURE

David Geneviève is a consultant for the Takeda Society. The society did not have any influence on the content of this report or its publication. The other authors declare no conflicts of interest.

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Henri Margot, MS¹, Guilaine Boursier, PharmD-PhD^{2,3}, Claire Duflos, MD⁴, Elodie Sanchez, MD^{2,3}, Jeanne Amiel, MD-PhD⁵, Jean-Christophe Andrau, PhD⁶, Stéphanie Arpin, MD⁷, Elise Brischoux-Boucher, MD⁸, Odile Boute, MD⁹, Lydie Burglen, MD-PhD¹⁰, Charlotte Caille, MS¹¹, Yline Capri, MD¹², Patrick Collignon, MD-PhD¹³, Solène Conrad, MS¹⁴, Valérie Cormier-Daire, MD-PhD⁵, Geoffroy Delplancq, MS⁸, Klaus Dieterich, MD-PhD¹⁵, Hélène Dollfus, MD-PhD¹⁶, Mélanie Fradin, MD¹⁷, Laurence Faivre, MD-PhD¹⁸, Helder Fernandes, PhD^{19,20,21}, Christine Francannet, MD²², Vincent Gatinois, PharmD-PhD^{2,3}, Marion Gerard, MD²³, Alice Goldenberg, MD-PhD²⁴, Jamal Ghoumid, MD-PhD⁹, Sarah Grotto, MD¹², Anne-Marie Guerrot, MD²⁴, Agnès Guichet, MD²⁵, Bertrand Isidor, MD-PhD¹⁴, Marie-Line Jacquemont, MD^{26,27}, Sophie Julia, MD²⁸, Philippe Khau Van Kien, MD-PhD²⁹, Marine Legendre, MD-PhD^{1,27}, K. H. Le Quan Sang, PhD³⁰, Bruno Leheup, MD-PhD³¹, Stanislas Lyonnet, MD-PhD⁵, Virginie Magry, MS^{10,22}, Sylvie Manouvrier, MD-PhD⁹, Dominique Martin, MD³², Godelieve Morel, MS³³, Arnold Munnich, MD-PhD⁵, Sophie Naudion, MD^{1,27}, Sylvie Odent, MD-PhD¹⁷, Laurence Perrin, MD¹², Florence Petit, MD-PhD⁹, Nicole Philip, MD-PhD³⁴, Marlène Rio, MD-PhD⁵, Julie Robbe, MS³⁴, Massimiliano Rossi, MD-PhD³⁵, Elisabeth Sarrazin, MD²⁷, Annick Toutain, MD-PhD⁷, Julien Van Gils, MD^{1,27,36}, Gabriella Vera, MS-PhD²⁴, Alain Verloes, MD-PhD¹², Sacha Weber, MD²³, Sandra Whalen, MD³⁷, Damien Sanlaville, MD-PhD³⁵, Didier Lacombe, MD-PhD^{1,27,36}, Nathalie Aladjidi, MD^{19,20,21} and David Geneviève, MD-PhD^{2,3}

¹Service de génétique médicale, CHU de Bordeaux, Bordeaux, France. ²Département de génétique médicale, Maladies rares et médecine personnalisée, CHU de Montpellier, Montpellier, France. ³INSERM U1183, Université de Montpellier, Montpellier, France. ⁴Département d'Information Médicale, CHU Montpellier, Montpellier, France. ⁵Fédération de Génétique et Institut Imagine, Hôpital Necker-Enfants Malades, AP-HP et INSERM UMR1163, Paris Descartes-Sorbonne Paris Cité University, Paris, France. ⁶Institut de Génétique Moléculaire de Montpellier (IGMM), Univ Montpellier, Montpellier, France. ⁷Service de génétique, CHU de Tours, UMR 1253, iBrain, Université de Tours, Inserm, Tours, France. ⁸Centre de Génétique Humaine, Université de Franche-Comté, Besançon, France. ⁹Centre de référence maladies rares pour les anomalies du développement Nord-Ouest, Clinique de Génétique médicale, CHU de Lille et EA7364, Université de Lille, Lille, France. ¹⁰Centre de référence des malformations et maladies congénitales du cerveau, département de génétique et embryologie médicale, APHP, GHUEP, Hôpital Trousseau, Paris, France. ¹¹Service de génétique médicale, CHU de Brest, Brest, France. ¹²Service de génétique médicale, AP-HP Robert-Debré, Paris, France. ¹³Service de génétique médicale, CHI de Toulon, Toulon, France. ¹⁴Service de génétique médicale, CHU de Nantes, Nantes, France. ¹⁵Service de génétique médicale, CHU de Grenoble, Grenoble, France. ¹⁶Service de génétique médicale, CHU de Strasbourg, Strasbourg, France. ¹⁷Service de génétique clinique, CHU de Rennes, Univ. Rennes, Institute of Genetics and Development of Rennes (IGDR) UMR6290 CNRS, Rennes, France. ¹⁸Service de génétique médicale et centre de référence Anomalies du Développement et Syndromes Malformatifs, CHU de Dijon, Dijon, France. ¹⁹Service d'onco hématologie pédiatrique, CHU de Bordeaux, Bordeaux, France. ²⁰Centre de référence des cytopénies auto-immunes de l'enfant, CHU de Bordeaux, Bordeaux, France. ²¹INSERM CICP, Université de Bordeaux, Bordeaux, France. ²²Service de génétique médicale, CHU de Clermont-Ferrand, Clermont-Ferrand, France. ²³Service de génétique médicale, CHU de Caen, Caen, France. ²⁴Normandie Univ, UNIROUEN, Inserm U1245 and Rouen University Hospital, Department of Genetics and Reference Center for Developmental Disorders, F 76000, Normandy Center for Genomic and Personalized Medicine, Rouen, France. ²⁵Service de génétique médicale, CHU d'Angers, Angers, France. ²⁶Service de génétique médicale, CHU de la Réunion, Saint-Pierre, France. ²⁷Centre de Référence Anomalies du développement et Syndromes malformatifs du Sud-Ouest Occitanie Réunion, Saint-Pierre, France. ²⁸Service de génétique médicale, CHU de Toulouse, Toulouse, France. ²⁹Service de génétique médicale, CHU de Nîmes, Nîmes, France. ³⁰Institut Imagine, Paris Descartes-Sorbonne Paris Cité University, Paris, France. ³¹Service de génétique médicale, CHU de Nancy, Nancy, France. ³²Service de génétique médicale, CH du Mans, Le Mans, France. ³³Service de génétique médicale, CHU de Nice, Nice, France. ³⁴Service de génétique médicale, CHU de

Marseille, Marseille, France. ³⁵Service de génétique médicale, CHU de Lyon, Lyon, France. ³⁶INSERM U1211, Université de Bordeaux, Bordeaux, France. ³⁷Service de génétique médicale, AP-HP Pitié Salpêtrière, Paris, France.