

# Racial and Ethnic Disparities in COVID-19 Treatments in the United States

Essy Mozaffari<sup>1</sup>, Aastha Chandak<sup>2</sup>, Alpesh N Amin<sup>3</sup>, Robert L Gottlieb<sup>4,5,6,7</sup>, Andre C Kalil<sup>8</sup>, Vishnudas Sarda<sup>9</sup>, Mark Berry<sup>1</sup>, Gina Brown<sup>1</sup>, Jason F Okulicz<sup>1</sup>, Chidinma Chima-Melton<sup>10</sup>

<sup>1</sup>Gilead Sciences, Foster City, CA, USA; <sup>2</sup>Certara, New York, NY, USA; <sup>3</sup>University of California Irvine, CA, USA; <sup>4</sup>Baylor University Medical Center, Dallas, TX, USA; <sup>5</sup>Baylor Scott & White Heart and Vascular Hospital, Dallas, TX, USA; <sup>6</sup>Baylor Scott & White The Heart Hospital, Plano, TX, USA; <sup>7</sup>Baylor Scott & White Research Institute, Dallas, TX, USA; <sup>8</sup>University of Nebraska Medical Center, Omaha, NE, USA; <sup>9</sup>Certara, Secunderabad, India; <sup>10</sup>University of California, Los Angeles, CA, USA

## Conclusions

- Black patients were less likely to receive COVID-19 treatment compared to White and Asian patients after accounting for differences in comorbidity burden and demographics
- The finding was consistent across all levels of COVID-19 severity, as measured using baseline supplemental oxygen requirement
- These findings are indicative of both under- and overtreatment according to patient race.
  - For example, White patients with NSOc were significantly more likely to receive corticosteroids and baricitinib than Black patients, despite clinical guidelines recommending against initiating these treatments in patients with NSOc
  - Conversely, White patients with LFO were more likely to receive remdesivir treatment than Black patients, which was indicative of undertreatment of Black patients.
- While remdesivir is selectively recommended for initiation among patients with HFO/NIV, growing evidence indicates its likely beneficial role among patients with and without hypoxemia<sup>3, 4</sup>
- The finding of inequitable delivery of life-saving treatments is likely to be an important contributor to the well-documented racial disparities in COVID-19 outcomes
- Few differences in the administration of COVID-19 treatments were observed between Hispanic and non-Hispanic patient populations, with the exception of tocilizumab
- This lack of consistent trend in treatment initiation according to ethnicity is likely to reflect the considerable heterogeneity of the non-Hispanic and Hispanic patient populations

### References:

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**Disclosures:** EM, MB, GB, JFO: employee and shareholder (Gilead Sciences, Inc.); AC, VS: employee of Certara (contracted by Gilead Sciences, Inc. to conduct the study); ANA: principal investigator or co-investigator (clinical trials sponsored by NIH/NIAID, NeuroRx Pharma, Pulmotect, Blade Therapeutics, Novartis, Takeda, Humanigen, Eli Lilly, PTC Therapeutics, OctaPharma, Fulcrum Therapeutics, Alexion), speaker and/or consultant (Pfizer, Salix, Alexion, AstraZeneca, Bayer, Ferring, Seres, Spero, Eli Lilly, Nova Nordisk, Gilead, Renibus, GSK, Dexcom, Reprieve, HeartRite, Aseptiscope)- these relationships are unrelated to the current work; RLG: advisor (AbbVie, Gilead Sciences, Inc., Eli Lilly, Roche, Johnson & Johnson), consultant (Eli Lilly, Gilead Sciences, Inc., Johnson & Johnson, Kinevart Sciences, Roche), de minimis investment (AbCellera), research contracts (Eli Lilly, Gilead Sciences, Inc., Johnson & Johnson, Pfizer), speaker's bureau (Pfizer); ACK: investigator (National Institutes of Health Adaptive COVID-19 Treatment Trial); CCM: advisor (AstraZeneca, Gilead Sciences, Inc.), speaker's bureau (AstraZeneca, Boehringer Ingelheim), consultant (Gilead Sciences, Inc.).

## Background

- Long-standing differential access to goods, services and opportunities according to race and ethnicity has led to widespread and well-documented disparities in health status and medical care in the United States (US).<sup>1,2</sup>
- The COVID-19 pandemic has magnified these racial and ethnic disparities, with heightened risk for COVID-19 incidence, morbidity and mortality demonstrated in many minority populations.<sup>2</sup>
- The rate of hospitalization of Black patients due to COVID-19 has been shown to be more than three times higher, and for Hispanic patients over four times higher, compared to White patients.<sup>2</sup>
- There is a necessity to examine and identify differences in the delivery of appropriate and potentially life-saving COVID-19 therapies according to race and ethnicity, so efforts can be made to ensure the delivery of equitable healthcare and lessen inequities in COVID-19 outcomes.
- The objective of the study was to characterize disparities in the administration of evidence-based COVID-19 treatments among patients hospitalized for COVID-19.

## Methods

### Study Design

- Retrospective cohort study
- **Study period:** May 2020-April 2022
- **Data source:** PINC AI Healthcare Database (formerly Premier Healthcare Database)
  - U.S. hospital-based, service-level, all-payer (Commercial, Medicare, Medicaid, others) database
  - Covers ~25% of all US hospitalizations from 48 states.
  - Includes information on billed services and activities for each day of the hospitalization
  - Underlying dataset included >700,000 COVID-19 hospitalizations with a primary diagnosis of COVID-19 from 954 hospitals during the study period, providing rich source of information and large sample size for each subgroup analysis considered

### Table 1. Study design

<b>Inclusion criteria</b>	<ul style="list-style-type: none"> <li>✓ First admission to the hospital between 1<sup>st</sup> May 2020 and 30<sup>th</sup> April 2022</li> <li>✓ <b>Primary discharge diagnosis of COVID-19</b> (ICD-10-CM: U07.1) flagged for being “present-on-admission”</li> <li>✓ Age ≥18 years old</li> </ul>
<b>Exclusion criteria</b>	<ul style="list-style-type: none"> <li>x Pregnant</li> <li>x Had incomplete data/ erroneous data fields</li> <li>x Transferred from hospice or another hospital</li> <li>x Unknown gender</li> <li>x Admitted for elective procedures</li> <li>x Admitted with no supplemental oxygen (NSOc) on admission to a hospital that did not report any low flow oxygen (LFO)</li> <li>x Patients with extracorporeal membrane oxygenation (ECMO) in the first two days of hospitalization</li> <li>x COVID-19 diagnosis not flagged as “Present on admission”</li> </ul>

- **Primary End Point:** Initiation of any guideline-recommended COVID-19 treatments within two days of hospitalization (corticosteroid, remdesivir, baricitinib and tocilizumab)
- Patients were categorized according to their recorded race (White, Black, Asian, Other) and their recorded ethnicity (Hispanic, Non-Hispanic, Unknown).

### Statistical analysis

- **Descriptive analyses:**
  - Baseline characteristics of study cohort overall and according to race
  - Baseline characteristics of study cohort overall and according to ethnicity
- **Multivariate logistic regression:**
  - Outcome: receiving any COVID-19 treatment within 2 days of hospitalization
  - Covariates: age, gender, hospital size, rural/urban, teaching hospital status, geographic region, primary payer, admission month/variant time period, renal disease, immunocompromised conditions, categories of CCI, and hospital ward upon admission

## Results

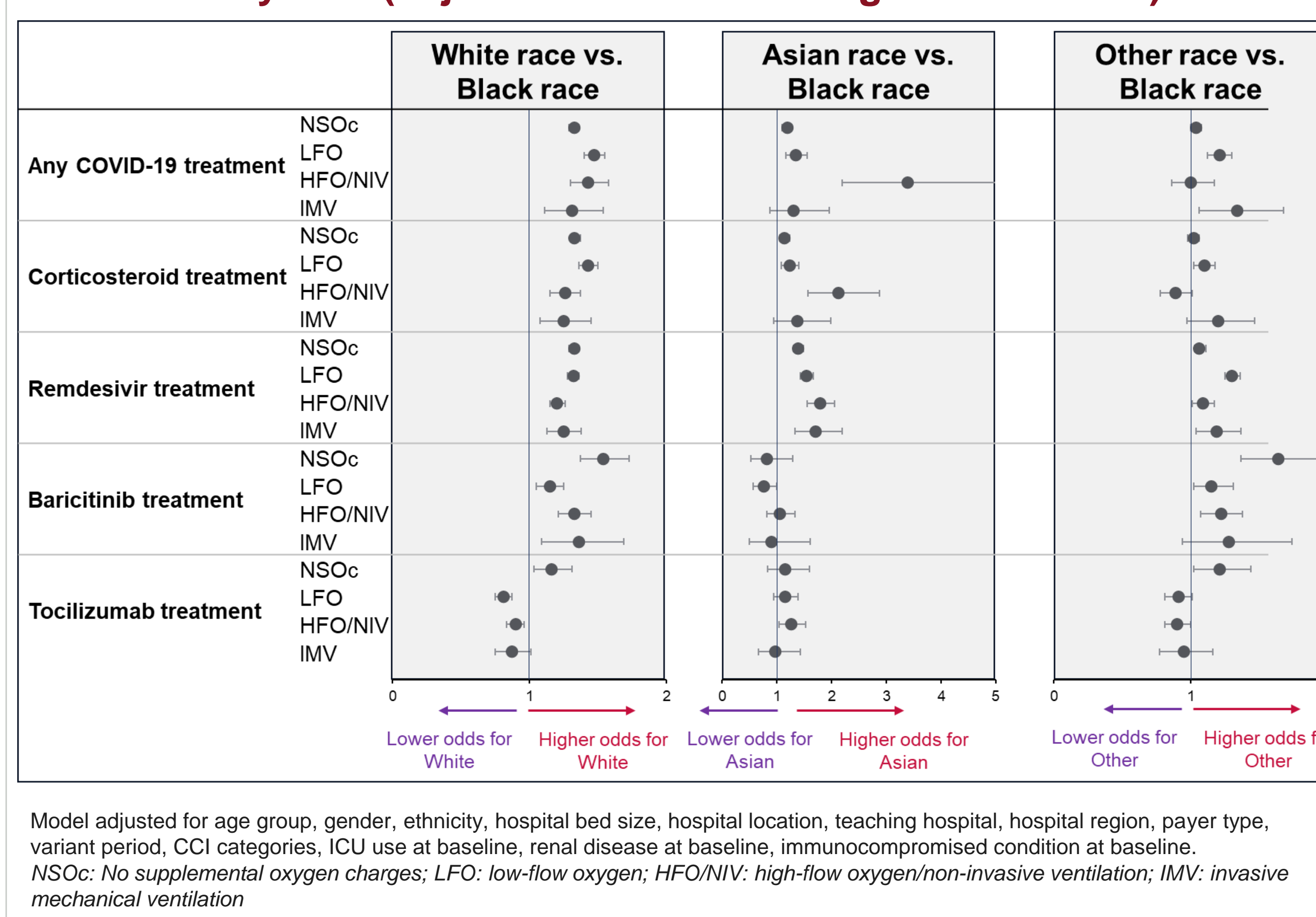
### Study population

- There were 454,761 adults hospitalized for COVID-19 between May 2020 and April 2022
  - White: 70%, Black: 17%, Asian: 2%, Other: 11%
  - Non-Hispanic: 72.6%, Hispanic: 16.3%, Unknown: 11.1%
- White patients were older, with a lower proportion admitted during the pre-Delta period, and had a lower comorbidity burden than Black, Asian, and other patients (**Table 2**)
- White patients were more likely to receive COVID-19 treatments, and specifically corticosteroids, compared to Black, Asian, and other patients (COVID-19 treatment: 87% vs. 81% vs. 85% vs. 84%, corticosteroids: 85% vs. 79% vs. 82% vs. 82%) (**Table 2**)
- White patients were less likely than Asian patients to have remdesivir (53% vs. 56%) initiated but considerably more likely than Black patients (53% vs. 43%) (**Table 2**)
- Hispanic patients were, younger, and had a lower comorbidity burden than non-Hispanic patients (**Table 2**)
- The proportion of patients receiving any COVID-19 treatment within 2 days of admission was similar across each ethnic grouping (Hispanic: 86%, Non-Hispanic: 86) (**Table 2**)

### Multivariable analyses

- White patients were significantly more likely to receive any COVID-19 treatment than Black patients across all supplemental oxygen levels
  - NSOc aOR: 1.33 (95% CI: 1.29 – 1.36), LFO aOR: 1.47 (1.40 – 1.55), HFO/NIV aOR: 1.43 (1.30 – 1.58), IMV aOR: 1.31 (1.11 – 1.54) (**Figure 1**)
- White patients were also statistically significantly more likely than Black patients to receive corticosteroids, remdesivir, and baricitinib treatment across all baseline supplemental oxygen levels, including among patients with NSOc (**Figure 1**)
- In contrast, White patients with LFO, HFO/NIV or IMV were less likely to receive tocilizumab treatment than Black patients (**Figure 1**)
  - LFO aOR: 0.81 (0.75 – 0.87), HFO/NIV aOR: 0.90 (0.83 – 0.96), IMV aOR: 0.87 (0.75 – 1.01)
- Asian patients were more likely to receive remdesivir treatment than Black patients across all supplemental oxygen (**Figure 1**)
  - NSOc aOR: 1.39 (1.30 – 1.48), LFO aOR: 1.54 (1.43 – 1.67), HFO/NIV aOR: 1.79 (1.56 – 2.06), IMV aOR: 1.71 (1.33 – 2.20)
- Asian patients with LFO or HFO/NIV were also more likely to receive any COVID-19 treatment and corticosteroids than Black patients (**Figure 1**)
- Differences in the administration of COVID-19 treatment between Hispanic and non-Hispanic patients varied considerably according to treatment type and baseline supplemental oxygen after multivariable adjustment (**Figure 2**)

**Figure 1. Likelihood of receiving COVID-19 treatment upon hospital admission by race (adjusted multivariable regression model)**



**Table 2. Baseline characteristics of study cohort overall and according to race and ethnicity**

	Overall	White	Black	Asian	Hispanic	Non-Hispanic	
# Patients	454,761	317,928	76,715	9,297	74,199	329,940	
Age (years)	Median (IQR)	64 (52-75)	66 (54-77)	60 (48-70)	63 (50-74)	57 (45-70)	65 (54-76)
Age group		21%	18%	28%	25%	33%	18%
		30%	29%	35%	30%	33%	30%
		49%	54%	38%	45%	34%	52%
Gender	Male	52%	53%	45%	54%	54%	51%
	White	70%	100%	0%	0%	65%	71%
	Black	17%	0%	100%	0%	2%	21%
	Asian	2%	0%	0%	100%	0%	2%
	Other <sup>1</sup>	11%	0%	0%	0%	33%	5%
	Hispanic	16%	15%	2%	2%	100%	0%
	Non-Hispanic	73%	74%	90%	86%	0%	100%
	Unknown	11%	11%	8%	12%	0%	0%
	Commercial	29%	28%	28%	34%	30%	29%
	Medicare	51%	55%	46%	40%	34%	54%
	Medicaid	11%	8%	17%	18%	16%	9%
	Other Payer	10%	9%	9%	8%	19%	8%
Hospital location	Urban	86%	84%	90%	92%	93%	85%
	Rural	14%	16%	10%	8%	7%	15%
	<100	7%	8%	4%	5%	4%	8%
	100-199	17%	18%	13%	16%	22%	17%
	200-299	20%	20%	19%	22%	15%	20%
	300-399	19%	18%	23%	19%	14%	20%
	400-499	10%	10%	6%	8%	11%	9%
	>=500	27%	25%	34%	29%	34%	26%
Hospital teaching status	Yes	39%	38%	45%	47%	44%	38%
	Midwest	21%	23%	19%	17%	8%	25%
	Northeast	9%	8%	9%	14%	8%	9%
	South	55%	54%	68%	29%	67%	56%
	West	14%	14%	5%	40%	17%	10%
	Pre-Delta	60%	58%	63%	70%	65%	57%
	Delta	33%	35%	31%	26%	31%	35%
	Omicron	7%	7%	6%	5%	5%	8%
CCI	0	30%	31%	26%	35%	38%	29%
	1 to 3	50%	51%	49%	49%	48%	51%
	≥4	19%	19%	25%	16%	14%	20%
Key comorbidities	Immunocompromised condition	10%	11%	11%	8%	8%	11%
	Renal disease	19%	18%	27%	18%	13%	21%
	NSOc	44%	42%	53%	47%	47%	42%
	LFO	38%	40%	31%	36%	38%	39%
	HFO/NIV	15%	15%	14%	13%	13%	15%
	IMV	3%	3%	3%	3%	3%	3%
Baseline ICU admission	Any COVID-19 treatment	86%	87%	81%	85%	86%	86%
	Corticosteroids	84%	85%	79%	82%	84%	84%
	Remdesivir	52%	53%	43%	56%	54%	52%
	Baricitinib	4%	4%	3%	2%	3%	4%
	Tocilizumab	4%	4%	4%	4%	4%	4%

<sup>1</sup>Other race includes race designations that have been rolled into “other” to ensure that the dataset conforms to regulatory requirements as well as race designations of “unable to determine”. CCI: Charlson Comorbidity Index; IQR: inter-quartile range; NSOc: supplemental oxygen charges; LFO: low-flow oxygen; HFO/NIV: high-flow oxygen/non-invasive ventilation; IMV: invasive mechanical ventilation

**Figure 2. Likelihood of receiving COVID-19 treatment upon hospital admission by ethnicity (adjusted multivariable regression model)**

