

BMJ Open Which parameters support disposition decision in suspected COVID-19 cases in the emergency department (ED): a German clinical cohort study

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ABSTRACT

Objectives One major goal of the emergency department (ED) is to decide, whether patients need to be hospitalised or can be sent home safely. We aim at providing criteria for these decisions without knowing the SARS-CoV-2 test result in suspected cases.

Setting Tertiary emergency medicine.

Participants All patients were treated at the ED of the Charité during the pandemic peak and underwent SARS-CoV-2 testing. Patients with positive test results were characterised in detail and underwent a 14-day-follow-up.

Primary and secondary outcome measures Logistic regression and classification and regression tree (CART) analyses were performed to identify predictors (primary endpoint), which confirm safe discharge. The clinical endpoint was all-cause mortality or need for mechanical ventilation during index stay or after readmission.

Results The primary test population of suspected COVID-19 consisted of n=1255 cases, 45.2% were women (n=567). Of these, n=110 tested positive for SARS-CoV-2 (8.8%). The median age of SARS-CoV-2-positive cases was 45 years (IQR: 33–66 years), whereas the median age of the group tested negative for SARS-CoV-2 was 42 years (IQR: 30–60 years) (p=0.096). 43.6% were directly admitted to hospital care.

CART analysis identified the variables oxygen saturation (<95%), dyspnoea and history of cardiovascular (CV) disease to distinguish between high and low-risk groups. If all three variables were negative, most patients were discharged from ED, and the incidence of the clinical endpoint was 0%. The validation cohort confirmed the safety of discharge using these variables and revealed an incidence of the clinical endpoint from 14.3% in patients with CV disease, 9.4% in patients with dyspnoea and 18.2% in patients with O₂ saturation below 95%.

Conclusions Based on easily available variables like dyspnoea, oxygen saturation, history of CV disease, approximately 25% of patients subsequently confirmed with COVID-19 can be identified for safe discharge.

Trial registration number DRKS00023117.

INTRODUCTION

The Coronavirus Disease 2019 (COVID-19) Pandemic poses unprecedented challenges

Strengths and limitations of this study

- This study constitutes a real-world cohort of patients tested positive for SARS-CoV-2 in the emergency department, and selection bias is, thus, very unlikely.
- All available clinical information were retrieved from patient files and, thus, a thorough analysis of clinical predictors of hospital admission was possible.
- The number of clinical endpoints like mechanical ventilation or death was low and, thus, the primary analysis of outcome prediction was not possible but described.
- Predictors of hospital admission were derived from a single-centre cohort, and thus generalisability might be questionable.
- The identified predictors for hospital admission were, however, validated in an independent, multicentre data set and results were similar, even though admission rates in general were higher in the multicentre cohort.

to modern healthcare worldwide. One of the particular challenges of COVID-19, the disease is caused by the most recently discovered coronavirus called SARS-CoV-2, is the possible presymptomatic or oligosymptomatic human-to-human transmission through small respiratory droplets through close-range contact¹ explaining its exponential spread. Additionally, while most patients develop mild disease, severe courses ranging to critical disease with acute respiratory distress syndrome are particularly prevalent in patients with risk factors, that is, pre-existing pulmonary diseases; age over 60 years or obesity.^{2–7} Due to widespread transmission of the virus with localised clusters and outbreaks in many countries, the management of patients with severe disease stretched healthcare systems in several countries, to their limits, regionally and beyond.

Emergency departments (ED) typically are the first point of contact with the health



system for patients with unclear and severe symptoms and disease states including COVID-19.⁸ On 1 March 2020, the first case in Berlin was diagnosed in our ED at Charité Universitätsmedizin Berlin located in the district Berlin Mitte. Charité university hospital as a tertiary care institution with 3000 beds is one of the largest university hospital centres in Europe. Compared with other European countries, Germany had the advantage of a longer period for health system contingency planning until more intense transmission occurred. This enabled healthcare providers to prepare for the pandemic by implementation of immediate-targeted action and a rapid, proactive and comprehensive approach. The early development and availability of diagnostic tests for virus detection by reverse transcription PCR (rt-PCR) from specimens,⁹ obtained from the upper respiratory tract (eg, oronasopharyngeal swabs), early establishment of testing capacities and the fact that the population initially affected in Germany was of younger age compared with other countries contributed to facilitating the country's response to the outbreak.

The public call for rapid sharing of research data resulted in a vast amount of high-ranking published papers from China and around the globe. Strategies for initial evaluation of suspected COVID-19 cases in the ED or fever outpatient clinic¹⁰ according to the original characteristics of the confirmed cases of COVID-19¹¹ were proposed. Modified clinical strategies were established in Germany. However, these proposals are based on clinical observations only and have not been backed up with data, yet.

For the present study, all patients tested for SARS-CoV-2 via rt-PCR from oronasopharyngeal swabs, obtained in the EDs of Charité Universitätsmedizin Berlin in the highly affected district Mitte, in the period from 1 March (case no. 1) to 15 April were analysed. The period covers the time of the first wave with the highest case numbers.

All patients with laboratory-confirmed SARS-CoV-2 infection were clinically characterised and followed up for at least 14 days. Factors associated with disposition and clinical outcome were determined and validated using independent multiple-centre cohort of confirmed cases of COVID-19 at the EDs of the University Hospital of Cologne, the University Hospital of Münster, the University Hospital of Kiel and the University Hospital of Essen, Germany between 1 March and 5 May 2020.

METHODS

Derivation cohort

ED patients at risk for infection were identified by a site-specific algorithm and isolated prior to the initial assessment of the clinical status and, depending on the severity of the illness, were directed to appropriately equipped and separate treatment areas. Oronasopharyngeal swabs for the detection of SARS-CoV-2 RNA via rt PCR were performed in all suspected cases. Berlin's first COVID-19 case was confirmed on 1 March 2020 at the ED at Charité

Universitätsmedizin Berlin in the Mitte district in the EDs of Charité Campus Virchow Klinikum and Campus Charité Mitte. From 1 March to 15 April, all primary suspected cases of COVID-19 or cases in differential diagnostic clarification according to the definition published by the German Robert Koch Institute¹² were included in the present study.

All following cases with confirmed SARS-CoV-2 infection were prospectively recorded and followed up in clinical routine. Hospital-wide and ED-specific protocols¹³ were established addressing establishment of a core team and key internal and external contact points, focusing on human, material and facility capacity, communication and data protection, training procedures, hand hygiene, personal protection equipment, waste management, triage, first contact and prioritisation, patient placement, moving of the patients in the facility and visitor access as well as environmental cleaning as proposed by international institutions.

Oligosymptomatic or asymptomatic self-presenter at our SARS-CoV-2 testing centre, which was set up as early as 3 March 2020, was not included in the analysis. Furthermore, confirmed cases tested positive prior to their presentation in the ED are not included in the series of primary suspected cases or cases in differential diagnostic clarification according to the definition published by the German Robert Koch Institute.¹²

In addition, a consecutive series of n=127 SARS-CoV-2-positive cases from the *Registry for Clinical Presentation and Management of Patients With COVID-19 in the Emergency Room (ReCovER)* and diagnosed in the ED of the University Hospitals of Cologne, Essen, Kiel and Münster between 1 March to 5 May 2020 were used for the validation of the multivariate model.

Data collection and endpoints

Clinical characteristics and in-hospital follow-up information of confirmed SARS-CoV-2 cases were extracted from electronic medical records. Particular attention was paid to vital parameters, certain chronic medical conditions, comorbidities and use of medication as ibuprofen, angiotensin converting enzyme inhibitors and angiotensin-II receptor antagonists.

Patients who did not meet the criteria for admission to the hospital and were discharged from ED were contacted by medical staff via telephone to inform them about their positive test result, educated on the importance of infection control and self-isolation and provided with instructions on the anticipated duration of isolation and warning signs that should prompt reevaluation. Another telehealth visit was scheduled within 14 days following the day of the ED presentation and confirmation of the infection at the University Hospitals of Berlin and Cologne.

The primary endpoint of the analyses was hospital admission after index presentation to the ED. Two further secondary combined clinical endpoints were used. The first clinical combined endpoint was intubation and death before discharge from the hospital or in case of initial

outpatient treatment after readmission to hospital. The second clinical endpoint included the components of the first clinical endpoint (intubation and death) and additionally all intensive care unit (ICU) admissions during the index stay and all inpatient readmissions.

Statistical analysis

The present analysis focuses on confirmed SARS-CoV-2 cases. The population of patients with suspected COVID-19 infection also included the SARS-CoV-2 negative cases and was characterised in terms of age and sex in the derivation cohort. In the subgroup of positive SARS-CoV-2 cases, the primary endpoint was hospital admission and cases were compared between primary outpatients and admitted patients using descriptive analysis of characteristics, clinical parameters and postemergency care. With regard to the derivation of clinical decision strategies, clinical characteristics were analysed in consideration of the published proposals with regard to the primary endpoint. Descriptive analyses included the calculation of relative and absolute frequencies as well as median and IQR. Statistical differences were calculated using the χ^2 test for categorical variables and the Mann-Whitney test for continuous variables. A p value of less than 0.05 was considered statistically significant. Due to the exploratory nature of the analysis, no corrections were made for multiple testing. Potential predictors for in-hospital treatment were first analysed univariately by above-mentioned statistical tests. All significant predictors were analysed by classification and regression analyses.¹⁴ For numeric variables, Youden optimised cut-off values were determined on basis of receiver operating characteristic curves. The identified predictors were then analysed as binary variables in logistic regression analysis and the best predicting variables were selected based on effect measure (OR) and model fit criteria (Cox & Snell R-Square and Nagelkerkes R-Square) after selection of the first best prediction variable and cut-off. The above-mentioned procedures were then repeated at each step in the resulting subgroups. All identified predictors, including cut-offs and resulting subgroups, were graphically illustrated by classification and regression trees (CART). All the above-mentioned analyses were performed in the derivation cohort of Charité Universitätsmedizin Berlin.

Validation cohort

The final CART model, that is, the identified predictors and cut-offs, which was developed in the Charité-cohort and can thus be regarded as optimised for this cohort, was then applied to the validation cohort. This validation cohort consisted of patients enrolled into the *ReCovER* of the EDs of the University Hospital of Cologne, Münster, Essen and Kiel. The *ReCovER* registry has been approved by the Ethics Committee of the Medical Faculty of the University of Cologne (EK 20–1198, NCT04351854). The Essen and Münster cohorts of confirmed-patients with SARS-CoV-2 were enrolled with approval of the respective Ethics committees (file numbers: 20-9310-BO,

2929-571-b-S). The principles of the basic data protection regulation apply.

Patient and public involvement statement

The development of the research question, study design and outcome measures was developed by a team of experienced ED doctors and researchers who also concerned patients' perceived preferences and priorities. Patients were not involved directly in these processes. The results of this research work are going to be published open access and disseminated to interested patients via the website of the institution.

RESULTS

Figure 1 shows the patient flowchart of the Berlin cohort with the corresponding case numbers.

The primary population of suspected COVID-19 cases which received testing in the ED consisted of n=1255 cases, 45.2% were women (n=567). The median age was 42 years (IQR: 31–60 years). The proportion of female patients was 39.1% (n=23) in the group of confirmed SARS-CoV2 cases, which was slightly lower than the proportion of female patients who tested negative for SARS-CoV2 at 45.3% (n=502; p=0.095). The median age of confirmed SARS-CoV-2 cases was in median 45 years (IQR: 33–66 years), whereas the median age of the group tested negative for SARS-CoV-2 was 42 years (IQR: 30–60 years) (p=0.096).

Figure 2 shows the daily test numbers during the study period and the proportion of positive cases that reached their maximum at the end of March. The online supplemental table 1S shows basic characteristics of the SARS-CoV-2 negatives (n=1070, nmiss=38).

Characteristics of patients with confirmed SARS-CoV-2 infection

Table 1 depicts the clinical characteristics of patients with confirmed SARS-CoV-2 infection stratified by the primary endpoint (outpatient care or admission to the hospital). The proportion of women was higher in the outpatient group and the age was lower compared with patients admitted to the hospital on ED presentation. Significant differences in vital parameters were observed for temperature, respiratory rate and oxygen saturation. The frequency of diarrhoea, dyspnoea and abdominal pain was higher in hospitalised patients compared with outpatients. Among common risk factors, pre-existing cardiovascular (CV) and hepatic diseases significantly associated with in-patient treatment.

Laboratory parameters at admission were in general comparable and showed a broad overlap between outpatients and hospitalised patients. However, there were significant differences in pCO₂, pH, glucose, lactate, lymphocytes, lactate dehydrogenases (LDH), C reactive protein (CRP) and procalcitonin (table 2).

Prediction of hospitalisation in patients with SARS-CoV-2-positive

Variables for classification and regression analyses were selected based on the bivariate association with

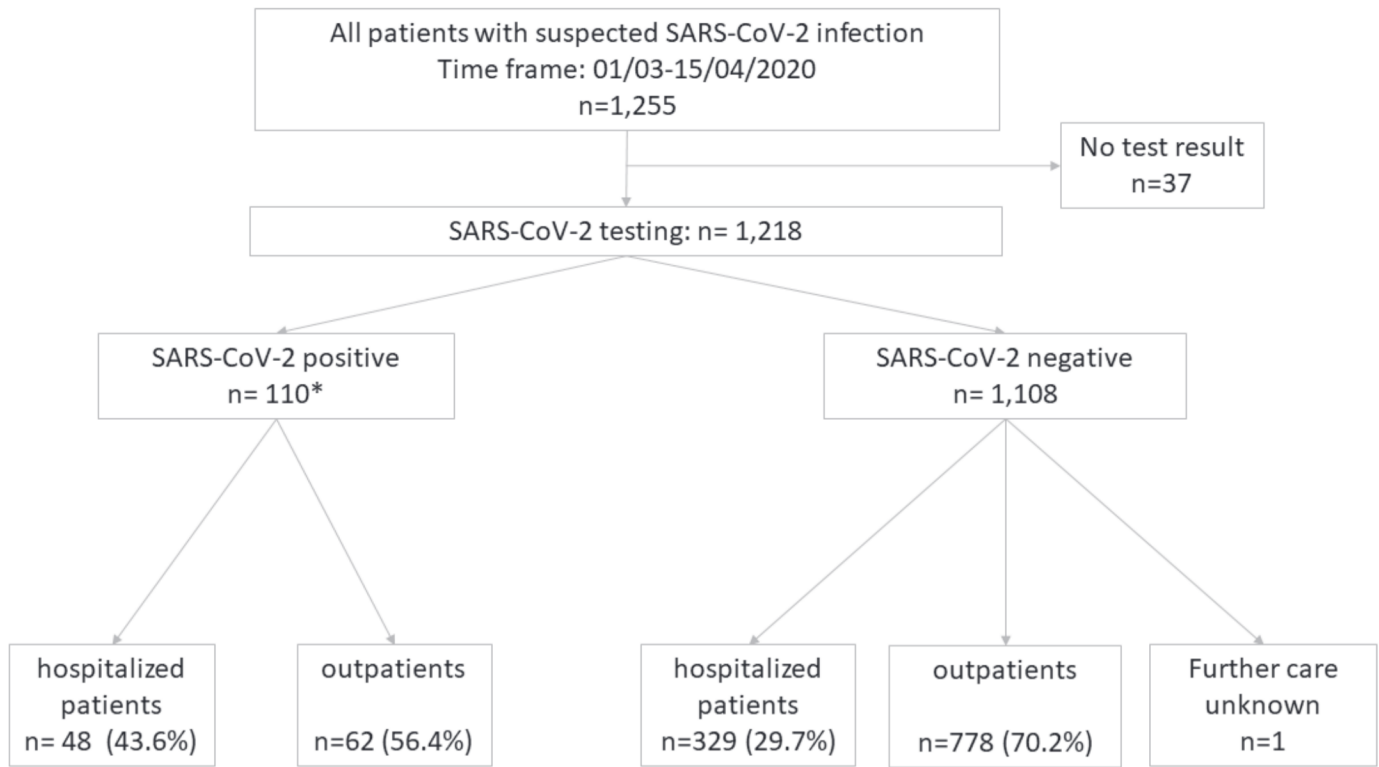


Figure 1 Patient flow diagram of the derivation cohort. *SARS-CoV-2 positive: n=106 positive tests performed at Charité laboratory, n=4 confirmed cases tested positive prior to their presentation in the ED (later confirmed in Charité laboratory) were also included in the analysis (total n=110). ED, emergency department.

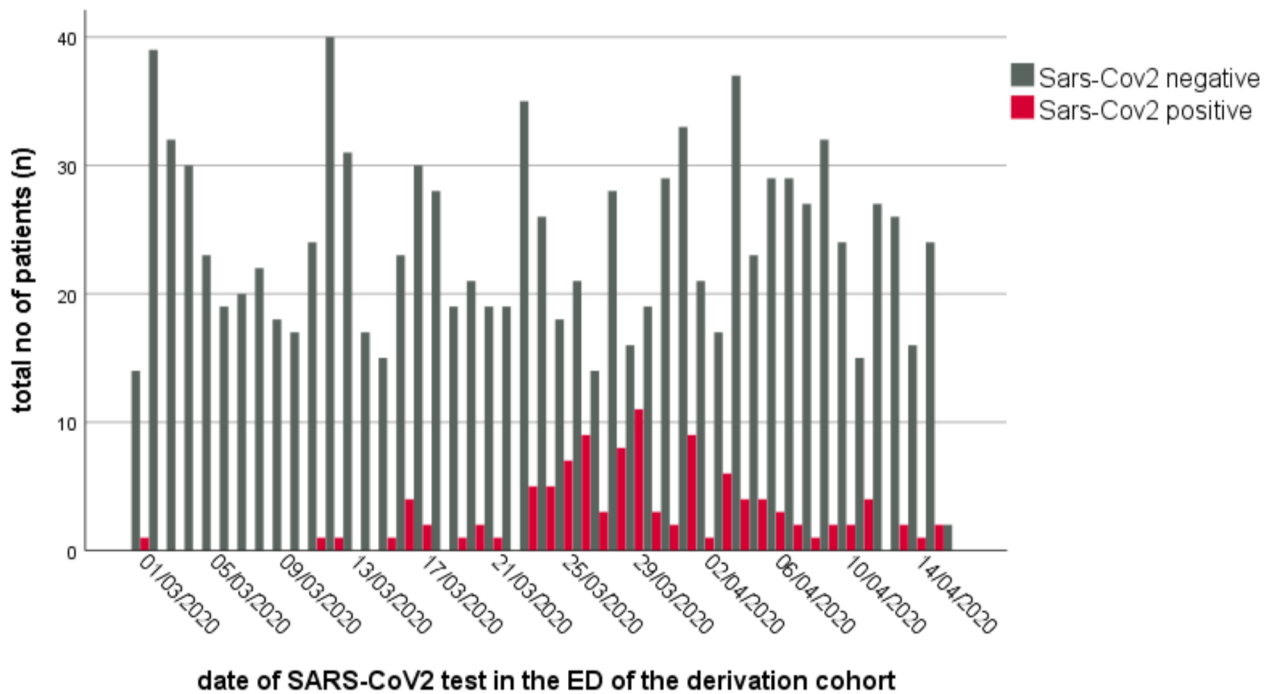


Figure 2 Absolute number of SARS-CoV-2 negative tests (blue) and confirmed SARS-CoV-2 cases (red) in patients with ED at Charité Universitätsmedizin Berlin (CVK, CCM). CCM, CampusCharité Mitte; CVK, CampusVirchow Klinikum; ED, emergency department.

Table 1 Demographic and clinical characteristics for patients with SARS-CoV2-positive with initial ambulatory treatment (outpatients) in the ED or inpatient treatment at Charité Universitätsmedizin Berlin

	SARS-CoV-2 positive hospitalised patients (n=48)	SARS-CoV-2 positive outpatient treatment (n=62)	P value
Women % (n)	31.3 (15)	50.0 (31)	0.048
Age (median. IQR)	56 (42–78)	38 (30–49)	<0.0001
BMI (median. IQR)	27 (24–31)	28 (22–30)	0.874
Vital signs (median. IQR)			
BP syst. mm Hg	135 (118–150)	133 (122–147)	0.912
BP diast. mm Hg	75 (69–86)	80 (74–90)	0.061
Heart rate/min	89 (79–99)	90 (80–101)	0.705
Temperature °C	37.8 (37.2–38.8)	37.3 (36.6–37.8)	0.004
Respiratory rate/min	18 (16–24)	16 (15–18)	0.009
Oxygen saturation %	95 (93–97)	99 (97–100)	<0.0001
Vital signs at established risk cut-offs			
BP syst. <90mm Hg/diast. ≤60mm Hg	10.4 (5)	12.9 (8)	0.012
Temperature >37.3°C	66.7 (32)	43.5 (27)	0.019
Respiratory rate >18/min	37.5 (18)	12.9 (8)	0.011
Respiratory rate >30/min	8.3 (4)	0	0.051
Oxygen saturation <90%*	12.5 (6)	0	0.009
Symptoms % (n)			
Fever	75.0 (36)	62.9 (39)	0.137
Cough	62.5 (30)	51.6 (32)	0.080
Haemoptysis	0	0	–
Sore throat	10.4 (5)	16.1 (10)	1.000
Rhinitis	4.2 (2)	4.8 (3)	0.584
Headache/muscle pain	25.0 (12)	46.8 (29)	0.707
Dyspnoea	58.3 (28)	19.4 (12)	<0.0001
GI-symptoms	12.5 (6)	6.5 (4)	0.064
Diarrhoea	27.1 (13)	9.7 (6)	0.004
Nausea/emesis	12.5 (6)	12.9 (8)	0.476
Loss of smell	2.1 (1)	3.2 (2)	0.783
Abdominal pain	10.4 (5)	3.2 (2)	0.029
Symptom onset time (days)	7.5 (5.0–10.8)	3.5 (2.0–6.0)	<0.0001
Risk factors % (n)			
Transplantation	4.2 (2)	1.6 (1)	0.532
Tumour	4.2 (2)	1.6 (1)	0.518
Cardiovascular disease	52.1 (25)	14.5 (9)	<0.0001
Respiratory disease	20.8 (10)	16.1 (10)	0.747
Renal disease	12.5 (6)	3.2 (2)	0.122
Hepatic disease	14.6 (7)	1.6 (1)	0.022
Pregnancy	4.2 (2)	0	0.290

Demographic and clinical characteristics for patients with SARS-CoV2-positive with initial ambulatory treatment (outpatients) in the ED or inpatient treatment at Charité Universitätsmedizin Berlin (CVK, CCM). The cut-offs presented in this table are previously reported risk cut-offs and were not derived from the current data analysis.

*The cohort consists of patients with and without oxygen supplementation.

BMI, body mass index; BP, blood pressure; CCM, Campus Charité Mitte; CVK, Campus Virchow Klinikum; ED, emergency department; GI, gastrointestinal.

the primary endpoint of hospital admission. [Table 3](#) summarises the results of the best predictors in bivariate analyses. Decreased oxygen saturation, older age,

presence of dyspnoea, longer duration of time since symptom onset, history of CV disease, elevated lactate, LDH and CRP were significantly associated with the

**Table 2** Laboratory parameters of patients with SARS-CoV2-positive with initial ambulatory treatment (outpatients) in the ED or inpatient treatment at Charité Universitätsmedizin Berlin (CVK, CCM)

	SARS-CoV-2 positive hospitalised patients (n=48)	SARS-CoV-2 positive outpatient treatment (n=62)	
pO ₂ (mm Hg) (median. IQR) (nmiss=38)	28.5 (20.4–34.8)	29.2 (23.5–36.2)	0.605
pCO ₂ (mm Hg) (median. IQR) (nmiss=32)	40.8 (37.2–45.2)	46.0 (41.1–49.5)	0.010
pH (median. IQR) (nmiss=32)	7.41 (7.38–7.45)	7.39 (7.36–7.41)	0.017
HCO ₃ ⁻ (mmol/L) (median. IQR) (nmiss=32)	26.0 (24.1–27.3)	26.9 (24.9–28.4)	0.132
BE (mmol) (median. IQR) (nmiss=35)	1.20 (-0.20–3.10)	2.35 (0.03–3.00)	0.483
Sodium (mmol/L) (median. IQR) (nmiss=32)	137 (134–140)	138 (136–141)	0.144
Potassium (mmol/L) (median. IQR) (nmiss=31)	4.0 (3.7–4.2)	4.0 (3.7–4.3)	0.684
Chloride (mmol/L) (median. IQR) (nmiss=33)	103 (101–107)	104 (101–107)	0.447
Glucose (mg/dl) (median. IQR) (nmiss=33)	130 (110–150)	108 (97–123)	0.001
Haemoglobin (g/L) (median. IQR) (nmiss=25)	138 (127–151)	139 (132–150)	0.790
Lactate (mg/dL) (median. IQR) (nmiss=35)	14.5 (12.0–19.0)	11.0 (9.5–13.5)	<0.0001
WBC (10exp9/L) (median. IQR) (nmiss=26)	6.9 (5.1–8.9)	5.7 (4.5–8.4)	0.406
Lymphocytes (%) (median. IQR) (nmiss=54)	0.92 (0.66–1.36)	1.41 (1.19–2.07)	0.003
CRP (mg/L) (median. IQR) (nmiss=31)	60.6 (27.1–118.8)	14.4 (3.9–30.4)	<0.0001
LDH (U/L) (median. IQR) (nmiss=44)	394 (291–501)	246 (212–326)	<0.0001
PCT (µg/L) (median. IQR) (nmiss=51)	0.11 (0.07–1.55)	0.05 (0.04–0.08)	0.002

Laboratory parameters are shown as median and IQR for patients with SARS-CoV2-positive with initial ambulatory treatment (outpatients) in the ED or inpatient treatment at Charité Universitätsmedizin Berlin (CVK, CCM).

BE, base excess; CCM, Campus Charité Mitte; CRP, C reactive protein; CVK, Campus Virchow Klinikum; ED, emergency department; HCO₃⁻, bicarbonate; LDH, lactate dehydrogenases; pCO₂, partial pressure of carbon dioxide; PCT, procalcitonin; PH, power of hydrogen; pO₂, partial pressure of oxygen; WBC, white cell count.

primary endpoint (admission to hospital). Oxygen saturation was the best predictor of hospital admission regarding area under the receiver operating

characteristics curve (0.822; 95% CI 0.735 to 0.909) and model fit criteria (R² values: Cox and Snell=0.237; Nagelkerkes=0.316).

Table 3 Bivariate analysis of parameters regarding the prediction of hospital admission in patients with SARS-CoV2-positive in the emergency department (ED) at Charité Universitätsmedizin Berlin (CVK, CCM)

	AUROC (95% CI)	P value ROC-analysis	Best cut-off value	OR (95% CI)	P value logistic regression	Cox and Snell R-square	Nagelkerkes R ²
Oxygen saturation (%)*	0.822 (0.735 to 0.909)	<0.0001	95%	36.6 (4.7 to 288.1)	0.001	0.237	0.316
Age (years)	0.775 (0.680 to 0.869)	<0.0001	55	9.5 (3.9 to 23.3)	<0.0001	0.225	0.302
Dyspnoea	NA	NA	NA	7.2 (2.9 to 17.7)	<0.0001	0.189	0.253
Time since onset of symptoms (days)	0.751 (0.645 to 0.858)	<0.0001	6.5	6.0 (2.3 to 15.5)	<0.0001	0.16	0.214
CV disease	NA	NA	NA	5.6 (2.2 to 14.2)	<0.0001	0.144	0.193
Lactate (mg/dL)	0.745 (0.632 to 0.857)	<0.0001	12.5	2.2 (1.0 to 5.2)	0.062	0.033	0.044
LDH (U/L)	0.798 (0.684 to 0.913)	<0.0001	272	2.4 (0.9 to 6.9)	0.092	0.028	0.038
CRP (mg/L)	0.786 (0.682 to 0.890)	<0.0001	30	2.0 (0.9 to 4.7)	0.099	0.026	0.035

Results of the receiver operating characteristics analysis (ROC), best identified cut-off values (optimised by Youden-Index) according to the ROC-curve and results of the univariate logistic regression analysis of dichotomised predictors. Also, goodness of fit criteria are reported (R²). The Cox and Snell R² is not standardised and can only be used to compare values between different models. The Nagelkerkes R² is standardised and produces values between 0 and 1.

*The cohort consists of patients with and without oxygen supplementation.

AUROC, area under the receiver operating characteristics curve; CCM, Campus Charité Mitte; CRP, C reactive protein; CV, cardiovascular; CVK, Campus Virchow Klinikum; LDH, lactate dehydrogenase.

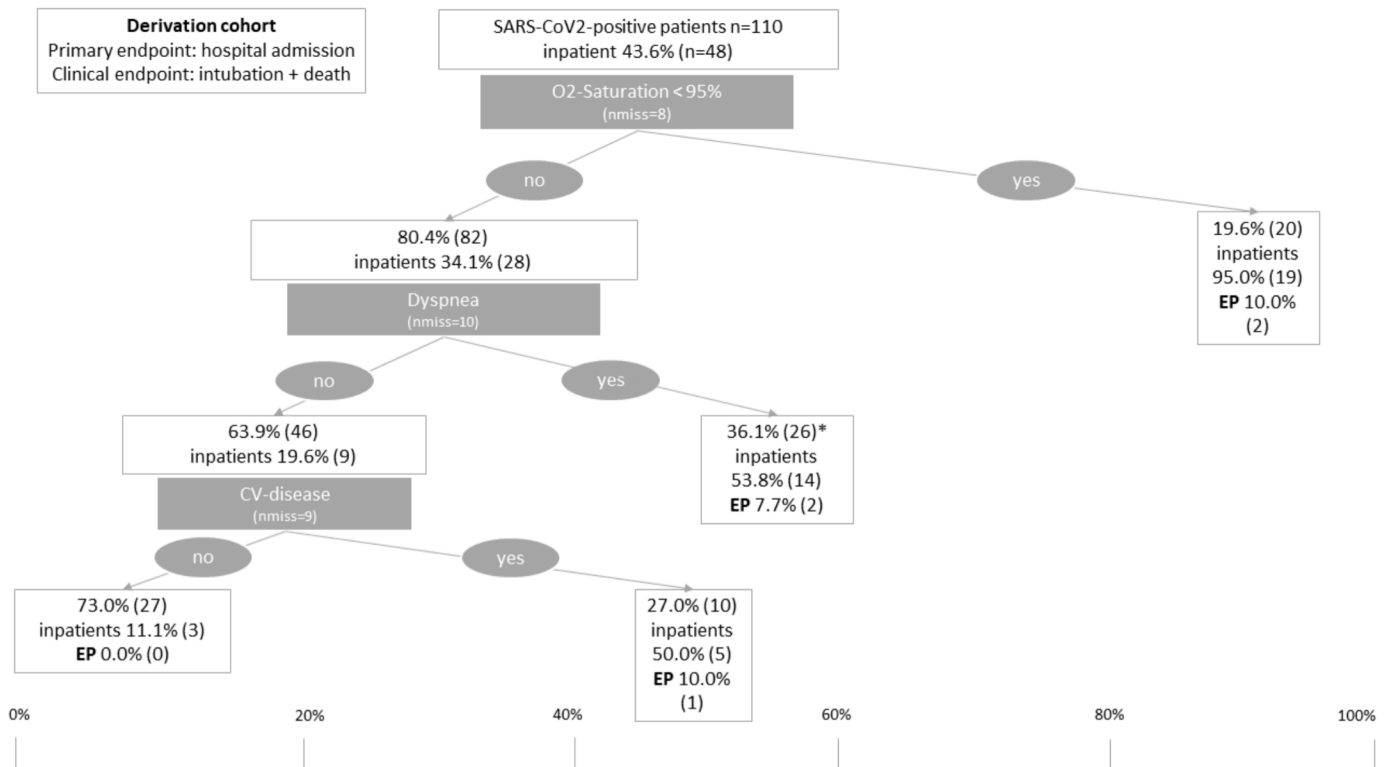


Figure 3 Classification and regression tree in the derivation cohort of Charité Universitätsmedizin Berlin. The position of the boxes on the x-axis illustrated the frequency of inpatient treatment in per cent while the size of the boxes is proportional to the size of the respective patient subgroup. Additionally to the proportion of patients who were admitted, the frequency of the clinical EP is displayed. *There was no further discriminating variable in the subgroup of patients with dyspnoea. CV, cardiovascular; EP, endpoint; nmiss, number of missing values; O₂, oxygen.

Ninety-five per cent was the best cut-off point for oxygen saturation to discriminate between patients who were admitted to hospital and those who were discharged home. Decreased oxygen saturation below 95% was observed in 19.6% (n=20) of confirmed cases with SARS-CoV-2 infection. There was no further distinct discriminating variable in this subgroup with an already high hospitalisation rate of 95% (n=19) in the group of SARS-CoV-2 cases with oxygen saturation below 95%. The clinical endpoint (intubation, death during index stay or after readmission) occurred in 10% (n=2) of this high-risk subgroup (figure 3). Report of dyspnoea and history of CV disease were the best further discriminating variables within the subgroup of patients with an oxygen saturation at or above 95% (80.4%; n=82). When the patient population was further divided into risk categories based on this information, patients with oxygen saturation at or above 95% but with reported dyspnoea or history of CV diseases showed a hospitalisation rate of 53.8% (n=14) and 50.0% (n=5), respectively. The risk for being intubated or death was 7.7% and 10%, respectively. No clinical endpoint occurred in the remaining patients with lower risk for hospitalisation (11.1%, n=3) and none of the above-mentioned conditions (n=27). The hospitalised patients with confirmed SARS-CoV-2 infection in the low risk group were diagnosed with cholangitis (n=1) or viral pneumonia (n=2) during their hospitalisation. All three hospitalised low-risk group patients were initially admitted to designated non-ICU COVID-19 wards. Both

patients with viral pneumonia were transferred to ICU over the course of their stay. After recovery, all three patients were discharged home.

The derivation cohort consisted of 39.1% female patients and the median age was 45 years. The demographic characteristics of the validation cohorts were Cologne: female 47.5%, median age 52 years; Essen: female 34.0%, median age 71 years; Münster: female 25%, median age 57 years; Kiel: female 41.7%, median age 63 years. In the validation cohort, consisting of 127 confirmed cases with SARS-CoV-2 infection who presented to the EDs of the University Hospital of Cologne, Münster, Essen and Kiel, the proportion of inpatients was in general higher and, thus, also higher in the respective lower risk groups (figure 4). In total, 66.9% (n=121) of SARS-CoV-2-positive patients were admitted to hospital in the validation cohort. The proportion of patients with an oxygen saturation at or above 95% was higher as compared with the derivation cohort (30.4%; n=55) but resulted in a similar risk of admission to hospital (94.5%; n=52). The clinical endpoint occurred in 18.2% (n=10) in this higher risk subgroup. The presence of symptoms of dyspnoea or pre-existing cardiovascular disease resulted in risk groups with a proportion of inpatients of 68.8% and 81.0%, respectively, and, thus, also higher as compared with the derivation cohort. Admission to hospital occurred in 63.2% in the low-risk group of the validation cohort (no dyspnoea and no CV disease).

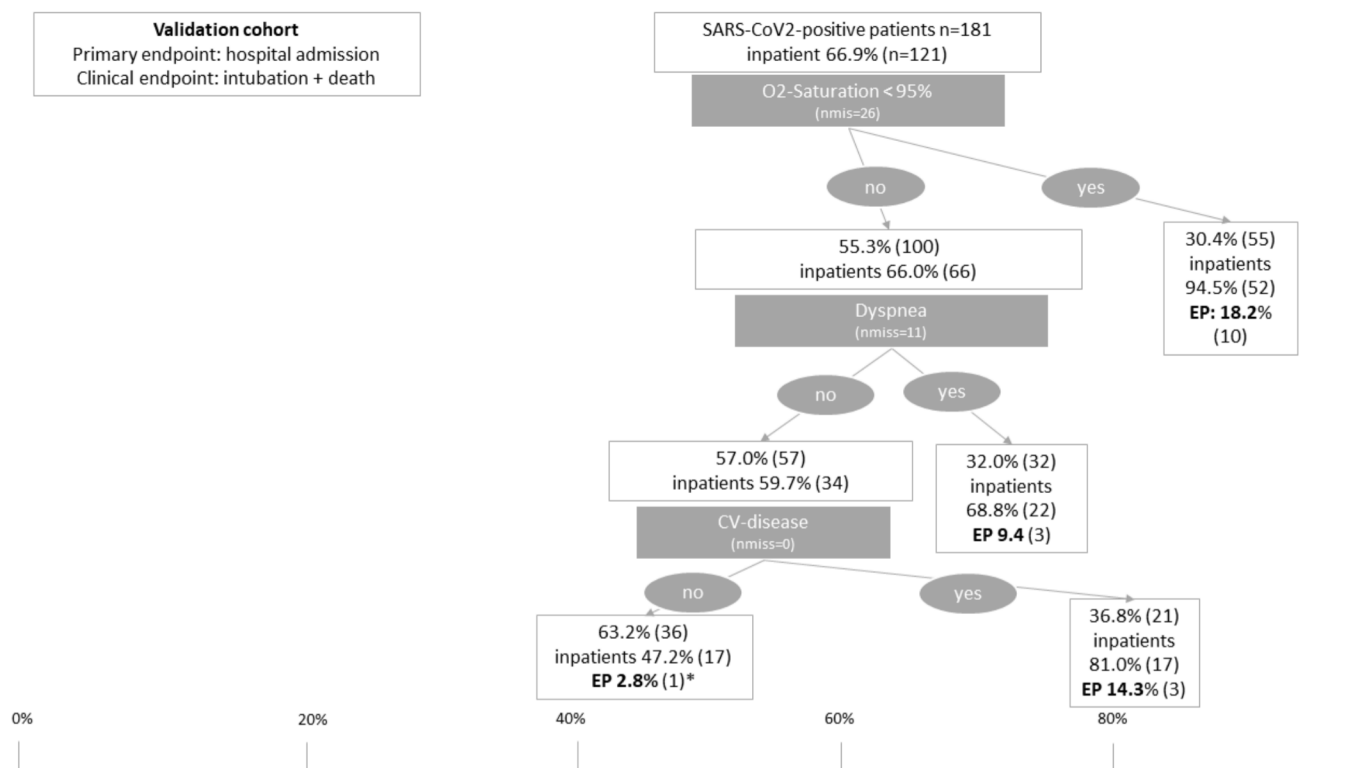


Figure 4 Classification and regression tree in the validation cohort of the EDs of the University Hospitals of Cologne, Münster, Essen and Kiel from the ReCover registry. The position of the boxes on the x-axis illustrated the frequency of inpatient treatment in per cent while the size of the boxes is proportional to the size of the respective patient subgroup. In the low risk group, one patient had a clinical endpoint. This patient was admitted to the ward directly from the ED and had a terminal oncological disease, which led to patient's death without ICU admission. ED, emergency department; ICU, intensive care unit.

Clinical endpoints in the derivation cohort

The first combined clinical endpoint (intubation, death during index stay or after readmission) occurred in 5% (n=5). The second combined clinical endpoint (intubation, death, inpatient readmission and all ICU admissions) was positive in 40% of all cases (n=44). Of all admitted patients, 37.5% (n=8) received oxygen supplementation. Of all 48 admitted patients, 23 patients were discharged during the follow-up period of 14 days. All other patients were still in hospital after 14 days and further length of hospital stay was not assessed. For these 23 patients, length of stay ranged between 1 and 12 days with a median length of stay (LOS) of 7 days (IQR: 4–10 days) (table 4).

DISCUSSION

In the present study, we provide evidence that the primary disposition of patients with suspected COVID-19 based on oxygen saturation, the cardinal symptom of dyspnoea and history of CV disease is safe and effective.

In contrast to most current strategies for risk stratification in COVID-19, which were developed focusing on severely ill hospitalised patients and the use of biomarkers like cardiac troponin and D-Dimer^{10 11 15–17}, our study investigates a cohort of suspected cases of COVID-19 in the ED for the first time. Approximately, 10% of the suspected cases were confirmed with COVID-19% and

43.6% were primarily admitted to hospital care (see figure 1).

Using a CART-model, we were able to identify a risk stratification strategy for suspected cases of COVID-19 at initial presentation in the ED that is based on the clinical items oxygen saturation, dyspnoea and history of CV disease. Our risk stratification strategy uses easy-to-examine criteria and, thus, may reveal its distinctive strength, especially when healthcare systems are challenged: first, these criteria are explicitly defined and can readily be assessed at the time of the initial presentation due to suspected COVID-19. Second, these criteria rely on information taken exclusively from the initial patient history and their physical examination. This may facilitate risk assessment for physicians in the outpatient setting, where ordering laboratory tests may be too time-consuming, costly and difficult to perform during the current pandemic. Third, our prediction strategy may also help reduce uncertainty and result in evidence-based use of patient admission to inpatient care, which may be crucial when healthcare systems are being stretched.

There were differences in the proportion of patients admitted to hospital, with an in general higher hospitalisation rate in the validation cohort. This is most likely caused by different standards for hospital admission and availability of beds in the respective hospitals. In addition, it could be influenced by prehospital patient selection.

Table 4 Clinical endpoints of patients with SARS-CoV2-positive with initial ambulatory treatment (outpatients) in the ED or inpatient treatment at Charité Universitätsmedizin Berlin (CVK, CCM)

	SARS-CoV-2-positive hospitalised patients (n=48)	SARS-CoV-2-positive outpatient treatment (n=62)
Primary hospitalised patients		
Intensive care unit % (n)	60.4% (n=29)	0
Intubation % (n)	10.4% (n=5)	0
ECMO % (n)	0.0% (n=0)	0
Death % (n)*	4.2% (n=2)	0
Rehospitalisation of patients with primary outpatient treatment		
Representation at Charité ED % (n)	DNA	17.7% (n=11)
Hospital admission Charité % (n)	DNA	6.5% (n=4)
ICU/intubation/ECMO % (n)	DNA	0.0% (n=0)
Discharged home % (n)	DNA	3.2% (n=2)†
Telehealth follow-up of patients with outpatient treatment during index stay		
Re-presentation other ED % (n)	DNA	4.8% (n=3)
Hospitalisation in other hospital % (n)	DNA	3.2% (n=2)
ICU in other hospital % (n)	DNA	0.0% (n=0)
Death % (n)	DNA	0.0% (n=0)

Clinical endpoints of patients with SARS-CoV2-positive with initial ambulatory treatment (outpatients) in the ED or inpatient treatment at Charité Universitätsmedizin Berlin (CVK, CCM). Patients were followed up during their index stay and patients with primary ambulatory (outpatient) treatment received a follow-up call to assess further clinical course and endpoints.

*Patients who died were intubated before death. Thus the first clinical endpoint occurred in n=5 patients.

†n=2 of cases with primary outpatient treatment were readmitted to hospital and were still in hospital at follow-up.

CCM, Campus Charité Mitte; CVK, Campus Virchow Klinikum; DNA, does not apply; ECMO, extracorporeal membrane oxygenation; ED, emergency department; ICU, intensive care unit.

However, the risk stratification of the above-mentioned criteria (oxygen saturation, dyspnoea and CV diseases) could be confirmed in the validation cohort. Especially the identification of patients of high risk of hospital admission based on oxygen saturation at admission resulted in comparable risk of about 95%. Of note, also other classification methods could have been used to identify predictors of hospital admission. The aim of the current analyses was to provide a robust risk classification algorithm, based

on routinely available clinical information which could be easily implemented in clinical routine. CART analysis is a robust statistical method to detect reliable predictors.¹⁴

However, we must address several limitations of our study: our study was conducted and validated in the EDs of five German University Hospitals in total. This leads to a selection bias, since the majority of asymptomatic to very mild suspected cases of COVID-19 may have been seen in the outpatient setting in testing centres or by their general practitioner and not in the ED. Nevertheless, in the early phase ('first wave') of the pandemic, the university EDs took care of many mild cases also, as general care facilities were in a delayed building up process. Furthermore, patients defined as being at low risk due to our simplified risk stratification strategy may have other medical or psychosocial contraindications to be treated in outpatient setting in the current COVID-19 pandemic. For example, patients may have pre-existing disorders, such as relevant oncological or neuromuscular diseases, which are not part of our risk assessment strategy, but which may lead to an increased likelihood of progression to severe illness during the course of COVID-19. Moreover, oxygen saturation might be influenced by early, undocumented oxygen supplementation in the ED. This could have masked some more severe cases with falsely high oxygen saturation and thus an potential underestimation of the discriminatory abilities of oxygen saturation as a risk predictor for hospitalisation in the current study. Additionally, patients with severe impaired cognitive function, underlying psychiatric diseases or with little social support, may require inpatient care regardless of the severity of underlying COVID-19. Furthermore, patients might be unable to adequately self-isolate themselves or are living in residential homes without adequate isolation management. In addition, our risk stratification does not apply to infants and children. The evaluation of infants and children needs to be developed separately. Another limitation is the inclusion of admitted patients only at the university hospitals of Essen, Kiel and Münster. Finally, during the pandemic, the ER utilisation was dramatically lower worldwide and it needs to be confirmed that the developed criteria are robust under regular circumstances.⁸

As a conclusion, we have developed an easy-to-determine risk stratification assay in a large set of suspected cases of COVID-19 that has been validated in an independent cohort from four separate German Emergency Medicine Centres. Our data provide preliminary evidence of a risk stratification strategy helping to determine whether hospital care is necessary in suspected cases of COVID-19. However, for a strong recommendation of our risk stratification and to confirm its safety and effectiveness, further trials comprising larger patient cohorts are warranted. For this purpose, we will make use of the already established ReCovER, an ongoing open retrospective survey platform facilitating anonymous data entry that is available on <http://www.covid-em.org/>.



Clinical significance

Among all patients presenting with respiratory symptoms and suspicion of COVID-19, approximately 10% are tested positive for SARS-CoV-2.

More than 40% of confirmed COVID-19 cases require hospital admission, mainly for symptomatic treatment and management of respiratory symptoms, hypoxia and other complications.

Factors for hospital admission are decreased oxygen saturation, dyspnoea and history of CV disease.

The primary disposition was confirmed safe by a very low event rate among those discharged home from the ED (0% ICU admission as well as 0% mortality).

Future studies may develop additional criteria to increase the number of patients who can safely be treated as outpatients.

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Which parameters support disposition decision in suspected COVID-19 cases in the Emergency Department (ED): a clinical cohort study

Martin Möckel, Miriam Songa Stegemann, Volker Burst, Philipp Kümpers, Joachim Risse, Felix Carlo Koehler, Domagoj Schunk, Jennifer Hitzek, Tamara Elene Dietrich and Anna Slagman

Supplement Table 1	SARS-CoV-2	SARS-CoV-2
	positive patients	negative patients
	(n=110)	(n=1,070)
Women % (n)	41.8 (46)	45.9% (491)
	(nmiss=0)	(nmiss=0)
Age (median. IQR)	45 (33-66)	43 (31-61)
	(nmiss=0)	(nmiss=1)
BMI (Median. IQR)	27.1 (24.0-30.1)	24.3 (21.5-27.9)
	(nmiss=82)	(nmiss=955)
	Vital signs (median; IQR)	
BP syst. mmHg	134 (120-150)	133 (121-148)
	(nmiss=10)	(nmiss=191)
BP diast. mmHg	79 (70-89)	80 (70-90)
	(nmiss=10)	(nmiss=191)
Heart rate / min	89 (80-100)	90 (80-103)
	(nmiss=14)	(nmiss=185)
Temperature °C	37.6 (36.7-38.3)	37.0 (36.5-37.8)
	(nmiss=6)	(nmiss=114)
Respiratory rate	16 (15-20)	16 (15-19)
/min	(nmiss=20)	(nmiss=350)
Oxygen saturation %	97 (95-100)	99 (97-100)
	(nmiss=8)	(nmiss=164)
	Symptoms % (n)	
Fever	68.2 (75)	36.2 (387)
	(nmiss=4)	(nmiss=102)
Cough	56.4 (62)	50.0 (535)
	(nmiss=0)	(nmiss=166)
Haemoptysis	0	1.2 (13)
	(nmiss=60)	(nmiss=728)
Sore throat	13.6 (15)	23.3 (249)
	(nmiss=50)	(nmiss=532)
Rhinitis	4.5 (5)	10.8 (116)
	(nmiss=59)	(nmiss=625)

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Headache/ muscle pain	37.3 (41) (nmiss=36)	24.2 (259) (nmiss=511)
Dyspnea	36.4 (40) (nmiss=13)	25.9 (277) (nmiss=340)
GI-Symptoms	9.1 (10) (nmiss=43)	6.4 (69) (nmiss=652)
Diarrhea	17.3 (19) (nmiss=31)	7.3 (78) (nmiss=482)
Nausea/emesis	12.7 (14) (nmiss=38)	11.0 (118) (nmiss=527)
Loss of smell	2.7 (3) (nmiss=61)	1.0 (11) (nmiss=711)
Abdominal pain	6.4 (7) (nmiss=49)	5.9 (63) (nmiss=605)
Symptom onset time [days]	6 (3-9) (nmiss=26)	3 (1-6) (n=454)
Risk factors % (n)		
Transplantation	2.7 (3) (nmiss=18)	1.3 (14) (nmiss=213)
Tumor	2.7 (3) (nmiss=21)	11.1 (119) (nmiss=211)
Cardiovascular disease	30.9 (34) (nmiss=19)	26.4 (283) (nmiss=172)
Respiratory disease	18.2 (20) (nmiss=16)	17.3 (185) (nmiss=185)
Renal disease	7.3 (8) (nmiss=18)	7.7 (82) (nmiss=205)
Hepatic disease	7.3 (8) (nmiss=18)	3.8 (4) (nmiss=212)
Pregnancy	1.8 (2) (nmiss=10)	1.0 (1) (nmiss=981)