

Clinical and laboratory characteristics in juvenile-onset Systemic Lupus Erythematosus across age groups

Short Title: Clinical and laboratory characteristics in JSLE

Massias J^{1,*}, Smith EMD^{2,3,*}, Eslam Al-Abadi⁴, Kate Armon⁵, Kathryn Bailey⁶, Coziana Ciurtin⁷, Joyce Davidson⁸, Janet Gardner-Medwin⁹, Kirsty Haslam¹⁰, Dan Hawley¹¹, Alice Leahy¹², Valentina Leone¹³, Flora McErlane¹⁴, Devesh Mewar¹⁵, Gita Modgil¹⁶, Robert Moots¹⁷, Clarissa Pilkington¹⁸, Athimalaipet Ramanan¹⁹, Satyapal Rangaraj²⁰, Phil Riley²¹, Arani Sridhar²², Nick Wilkinson²³, Beresford MW^{2,3,\$}, Hedrich CM^{2,3,\$}, on behalf of the UK JSLE Study Group

* and ^{\$} Both authors contributed equally

¹School of Medicine, University of Liverpool, UK.

²Department of Women's & Children's Health, Institute of Translational Medicine, University of Liverpool, UK.

³Department of Paediatric Rheumatology, Alder Hey Children's NHS Foundation Trust Hospital, UK.

⁴Department of Rheumatology, Birmingham Children's Hospital, Birmingham, UK.

⁵Department of Paediatric Rheumatology, Cambridge University Hospitals, Cambridge, UK.

⁶Department of Paediatric Rheumatology, Oxford University Hospitals, Oxford, UK.

⁷Department of Rheumatology, University College London Hospitals NHS Foundation Trust, London, UK.

⁸Department of Paediatric Rheumatology, Royal Hospital for Sick Children, Edinburgh, UK.

⁹Department of Paediatric Rheumatology, NHS Greater Glasgow and Clyde (Yorkhill Division), Glasgow, UK.

¹⁰Department of Paediatrics, Bradford Royal Infirmary, Bradford, UK.

¹¹Sheffield Children's Hospital, Sheffield, UK.

¹²Department of Paediatric Rheumatology, Southampton General Hospital, Southampton, UK.

¹³Department of Paediatric Rheumatology, Leeds General Infirmary, Leeds, UK.

¹⁴Department of Paediatric Rheumatology, Royal Victoria Infirmary, Newcastle upon Tyne, UK.

¹⁵Department of Rheumatology, Royal Liverpool University Hospital, Liverpool, UK.

¹⁶Department of Paediatrics, Musgrove Park Hospital, Taunton, UK.

¹⁷Department of Rheumatology, University Hospital Aintree, Liverpool, UK.

¹⁸Department of Paediatric Rheumatology, Great Ormond Street Hospital, London, UK.

¹⁹Department of Paediatric Rheumatology, Bristol Royal Hospital for Children, Bristol, UK.

²⁰Department of Paediatric Rheumatology, Nottingham University Hospitals Nottingham, UK.

²¹Department of Paediatric Rheumatology, Royal Manchester Children's Hospital, Manchester, UK.

²²Department of Paediatrics, Leicester Royal Infirmary, Leicester, UK.

²³Guy's & St Thomas's NHS Foundation Trust, Evelina Children's Hospital, London, UK.

Address correspondence to:

Christian M Hedrich, Institute in the Park, University of Liverpool, Alder Hey Children's Hospital, Eaton Rd, Liverpool, L12 2AP, chedrich@liverpool.ac.uk

Or

Eve MD Smith, Institute in the Park, University of Liverpool, Alder Hey Children's Hospital, Eaton Rd, Liverpool, L12 2AP, esmith8@liverpool.ac.uk

Abstract

Background: Systemic lupus erythematosus (SLE) is a systemic autoimmune/inflammatory condition. Approximately 15-20% of patients develop symptoms before their 18th birthday and are diagnosed with juvenile-onset SLE (JSLE). Gender distribution, clinical presentation, disease courses and outcomes vary significantly between JSLE patients and individuals with adult-onset SLE. This study aimed to identify age-specific clinical and/or serological patterns in JSLE patients enrolled to the UK JSLE Cohort Study.

Methods: Patient records were accessed and grouped based on age at disease-onset: pre-pubertal (≤ 7 years), peri-pubertal (8-13 years) and adolescent (14-18 years). The presence of ACR classification criteria, laboratory results, disease activity (BILAG score) and damage (SLICC damage index) were evaluated at diagnosis and last follow-up.

Results: A total of 418 JSLE patients were included in this study: 43 (10.3%) with pre-pubertal disease onset; 240 (57.4%) with peri-pubertal onset, and 135 (32.3%) were diagnosed during adolescence. At diagnosis, adolescent JSLE patients presented with a higher number of ACR criteria when compared to pre-pubertal and peri-pubertal patients (pBILAG2004 scores: 9[4-20] vs. 7[3-13] vs. 7[3-14] respectively, $p=0.015$) with increased activity in the following BILAG domains: mucocutaneous ($p=0.025$), musculoskeletal ($p=0.029$), renal ($p=0.027$), and cardiorespiratory ($p=0.001$). Furthermore, adolescent JSLE patients were more frequently ANA positive ($p=0.034$) and exhibited higher anti-dsDNA titres ($p=0.001$). Pre-pubertal individuals less frequently presented with leukopenia ($p=0.002$), thrombocytopenia ($p=0.004$) or low complement ($p=0.002$) when compared to other age groups. No differences were identified in disease activity (pBILAG2004 score), damage (SLICC damage index) and the number of ACR criteria fulfilled at last follow-up.

Conclusions: Disease presentations and laboratory findings vary significantly between age groups within a national cohort of JSLE patients. Patients diagnosed during adolescence exhibit greater disease activity and “classic” autoantibody, immune cell and complement patterns when compared to younger patients. This supports the hypothesis that pathomechanisms may vary between patient age groups.

Introduction

Systemic lupus erythematosus (SLE) is a systemic autoimmune/inflammatory condition that can affect any organ system and result in significant damage and organ failure^{1, 2}. Clinical characteristics, underlying pathomechanisms, disease progression and outcomes vary between individuals, age groups and races. Approximately 15-20% of SLE patients develop the disease before their 18th birthday and are therefore diagnosed with juvenile-onset SLE (JSLE)^{1, 2}. Juvenile onset-disease is associated with more severe organ involvement (including renal and CNS disease), increased disease activity, presence of greater damage at the time of diagnosis, and higher steroid burden, contributing to the increased morbidity and mortality when compared to adult-onset SLE³⁻⁵. Even within the JSLE population, very early disease onset (before the 5th birthday) may be associated with atypical presentations (including fewer autoantibodies), more severe disease courses and poor prognosis^{1, 6-8}. However, assumptions on variable disease presentation and progression within different JSLE age sub-groups are generally based on case reports, case series or relatively small cohorts^{7, 8} and currently lack scientific evidence from longitudinal national or international studies.

Preliminary datasets suggest that clinical differences may reflect variable pathomechanisms and that patients with JSLE may have increased genetic burden when compared to individuals with adult-onset disease, contributing to early disease onset and more severe presentations^{1, 9}. Very early disease onset, atypical disease presentation and severe manifestations may be the result of (very rare) disease-causing mutations in single genes or the combination of multiple genomic variants that individually increase an individual's risk for the development of SLE^{1, 9-11}. To date, evidence still remains weak and it is largely unclear whether distinct clinical and laboratory differences exist between age groups within the paediatric population^{1, 12, 13}.

This study aimed to assess if there are differential clinical and laboratory characteristics in patients presenting with JSLE at different ages, sub-dividing patients into three groups: pre-pubertal (≤ 7 years), peri-pubertal (8-13 years) or adolescence (14-18 years). To achieve this, prospectively collected data from a national cohort of JSLE patients (the UK JSLE Cohort Study) was interrogated.

Methods

Patients

Participants of the UK JSLE Cohort Study¹⁴, followed between 2006-2018, aged ≤ 16 years at the time of diagnosis and with ≥ 4 American College of Rheumatology (ACR) classification criteria for SLE¹⁵ were included in this study. Participants were excluded from the study if they did not have a diagnosis date recorded, as this precluded them from being categorized on the basis on their age at disease-onset (pre-pubertal (≤ 7 years), peri-pubertal (8-13 years) or adolescent (14-18 years)). Patient/family reported ethnicity information was collected using the UK National Census categorisations¹². Data of patients who were of mixed race were grouped with those of the associated ethnic minority group (e.g. Asian if mixed Asian and Caucasian race).

Data collected

The following clinical and laboratory data were collected: 1) total ACR score with its individual domains¹⁵; 2) anti-nuclear antibody (ANA) positivity and titre; 3) Systemic Lupus International Collaborating Clinics standardised damage index (SLICC-SDI) score¹⁶; 4) paediatric British Isles Lupus Assessment Grade 2004 numerical scores (pBILAG2004) with individual organ/system domains (alphabetical score A-E)¹⁷; 5) key laboratory findings, including haemoglobin levels, white cell count and differentiation, platelets, erythrocyte sedimentation rate (ESR), complement levels (C3, C4) and anti-double-stranded DNA (dsDNA) titres

The SLICC-SDI tool records permanent damage that occurs as a result of SLE activity, and is composed of 41 different components¹⁶. The pBILAG2004 score is a composite disease activity measure focusing on nine organ/system domains (constitutional, mucocutaneous, neurological, musculoskeletal, cardiovascular/respiratory, renal, gastrointestinal, ophthalmic and haematological). Each organ domain is graded A-E and defined as follows; pBILAG2004 grade A/B: severe and moderate disease respectively, grade C patients: mild/improving renal disease, grade D: inactive disease but previous system involvement, grade E: system has never been involved^{18, 19}. For each organ/system domain, an alphabetical (A-E grade) is determined, equating to a numerical value for each organ/system domain. These can be combined to give the total numerical pBILAG2004 score¹⁷. Within these analyses, presence of pBILAG2004 domains A and B was taken to signify active organ/system involvement, in-keeping with previous studies¹⁷. All data items (1-5 listed above) were collected at the time of initial diagnosis. At the patients' last follow-up visit, data from items 1 and 3 were collected.

Furthermore, data from item 4 were collected from patients as their cumulative maximum disease activity level (for each individual organ/system domain) throughout the disease course.

Statistical analysis

Laboratory findings, total number of ACR criteria, SLICC and pBILAG2004 scores were compared between groups using Kruskal Wallace tests. Median values and interquartile ranges (IQRs) are displayed within tables. Categorical pBILAG2004 domain data is presented as a percentage of patients with active organ/system involvement for each age group along with 95% confidence intervals. Individual domains of the pBILAG2004 score were compared between groups using Chi-square and Fisher's exact tests. Analyses were completed using SPSS software, version 25 (IBM SPSS).

Power analysis revealed that the three patient groups should all have approximately 700 patients each to reach sufficient statistical power. Limited by the rarity of JSLE and resulting number of patients included in the national UK JSLE cohort study since 2006, these numbers are extremely difficult to obtain in national or even international cohorts. Thus, p values of statistical tests should be interpreted with caution, based upon the limited statistical power of this study.

Results

Demographics

A total of 418 eligible patients enrolled in the UK JSLE Cohort Study were included in this study; five JSLE patients were excluded due to unknown age at diagnosis. The mean age at diagnosis was 12.1 years (range: 0.17-17.91), with 43/418 (10.3%) participants presenting in the pre-pubertal period, 240/418 (57.4%) were peri-pubertal, and 135/418 (32.3%) were in the adolescent age group. The overall female:male ratio was 5.4:1 and increased with age (pre-pubertal=3.3:1; peri-pubertal=5.24:1; adolescent=7.25:1). No statistically significant differences were demonstrated between groups in relation to ethnicity ($p>0.05$) (supplementary table 1).

Clinical features

At diagnosis, adolescent JSLE patients exhibited higher median ACR scores when compared to younger JSLE patients (pre-pubertal: median 4[IQR 4-5] vs. peri-pubertal: 4[4-5] vs. adolescent: 5[4-6], $p=0.004$). Similarly, pBILAG2004 disease activity scores were higher in

newly diagnosed adolescent JSLE patients (pBILAG2004: 9[4-20]) when compared to younger JSLE patients (pre-pubertal: 7[3-13]; peri-pubertal: 7[3-14], $p = 0.015$) (Table 1). Furthermore, adolescents with a new diagnosis of JSLE exhibited more activity in the following pBILAG domains when compared to new peri-pubertal and pre-pubertal JSLE patients: mucocutaneous ($p=0.025$), musculoskeletal ($p=0.029$), cardiorespiratory ($p=0.001$) and renal ($p=0.027$) (Table 1).

At the time of last follow-up differences were not apparent between age groups in terms of total ACR scores (median of 5[IQR 4-7] in all groups) and the proportion of patients that were ANA positive. Over the disease course active organ/system involvement (as defined by the pBILAG2004 score) also did not differ significantly between age groups (Table 2). There was little variance in SLICC-SDI defined damage at diagnosis ($p=0.410$) or last follow-up ($p=0.284$) between age groups (Tables 1 and 2).

Laboratory features

Laboratory findings varied between JSLE patients from different age groups at diagnosis (Table 3). White blood cell and platelet counts reduced with growing age across the JSLE cohort; with pre-pubertal patients exhibiting median white cell counts of $6.7 \times 10^9/L$ [4.69 – 9.53] vs. $6.09 \times 10^9/L$ [4.16-8.67] in peri-pubertal vs. $4.69 \times 10^9/L$ [3.7-6.54] in the adolescent age group ($p=0.002$). Median platelet counts were within the normal range, but followed a similar pattern to the white cell count, with $293 \times 10^9/L$ [212-426] in the pre-pubertal group vs. $271 \times 10^9/L$ [191-388] in the peri-pubertal vs. $242 \times 10^9/L$ [168-298] in the adolescent group ($p=0.004$). Median levels of haemoglobin ($p=0.404$) and ESR ($p=0.2$) did not differ between age groups (Table 3).

Serum complement is a measure of disease activity in SLE as it indicates activation and consumption of complement components²⁰. Median complement levels differed significantly between age groups, with higher complement levels in younger patients (C3: $0.95g/L$ [0.73-1.11] in pre-pubertal patients vs. $0.81g/L$ [0.50-1.22] in peri-pubertal vs. $0.69g/L$ [0.28-0.98] in adolescent patients ($p=0.002$); C4: $0.13g/L$ [0.08-0.28] in pre-pubertal patients vs. $0.11g/L$ [0.06-0.19] peri-pubertal patients vs. $0.08g/L$ [0.04-0.14] in adolescent patients ($p=0.002$)) (Table 3).

In the UK JSLE cohort, patients with disease-onset during adolescence were more frequently ANA positive 131/135 (97.0%) at diagnosis, when compared to the other age groups; 37/43 (86.0%) with pre-pubertal onset, and 223/240 (92.9%) in peri-pubertal onset ($p=0.034$). Anti-dsDNA antibody titres were higher in older patients than younger patients; pre-pubertal onset 15 IU/L [0.25-89] vs. 67 IU/L [19-200] in peri-pubertal group vs. 111 IU/L [15-300] in adolescents ($p=0.001$) (Table 3).

Discussion

While clinical and laboratory differences between JSLE and adult-onset SLE have been acknowledged⁸, only few short reports discuss differences within the paediatric age group^{12, 13}. The 418 JSLE patients included in this study allow for more reliable assessment of clinical and laboratory features between the paediatric age groups. When compared to younger children, adolescents exhibit an increased number of ACR criteria, and show typical autoantibody patterns (ANA and anti-dsDNA positivity), haematological involvement (leukopenia, thrombocytopenia) and immunological characteristics (hypocomplementaemia) reflecting “classical” SLE. Of note, adolescents also present with higher disease activity at diagnosis when compared to younger children (total numerical BILAG score; $p=0.015$). At diagnosis, differences were also seen in the organ domains involved across age groups, including increased mucocutaneous, musculoskeletal, cardiorespiratory and renal system involvement in adolescents when compared to other age groups. Notably, previous studies did not consider pre-pubertal (≤ 7) JSLE patients as a distinct age group^{3, 6-8}.

One of the most interesting differences between JSLE patients of three age groups relates to laboratory findings. Patients diagnosed in early childhood (≤ 7 years) had lower rates of ANA positivity; with 14% of the pre-pubertal JSLE patients being ANA negative vs. 3% of the adolescent JSLE group ($p=0.034$). Pre-pubertal children also displayed lower median anti-dsDNA titres, than the other age groups ($p=0.001$). These laboratory differences may reflect differences in pathophysiology at varying ages, and a potentially more “innate” disease phenotype in at least a subset of early-onset JSLE patients¹.

Of note, previous studies failed to identify serological differences between paediatric and adult SLE populations, which may be due to them not discriminating between age groups within the JSLE population^{3, 8, 21}. This potential explanation is supported by the observation that differences in immunological patterns (ANA positivity) disappeared by the time of last follow

up prior to transition into adult care ($p=0.559$). Most patients who were initially autoantibody negative in the pre-pubertal (11.7%) and peri-pubertal age groups (2.9%), eventually developed ANA positivity (pre-pubertal group: 14% at diagnosis vs. 2% at last follow up) between the time of initial diagnosis and last follow-up. Why this is can only be speculated, but it has previously been discussed that early-onset JSLE patients, who may have a higher genetic risk when compared to older SLE patients or have a more monogenic disease phenotype, can develop autoantibodies over time as a result of tissue damage and subsequent presentation of physiologically nuclear components to the immune system^{1, 22}.

This study also found increased frequencies of ANA positivity to coincide with an increased prevalence of likely autoantibody-mediated symptoms, e.g. renal, musculoskeletal and haematological anomalies (thrombocytopenia, lymphopenia and low complement levels, all $p<0.05$). Autoantibodies (particularly anti-dsDNA antibodies) indeed contribute to renal disease and immune complex deposition, which may also partially cause the pathologically reduced complement levels observed with increasing age^{2, 20, 23}. Also, increased musculoskeletal involvement in adult-onset SLE vs. JSLE patients has been previously demonstrated^{3, 8, 13}. Tavangar-Rad *et al.* studied 120 Iranian children with JSLE and compared age groups in a similar way to the current study (<7, 7-14, and >14 years) and reported more joint involvement with increasing age¹³. While it remains unclear why this is, musculoskeletal involvement is another example of a clinical feature that may be auto-antibody driven, thus becoming more prevalent with advancing age at presentation.

Findings from this study also suggest that disease activity within the paediatric age group may (at diagnosis) be more severe in individuals diagnosed in adolescence, while disease severity increases over time in children diagnosed at a younger age. This is indicated by comparable disease activity and damage scores at last follow up. Based on variable clinical patterns over time that coincide with increased disease activity, autoantibodies, immune complex deposition, and complement activation may likely be involved in this process^{2, 20, 23}. Differences between the present study and previous reports suggesting increased disease severity in very early-onset SLE when compared to “older” children with JSLE, may be due to the character of previous reports¹². Small case series and individual case reports tend to over-report particularly severe, interesting and/or complicated presentations and disease courses.

The absence of ANA antibodies in 14% of pre-pubertal JSLE patients is interesting when considering the classification criteria for SLE. Recently proposed “new” ACR/EULAR criteria for SLE include ANA titres of $\geq 1:80$ as entry criterion²⁴. While application of these criteria would only affect a relatively small number of peri-pubertal or adolescent JSLE patients, 14% of patients with early disease-onset could potentially remain without a diagnosis, as classification criteria are frequently (incorrectly) used by colleagues (not necessarily specialized in paediatric rheumatology) to diagnose SLE and refer to tertiary care. One may argue that very early disease-onset in the absence of autoantibodies can indicate genetic conditions (“monogenic SLE-like disease”, such as complement deficiencies, primary type I interferonopathies) and that it is beneficial for patients to not be classified as “classical” SLE. However, this may result in diagnostic delays and that young patients not being seen by paediatric rheumatologists²⁵.

Although this study involves one of the largest national JSLE cohorts available, it is still limited by JSLE being a rare disease and patient numbers. A power analysis performed prior to this study suggested that around 700 patients were required per group for the analysis to be statistically reliable. Since the UK JSLE cohort study is the largest JSLE cohort across Europe and one of the largest in the world, this limitation can unfortunately currently not be addressed. International collaboration is therefore warranted in the future. The variable duration of follow-up from initial evaluation to last visit between the three age groups may also be seen as a potential limitation. This was mainly caused by the time of transition to adult care.

Conclusion

This is the largest study to date comparing clinical and laboratory features of JSLE patients diagnosed during the pre- (≤ 7), peri-pubertal (8-13) and adolescent (14-18) periods. Distinct clinical and laboratory differences between age groups support the hypothesis that variable pathomechanisms may contribute to differences in clinical presentations, treatment responses and disease outcomes not only between adult and paediatric patients but also within the cohort of JSLE patients. Based on the presence of autoantibodies and higher prevalence of antibody-mediated features (including thrombocytopenia, lymphopenia, hypocomplementaemia), adaptive immune mechanisms may play an increasing role with growing age. Though the largest study of its kind, it is still limited by patient numbers, due to the rarity of JSLE. Thus, international collaborations are warranted to address age-specific differences in JSLE in more detail.

Conflict of interest

The authors declare no conflict of interest relevant to this work.

Acknowledgements

The authors acknowledge all participants of the study and their families. Authors would also like to acknowledge specialist pediatric centers who support the collection of data for the UKJSLE cohort (<https://www.liverpool.ac.uk/translational-medicine/research/ukjsle/jsle/>). Special recognition also goes to Duncan Appleby for database and information technology support and Carla Roberts for co-ordination of the UK JSLE Cohort study. This work was supported by the UK's Experimental Arthritis Treatment Centre for Children (<https://www.liverpool.ac.uk/eatc-for-children/>; supported by Versus Arthritis, Alder Children's NHS Foundation Trust, the National Institute for Health Research (NIHR) Alder Hey Clinical Research Facility, the Alder Hey Charity, and the University of Liverpool) and partially carried out at the National Institute for Health Research (NIHR) Alder Hey Clinical Research Facility. Author JM would also like to thank Steven Lane (University of Liverpool, Department of Biostatistics) for his advice on statistical analysis, and Dr A Frankel (Imperial College Healthcare, Department of Nephrology) who's passion for research and bedside manner inspired him to pursue a career in medicine.

Funding

This work was supported by Lupus UK, who provide financial support for co-ordination of the UK JSLE Cohort Study [grant numbers: LUPUS UK: JXR10500, JXR12309]. The bodies detailed above were not involved in the design, collection, analysis, and interpretation of data; in the writing of the manuscript; and in the decision to submit the manuscript for publication.

Authors contributions

EMDS, CMH and JM were involved in study design. JM performed statistical analysis. MWB is Chief Investigator of the UK JSLE Cohort Study and supported all steps of the work presented. All authors participated in the interpretation of data, and have since revised drafts. They have also read and given final approval for the version to be published.

1. References

1. Hedrich CM, Smith EMD and Beresford MW. Juvenile-onset systemic lupus erythematosus (jSLE) - Pathophysiological concepts and treatment options. *Best Pract Res Clin Rheumatol*. 2017; 31: 488-504.
2. Tsokos GC. Systemic lupus erythematosus. *N Engl J Med*. 2011; 365: 2110-21.
3. Ambrose N, Morgan TA, Galloway J, et al. Differences in disease phenotype and severity in SLE across age groups. *Lupus*. 2016.
4. Brunner HI, Gladman DD, Ibañez D, Urowitz MD and Silverman ED. Difference in Disease Features Between Childhood-Onset and Adult-Onset Systemic Lupus Erythematosus. *Arth & Rheum*. 2008; 58: 556-62.
5. Tucker LB, Uribe AG, Fernandez M, et al. Adolescent onset of lupus results in more aggressive disease and worse outcomes: results of a nested matched case-control study within LUMINA, a multiethnic US cohort (LUMINA LVII). *Lupus*. 2008; 17: 314-22.
6. Chen YM, Lin CH, Chen HH, et al. Onset age affects mortality and renal outcome of female systemic lupus erythematosus patients: a nationwide population-based study in Taiwan. *Rheumatology (Oxford)*. 2014; 53: 180-5.
7. Descloux E, Durieu I, Cochat P, et al. Influence of age at disease onset in the outcome of paediatric systemic lupus erythematosus. *Rheumatology (Oxford)*. 2009; 48: 779-84.
8. Fonseca R, Aguiar F, Rodrigues M and Brito I. Clinical phenotype and outcome in lupus according to age: a comparison between juvenile and adult onset. *Reumatol Clin*. 2018; 14: 160-3.
9. Webb R, Kelly JA, Somers EC, et al. Early disease onset is predicted by a higher genetic risk for lupus and is associated with a more severe phenotype in lupus patients. *Ann Rheum Dis*. 2011; 70: 151-6.
10. Alperin JM, Ortiz-Fernandez L and Sawalha AH. Monogenic Lupus: A Developing Paradigm of Disease. *Front Immunol*. 2018; 9: 2496.
11. Lo MS and Tsokos GC. Recent developments in systemic lupus erythematosus pathogenesis and applications for therapy. *Curr Opin Rheumatol*. 2018; 30: 222-8.
12. Hui-Yuen JS, Imundo LF, Avitabile C, Kahn PJ, Eichenfield AH and Levy DM. Early versus later onset childhood-onset systemic lupus erythematosus: Clinical features, treatment and outcome. *Lupus*. 2011; 20: 952-9.
13. Tavangar-Rad F, Ziaee V, Moradinejad MH and Tahghighi F. Morbidity and Mortality in Iranian Children with Juvenile Systemic Lupus erythematosus. *Iran J Pediatr*. 2014; 24: 365-70.
14. Watson L, Leone V, Pilkington C, et al. Disease activity, severity, and damage in the UK Juvenile-Onset Systemic Lupus Erythematosus Cohort. *Arthritis Rheum*. 2012; 64: 2356-65.
15. Hochberg MC. Updating the American College of Rheumatology revised criteria for the classification of systemic lupus erythematosus. *Arthritis Rheum*. 1997; 40: 1725.
16. Gladman D, Ginzler E, Goldsmith C, et al. The development and initial validation of the Systemic Lupus International Collaborating Clinics/American College of Rheumatology damage index for systemic lupus erythematosus. *Arthritis Rheum*. 1996; 39: 363-9.
17. Stoll T, Stucki G, Malik J, Pyke S and Isenberg DA. Association of the Systemic Lupus International Collaborating Clinics/American College of Rheumatology Damage Index with measures of disease activity and health status in patients with systemic lupus erythematosus. *J Rheumatol*. 1997; 24: 309-13.
18. Marks SD, Pilkington C, Woo P and Dillon MJ. The use of the British Isles Lupus Assessment Group (BILAG) index as a valid tool in assessing disease activity in childhood-onset systemic lupus erythematosus. *Rheumatology (Oxford)*. 2004; 43: 1186-9.
19. Isenberg DA, Rahman A, Allen E, et al. BILAG 2004. Development and initial validation of an updated version of the British Isles Lupus Assessment Group's disease activity index for patients with systemic lupus erythematosus. *Rheumatology (Oxford)*. 2005; 44: 902-6.

20. Ricklin D, Reis ES and Lambris JD. Complement in disease: a defence system turning offensive. *Nat Rev Nephrol.* 2016; 12: 383-401.
21. Carreno L, Lopez-Longo FJ, Monteagudo I, et al. Immunological and clinical differences between juvenile and adult onset of systemic lupus erythematosus. *Lupus.* 1999; 8: 287-92.
22. Hedrich CM. Shaping the spectrum - From autoinflammation to autoimmunity. *Clin Immunol.* 2016; 165: 21-8.
23. Alba P, Bento L, Cuadrado MJ, et al. Anti-dsDNA, anti-Sm antibodies, and the lupus anticoagulant: significant factors associated with lupus nephritis. *Ann Rheum Dis.* 2003; 62: 556-60.
24. Aringer M, CK, Brinks R., Boumpas D., Daikh D., Jayne D., Kamen D.L., Mosca M., Ramsey-Goldman R., Smolen J.S., Wofsy D., Diamond B., Jacobsen S., McCune W.J., Ruiz-Irastorza G., Schneider M., Urowitz M., Bertsias G., Hoyer B., Leuchten N., Tani C., Tedeschi S.K., Touma Z., Anic B., Assan F., Chan T.M., Clarke A.E., Crow P., Czirják L., Doria A., Graninger W., Halda-Kiss B., Hasni S.A., Izmirly P.M., Jung M., Kumanovics G., Mariette ., Padjen I., Pego-Reigosa J.M., Romero-Díaz J., Rúa-Figueroa I., Seror R., Stummvoll G., Tanaka Y., Tektonidou M., Vasconcelos C., Vital E.M., Wallace D.J., Yavuz S., Naden R.P., Dörner T., Johnson S. Validation of New Systemic Lupus Erythematosus Classification Criteria. *Arthritis Rheumatol* 2018; 70 (suppl 10). .
25. Johnson SR, Khanna D, Daikh D, et al. Use of Consensus Methodology to Determine Candidate Items for Systemic Lupus Erythematosus Classification Criteria. *J Rheumatol.* 2018.

Supplementary table 1 – Ethnicity of participants

Ethnicity	Pre-pubertal disease-onset (n=43)	Peri-pubertal disease-onset (n=240)	Adolescence (n=135)	P value
British or Irish	19/43 (44.2%) [29.3%, 59.0%]	104/240 (43.3%) [37.0%, 49.6%]	77/135 (57.0%) [48.7%, 65.4%]	0.100
Asian	13/43 (30.2%) [16.5%, 44.0%]	73/240 (30.4%) [24.6%, 36.2%]	41/135 (30.4%) [22.6%, 38.1%]	
African/Caribbean	8/43 (18.6%) [7.0%, 30.2%]	45/240 (18.8%) [13.8%, 23.7%]	14/135 (10.4%) [5.2%, 15.5%]	
Other Caucasian origin	1/43 (2.3%) [-2.2%, 6.8%]	7/240 (2.9%) [0.8%, 5.0%]	2/135 (1.5%) [-0.6%, 3.5%]	
Any other mixed/black background	0/43 (0%) [0%, 0%]	7/240 (2.9%) [0.8%, 5.0%]	0/135 (0%) [0%, 0%]	
Ethnicity not stated	2/43 (4.7%) [-1.6%, 10.9%]	4/240 (1.67%) [0.0%, 3.3%]	1/135 (0.7%) [-0.7%, 2.2%]	

Ethnicities were compared between age groups using the Chi Square test of independence. For each ethnic group, the total number of patients is provided along with the percentage (in curved brackets), and 95% confidence intervals for the percentage [in square brackets].

Table 1 – Clinical features of JSLE subgroups at diagnosis

Item	Pre-pubertal disease-onset (n=43)	Peri-pubertal disease-onset (n=240)	Adolescence (n=135)	P value
<i>Female:Male ratio</i>	<i>3.3:1</i>	<i>5.24:1</i>	<i>7.25:1</i>	<i>0.347</i>
<i>Total ACR score</i>	<i>4 [4-5]</i>	<i>4 [4-5]</i>	<i>5 [4-6]</i>	<i>0.004</i>
SLICC-SDI	0 [0]	0 [0]	0 [0]	0.410
<i>Total numerical pBILAG2004 score</i>	<i>7 [3-13]</i>	<i>7 [3-14]</i>	<i>9 [4-20]</i>	<i>0.015</i>
Active organ/system involvement at diagnosis (pBILAG2004 defined)				
• Constitutional	13 (30.2%) [16.5%, 43.9%]	67 (27.9%) [22.2%, 33.6%]	51 (37.8%) [29.6%, 46.0%]	0.140
• <i>Mucocutaneous</i>	<i>19 (44.2%)</i> <i>[29.5%, 59.0%]</i>	<i>78 (32.5%)</i> <i>(26.6%, 38.4%)</i>	<i>62 (45.9%)</i> <i>[37.5%, 54.3%]</i>	<i>0.025</i>
• Neuropsychiatric	6 (40.0%) [25.4%, 54.6%]	20 (8.3%) [4.8%, 11.8%]	14 (10.4%) [5.3%, 15.3%]	0.477
• <i>Musculoskeletal</i>	<i>7 (16.3%)</i> <i>[5.3%, 27.3%]</i>	<i>66 (27.5%)</i> <i>[21.9%, 33.1%]</i>	<i>49 (36.3%)</i> <i>[28.2%, 44.4%]</i>	<i>0.029</i>
• <i>Cardiorespiratory</i>	<i>4 (9.3%)</i> <i>[0.6%, 18%]</i>	<i>18 (7.5%)</i> <i>[4.2%, 10.8%]</i>	<i>27 (20%)</i> <i>[13.3%, 26.7%]</i>	<i>0.001</i>
• Gastrointestinal	2 (4.7%) [0%, 11%]	15 (6.3%) [3.2%, 9.4%]	4 ((3.0%) [0.1%, 5.9%]	0.442
• Ophthalmic	1 (2.3%) [0%, 6.8%]	2 (0.8%) [0%, 1.9%]	1 (0.7%) [0%, 2.1%]	0.548
• <i>Renal</i>	<i>9 (20.9%)</i> <i>[8.7%, 33.1%]</i>	<i>73 (30.4%)</i> <i>[24.6%, 36.2%]</i>	<i>55 (40.7%)</i> <i>[32.4%, 49.0%]</i>	<i>0.027</i>
• Hematological	11 (25.6%) [12.6%, 38.6%]	58 (24.1%) [18.7%, 29.5%]	37 (27.4%) [19.9%, 34.9%]	0.786

Total ACR, SLICC-SDI, and pBILAG2004 scores, and key laboratorial findings are reported as median values and interquartile ranges [in square brackets]. For individual pBILAG2004

organs/systems involved, the total number of patients with active involvement (defined as pBILAG2004 domain score of A or B within a given organ domain/system) is provided along with the percentage (in curved brackets) and 95% confidence intervals for the percentage [in square brackets].

Table 2 – Clinical features of the different age groups over time

Items on last follow up	Pre-pubertal disease-onset (n=43)	Peri-pubertal disease-onset (n=240)	Adolescence (n=135)	P value
<i>Length of follow up in years</i>	6 [3-9]	4 [2-6]	2 [1-4]	<0.001
Total ACR score	5 [4-7]	5 [4-7]	5 [4-7]	0.686
SLICC-SDI score	0 [0-1]	0 [0-1]	0 [0-1]	0.284
ANA positivity	42 (97.7%) [93.2%, 102.2%]	230 (95.8%) [93.3%, 98.3%]	132 (97.8%) [95.3%, 100.0%]	0.559
pBILAG2004 defined organ/system domain involvement throughout the disease course				
Constitutional	18 (41.9%) [27.2%, 56.6%]	98 (40.8%) [34.1%, 46.5%]	56 (41.5%) [33.2%, 49.8%]	0.988
Mucocutaneous	33 (76.7%) [64.1%, 89.3%]	157 (65.4%) [59.4%, 71.4%]	90 (66.7%) [58.7%, 74.7%]	0.346
Neuropsychiatric	11 (25.6%) [12.6%, 38.6%]	57 (27.9%) [22.2%, 33.6%]	28 (20.7%) [13.9%, 27.5%]	0.731
Musculoskeletal	18 (41.9%) [16.5%, 43.9%]	121 (50.4%) [44.1%, 56.7%]	73 (54.1%) [45.7%, 62.5%]	0.374
Cardiorespiratory	13 (30.2%) [16.5%, 43.9%]	46 (19.2%) [14.2%, 24.2%]	35 (25.9%) [18.5%, 33.3%]	0.141
Gastrointestinal	8 (18.6%) [7.0%, 30.2%]	28 (11.7%) [7.6%, 15.8%]	10 (7.4%) [3.0%, 11.8%]	0.107
Ophthalmic	3 (7.0%) [0%, 14.6%]	12 (5.0%) [2.2%, 7.8%]	4 (3.0%) [0.1%, 5.9%]	0.467
Renal	28 (65.1%) [50.9%, 79.3%]	153 (63.8%) [57.7%, 69.9%]	94 (69.6%) [61.8%, 77.4%]	0.513
Hematological	26 (60.5%) [45.9%, 75.1%]	114 (47.5%) [41.2%, 53.8%]	55 (40.7%) [32.4%, 49.0%]	0.072

Total ACR, SLICC-SDI, pBILAG2004 scores are reported as median values and interquartile ranges. For individual pBILAG2004 domains, the total number of patients with activity involvement (defined as a pBILAG2004 domain score of A or B in a given organ domain/system) are provided along with percentage (in curved brackets), and 95% confidence

intervals for the percentage [square brackets]. SLICC-SDI and ACR scores are provided from the last follow-up visit.

Table 3 – Laboratory features of JSLE subgroups at diagnosis

Key laboratory findings	Pre-pubertal disease-onset (n=43)	Peri-pubertal disease-onset (n=240)	Adolescence (n=135)	P value
Haemoglobin level (g/dL)	11 [9-11.9]	11.3 [9.9-12.6]	11.08 [9.7-12.53]	0.404
White cell count (x 10⁹/L)	6.7 [4.69-9.53]	6.09 [4.16-8.67]	4.69 [3.7-6.54]	0.002
Platelets (x 10⁹/L)	293 [212-426]	271 [191-338]	242 [168-298]	0.004
ESR (mm/hr)	18 [11-72]	36 [12-76]	42.5 [19-86.75]	0.200
C3 median (g/L)	0.95 [0.73-1.11]	0.81 [0.50-1.22]	0.69 [0.28-0.98]	0.002
C4 median (g/L)	0.13 [0.08-0.28]	0.11 [0.06-0.19]	0.08 [0.04-0.14]	0.002
ANA positive	37 (86.0%) [80.7%, 91.3%]	223 (92.9%) [89.7%, 96.2%]	131 (97.0%) [94.1%, 99.9%]	0.034
ANA titre median	1:640 [1:320-1:960]	1:640 [1:320-1:1280]	1:640 [1:320-1:2560]	0.565
dsDNA levels (IU/L)	15 [0.25-89]	67 [19-200]	111 [15-300]	0.001

Haemoglobin, white cell count, platelets, ESR, C3, C4, ANA titre and dsDNA titre are reported as median values and interquartile ranges [in square brackets].