Oxygen provision to severely ill COVID-19 patients at the peak of the 2020 pandemic in a Swedish district hospital

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Abstract

Oxygen is a low-cost and life-saving therapy for patients with COVID-19. Yet, it is a limited resource in many hospitals in low income countries and in the 2020 pandemic even hospitals in richer countries reported oxygen shortages. An accurate understanding of oxygen requirements is needed for capacity planning. The World Health Organization estimates the average flow-rate of oxygen to severe COVID-19-patients to be 10 l/min. However, there is a lack of empirical data about the oxygen provision to patients. This study aimed to estimate the oxygen provision to COVID-19 patients with severe disease in a Swedish district hospital.

A retrospective, medical records-based cohort study was conducted in March to May 2020 in a Swedish district hospital. All adult patients with severe COVID-19 – those who received oxygen in the ward and had no ICU-admission during their hospital stay – were included. Data were collected

NOTE: This preprint reports new research that has not been certified by peer review and should not be used to guide clinical practice.
on the oxygen flow-rates provided to the patients throughout their hospital stay, and summary measures of oxygen provision calculated.

One-hundred and twenty six patients were included, median age was 70 years and 43% were female. On admission, 27% had a peripheral oxygen saturation of ≤91% and 54% had a respiratory rate of ≥25/min. The mean oxygen flow-rate to patients while receiving oxygen therapy was 3.0 l/min (SD 2.9) and the mean total volume of oxygen provided per patient admission was 16,000 l (SD 23,000).

In conclusion, the provision of oxygen to severely ill COVID-19-patients was lower than previously estimated. Further research is required before global estimates are adjusted.

Introduction

The world has been tackling the COVID-19 pandemic since early 2020 (1). The pandemic has led to high demands on health systems and many lives have been lost because of the disease (2). The severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) causing COVID-19 can affect most organs but is primarily a respiratory virus (3). Viral pneumonia is the most common serious manifestation and can result in hypoxemia and acute respiratory distress syndrome (ARDS) (3,4).

To treat severe and critical COVID-19, much global focus has been on expanding intensive care capacity including the use of mechanical ventilation (5). Whilst many critically ill patients can benefit from mechanical ventilation if it is administrated by experienced personnel, it is a staff intensive measure and requires training to be effective and avoid harm (6).

Oxygen therapy is a low cost treatment, less complex than mechanical ventilation and saves lives in COVID-19 (4,7–9). It is the first-line treatment of hypoxemia and has been listed as a World Health Organization (WHO) essential medicine (10). Oxygen is a limited resource in many hospitals in low income countries (11–14), and during the peaks of the pandemic waves there have been reports
of hospitals running out of oxygen in high and middle income countries such as the UK, USA, South Africa, Portugal, Egypt and Brazil (15–21). In addition, sudden failures of the oxygen systems requiring emergency transport from other sites may happen to hospitals anywhere in the world.

Capacity planning is needed to optimise the distribution of oxygen and reach highest positive impact on patient outcomes. This requires an accurate understanding of oxygen requirements for patients with COVID-19. The WHO estimates the average flow-rate of oxygen to severely ill COVID-19-patients (referring to those requiring oxygen but not intensive care unit treatment) to be 10 l/min (22). This estimate is not based on empirical findings and there is a lack of quantitative data on oxygen provision to patients. This study aims to estimate oxygen provision to severely ill COVID-19 patients in a Swedish district hospital.

**Method**

A retrospective, electronic medical records-based cohort study in the medical department in Nyköping Hospital, Sweden.

**Setting**

Nyköping Hospital is a first-line district hospital in Sörmland Region in Sweden with a catchment area of 90,000 people. The medical department in the hospital has a usual capacity for managing 35 inpatients at a time. Sörmland was one of the first Swedish regions to be significantly affected by COVID-19 in 2020 (23) – the first COVID-19 patient was admitted to the department in early March. Less than a month later, the number of in-patients with COVID-19 was 43, with an average of seven new COVID-19 admissions per day.

In the medical department, 40 beds on two new wards were opened within two weeks of the first COVID-19-admission and extra staff were drawn from other areas of the hospital. The patients’
respiratory rate and peripheral oxygen saturation (SpO2) were assessed at least every two hours and oxygen therapy adjusted to maintain SpO2 within an individualized target range (standard 92-96%, lower to patients with assumed hypercapnic respiratory failure). The hospital has an intensive care unit (ICU), which was expanded from the usual 5-beds to 14-beds during the pandemic. There was no shortage of oxygen during the study period.

Before rapid polymerase chain reaction (PCR) testing capacity was widely available, an initial diagnosis of COVID-19 was based on the clinical picture together with typical findings on a thoracic computed tomography (CT) scan (24). Often the initial diagnosis was confirmed by later PCR-testing. This early diagnostic reliance on CT scans shifted towards PCR-testing during the time of the study.

Standard treatments protocols evolved during the study period and included at different times chloroquine, antibiotics and anticoagulation prophylaxis. Corticosteroids, Remdesivir and other COVID-therapeutics were not included as standard treatment during the time of the study. High-flow nasal oxygen and non-invasive ventilation were neither recommended for ARDS in COVID-19, nor available in the medical wards at this time. Patients that needed more respiratory support than low-flow oxygen were transferred to the ICU where most received invasive mechanical ventilation. As many of the admitted patients were elderly and frail, in which ICU care may cause harm or would not be in the patient’s best interest, a policy was introduced to make patient-centred decisions for every admitted patient about the appropriateness of care-escalation to ICU in the event of clinical deterioration. The decision of no escalation of care to ICU (no-ICU) was documented in the patient’s notes and, importantly, was not regarded as synonymous with end-of-life or palliative care. Patients for whom a no-ICU decision had been made received all other therapies, including oxygen therapy when indicated, unless an additional clinical decision was made to provide end-of-life palliative care.
Study cohort

The Sörmland Region database of COVID-19 patients was used to identify participants. All adult patients (age ≥18) admitted to the department of medicine in Nyköping Hospital from March 13 to May 10 who had been diagnosed with COVID-19 during their admission and who fulfilled the criteria for “severe” disease were included. Severe patients were those who received oxygen at some point during their care in the ward and had no ICU-admission during their hospital-stay, in-line with the WHO’s classification (22). Non-severe COVID-19 patients — patients with either moderate disease (never received oxygen) or critical disease (admitted to the ICU, either from the wards or directly from the emergency department) (22) — were not included in the study cohort. However, to describe the study cohort in context, data about admission findings, the most advanced mode of respiratory support and outcomes were collected for all COVID-19 patients admitted to the department during the study period. For patients with more than one admission, only data from the first admission were included.

Sub-groups

Two a-priori defined sub-groups were analysed, as it was hypothesised that their oxygen provision may differ substantially from other patients. The first group were those patients for whom a no-ICU-decision had been made. The second group consisted of patients younger than 70 years old.

Data extraction

Data were extracted in two ways. Data on vital signs and oxygen treatment on admission, patient characteristics including Charlson’s Age Adjusted Comorbidity score (CACI) (25,26), clinical and laboratory findings, pharmaceutical treatments and the presence of a no-ICU decision were extracted manually from the patients’ electronic medical records. Other data were collected through a computerised search in the electronic medical records system, including for dates of admission,
discharge, outcomes, and all registered SpO2 values and oxygen flow-rates throughout the patients’ care. The data extracted with a computerised search were validated by manual cross-checking a 10% sample of the data with the medical records. Data were used from the patients’ entire stay in the medical department. The patients’ outcome in the medical department were noted as either transfer to another department, discharge home or died, with an additional outcome of died within 60 days.

Analysis

All data were anonymised before analysis. Oxygen provision to each patient was estimated by multiplying the oxygen flow-rate during each assessment of the patient (l/min) and the time since the previous assessment (minutes). Thereby, the volumes of oxygen provided to the patient over time were generated, and the sum of these made up the total volume of oxygen provided to the patient (formula 1).

\[
\text{Formula 1, \quad Total volume of oxygen provided to a patient:} \\
\sum_n X_n(T_{n+1} - T_n)
\]

\[n\]: one time point in a series of consecutive time points, \(T_n\): time at the time point \(n\), \(X_n\): oxygen flow (l/min) at the time point \(n\)

The mean oxygen flow-rate for each patient while receiving oxygen was calculated as the total volume of oxygen provided divided by time spent receiving oxygen treatment. An additional analysis of the mean oxygen flow-rate for each patient during their care in the ward used the total oxygen volume provided divided by total time in the ward. Means were used to provide an estimate of the total amount of oxygen that needs to be supplied to such a patient cohort. Medians were also
calculated for both analyses to provide an estimate of a “typical” patient, given the non-normally
distributed data. For other variables, median and interquartile range (IQR) were used for non-
normally distributed values and mean and standard deviation (SD) for normally distributed values.
For all admitted patients, the clinical progression score was determined as has been recently defined
(27). Admission vital signs were analysed as the proportion of vital signs corresponding to a red
NEWS-2 parameter (28). Missing data was omitted from analysis. STATA IC/15.1 (StataCorp LLC) was
used for the analysis.

Ethical considerations
The study was approved by the Ethical Review Board in Lund, reference number 2020-04012. As the
study did not alter patient care and data were anonymised before analysis, individual patient consent
was waived.

Results
Patient characteristics
In total, 206 COVID-19 patients were admitted to the department during the study period (Table 1,
Supplementary table 1). Of these, 126 had severe disease and were included as the study cohort. The
median age was 70 years (IQR 57-82), 54 (43%) were female and 42% had a body mass index (BMI) of
30 or above. CACI scores were 4 or above (indicating <50% estimated 10-years survival) in 50% of the
patients (26,27). A no-ICU decision was documented for 48% of the patients. On admission, 34 (27%)
had an SpO2 of 91% or below and 68 (54%) had a respiratory rate of 25 or above. The length of stay
was a median of 4.9 days (IQR 2.8-7.8). Eight (6.4%) patients were transferred to another
department. The in-hospital mortality was 26% and the 60-day mortality was 32%. For the subset of


patients aged <70, in-hospital and 60-day mortalities were 6.5% and 8.1% respectively and for
patients with a documented no-ICU-decision 55% and 65% respectively. The clinical progression
scores for all admitted patients are presented in supplementary table 2.

| Table 1 Patient characteristics and outcomes |
|----------------------------------------------|
| Study cohort | All patients admitted |
| % (n/N), unless otherwise stated | to the department % (n/N), unless otherwise stated |
| Age (years), median (IQR) | 70 (57-82) | 65 (54-78) |
| Female | 43% (54/126) | 42% (87/206) |
| Diagnosis of COVID-19 confirmed by PCR | 90% (114/126) | 89% (184/206) |
| BMI ≥30 | 42% (38/91) | 40% (56/141) |
| CACI ≥4 | 50% (63/126) | 43% (88/206) |
| No-ICU-decision documented | 48% (60/126) | 37% (77/206) |

Red NEWS-2 (28) parameter on first measurements of vital signs

| | Study cohort | All patients admitted |
| | % (n/N), unless otherwise stated | to the department % (n/N), unless otherwise stated |
| SpO2 (≤91%) | 27% (34/126) | 28% (57/206) |
| Respiratory rate (≤8 or ≥25 breaths/min) | 54% (68/126) | 49% (100/203) |
| Heart rate | 3.2% (4/125) | 2.4% (5/205) |
|                        | Column 1 | Column 2 |
|------------------------|----------|----------|
| Systolic blood pressure| 2.4% (3/125) | 2.0% (4/205) |
| (≤90 or ≥220 mmHg)     |          |          |
| Consciousness          | 13% (17/126) | 10% (20/204) |
| (Non-alert)            |          |          |
| Temperature            | 14% (18/125) | 16% (32/205) |
| (≤35.0 or ≥39.1°C)     |          |          |

*Treatments during hospital-stay*

| Antibiotics           | 79% (99/126) | 77% (158/206) |
| Chloroquine           | 12% (15/126) | 14% (28/206) |
| Anticoagulants        | 52% (65/126) | 55% (114/206) |

*Patient outcomes*

| Length of stay (days), median (IQR) | 4.9 (2.8-7.8) | 4.3 (2.2-9.0) |
| Transfer to another department    | 6.4% (8/126)  | 13% (27/206)  |
| Dead in-hospital                  | 26% (33/126)  | 19% (40/206)  |
| Dead at 60 days                   | 32% (40/126)  | 23% (48/206)  |

*Abbreviations: PCR: polymerase chain reaction, BMI: body mass index, CACI: Charlson’s age adjusted comorbidity score, ICU: intensive care unit, SpO2: peripheral oxygen saturation*

**Oxygen provision**

The mean oxygen flow-rate to the patients while receiving oxygen therapy was 3.0 l/min (SD 2.9) and the median was 2.0 l/min (IQR 1.3-3.5). Results for oxygen provision to the study cohort are shown in Table 2 and for patients that were initially cared for in the wards but later transferred to ICU (therefore not part of the study cohort) in supplementary table 3. The highest oxygen flow-rate provided to the patients was median 4.0 l/min (IQR 2.0-8.0) and mean 5.4 l/min (SD 4.1) (figure 1).
Table 2 Oxygen provision

|                        | Study cohort (n=126) | Subgroups                         |
|------------------------|----------------------|-----------------------------------|
|                        | Patients aged <70 (n=62) | Patients with a no-ICU-decision (n=60) |
| Age, median (IQR)      | 70 (57-82)           | 57 (48-61)                        | 83 (75-88) |
| Days on oxygen         | 2.3 (0.68-4.2)       | 1.8 (0.68-3.9)                    | 2.5 (0.77-4.8) |
| treatment, median      |                      |                                  |            |
| (IQR)                  |                      |                                  |            |
| Oxygen flow to         |                      |                                  |            |
| patients during        |                      |                                  |            |
| oxygen therapy         |                      |                                  |            |
| (l/min)                |                      |                                  |            |
| - Mean (SD)            | 3.0 (2.9)            | 2.6 (2.3)                        | 4.0 (3.7)  |
| - Median (IQR)         | 2.0 (1.3-3.5)        | 1.9 (1.3-2.9)                    | 2.9 (1.6-5.6) |
| Oxygen flow to         |                      |                                  |            |
| patients during time   |                      |                                  |            |
| in the ward (l/min)    |                      |                                  |            |
| - Mean (SD)            | 2.2 (2.9)            | 1.8 (2.3)                        | 3.1 (3.7)  |
| - Median (IQR)         | 1.2 (0.32-2.6)       | 1.1 (0.43-2.0)                   | 1.6 (0.28-4.6) |
| Total volume of        |                      |                                  |            |
| oxygen provided per    |                      |                                  |            |
| patient admission (l)  |                      |                                  |            |
| - Mean (SD)            | 16,000 (23,000)      | 12,000 (16,000)                  | 22,000 (26,000) |
- Median (IQR)  
  7,400 (1,200-21,000)  
  4,8000 (1,800-17,000)  
  12,000 (2,000-32,000)

**Figure 1** Highest oxygen flow-rate received by patients with severe COVID-19 during their care in the medical wards

**Discussion**

We have found that a mean of 3.0 l/min (median 2.0 l/min) oxygen were provided to severely ill COVID-19 patients while receiving oxygen in a Swedish district hospital during the first peak of the pandemic in 2020. This calculated oxygen flow-rate is lower than the 10 l/min estimated by the WHO for COVID-19 patients with severe disease (22).

A strength of this study is the use of a method for calculating oxygen provision to hospitalized patients by summing up documented flow-rates in the medical records. Previously, findings of the proportions of patients receiving oxygen (8,29–32) and assumptions on required oxygen flow (22) have been presented. To the best of our knowledge, this study is the first to use all the necessary data to be able to calculate total oxygen provision to patients.

The 206 patients admitted to the medical department in Nyköping with COVID-19 during this study period had similar age, gender proportions and in-hospital mortality as other described cohorts of hospitalised COVID-19 patients (8,29,31,33). The median length of stay of 4.3 days was shorter in the Nyköping patient group than in two cohorts from New York and Madrid described during the early pandemic, as well as in the multinational ISARIC survey, with stays of seven to nine days (31,33,34). This could be explained by Sweden’s early discharge policies (35,36) and a considerable
amount of patient transfers in this study (13% of all admitted patients and 6.4% of the study cohort) due to overcrowding.

The lower flow-rates of oxygen to patients with severe COVID-19 in this study compared to previous estimates are interesting but, for several reasons, should be interpreted with caution when planning health care services. Firstly, it is treacherous for health care systems to make plans based on the COVID-19 patient categories “severe” and “critical” as there are currently several definitions in use and many of them categorise patients based on the care they receive rather than the care they require (22,27,37). Since resources and practices for ICU-admission vary greatly around the world, for example ICU beds per 100,000 population vary from 29 in Germany to 6 in Sweden and <1 in Uganda (38), very few COVID-19 patients would be classified as critical in Uganda according to the definition used by WHO and in this study. Indeed, in settings with less availability of ICU beds, sicker patients will be classified as “severe” and require higher flows of oxygen than the cohort in this study. In addition, suboptimal respiratory support might cause shorter survival or survival with prolonged time to recovery – hence it is difficult to make assumptions on the quantity of oxygen they need without available data. This study aimed to study a patient group that was truly severe according to the WHO classification, and due to the well-resourced hospital system in Sweden succeeded with this aim, but a similar group may not be easy to delineate in other settings.

Secondly, decisions on treatment limitations were important to the assignment of patients to the severe group in this study and such practices vary largely across countries. Of the study patients, 48% had a no-ICU-decision meaning that even if they deteriorated, they were kept in the general wards and received targeted oxygen treatment. Although oxygen needs for this subgroup was also considerably lower than the WHO estimate (4.0 l/min), it is possible that in settings with other norms around treatment limitations, these patients may, at some point, have received ICU-treatments such as mechanical ventilation with substantially higher oxygen flow-rates.
Thirdly, the flow-rates of oxygen provided to patients depend on target saturation and duration between treatment modifications. The patients in this study were cared for with defined targets and frequent saturation controls. In wards where oxygen flow is not—or cannot be—adjusted as frequently, oxygen flow-rates and target saturations may be higher to provide patients with a safe margin for avoiding hypoxia (39). While the optimal target saturation for hospitalised patients receiving oxygen is debated (40,41), the target range in Nyköping was set following the Surviving Sepsis Campaign guidelines (42).

Our findings suggest that the oxygen need for severely ill COVID-19 patients may be lower than previously estimated. Future research using the methods described in this study in larger cohorts and from other settings would be useful to inform capacity planning with updated estimates of oxygen need. Additionally, as oxygen is essential in many other conditions such as sepsis, trauma (21,43–45) and notably child pneumonia, a disease killing 800,000 children under 5 each year (14,46), estimates for the oxygen needs for treating these conditions would be beneficial.

Conclusion

The provision of oxygen to severely ill COVID-19-patients was lower than previously estimated. Further research is required before global estimates are adjusted.

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Highest oxygen flow-rate (n=126)

- ≤2: 34
- >2 to ≤5: 52
- >5 to ≤10: 29
- >10: 11