



**HAL**  
open science

## Different phenotypes in dermatomyositis associated with anti-MDA5 antibody

Yves Allenbach, Yurdagul Uzunhan, Ségolène Toquet, Gaëlle Leroux, Laure Gallay, Alicia Marquet, Alain Meyer, Constance Guillaud, Nicolas Limal, Frédéric Gagnadoux, et al.

### ► To cite this version:

Yves Allenbach, Yurdagul Uzunhan, Ségolène Toquet, Gaëlle Leroux, Laure Gallay, et al.. Different phenotypes in dermatomyositis associated with anti-MDA5 antibody. *Neurology*, 2020, 95 (1), pp.e70-e78. 10.1212/WNL.0000000000009727 . hal-03425167

**HAL Id: hal-03425167**

**<https://hal.univ-reims.fr/hal-03425167>**

Submitted on 12 Jan 2022

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

# Different phenotypes in dermatomyositis associated with anti-MDA5 antibody

## Study of 121 cases

Yves Allenbach, MD, PhD,\* Yurdagul Uzunhan, MD, PhD,\* Ségolène Toquet, MD,\* Gaëlle Leroux, MD, Laure Gallay, MD, Alicia Marquet, MD, Alain Meyer, MD, PhD, Constance Guillaud, MD, Nicolas Limal, MD, Frédéric Gagnadoux, MD, PhD, Baptiste Hervier, MD, PhD, Raphaël Borie, MD, PhD, Christophe Deligny, MD, PhD, Benjamin Terrier, MD, PhD, Alice Berezne, MD, Sylvain Audia, MD, PhD, Nicolas Champiaux, MD, PhD, Hervé Devilliers, MD, PhD, Nicol Voermans, MD, Elizabeth Diot, MD, Amélie Servettaz, MD, PhD, Thierry Marhadour, MD, Vincent Castelain, MD, PhD, Sébastien Humbert, MD, Claire Blanchard-Delaunay, MD, Nathalie Tieulie, MD, PhD, Pierre Charles, MD, Magdalena Gerin, MD, Arsène Mekinian, MD, PhD, Pascaline Priou, MD, Jean Claude Meurice, MD, PhD, Abdellatif Tazi, MD, PhD, Vincent Cottin, MD, PhD, Makoto Miyara, MD, PhD, Benjamin Grange, MD, Dominique Israël-Biet, MD, PhD, Sophie Phin-Huynh, MD, Camille Bron, MD, Luc De Saint Martin, MD, Nicole Fabien, MD, Kubéraka Mariampillai, MD, Hilario Nunes, MD, PhD,† and Olivier Benveniste, MD, PhD,† the French Myositis Network

### Correspondence

Dr. Allenbach  
yves.allenbach@aphp.fr

*Neurology*® 2020;95:1-9. doi:10.1212/WNL.0000000000009727

## Abstract

### Objectives

The predominance of extramuscular manifestations (e.g., skin rash, arthralgia, interstitial lung disease [ILD]) as well as the low frequency of muscle signs in anti-melanoma differentiation-associated gene 5 antibody-positive (anti-MDA5+) dermatomyositis caused us to question the term myositis-specific antibody for the anti-MDA5 antibody, as well as the homogeneity of the disease.

### Methods

To characterize the anti-MDA5+ phenotype, an unsupervised analysis was performed on anti-MDA5+ patients (n = 83/121) and compared to a group of patients with myositis without anti-MDA5 antibody (anti-MDA5−; n = 190/201) based on selected variables, collected retrospectively, without any missing data.

### Results

Within anti-MDA5+ patients (n = 83), 3 subgroups were identified. One group (18.1%) corresponded to patients with a rapidly progressive ILD (93.3%;  $p < 0.0001$  across all) and a very high mortality rate. The second subgroup (55.4%) corresponded to patients with pure dermatorheumatologic symptoms (arthralgia; 82.6%;  $p < 0.01$ ) and a good prognosis. The third corresponded to patients, mainly male (72.7%;  $p < 0.0001$ ), with severe skin vasculopathy, frequent signs of myositis (proximal weakness: 68.2%;  $p < 0.0001$ ), and an intermediate prognosis. Raynaud phenomenon, arthralgia/arthritis, and sex permit the cluster apparition (83.3% correct estimation). Nevertheless, an unsupervised analysis confirmed that anti-MDA5 antibody delineates an independent group of patients (e.g., dermatomyositis skin rash, skin ulcers, calcinosis, mechanic's hands, ILD, arthralgia/arthritis, and high mortality rate) distinct from anti-MDA5− patients with myositis.

### Conclusion

Anti-MDA5+ patients have a systemic syndrome distinct from other patients with myositis. Three subgroups with different prognosis exist.

\*These authors contributed equally to the work as co-first authors.

†These authors contributed equally to the work as co-last authors.

From the Departments of Clinical Immunology and Internal Medicine (Y.A., S.T., G.L., B.H., N.C., K.M., O.B.), Immunobiology (M.M.), and Biostatistics, Public Health, and Medical Information (B.G.), Pitié-Salpêtrière Hospital, APHP, Sorbonne Universities; Myology Research Centre (Y.A., O.B.), INSERM, UMRS 974, Paris; Department of Pneumology (Y.U., H.N.), APHP, Avicenne Hospital, INSERM U1272, Université Sorbonne Paris Nord, Bobigny; Department of Internal Medicine (L.G., A. Marquet), Lyon Sud Hospital, Lyon Sud University, Paris; Reference Centre for Rare Autoimmune Diseases (A. Meyer) and Medical Intensive Care Unit, Hautepierre Hospital (V. Castelain), Hôpitaux Universitaires de Strasbourg, Strasbourg University; Division of Pneumology (R.B.), Bichat Hospital, AP-HP, Paris-Diderot University, Paris; Department of Pneumology (F.G., P.P.), Centre Hospitalier Universitaire, Angers University; Department of Internal Medicine Infectieuses (C.D.), CHU de Martinique, Fort de France; Departments of Internal Medicine and Clinical Immunology (S.A.) and Internal Medicine and Systemic Diseases (H.D.), François-Mitterrand University Hospital, University of Bourgogne-Franche-Comté, Dijon; Internal Medicine Department (B.T., A.B.), Cochin Hospital, AP-HP, René Descartes University, Paris; Department of Internal Medicine (C.G., N.L.), Henri Mondor Hospital, Paris Est Créteil University, Créteil, France; Department of Neurology (N.V.), Donders Institute for Brain, Cognition, and Behaviour, Radboud University Medical Centre, Nijmegen, the Netherlands; Department of Internal Medicine (E.D.), Hôpital Bretonneau Tours, Tours University; Internal Medicine Department (A.S.), Hôpital Robert-Debré, Reims University; Departments of Rheumatology (T.M.) and Internal Medicine (L.D.S.M.), Bretagne Occidentale University, Centre Hospitalo-Universitaire de Brest; Department of Internal Medicine (S.H.), CHU de Besançon; Department of Internal Medicine (C.B.-D.), Centre Hospitalier Général de Niort; Department of Internal Medicine (N.T.), Nice University, CHU Nice; Department of Internal Medicine (P.C.), Institut Mutualiste Montsouris, Paris; Department of Internal Medicine (M.G.), Jean Verdier Hospital, Paris Seine-Saint-Denis University, Bondy; Department of Internal Medicine (A. Mekinian), Saint Antoine Hospital, APHP, Sorbonne Universities, Paris; Department of Pneumology CHU (J.C.M.), Poitiers University; Department of Pneumology (A.T.), Hôpital Saint-Louis, APHP, Sorbonne Paris Cité, University Paris-Diderot, Paris; Department of Respiratory Medicine (V. Cottin), National Reference Center for Rare Pulmonary Diseases, Hospices Civils de Lyon, Claude Bernard University Lyon 1, Lyon University; Department of Pneumology (D.I.-B.), European George Pompidou Hospital, APHP, Paris V University; Department of Pneumology (S.P.-H.), Sainte-Camille Hospital, Bry-sur-Marne; Department of Pneumology (C.B.), Foch Hospital, Saint Quentin en Yveline University, Suresnes; and Department of Immunology (N.F.), Lyon-Sud Hospital, Hospices Civils-de-Lyon, Pierre-Bénite, France.

Go to [Neurology.org/N](https://www.neurology.org/N) for full disclosures. Funding information and disclosures deemed relevant by the authors, if any, are provided at the end of the article.

## Glossary

**CK** = creatine kinase; **DM** = dermatomyositis; **ENMC** = European NeuroMuscular Center; **ICU** = intensive care unit; **ILD** = interstitial lung disease; **MDA5** = melanoma differentiation-associated gene 5; **RP-ILD** = rapidly progressive interstitial lung disease.

Dermatomyositis (DM) is a heterogeneous group of autoimmune diseases, including disorders limited to the skin of patients, with extracutaneous manifestations, such as muscle, articular or pulmonary lesions, and sometimes with an association with malignancy.<sup>1</sup> Myositis-specific autoantibodies permit the delineation of homogenous subgroups of DM.<sup>1,2</sup> DM associated with anti-melanoma differentiation-associated gene 5 antibody (anti-MDA5+) is typically characterized by the presence of a DM skin rash and polyarthralgia and interstitial lung disease (ILD), whereas the clinical signs of myositis are frequently absent.<sup>1,3–5</sup> Anti-MDA5+ DM has a high mortality rate related to the presence of rapidly progressive ILD (RP-ILD).<sup>1</sup>

The predominance and the variety of the extramuscular manifestations, as well as the absence of muscle symptoms, calls into question the term myositis-specific antibody for anti-MDA5 antibodies.

The anti-MDA5 antibody was identified in 2009,<sup>5</sup> and a limited number of case reports and case series have been reported. Knowledge of the precise clinical phenotype and the prognosis of anti-MDA5+ patients is necessary to improve the management of this potentially severe disease.

To characterize the anti-MDA5+ phenotype, we analyzed the characteristics of a large group of patients (n = 121) and performed unsupervised analysis to detect subgroups with different prognoses among anti-MDA5+ patients. We also compared anti-MDA5+ patients with a group of myositis patients without anti-MDA5 antibody (anti-MDA5–; n = 201) to confirm the specificity of the phenotype.

## Methods

### Patients

Anti-MDA5+ patients were included in the study if they presented any of the following or a combination thereof: a DM skin rash compatible with DM, according to the European NeuroMuscular Center (ENMC) criteria<sup>6</sup> or Sontheimer criteria<sup>7</sup>; myositis (pathologic features showing the presence of inflammatory infiltrates); arthralgia; or ILD, without other etiology. Anti-MDA5 antibody detection was performed using line-immunoassays (Euroimmun [Germany] or D-Tek [Belgium]). This multicenter observational study was performed on data available from 37 medical centers in France from 2011 to 2017. Medical records were reviewed retrospectively (Y.A., S.T., G.L., and Y.U.) to collect clinical, laboratory, and imaging data. ILD was defined based on CT

imaging studies and RP-ILD was defined by the acute onset and rapid worsening within 3 months of the onset of respiratory symptoms leading to severe hypoxia  $\leq 60$  mm Hg. Early mortality was defined by death within 3 months of the diagnosis. As a control, a cohort of anti-MDA5– patients with myositis was used. All of the controls had myositis defined based on ENMC<sup>6</sup> or Bohan and Peter criteria<sup>8</sup> and were followed in one center (Pitié-Salpêtrière Hospital). The controls' characteristics are detailed in table e-1 (doi.org/10.5061/dryad.t39c4k1).

### Standard protocol approvals, registrations, and patient consents

Agreement for the study was obtained from the French Ministry of Research (CCTIRS no. 14.323 and AC-2013-1868) and the study was approved by the Research Ethics Committee of the Pitié-Salpêtrière Hospital (Paris, France).

### Statistics

Quantitative data (median [interquartile range]) and qualitative data (frequency and percentage) were described. Unsupervised descriptive methods of statistical learning were used to analyze either the anti-MDA5+ patients or the global cohort of patients with myositis (anti-MDA5+ and anti-MDA5–). A multiple correspondence analysis and hierarchical cluster analysis were used to resume the dataset and aggregate patients in subgroups, as previously reported.<sup>9</sup> Only patients with an exhaustive set of data were included for the unsupervised analysis. The clustering of patients was unsupervised using Euclidean distance and the Ward agglomerative method. *V* test *p* values are represented for the variables that participated in the multidimensional analyses (see supplemental tables, doi.org/10.5061/dryad.t39c4k1).<sup>10</sup> In addition, the Wilcoxon test was used to compare quantitative data and anti-MDA5 status (positive or negative), and Fisher exact test was used for qualitative data in the different groups (anti-MDA5 status [positive or negative]) or global association with the clusters obtained by hierarchical cluster analysis (tables 1 and 2).

To construct a decisional algorithm tree to position the patients in a cluster, we used classification and regression trees. A *p* value  $< 0.05$  was considered significant. For the survival analysis, Kaplan-Meier curves were performed on different patient subgroups and compared using log-rank tests. Statistical analyses were performed (R version 3.4.0 software and GraphPad Prism software).

### Data availability

All supplementary data are available from the Dryad digital repository (doi.org/10.5061/dryad.t39c4k1). Further anonymized

**Table 1** Characteristics of 3 groups of anti-melanoma differentiation-associated gene 5 antibody-positive (anti-MDA5+) patients

	Cluster 1 (n = 15)	Cluster 2 (n = 46)	Cluster 3 (n = 22)	p Value
<b>Sex<sup>a</sup></b>				
Women	11 (73.3)	38 (82.6)	6 (27.3)	<0.0001
Men	4 (26.7)	8 (17.4)	16 (72.7)	
<b>Ethnicity<sup>a</sup></b>				
Caucasian	6 (40)	23 (50)	12 (54.5)	0.67
African	7 (46.7)	20 (43.5)	10 (45.4)	
Asian	2 (13.3)	2 (4.5)	0 (0)	
Hispanic	0 (0)	1 (2.2)	0 (0)	
<b>General condition/inflammation</b>				
Fever	6 (40)	14 (30.4)	9 (42.9)	0.57
Deterioration of general condition <sup>a</sup>	14 (93.3)	30 (65.2)	10 (45.4)	0.008
CRP or ESR increase	7 (58.3)	18 (41.9)	9 (47.4)	0.62
<b>Skin changes</b>				
Skin lesions <sup>a</sup>	15 (100)	38 (82.6)	21 (95.4)	0.16
Mechanic's hands <sup>a</sup>	11 (73.3)	20 (43.5)	3 (13.6)	0.0011
Typical DM skin rash <sup>a</sup>	12 (80)	31 (67.4)	17 (77.3)	0.63
Raynaud phenomenon <sup>a</sup>	1 (6.7)	5 (10.9) <sup>a</sup>	18 (81.8)	<0.0001
Skin ulcers <sup>a</sup>	4 (26.7)	17 (37)	17 (77.3)	0.002
Digital necrosis <sup>a</sup>	0 (0)	2 (4.3)	7 (31.8)	0.002
Calcinosis <sup>a</sup>	0 (0)	2 (4.3)	5 (22.7)	0.02
Skin sclerosis <sup>a</sup>	1 (6.7)	2 (4.3)	5 (22.7)	0.05
<b>Muscular manifestations</b>				
Proximal weakness <sup>a</sup>	4 (26.7)	7 (15.2) <sup>a</sup>	15 (68.2)	0.0001
MRC ≤3 <sup>a</sup>	0 (0)	1 (2.3)	5 (26.3)	0.003
Increased CK level <sup>a</sup>	10 (66.7)	10 (21.7)	14 (63.6)	0.0004
CK >160 IU/L and <800 IU/L <sup>a</sup>	10 (66.7)	11 (24)	6 (27.3)	
>800 IU/L <sup>a</sup>	1 (6.7)	0 (0)	8 (36.4)	
Myositis on muscle biopsy	2 (50)	12 (52.2)	9 (56.2)	1
<b>Lung manifestations</b>				
Dyspnea <sup>a</sup>	15 (100)	28 (60.9)	10 (45.4)	0.0008
ILD <sup>a</sup>	15 (100)	38 (82.6)	11 (50)	0.0008
RP-ILD <sup>a</sup>	14 (93.3)	8 (17.4)	5 (22.7)	<0.0001
PaO <sub>2</sub> <60 mm Hg	13 (92.8)	8 (23.6)	8 (57.1)	<0.0001
<b>Rheumatologic manifestations</b>				
Arthritis/arthralgia <sup>a</sup>	4 (26.7)	38 (82.6)	17 (77.3)	0.0003
Synovitis <sup>a</sup>	1 (6.7)	17 (37)	9 (41)	0.0479
RF or anti-CCP antibody	3 (27.7)	5 (13.5)	4 (21)	0.46

Continued

**Table 1** Characteristics of 3 groups of anti-melanoma differentiation-associated gene 5 antibody-positive (anti-MDA5+) patients (continued)

	Cluster 1 (n = 15)	Cluster 2 (n = 46)	Cluster 3 (n = 22)	p Value
<b>Outcomes</b>				
ICU <sup>a</sup>	13 (86.7)	5 (10.9) <sup>a</sup>	8 (36.4)	<0.0001
Death within 3 months <sup>a</sup>	12 (80)	0 (0) <sup>a</sup>	1 (4.5)	<0.0001
<b>Miscellaneous</b>				
Malignancy <sup>a</sup>	1 (6.7)	2 (4.3)	1 (4.5)	1
Lymphadenopathy	5 (38.5)	15 (33.3)	4 (19)	0.4
Seritis	0 (0)	7 (16.7)	4 (18.2)	0.27
Increased liver enzymes	8 (66.7)	16 (40)	11 (61.1)	0.18
Leukopenia	2 (16.7)	13 (31.7)	3 (15)	0.32
ANA	5 (41.7)	21 (55.3)	11 (57.9)	0.72

Abbreviations: ANA = antinuclear antibody; anti-CCP = anti-cyclic citrullinated peptide antibody; CK = creatine kinase; CRP = C-reactive protein; DM = dermatomyositis; ESR = erythrocyte sedimentation rate; ICU = intensive care unit; ILD = interstitial lung disease; MRC = Medical Research Council scale; RF = rheumatoid factor; RP-ILD = rapidly progressive interstitial lung disease. Values are n (%).

<sup>a</sup> Variables used in the unsupervised analysis permitting us to identify the 3 clusters. The other variables (not used for the clustering, because not available for all anti-MDA5+ patients) were positioned in the different clusters.

data can be made available to qualified investigators upon request to the corresponding author.

## Results

### Three MDA5+ patient subgroups

Anti-MDA5+ patients (n = 121) were mainly female, 49 years old (34–58), and Caucasian, but African origin was also frequent. As expected, most of the patients had a DM skin rash (71.2%), ILD (76.5%), and arthralgia (69%), whereas only one-third of patients had muscle weakness (table e-2, doi.org/10.5061/dryad.t39c4k1). Of note, 2 patients had concomitant myocarditis with severe cardiac failure.

We aimed to identify the phenotype variations within the anti-MDA5+ group (n = 83/121). The unsupervised hierarchical analysis tree that we positioned on the factorial map showed 3 clusters (figure 1, A and B). The characteristics of each cluster are reported in table 1.

Cluster 1 (18.1%; n = 15) corresponded to patients with very severe lung disease (RP-ILD cluster). All patients had ILD, and all but one had RP-ILD (93.3%; V test  $p < 0.0001$ ). Patients in this cluster were admitted to the intensive care unit (ICU) in 86.7% of cases ( $p < 0.0001$ ). The early mortality rate was high (80%; V test  $p < 0.0001$ ). The other main characteristic was that these patients frequently had mechanic's hands (73.3%; V test  $p = 0.006$ ).

Cluster 2 (55.4%; n = 46) corresponds to a pure dermatorheumatologic pattern (rheumatoid cluster). As for the RP-ILD

cluster, patients in cluster 2 were mainly women and less frequently had skin lesions (82.6%; V test  $p = 0.03$ ), digital necrosis (4.35%;  $p = 0.04$ ), and the Raynaud phenomenon (10.9%; V test  $p < 0.0001$ ) (table 1). Most patients in cluster 2 complained of arthralgia or arthritis (82.6%; V test  $p = 0.01$ ). Signs of myositis (muscle weakness 15.2%; V test  $p = 0.0005$  and increased creatine kinase [CK] level 21.7%; V test  $p < 0.0001$ ), as well as RP-ILD (17.4%; V test  $p = 0.001$ ), were infrequent, and no early death occurred in this group (V test  $p < 0.0001$ ).

Cluster 3 (26.5%; n = 22) corresponded to patients with severe skin vasculopathy and an intermediate prognosis (vasculopathic cluster). Patients here were mainly male (72.7%; V test  $p < 0.0001$ ). In addition to the classical DM skin rash, those patients frequently harbored signs of skin vasculopathy with the Raynaud phenomenon (81.8%; V test  $p < 0.0001$ ), skin ulcers (77.3%; V test  $p = 0.0006$ ), digital necrosis (31.8%;  $p = 0.001$ ), and calcinosis (22.7%; V test  $p = 0.01$ ). In addition, most patients had proximal weakness (68.2%; V test  $p < 0.0001$ ) with increased CK levels in 63.6% (V test;  $p < 0.0001$ ). Some of the patients in cluster 3 had RP-ILD (22.7%), and the early mortality rate was low (4.5% in the vasculopathic cluster).

### Three different MDA5+ patients prognoses

We next analyzed the long-term survival of anti-MDA5+ patients depending on the 3 clusters (figure 2A). The analysis clearly confirmed a different prognosis between the clusters. The RP-ILD cluster had a very high mortality rate compared to patients in the rheumatic and vasculopathic clusters ( $p <$

**Table 2** Comparison of anti-melanoma differentiation-associated gene 5 antibody–positive (anti-MDA5+) vs anti-MDA5– patients with myositis

	Anti-MDA5– (n = 201)	Anti-MDA5+ (n = 121)	p Value
<b>Sex</b>			
Male	56 (27.9)	40 (33.1)	0.3889
<b>Ethnicity</b>			
Caucasian	144 (71.6)	56 (48.3)	<0.0001
African	30 (14.9)	50 (43.1)	
Asian	6 (3)	8 (6.9)	
Others	21 (10.4)	2 (1.7)	
<b>Skin</b>			
Skin lesions	129 (64.2)	105 (87.5)	<0.0001
Typical DM skin rash	54 (26.9)	84 (70)	<0.0001
Mechanic's hands	35 (17.4)	50 (42.7)	<0.0001
Raynaud phenomenon	70 (35.2)	34 (29.8)	0.3995
Skin ulcers	11 (5.5)	49 (41.5)	<0.0001
Calcinosis	4 (2)	12 (10.2)	0.003
<b>Muscle</b>			
Proximal weakness	180 (90)	35 (30.7)	<0.0001
Psoas MRC ≤3	107 (57.53)	8 (8.2)	<0.0001
Increased CK level	190 (94.5)	46 (41.4)	<0.0001
CK level >800 IU/L	162 (80.6)	13 (10.7)	<0.0001
Myositis on biopsy	170 (90.9)	30 (55.6)	<0.0001
<b>Joints</b>			
Arthritis/arthralgia	92 (45.8)	80 (69)	<0.0001
<b>Lungs</b>			
Dyspnea	137 (70.3)	80 (66.7)	0.5871
ILD	90 (46.4)	91 (76.5)	<0.0001
Malignancy	24 (12)	9 (7.6)	0.296
Death	13 (6.8)	33 (27.3)	<0.0001

Abbreviations: CK = creatine kinase; DM = dermatomyositis; ILD = interstitial lung disease; MRC = Medical Research Council scale. Values are n (%).

0.0001). Comparing patients in the rheumatic cluster to those in the vasculopathic cluster, it appeared that the prognosis was better (very good) in the rheumatic cluster than in the vasculopathic cluster ( $p = 0.01$ ).

Finally, we aimed to easily position a participant in one of the 3 clusters. Classification and regression tree analysis was used as a predictive model to determine the items

permitting us to classify a patient (figure 2B). Because we aimed to predict patient outcome, the variables related to obvious severity were removed (RP-ILD and ICU). The models including 3 variables Raynaud phenomenon, arthralgia/arthritis, and sex permits 83.3% correct estimation. Of note, adding the variable RP-ILD permits us to reach 91.6% correct estimation.

### Comparison of anti-MDA5+ vs anti-MDA5– patients

Because we observed the different subgroups of anti-MDA5+ patients, we aimed to verify if anti-MDA5+ disease remains a separate entity within myositis. We compared anti-MDA5+ patients ( $n = 121$ ) to anti-MDA5– myositis patients ( $n = 201$ ) (table 2).

Compared to anti-MDA5– patients, anti-MDA5+ patients were more frequently African, more frequently had a DM skin rash, skin ulcers, calcinosis, mechanic's hands, ILD, and arthralgia/arthritis, and had a higher mortality rate (table 2). Anti-MDA5– patients had more severe myositis (based on muscle weakness and CK level).

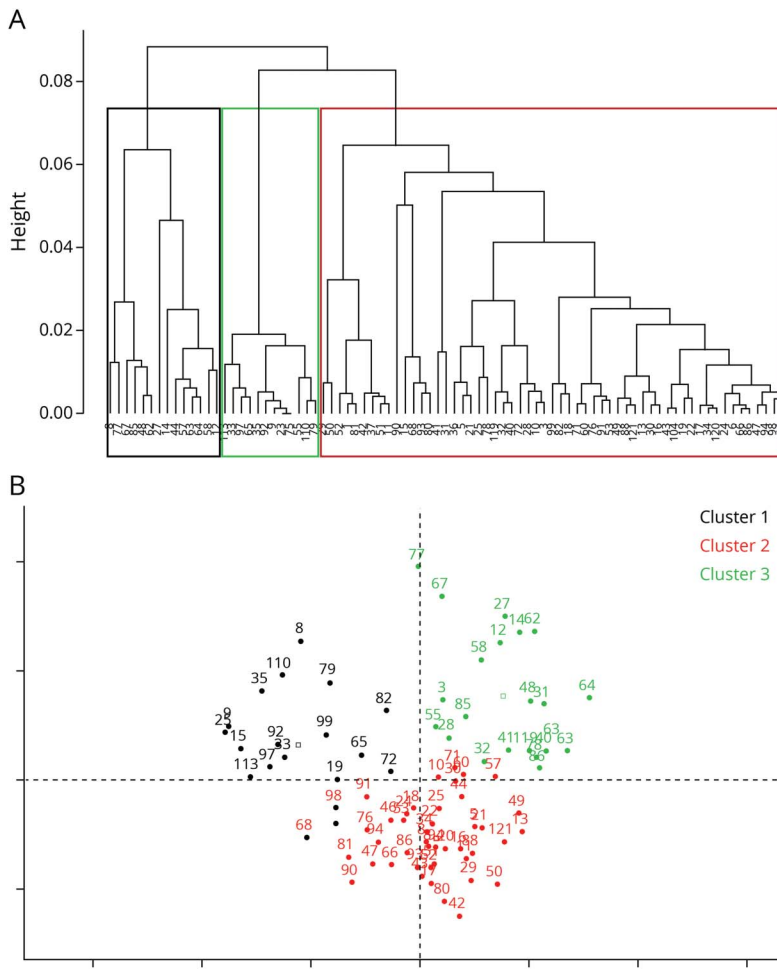
Next, we performed an unsupervised hierarchical clustering analysis in a large group of patients with myositis (anti-MDA5+ and anti-MDA5– patients;  $n = 274/322$ ) with 20 variables, including long-term mortality (table e-2, doi.org/10.5061/dryad.t39c4k1), but importantly not including anti-MDA5 antibody status. We clearly observed 2 clusters (figure 3 and figure e-1, doi.org/10.5061/dryad.t39c4k1). The hierarchical tree positioned on the factorial map showed that cluster 2 regrouped mainly anti-MDA5+ patients (87.5%;  $n = 84/96$  vs 7.3%;  $n = 13/178$ ;  $p < 0.0001$ ) after having positioned the variable anti-MDA5. The anti-MDA5 antibody delineates an independent subgroup of patients from a large cohort of patients with myositis.

### Discussion

We report the phenotype of a large group of anti-MDA5+ patients. Overall, our observation confirmed previous studies showing that anti-MDA5+ patients present a DM skin rash with frequent signs of skin vasculopathy,<sup>3</sup> frequent arthralgia/arthritis,<sup>4</sup> and ILD with a high mortality rate.<sup>1</sup>

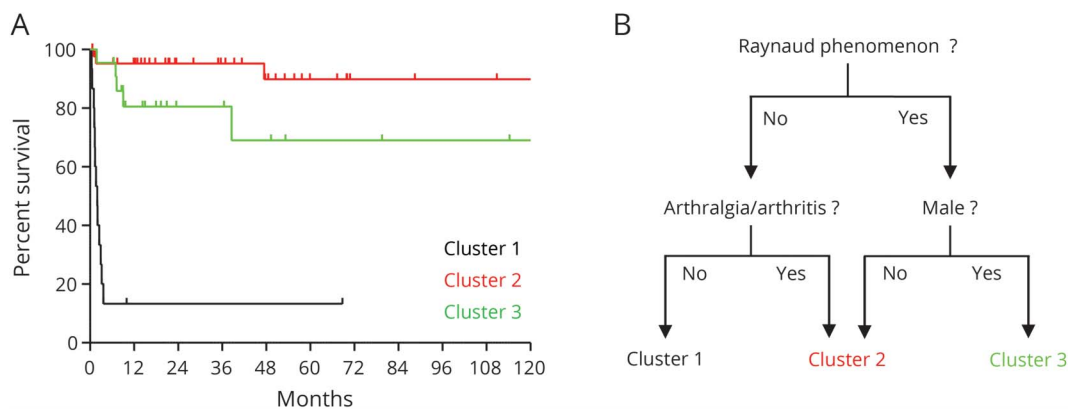
Using unsupervised analyses, we observed for the first time 3 distinct subgroups with different prognoses. The RP-ILD cluster with severe lung involvement and a poor prognosis corresponds to well-recognized anti-MDA5+ RP-ILD.<sup>1,5</sup> In addition, we were able to isolate 2 new forms: anti-MDA5+ rheumatic DM, with a good prognosis, and anti-MDA5+ vasculopathic DM, with an intermediate prognosis. The algorithm decisional algorithm showed that only 3 variables (Raynaud phenomenon, arthralgia/arthritis, and sex) are good predictors for cluster appartenance.

**Figure 1** Unsupervised analysis of anti-melanoma differentiation-associated gene 5 antibody-positive (anti-MDA5+) patients



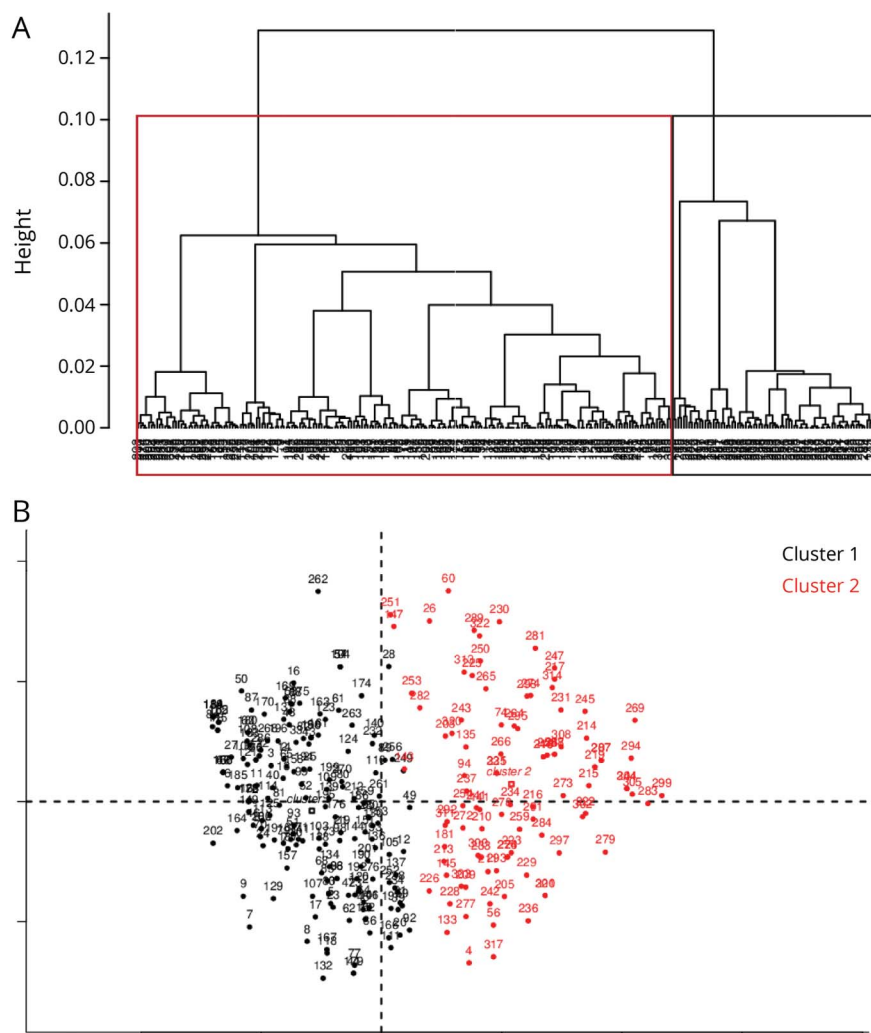
(A) The hierarchical cluster analysis of anti-MDA5+ patients showed 3 clusters (the variables included in the unsupervised analysis were sex, ethnicity, skin changes, typical dermatomyositis skin rash, mechanic's hands, skin ulcers, calcinosis, digital necrosis, sclerosis, Raynaud phenomenon, abnormal creatine kinase [CK] level, CK level, proximal muscle weakness, manual muscle testing score of the psoas [Medical Research Council score], arthritis/arthralgia, synovitis, dyspnea, interstitial lung disease [ILD], rapidly progressive ILD, resuscitation, deterioration of general status, malignancy, intensive care unit admission, malignancy within 3 years before or after the myositis diagnosis, and early mortality). (B) Multiple correspondence analysis confirmed the presence of 3 groups of anti-MDA5+ patients. All above-mentioned variables except intensive rapidly progressive ILD, care unit admission, and mortality were included in the classification algorithm regression tree analysis.

**Figure 2** Anti-melanoma differentiation-associated gene 5 antibody-positive (anti-MDA5+) prognosis



(A) The survival curves showed that cluster 1 (black) had a poor prognosis, cluster 2 (red) had a good prognosis, and cluster 3 (green) had a mild prognosis. (B) Pruned model of prediction without obvious signs of severity (the variables included in the classification algorithm regression tree analysis did not encompass obvious signs of severity [rapidly progressive interstitial lung disease (ILD), intensive care unit admission, and mortality], sex, ethnicity, skin changes, typical dermatomyositis skin rash, mechanic's hands, skin ulcers, calcinosis, digital necrosis, sclerosis, Raynaud phenomenon, abnormal creatine kinase [CK] level, CK level, proximal muscle weakness, manual muscle testing score of the psoas [Medical Research Council score], arthritis/arthralgia, synovitis, dyspnea, ILD, deterioration of the general status, malignancy, and malignancy within 3 years before or after the myositis diagnosis).

**Figure 3** Unsupervised analysis of anti-melanoma differentiation-associated gene 5 antibody-positive (anti-MDA5+) and anti-MDA5- patients



(A) Hierarchical cluster analysis of anti-MDA5+ patients and anti-MDA5- myositis patients without including the anti-MDA5 antibody status as variable (the variables included for the unsupervised analysis were sex, ethnicity, proximal muscle weakness, abnormal creatine kinase [CK] level, CK level, skin changes, typical dermatomyositis skin rash, mechanic's hands, skin ulcers, calcinosis, Raynaud phenomenon, arthritis/arthralgia, dyspnea, interstitial lung disease, and malignancy within 3 years before or after the myositis diagnosis). The analysis permitted the identification of 2 main clusters. (B) Using multiple correspondence analysis, both clusters were positioned on the factor map of individuals, which confirmed that patients were segregated into 2 groups. When the variable anti-MDA5 was next positioned, it appeared that cluster 2 was mainly composed of anti-MDA5+ patients (87.1%).

These observations and the large spectrum of disease manifestations suggest that the so-called anti-MDA5+ DM is rather a systemic syndrome (anti-MDA5+ syndrome) than a musculoskeletal disease.

This finding underlines the importance of anti-MDA5 antibody screening in patients with seronegative polyarthritis and patients with interstitial pneumonia, with a suspicion of autoimmune features,<sup>11</sup> or ICU patients with acute respiratory distress syndrome of unknown origin. The prognosis related to the different clusters is also important regarding the treatment strategy and may suggest different pathophysiological mechanisms.

The comparison of anti-MDA5+ patients with a large cohort of MDA5- patients showed that anti-MDA5+ syndrome remains a separate entity. Using unsupervised clustering analysis, we showed that the anti-MDA5+ antibody was independently and strongly associated with this phenotype. This result emphasizes

the crucial role of myositis-specific antibodies to diagnose and delineate a subgroup of patients. However, the term myositis-specific antibody seems inappropriate, because many anti-MDA5+ patients are amyopathic.

The nationwide retrospective cohort study design we used is the main limitation of our work. The low frequency of the reported rheumatologic signs in the RP-ILD cluster could be due to a detection bias since the majority of patients were in the ICU. Nevertheless, patients in the RP-ILD cluster were also characterized by an increased CK level, whereas this was not the case for patients with the rheumatologic form in cluster 2.

Together, these data show that anti-MDA5+ patients are a distinct group from patients with myositis and have a systemic syndrome composed of 3 different entities with different prognoses. The anti-MDA5+ antibody is a key biomarker to define this syndrome.



## Study funding

No targeted funding reported.

## Disclosure

The authors report no relevant disclosures. Go to [Neurology.org/N](https://www.neurology.org/N) for full disclosures.

## Publication history

Received by *Neurology* February 14, 2019. Accepted in final form December 12, 2019.

## Appendix Authors

Name	Location	Contribution
<b>Yves Allenbach, MD, PhD</b>	Sorbonne Université, Paris	Designed and conceptualized the study, analyzed the data, drafted the manuscript
<b>Yurdagul Uzunhan, MD, PhD</b>	APHP, Avicenne hospital, INSERM U1272, Université Sorbonne Paris Nord, Bobigny	Designed and conceptualized the study, analyzed the data, drafted the manuscript
<b>Ségolène Toquet, MD</b>	Sorbonne Université, Paris	Major role in data acquisition, revised the manuscript for intellectual content
<b>Gaëlle Leroux, MD</b>	Sorbonne Université, Paris	Major role in data acquisition, revised the manuscript for intellectual content
<b>Laure Gallay, MD</b>	Lyon Université	Major role in data acquisition, revised the manuscript for intellectual content
<b>Alicia Marquet, MD</b>	Lyon Université	Major role in data acquisition
<b>Alain Meyer, MD, PhD</b>	Strasbourg Université	Major role in data acquisition
<b>Constance Guillaud, MD</b>	Paris Est Université, Créteil	Major role in data acquisition
<b>Nicolas Limal, MD</b>	Paris Est Université, Créteil	Major role in data acquisition
<b>Frédéric Gagnadoux, MD, PhD</b>	Angers Université	Major role in data acquisition
<b>Baptiste Hervier, MD, PhD</b>	Sorbonne Université, Paris	Major role in data acquisition
<b>Raphaël Borie, MD, PhD</b>	Paris Diderot Université, Paris	Major role in data acquisition
<b>Christophe Deligny, MD, PhD</b>	Centre Hospitalier Universitaire, Martinique	Major role in data acquisition
<b>Benjamin Terrier, MD, PhD</b>	Descartes Université, Paris	Major role in data acquisition
<b>Alice Berezne, MD</b>	Hôpital Annecy, Annecy	Major role in data acquisition

## Appendix (continued)

Name	Location	Contribution
<b>Sylvain Audia, MD, PhD</b>	Université Bourgogne Franche Comté, Dijon	Major role in data acquisition
<b>Nicolas Champetiaux, MD, PhD</b>	Sorbonne Université, Paris	Major role in data acquisition
<b>Hervé Devilliers, MD, PhD</b>	Université Bourgogne Franche Comté, Dijon	Major role in data acquisition
<b>Nicol Voernmans, MD</b>	Radboud Université, Nijmegen	Major role in data acquisition
<b>Elizabeth Diot, MD</b>	Tour Université, Tour	Major role in data acquisition
<b>Amélie Servettaz, MD, PhD</b>	Reims Université	Major role in data acquisition
<b>Thierry Marhadour, MD</b>	Bretagne Université, Reims	Major role in data acquisition
<b>Vincent Castelain, MD, PhD</b>	Strasbourg Université	Major role in data acquisition
<b>Sébastien Humbert, MD</b>	Besançon Université	Major role in data acquisition
<b>Claire Blanchard-Delaunay, MD</b>	Niort Hôpital	Major role in data acquisition
<b>Nathalie Tieulie, MD, PhD</b>	Nice Université	Major role in data acquisition
<b>Pierre Charles, MD</b>	Institut Mutualiste Montsouris, Paris	Major role in data acquisition
<b>Magdalena Gerin, MD</b>	Saint Denis Université	Major role in data acquisition
<b>Arsène Mekinian, MD, PhD</b>	Sorbonne Université, Paris	Major role in data acquisition
<b>Pascaline Priou, MD</b>	Angers Université	Major role in data acquisition
<b>Jean Claude Meurice, MD, PhD</b>	Poitier Université	Major role in data acquisition
<b>Abdellatif Tazi, MD, PhD</b>	Paris Diderot Université	Major role in data acquisition
<b>Vincent Cottin, MD, PhD</b>	Lyon Université	Major role in data acquisition
<b>Makoto Miyara, MD, PhD</b>	Sorbonne Université, Paris	Major role in data acquisition
<b>Benjamin Grange, MD</b>	Sorbonne Université, Paris	Interpreted the data and statistical analysis
<b>Dominique Israël-Biet, MD, PhD</b>	Descartes Université, Paris	Major role in data acquisition

## Appendix *(continued)*

Name	Location	Contribution
<b>Sophie Phin-Huynh, MD</b>	Sainte Camille Hôpital, Bry-sur-Marne	Major role in data acquisition
<b>Camille Bron, MD</b>	Saint Quentin Université, Suresnes	Major role in data acquisition
<b>Luc De Saint Martin, MD</b>	Brest Université	Major role in data acquisition
<b>Nicole Fabien, MD</b>	Lyon Université	Major role in data acquisition
<b>Kubéraka Mariampillai, MD</b>	Sorbonne Université, Paris	Interpreted the data and statistical analysis, revised the manuscript for intellectual content
<b>Hilario Nunes, MD, PhD</b>	APHP, Avicenne hospital, INSERM U1272, Université Sorbonne Paris Nord, Bobigny	Major role in data acquisition, interpreted the data, revised the manuscript for intellectual content
<b>Olivier Benveniste, MD, PhD</b>	Sorbonne Université, Paris	Major role in data acquisition, interpreted the data, revised the manuscript for intellectual content

## References

1. Hamaguchi Y, Kuwana M, Hoshino K, et al. Clinical correlations with dermatomyositis-specific autoantibodies in adult Japanese patients with dermatomyositis: a multicenter cross-sectional study. *Arch Dermatol* 2011;147:391–398.
2. Benveniste O, Stenzel W, Allenbach Y. Advances in serological diagnostics of inflammatory myopathies. *Curr Opin Neurol* 2016;29:662–673.
3. Fiorentino D, Chung L, Zwerner J, Rosen A, Casciola-Rosen L. The mucocutaneous and systemic phenotype of dermatomyositis patients with antibodies to MDA5 (CADM-140): a retrospective study. *J Am Acad Dermatol* 2011;65:25–34.
4. Hall JC, Casciola-Rosen L, Samedy LA, et al. Anti-melanoma differentiation-associated protein 5-associated dermatomyositis: expanding the clinical spectrum: clinical features of anti-MDA-5-positive patients. *Arthritis Care Res* 2013;65:1307–1315.
5. Sato S, Hoshino K, Satoh T, et al. RNA helicase encoded by melanoma differentiation-associated gene 5 is a major autoantigen in patients with clinically amyopathic dermatomyositis: association with rapidly progressive interstitial lung disease. *Arthritis Rheum* 2009;60:2193–2200.
6. Hoogendijk JE, Amato AA, Lecky BR, et al. ENMC international workshop: trial design in adult idiopathic inflammatory myopathies, with the exception of inclusion body myositis, 10–12 October 2003, Naarden, The Netherlands. *Neuromuscul Disord* 2004;14:337–345.
7. Sontheimer RD. Would a new name hasten the acceptance of amyopathic dermatomyositis (dermatomyositis sine myositis) as a distinctive subset within the idiopathic inflammatory dermatomyopathies spectrum of clinical illness? *J Am Acad Dermatol* 2002;46:626–636.
8. Bohan A, Peter JB. Polymyositis and dermatomyositis (first of two parts). *N Engl J Med* 1975;292:344–347.
9. Mariampillai K, Granger B, Amelin D, et al. Development of a new classification system for idiopathic inflammatory myopathies based on clinical manifestations and myositis-specific autoantibodies. *JAMA Neurol* 2018;75:1528–1537.
10. Lebart L, Morineau A, Warwick KM. *Multivariate Descriptive Statistical Analysis (Correspondence Analysis and Related Techniques for Large Matrices)*. Chichester: Wiley; 1984.
11. Fischer A, Antoniou KM, Brown KK, et al. An official European Respiratory Society/American Thoracic Society research statement: interstitial pneumonia with autoimmune features. *Eur Respir J* 2015;46:976–987.

# Neurology<sup>®</sup>

## Different phenotypes in dermatomyositis associated with anti-MDA5 antibody: Study of 121 cases

Yves Allenbach, Yurdagul Uzunhan, Ségolène Toquet, et al.  
*Neurology* published online June 2, 2020  
DOI 10.1212/WNL.00000000000009727

**This information is current as of June 2, 2020**

<b>Updated Information &amp; Services</b>	including high resolution figures, can be found at: <a href="http://n.neurology.org/content/early/2020/06/02/WNL.00000000000009727.full">http://n.neurology.org/content/early/2020/06/02/WNL.00000000000009727.full</a>
<b>Permissions &amp; Licensing</b>	Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at: <a href="http://www.neurology.org/about/about_the_journal#permissions">http://www.neurology.org/about/about_the_journal#permissions</a>
<b>Reprints</b>	Information about ordering reprints can be found online: <a href="http://n.neurology.org/subscribers/advertise">http://n.neurology.org/subscribers/advertise</a>

*Neurology*® is the official journal of the American Academy of Neurology. Published continuously since 1951, it is now a weekly with 48 issues per year. Copyright © 2020 American Academy of Neurology. All rights reserved. Print ISSN: 0028-3878. Online ISSN: 1526-632X.

