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Remote Biometric Monitoring of Patients With COVID-19 With Exertional Hypoxia Treated With Supplemental Oxygen

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Hospitalizations to treat patients infected with COVID-19 have strained health systems worldwide. Monitored outpatient treatment of select low-risk patients with COVID-19 could help preserve hospital resources and reduce costs while aiming to maintain a high standard of care. However, criteria to identify patients appropriate for outpatient monitoring do not exist. Isolated exertional hypoxia may be an early predictor of moderate-to-severe COVID-19 and could be an indication for remote monitoring or inpatient admission; however, data to support its use are

Patients and Methods

The cRPM-O2 algorithm was implemented across 23 of Intermountain Healthcare's integrated hospitals from November 2020 to April 2021, during the COVID-19 alpha variant surge. Included patients had an ED encounter within 14 days after a confirmed COVID-19 positive test, received a COVID exercise tolerance test (cETT) to assess exertional hypoxia per protocol, and had a cHIS <2 by laboratory results that were obtained during the index ED encounter. cETT consisted of either briskly walking for 1 min or sit-to-stand for 1 min. Any SpO2 result <90% or severe dyspnea that prevented completion of the test was considered a positive cETT. The clinical criteria for cRPM-O₂ inclusion and triage process are presented in Figure 1. Patients who met the criteria for cRPM-O₂ participation from the ED, but with an inpatient admission, constituted the control arm. The primary outcome was inpatient hospital days associated with the first admission within 14 days after the index ED encounter modeled by ordinary least squares regression (covariates are listed in Table 1). Secondary outcomes were ICU length of stay and combined ICU admission or all-cause death within 28 days. ICU length of stay was modeled in the same fashion as the primary outcome. Death was identified by state death database review to ascertain any death that occurs outside of our system. The composite

Results

Of the 755 patients identified in the study cohort, 522 patients comprised the control group, and 233 patients were in the cRPM-O₂ group. The cRPM-O₂ group was younger and had fewer comorbidities (Table 1). After

limited.¹⁻³ Our institution implemented a 61 standardized ED triage process that included 62 exertional hypoxia and the COVID-19-associated 63 hyperinflammation (cHIS) score.⁴ The cHIS scoring 64 65 includes fever, degree of cytopenia, hyperferritinemia, 66 D-dimer, markers of hepatic injury (lactate 67 dehydrogenase or aspartate aminotransferase), and 68 markers of cytokinemia (c-reactive protein, IL-6, or 69 hypertriglyceridemia) and is associated with 70 progression to mechanical ventilation and death. This 71 triage process identified patients who were considered 72 to be at an acceptable morbidity and mortality risk 73 for discharge to home on low-flow home oxygen with 74 remote SpO₂ and vital sign monitoring. In this 75 retrospective cohort analysis, we report the safety and ⁷⁶ 77 effectiveness of the COVID-19 remote patient 78 monitoring with home oxygen (cRPM-O₂) program 79 as a strategy to reduce hospitalization days and 80 preserve resource utilization. 81

binary outcome of ICU admission or death was modeled with the 84 use of a logistic regression. Patient COVID-19 vaccination status was not available. The Intermountain Healthcare Institutional Review Board approved the analysis.

Patients who enrolled in cRPM-O2 received a Bluetooth-enabled pulse 88 oximeter paired to their smart phone via an application downloaded 89 and confirmed by the ED nurse or technician prior to ED discharge. Patients were asked to record SpO2/heart rate biometrics at least 90 twice daily, or as often as they preferred, for 14 days. Submitted 91 biometric data from enrolled patients were monitored 24 hours per 92 day, 7 days per week. Patients without transmitted vital signs for 93 >18 hours or with $\text{SpO}_2 < 88\%$ and/or pulse rate >120 beats/min 94 were contacted by a monitoring technician. If new hypoxemia was identified despite supplemental oxygen, the patient was instructed to 95 return to the ED. Patients with concerns about their symptoms or 96 the device could contact a monitoring technician via the application 97 at any point, and concerns could be escalated to the monitoring 98 center critical care nurse or nurse practitioner. The application 99 included built-in text and video visit functionality to facilitate remote patient assessment. The pulse oximeter device, application, and 100 monitoring service were offered to patients without out-of-pocket 101 charges. 102

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adjustment for age, sex, ethnicity/race identifiers, 105 insurance provider, Charlson Comorbidity Index, and 106 lowest recorded oxygen saturation during index ED visit, 107 cRPM-O₂ participation was associated with a reduction 108 of 2.27 hospital days per patient (95% CI, 1.53 to 3.01; 109 110

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Figure 1 – COVID-19-Associated Hyperinflammation Score Assessment Flow Diagram. AST = aspartate aminotransferase; cETT = COVID exercise tolerance test; cHIS = COVID-19-associated hyperinflammation score; cRPM = COVID remote patient monitoring; Hgb = hemoglobin; Q13 LDH = lactate dehydrogenase.

14**@** P < .001). Of the 233 patients who received cRPM-O₂, 147 39 patients (16.7%) subsequently were admitted with a 148 median of 4 days (interquartile range, 2 to 6) from ED 1495 visit to admission. cRPM-O₂ participation was 150 associated with a numeric increase in combined ICU 151 admission and deaths (OR, 1.93; P = .07; 95% CI, 0.94 to 152 3.95); ICU days were significantly increased in cRPM-O₂ 153 154 participants (0.33 days; 95% CI, 0.00 to 0.65; P = .05). In 155 the preplanned subgroup analysis that was limited to 156 patients \geq 70 years old, hospital length of stay did not 157 differ with cRPM-O₂ participation (-0.50 days; 95% CI, 158 -3.44 to 2.44; P = .74) and participation in cRPM-O₂ 159 was associated with increased ICU length of stay 160 (1.05 days; 95% CI, 0.07 to 2.0; P = .04) and a 161 numerically increased odds of ICU admission or death 162 (OR, 5.44; 95% CI, 0.93 to 31.75; P = .06). In this 163 subgroup, there were four deaths in the control group 164 165

and no deaths in the cRPM-O₂ group. In contrast, cRPM-O₂ for patients <70 years of age was associated with hospital length of stay reduction of 2.56 days (95% CI, 1.84 to 3.30; P < .001), no change in ICU length of stay (+0.18 days; 95% CI, -0.16 to 0.53; P = .296), and no difference in the odds of combined ICU admission or death (OR, 1.40; 95% CI, 0.63 to 3.14; P = .410).

Discussion

This ED program to triage low-risk patients with213COVID-19 with exertional hypoxia to treatment at214home with remote monitoring and low-level216supplemental oxygen was associated with fewer217hospital days but with an increase in ICU days and218trend toward an increase in the composite outcome of219

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Model Input and Patient Characterist	ED Admission to Floor (n = 522)	COVID Remote Patient Monitoring With Home Oxygen (n = 233)	<i>P</i> value
Age, mean (SD), y	61.5 (47.5-71.5)	55.1 (43.6-64.9)	<.001
Sex, No. (%)			.82
Male	251 (48.1)	110 (47.2)	
Female	271 (51.9)	123 (52.8)	
Insurance, No. (%)			<.001
Private or commercial	253 (48.5)	167 (71.7)	
Medicaid	46 (8.8)	11 (4.7)	
Medicare	204 (39.1)	45 (19.3)	
Other	19 (3.6)	10 (4.3)	
Collapsed race/ethnicity, No. (%)		.28
White, not Hispanic	370 (70.9)	174 (74.7)	
Hispanic or other	152 (29.1)	59 (25.3)	
Charlson Comorbidity Index	1.6 (2.7)	0.8 (2.0)	<.0001
Lowest oxygen saturation durin	ng ED visit 87.6 (4.4)	87.3 (3.2)	.38
Unadjusted patient outcomes ^b			
Inpatient hospital days, No.	%) 4.0 (3.2)	1.5 (4.7)	<.001
ICU length of stay, d (%)	0.2 (1.8)	0.5 (2.1)	.079
ICU level care or death, No.	(%) 22 (4.2)	15 (6.4)	.19
28-Day all-cause deaths, No.	(%) 5 (1.0)	2 (0.9)	.90
cRPM-O2 adjusted outcomes $^{\circ}$	Coefficient/OR	95% CI	
Hospital days (coefficient)	-2.27	-3.01 to -1.53	<.001
<70 Years old	-2.57	-3.30 to -1.84	<.001
\geq 70 Years old	-0.05	-3.44 to 2.44	.74
ICU days (coefficient)	0.33	0.00 - 0.65	.048
<70 Years old	0.18	-0.16 - 0.53	.30
\geq 70 Years old	1.05	0.07 - 2.031	.04
ICU admission of death (OR)	1.93	0.94 - 3.95	.074
<70 Years old	1.40	0.63 - 3.14	.41
≥70 Years old	5.44	0.93 - 31.75	.06

^aCovariates adjusted for in our model for all 3 outcomes.

²⁵⁹ ^bUnadjusted outcome measures.

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260 ^cAdjusted outcomes (ordinary least squares and logistic regression).

262 death or ICU for patients triaged to home with 263 cRPM-O₂, driven by events that occurred among 264 patients \geq 70 years old. In total, 83.3% of patients with 265 cRPM-O₂ were treated safely without hospitalization; 266 267 however, only 67% in those patients aged \geq 70 years. 268 Inclusion in cRPM-O₂ significantly increased ICU 269 days, but by a clinically insignificant 0.33 days in our 270 main analysis. When assessing only those patients 271 admitted to the ICU in each group, we observed the 272 adjusted ICU length of stay is not statistically 273 different, which suggests that trends toward increased 274 risk of ICU admission between groups accounts for 275

317 some of the increased ICU length of stay finding. 318 When assessing only patients who were admitted from 319 the cRPM-O2 , we found that patients who required 320 hospitalization are more likely to be older and male 321 and to have more comorbidities than those patients 322 who remain at home. Inpatient interventions such as 323 324 Remdesivir and steroids are considered confounders 325 that may have influenced the differences in outcomes 326 between groups in addition to expected confounders 327 of a retrospective analysis. Additionally, this 328 implementation from a single integrated health care 329 system may not be generalizable externally. 330

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331 ED triage of select patients with COVID-19 with isolated 332 exertional hypoxia to home with low-flow supplemental 333 oxygen plus remote monitoring is associated with fewer 334 hospital days, and a net clinical benefit especially may be 335 realized among patients <70 years old. Appropriate risk 336 assessment and treatment with cRPM-O₂ may be a 337 strategy to improve hospital capacity safely. Future study 338 of subgroups is warranted to ensure the safety of this 339 approach. 340 341 Harris L. Carmichael <mark>3</mark>42

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4 Research Letter