

DONALD AND BARBARA ZUCKER SCHOOL of MEDICINE AT HOFSTRA/NORTHWELL

Association Between Skin-Related Quality of Life and Race in Patients with Hidradenitis Suppurativa

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Background

Hidradenitis suppurativa (HS) disproportionately affects Black patients. Patients with HS experience significant decrease in quality of life (QoL) due to pain, drainage, disfigurement, and shame. In a Global Survey Of Impact and Healthcare Needs (Global VOICE) analysis, Black patients with HS experienced lower disease-related QoL compared to those who were White.¹ The Dermatology Life Quality Index (DLQI) is commonly used in clinical trials to measure QoL. However, HS-related life impact according to race has yet to be investigated using DLQI data. Evaluating life impact in HS patients with highquality evidence is critical to understand potential racerelated differences in disease burden.

Objective

To compare patient-reported quality of life at baseline between White and Black patients in two Phase III clinical trials of HS patients using the Dermatology Life Quality Index (DLQI).

Methods

- Study design: Pooled analysis of baseline data from two phase 3 multicenter trials of adalimumab for HS: PIONEER I and PIONEER II.
- Inclusion Criteria: Self-reported Black or White race, United States residence, and available baseline DLQI data in addition to main trial eligibility criteria.
- > Primary Outcome: Baseline DLQI score
- Statistical Analysis: Multiple linear regression was used to compare mean baseline DLQI at baseline between White and Black patients while adjusting for age, sex, smoking status, baseline abscess/nodule (AN) count, and baseline draining fistula count. Results were pooled using fixed-effect inverse-variance weighted metaanalysis.

Results

 Table 1. Baseline Characteristics by Race Among US

 Patients

Characteristic	PIONEER I (N = 147)		PIONEER II (N = 94)	
	White (n = 90)	Black (n = 57)	White (n = 71)	Black (n = 23)
Female sex, n (%)	63 (70.0)	45 (78.9)	56 (78.9)	14 (60.9)
Age, years, mean (SD)	35.1 (10.1)	35.5 (10.8)	35.2 (9.8)	33.9 (9.9)
Body mass index				
Mean (SD)	35.0 (8.0)	34.6 (7.0)	36.8 (9.0)	32.5 (6.9)
Current smoker, n (%)	53 (58.9)	14 (24.6)	42 (59.1)	8 (34.8)
Hurley stage, n (%)				
П	57 (63.3)	28 (49.1)	36 (50.7)	7 (30.4)
111	33 (36.7)	29 (50.9)	35 (49.3)	16 (69.6)
Disease duration, years				
Mean (SD)	10.8 (8.9)	10.2 (9.2)	11.7 (8.3)	11.8 (10.1)
Median (range)	8.2 (1.0- 42.8)	6.8 (1.2- 34.1)	9.9 (1.4- 35.4)	7.1 (1.3- 41.6)
Prior surgery for HS, n (%)	6 (6.7)	14 (24.6)	6 (8.5)	3 (13.0)
Lesion counts				
Total # of abscess and inflammatory nodules	14.1 (12.3)	11.3 (7.5)	13.2 (11.5)	10.2 (6.4)
# of abscesses	2.9 (3.6)	2.8 (2.8)	2.6 (3.3)	3.0 (2.4)
# of inflammatory nodules	11.2 (10.8)	8.6 (6.9)	10.6 (10.2)	7.2 (5.0)
# of draining fistulas	3.6 (5.2)	4.4 (4.9)	3.0 (5.0)	6.5 (7.0)
High-sensitivity C- reactive protein				
Mean (SD), mg/L	16.1 (24.2)	21.6 (19.7)	14.4 (15.4)	33.9 (38.7)

Abbreviations: SD, standard deviation; HS, hidradenitis suppurativa

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Results

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Table 2. Comparison of baseline DLQI between White and Black patients, US

	PIONEER I (N = 147)		PIONEER II (N = 94)	
Baseline DLQI	White	Black	White	Black
	(n = 90)	(n = 57)	(n = 71)	(n = 23)
Mean (SD)	15.6 (6.3)	18.5 (6.4)	15.8 (7.5)	17.3 (7.9)
Median (Q1, Q3)	15 (10, 20)	20 (15, 23)	16 (10, 21)	19 (12, 23)
Unadjusted mean				
difference	Reference	2.9 (0.8-	Reference	1.6 (-2.1,
(Black – White)		5.0)		5.2)
Adjusted mean difference ^a	Reference	3.5 (1.3-	Reference	1.9 (-1.8,
(Black – White)		5.8)		5.6)
p-value (Adjusted mean difference)	Reference	.00226	Reference	.31794

Pooled mean DLQI difference from both studies after covariate adjustment was 3.1 (95% CI 1.2,5.0).

Conclusions

- Black patients with HS experienced lower skin-related quality of life compared to those who were White. QoL burden remained higher among Black patients even when controlling for disease severity.
- ➢ Prior studies have reported increased disease severity, higher healthcare utilization, and greater pain severity among Black patients with HS.^{2,3} Differences in QoL may have been underestimated as DLQI is less sensitive to HS-related factors. Understanding these factors can mitigate race-related differences and reduce diseaserelated life impact in patients with HS.

References

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