

IMPACT OF ACTIVE LUPUS NEPHRITIS ON THE QUALITY OF LIFE OF PATIENTS FROM A LATIN AMERICAN LUPUS COHORT

Romina Nieto; Rosana Quintana; Diana Carolina Fernández Ávila; Rosa Serrano; Guillermina Harvey; Lucia Hernandez; Karen Roberts; Erika S. Palacios Santillan; Nidia Meras; Cintia Otaduy; Elisa Novatti; Valeria Arturi; Boris Kisluk; Luciana González Lucero; Eduardo Kerzberg; Nicolás Pérez; Cecilia Pisoni; Paola Pirruccio; María E. Crespo; Ana Carolina Montandon; Andrese A. Gasparin; Angela Duarte; Laissa C. Alves Alvino; Eloisa Bonfa; Emily Figuereido Neves; Lucas Victoria de Oliveira Martins; Iris Guerra; Milena Mimica Davet; Lizeth De la Hoz Rueda; Andrés Cadena Bonfanti; Roberth Rivera; Paola Coral Alvarado; John Fredy Jaramillo; José Martínez; Mario Pérez Cristóbal; Eduardo Martin Nares; Yaneli Juárez-Vicuña; Yelitza González Bello; Octavio González; Leonardo R. Aguilar Rivera; Margarita Duarte; Patricia Langjarth; Wilkerson Pérez Medina; Armando Calvo; Teresandris Polanco; Carina Pizzarossa; Gonzalo Silveira, Cristina Reategui; Graciela S. Alarcón; Urbano Sbarigia; Federico Zazzetti; Ashley Orillion; Guillermo Pons-Estel^{*}; Bernardo Pons-Estel; on behalf of Grupo Latino Americano de Estudio del Lupus (GLADEL)

BACKGROUND

1510

- Systemic lupus erythematosus (SLE) has a negative impact on patients' quality of life and is associated with an increased economic
- Lupus nephritis (LN) is present in up to 65% of patients with SLE, and approximately 13% of patients with LN will progress to end-stage renal disease within 10 years³
- There are limited data on the impact of active LN on the health-related quality of life (HRQoL) of patients with LN and on whether achieving renal clinical response with current treatments can improve HRQoL
- The Latin American Lupus Study Group (Grupo Latino Americano De Estudio del Lupus [GLADEL]) was created to explore disease features, the clinical course, and outcomes in Latin American patients with SLE⁴

- GLADEL 2.0 is an observational prevalent and incident cohort that was initiated in 2019 in Argentina, Brazil, Chile, Colombia, the Dominican Republic, Ecuador, Mexico, Paraguay, Peru, and Uruguay⁵

• The study aimed to evaluate HRQoL in patients with active LN in the GLADEL 2.0 cohort at baseline and 12 months after treatment and to assess the impact of renal response on HRQoL

METHODS

Study population

- A total of 44 centers from 10 Latin American countries enrolled patients aged ≥18 years who fulfilled the 1982/1997 American College of Rheumatology (ACR) and/or 2012 Systemic Lupus International Collaborating Clinics (SLICC) classification criteria
- Patients were categorized into 4 subsets according to the presence of LN, as follows:
- Group I: no LN
- Group II: prevalent and inactive LN
- Group III: prevalent and active LN
- Group IV: incident LN with an onset of <3 months and renal biopsy
- For this analysis, patients in Groups II, III, and IV with sufficient follow-up data at 12 months were included

Study assessments

- Baseline demographics, clinical manifestations, and disease activity based on the Systemic Lupus Erythematosus Disease Activity Index (SLEDAI) and SLICC/ACR Damage Index were assessed
- At baseline, HRQoL was assessed based on the Lupus Quality of Life (LupusQoL) questionnaire and was stratified by the presence of active or inactive LN
- At the 12-month follow-up, LupusQoL responses were assessed in patients with active LN and stratified by their renal response
- Renal response at the 12-month follow-up was assessed and categorized according to European Alliance of Associations for Rheumatology/Kidney Disease: Improving Global Outcomes criteria, as follows:
- Complete response (CR): <0.5 g/g reduction in proteinuria, measured as urine protein-to-creatinine ratio (UPCR) from a 24-hour urine collection
- Partial response (PR): ≥50% reduction in proteinuria, measured as UPCR from a 24-hour urine collection
- No response (NR): <50% reduction in proteinuria

Statistical analysis

- Continuous variables were reported as medians (interquartile ranges), and categorical variables were reported as frequencies (percentages)
- Baseline LupusQoL responses were compared between patients in Groups II and Groups III + IV using a Kruskal-Wallis test
- LupusQoL responses at the 12-month follow-up were compared between patients with NR and PR + CR using a Kruskal-Wallis test

RESULTS

Patient characteristics

- Of the 1081 patients included in the GLADEL 2.0 cohort, 651 with a history of LN were evaluated (423 with active LN and 228 with inactive LN)
- Patients with active LN were predominantly female (369/423 [87.2%]) and had a lower socioeconomic status, a higher unemployment rate, and a higher SLEDAI score than patients with inactive LN

Baseline LupusQoL responses

• Patients with active LN had significantly worse baseline LupusQoL responses across all domains compared to those with inactive LN (**Table 1**)

TABLE 1: LupusQoL responses by domain at baseline in patients with inactive LN versus active + incident LN

LupusQoL domain, median (IQR)	Total (N = 651)	Group II: prevalent and inactive LN (n = 228)	Groups III + IV: prevalent and active + incident LN (n = 423)	<i>P</i> value
Physical health	78.1 (56.3-93.8)	87.5 (71.9-96.9)	71.9 (50.0-87.5)	<0.0001
Pain	83.3 (58.3-100.0)	91.7 (75.0-100.0)	75.0 (50.0-91.7)	<0.0001
Planning	83.3 (58.3-100.0)	91.7 (75.0-100.0)	75.0 (41.7-91.7)	<0.0001
Intimate relationship	100.0 (62.5-100.0)	100.0 (62.5-100.0)	75.0 (50.0-100.0)	0.001
Burden to others	58.3 (25.0-83.3)	75.0 (41.7-91.7)	50.0 (16.7-75.0)	<0.0001
Emotional health	75.0 (50.0-87.5)	83.3 (62.5-95.8)	66.7 (45.8-83.3)	<0.0001
Body image	80.0 (55.0-100.0)	90.0 (70.0-100.0)	75.0 (50.0-95.0)	<0.0001
Fatigue	68.8 (43.8-87.5)	81.3 (56.3-93.8)	62.5 (37.5-81.3)	<0.0001

IQR=Interquartile range; LN=Lupus nephritis; LupusQoL=Lupus Quality of Life questionnaire.

Impact of achieving renal response on LupusQoL responses

• At the 12-month follow-up, no differences were found between patients who achieved CR or PR and those who did not (Table 2)

TABLE 2: Impact of achieving renal response at 12 months on LupusQoL responses in patients with active LN

LupusQoL domain, median (IQR)	Total (N = 328)	NR (n = 113)	PR + CR (n = 215)	P value		
Physical health	84.4 (68.8-93.8)	81.3 (65.6-93.8)	84.4 (68.8-93.8)	0.777		
Pain	83.3 (66.7-100.0)	83.3 (66.7-100.0)	83.3 (75.0-100.0)	0.407		
Planning	83.3 (66.7-100.0)	83.3 (66.7-100.0)	83.3 (66.7-100.0)	0.704		
Intimate relationship	87.5 (75.0-100.0)	87.5 (75.0-100.0)	87.5 (62.5-100.0)	0.948		
Burden to others	58.3 (33.3-83.3)	58.3 (33.3-83.3)	58.3 (33.3-75.0)	0.657		
Emotional health	77.1 (62.5-91.7)	75.0 (50.0-91.7)	79.2 (66.7-87.5)	0.310		
Body image	85.0 (70.0-95.0)	85.0 (65.0-100.0)	85.0 (70.0-95.0)	0.629		
Fatigue	75.0 (56.3-87.5)	75.0 (56.3-90.6)	75.0 (56.3-87.5)	0.938		

CR=Complete response; IQR=Interquartile range; LN=Lupus nephritis; LupusQoL=Lupus Quality of Life questionnaire; NR=No response; PR=Partial response.

PRESENTED AT: AMERICAN COLLEGE OF RHEUMATOLOGY (ACR) CONVERGENCE; WASHINGTON, D.C., USA; NOVEMBER 14–19, 2024.

HRQoL in patients with LN



- response at the 12-month follow-up

REFERENCES

- 1. Cornet A, et al. *Lupus Sci Med.* 2021;8(1):e000469.
- 2. Katz P, et al. *J Manag Care Spec Pharm.* 2020;26(3):275-283.
- 3. Hocaoğlu M, et al. *Arthritis Rheumatol.* 2023;75(4):567-573.
- 4. Pons-Estel BA, et al. *Medicine* (*Baltimore*). 2004;83(1):1-17.
- 5. Gómez-Puerta JA, et al. *Lupus.* 2021;30(4):630-640.

ACKNOWLEDGMENTS

This study was sponsored by Janssen Research & Development, LLC. Medical writing support was provided by Panita Maturavongsadit, PhD, of Lumanity Communications Inc., and was funded by Janssen Global Services, LLC. Layout design and reformatting for this encore presentation was provided by Sandeep Chavan of Siro Clinpharm Pvt Ltd, Thane, Maharashtra, India.

DISCLOSURES

US, FZ, and AO are employees of Janssen and may hold stock/stock options from Johnson & Johnson. GP-E received grants and consulting fees from and participated as a speaker, as an advisor, and/or on a steering committee for AstraZeneca, Boehringer Ingelheim, GSK, Janssen, Novartis, Pfizer. RemeGen. Sanofi, and Werfen Diagnostics. BP-E served as a speaker and/or advisor for AstraZeneca, GSK, and Janssen. RN, RQ, DCFÁ, RS, GH, LH, KR, ESPS, NM, CO, EN, VA, BK, LGL, EK, NP, C Pisoni, PP, MEC, ACM, AAG, AD, LCAA, EB, EFN, LVdOM, IG, MMD, LDIHR, ACB, RR, PCA, JFJ, JM, MM, RESB, MPC, EMN, YJ-V, YGB, OG, LRAR, MD, PL, WPM, AC, TP, C Pizzarossa, GS, CR, and GSA declared no conflicts of interest. Previously presented at PANLAR 2024; Barranguilla, Colombia; April 10-13, 2024.



*Presenting author.

KEY TAKEAWAY

• In the GLADEL 2.0 cohort, active LN had a significant impact on the HRQoL of patients; further evaluation is warranted to confirm whether achieving renal clinical response can improve



• Patients with active LN showed a worse HRQoL compared to those with inactive LN

• No differences in HRQoL were found between patients who achieved and did not achieve renal

• Future analyses with a larger number of patients are necessary to provide conclusive data