Disparities in NMOSD Hospitalization Outcomes: A 5-Year Analysis of the Impact of Sociodemographic Factors On Healthcare Utilization and Charges

Background

- Neuromyelitis Optica Spectrum Disorder (NMOSD) is a rare autoimmune disease, in which the immune system attacks nervous system cells, mainly resulting in inflammation of the optic nerve.^{1,2,3} This may lead to severe vision and movement problems.
- The prevalence of NMOSD was notably elevated among different racial groups, with the highest rates observed in Blacks (12.99/100,000), constituting 27.7% of NMOSD patients.¹
- In the female population, the prevalence is 9.48/100,000, with Black and Asian females exhibiting 2.65- and 1.94-times higher prevalence rates, respectively, compared to White females.¹
- Total healthcare costs of managing NMOSD and relapses are estimated to be about \$60,500 annually, with a relapse episode lasting for 10-15 days. Due to the severe impact of the relapses, many relapses may require emergency department visits.⁴
- Limited research has evaluated the impact of sociodemographic factors on NMOSD hospitalizations.

Objectives

- 1. To examine the hospitalization characteristics and outcomes [length of stay (LOS), hospitalization charges] of NMOSD patients across 2016-2020.
- 2. To determine the impact of social determinants of health (age, gender, race, income) on NMOSD hospitalization outcomes.

Methods

- We used the Healthcare Cost and Utilization Project (HCUP) National Inpatient Sample (NIS) database (2016-2020).
- We used International Classification of Diseases-10 (ICD-10) codes to identify hospitalizations associated with NMOSD (ICD10: G36.0).
- Sociodemographic factors, hospital characteristics, and outcomes were pooled for analyses.
- We conducted multivariate analyses adjusted for income, region, procedures, and hospital characteristics.
- Total charges were subjected to regression analysis, with log transformation applied to the values for interpretation and statistical analysis.

- Over 5 years there were 1,686 NMOSD hospitalizations, with a mean age at hospitalizations of 45.9 ±15.4 years (Table 1).
- NMOSD hospitalizations were predominantly for women patients (79.6%)
- Black patients accounted for 39.7% of the hospitalizations, White 37.3% and Hispanic patients for 11.9%
- hospitalizations (Table 2).

Table 1. NMOSD Overall Hospitalization Characteristics				
	Age	Male	Female	Total
Average	45.9	44.35	45.12	
Age (SD)	(15.4)	(15.43)	(15.46	
Race	Mean (SD)	N (%)	N (%)	N (%)
White	47.9	170	459	629
	(16.4)	(49.4%)	(34.2%)	(37.3%)
Black	42.9	103	566	669
	(14.0)	(30.0%)	(42.2%)	(39.7%)
Hispanic	42.7	45	156	201
	(14.2)	(13.1%)	(11.6%)	(11.9%)
Other	45.0	19	118	137
	(16.1)	(5.5%)	(8.8%)	(8.1%)
Missing		7	43	50
		(2.0%)	(3.2%)	(3.0%)
Total		344	1,342	
		(20.4%)	(79.6%)	

Table 2. NMOSD Hospitalization Outcomes Overall Gender Male Female

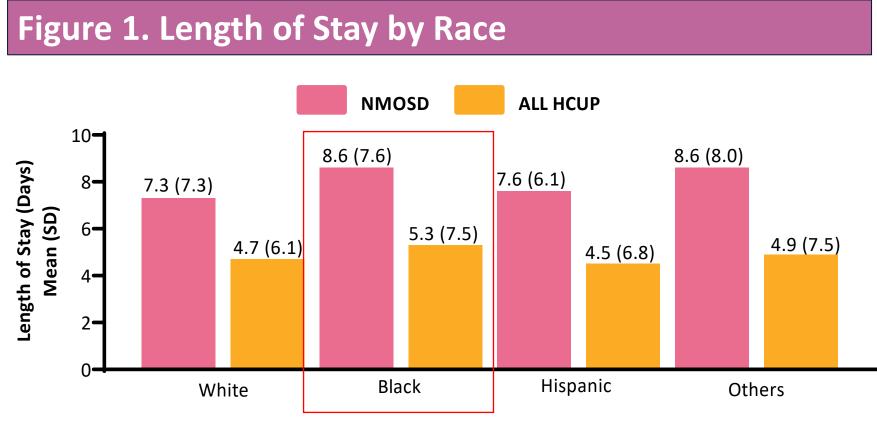
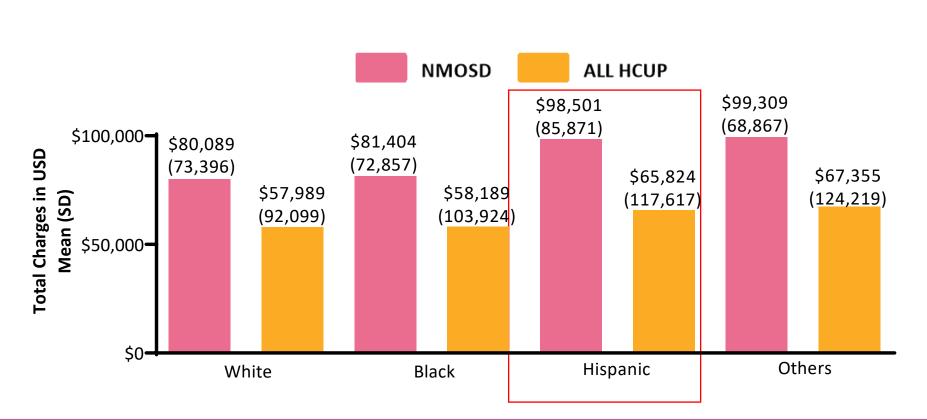


Figure 2. Hospitalization Charges by Race



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• There were minimal differences between Males and Females in terms of LOS and Total Charges for NMOS

Length of Stay	Total Charges
Mean (SD)	Mean (SD)
8.0 (7.3)	\$84,171 (74,294)
7.8 (7.6)	\$85,550 (74,899)
8.0 (7.3)	\$83,818 (74,162)

Results

NMOSD Hospitalizations Compared to All Hospitalizations

NMOSD Hospitalizations Length of Stay (LOS)

- NMOSD LOS was longer (8 ± 7.3 days) compared to HCUP average (4.8 ± 6.5) days).
- Hospitalizations in Black patients had longer LOS (8.6 ± 7.6 days) compared to White $(7.3 \pm 7.3 \text{ days})$ and Hispanic (7.6)± 6.1 days) patients (Figure 1).

NMOSD Hospitalizations Total Charges

- Total charges were higher among NMOSD hospitalizations (\$84,171 ± 74,294) compared to HCUP average $($57,419 \pm 96,851).$
- NMOSD hospitalizations for Hispanic patients incurred the highest total charges (\$98,501 ± 85,871), followed by Black (\$81,404 ± 72,857) and White (\$80,089 ± 73,396) patients (Figure 2).

Regression Analysis Model

The model used in the analysis is a multiple linear regression model, specified as follows:

Length of Stay / log(Total Charges) = B0 (Intercept) + β 1(Age) + β2(Female) + β3(Black) + β4(Hispanic)+ β5(Other Race)+ β6(Income Zip Quartile) + β7(Large Metro) + β8(Med Metro)+ β 9(Mid West)+ β 10(South) + β 11(West) + β 12(Medium Hospital) + β13(Large Hospital) + β14(Urban Non-Teaching) + β 15(Urban Teaching) + β 16(Number of Procedure Codes) + β 17(Number of Diseases) + β 18(Medicare)+ β 19(Private including HMO) +

 β 20(Other Insurance) + ϵ

Where:

β0 is the intercept

 $\beta 1 + \beta 2 + \beta 3$ etc. are the coefficients for each independent variable

ε is the error term, capturing the variability in Total charges or Length of stay

References

Table 1: Regression Ana	alysis			
	Length of Stay R2=36		Log Total Cha B) R2=3	•
Variables	Coefficient	p value	Coefficient	p
Patient Age (years)	0.023	0.046	0.003	
Gender (Reference Male)				
Female	-0.268	0.464	-0.062	
Race (Reference White)				
Black	1.415	<0.001	0.120	
Hispanic	0.836	0.111	0.165	
Other	1.816	0.002	0.184	
Quartile Income	0.032	0.828	0.002	
Patient's Location (Reference	Small Metro)			
Large Metro	-0.245	0.670	-0.065	
Medium Metro	0.252	0.663	-0.065	
Hospital Region (Reference No	ortheast)			
Midwest	-0.686	0.148	-0.353	
South	-0.346	0.407	-0.432	
West	-0.707	0.170	0.023	
Hospital Size (Reference Smal	I)			
Medium	0.122	0.829	-0.008	
Large	0.468	0.352	0.140	
Hospital Location/Type (Refer	ence Rural)			
Urban Non-Teaching	-0.469	0.729	0.700	
Urban Teaching	0.228	0.858	0.787	
Hospital Ownership (Reference	e Governmen	t Owned)		
Private, Non-Profit	-0.417	0.286	-0.026	
Private, Investor own	-1.141	0.089	0.365	
Expected Payer Type (Referen	ce Medicaid)			
Medicare	-1.059	0.020	-0.010	
Private including HMO	-0.811	0.040	0.043	
Other	0.653	0.246	0.028	
Number of Procedures performed	1.313	<0.001	0.177	
Number of Diseases	0.392	<0.001	0.030	

Length of Stay (Model A; Table 3)

- Age: Each additional year of age is associated with a longer stay of approximately 0.023 days (p < 0.05).
- **Race:** Compared with the reference group (White), African American patients have, on average, a longer hospital stay by 1.4 days (p < 0.001).

Total Charges (Model B; Table 3)

- Age: For every additional year of age, charges increase by 0.3% (p < 0.05).
- **Race:** On average, hospitalizations for Black patients incurred charges 12% higher than those for White patients (p < 0.05), while hospitalizations for Hispanic patients incurred charges 16.5% higher (p < 0.05).

1. Kim et al. Differential diagnosis of neuromyelitis optica spectrum disorders. Ther Adv Neurol Disord. 2017 Jul;10(7):265-289. 2. Briggs et al. Prevalence of neuromyelitis optica spectrum disorder in the United States. Mult Scler. 2024 Jan 27:13524585231224683. 3. Kim et al. Racial differences in neuromyelitis optica spectrum disorder. Neurology. 2018 Nov 27; 91(22): e2089–e2099. 4. Royston et al. Neuromyelitis optica spectrum disorder: Clinical burden and cost of relapses and disease-related care in US clinical practice

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s (Model I%	
o value	
0.029	
0.164	
0.007	
0.010	
0.010	
0.932	
0.362	
0.362	
<0.001	
<0.001	
0.718	
0.910	
0.022	
<0.001	
<0.001	
0.590	
<0.001	
0.856	
0.366	
0.687	
<0.001	
\U.UU1	
<0.001	



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Conc	lusions

- This is the first US study of NMOSD hospitalizations over 5 years.
- Disproportionately more women (79%) and Blacks (40%) are hospitalized due to the higher prevalence of the disease in women and Black Americans.
- On average, the NMOSD hospitalizations were almost twice as long as average HCUP hospitalizations.
- Furthermore, NMOSD hospitalizations were ~50% more expensive than an average HCUP hospitalization.
- Black patients had significantly longer LOS (1 extra day) and **Hispanic patients incurred** significantly higher charges (~\$17,500 more).
- Effective treatment of NMOSD in the community is crucial to ensure that these vulnerable patients are not hospitalized, where they may bear disproportionately higher healthcare burden.

Limitations

- L. HCUP data which is based on total charges may not accurately reflect the actual cost structure related to hospital services.
- 2. HCUP primarily focuses on inpatient care and hence no extrapolation can be made about NMOSD in nonhospital settings, which will require further study

Author Disclosures

- JA served on scientific advisory boards for of EMD Serono, Genentech, Horizon Therapeutics/Amgen & TG Therapeutics; has received research support from Horizon Therapeutics/Amgen
- AA and MR are employees of APPERTURE LLC and are HEOR consultants to several Biopharmaceutical companies
- KP, HP, JP are employees of Amgen (Horizon Therapeutics) and are stock-holders of Amgen