The impact of the COVID-19 pandemic on recovery from cardiac surgery: 1-year outcomes

Julie Sanders 1,2*, Teo Bueser 3,4, Emma Beaumont 5, Matthew Dodd 5, Sarah E Murray 6, Gareth Owens 7, Alan Berry 8, Edward Hyde 8, Tim Clayton 9, and Aung Ye Oo 10

1Director of Clinical Research, St Bartholomew’s Hospital, Barts Health NHS Trust, London EC1A 7DN, UK; 2Honorary Clinical Professor of Cardiovascular Nursing, William Harvey Research Institute, Queen Mary University of London, London EC1M 6BQ, UK; 3Associate Director of Clinical Research, St Bartholomew’s Hospital, Barts Health NHS Trust, London EC1A 7BE, UK; 4NIHR 70@70 Senior Nurse/Midwife Research Leader programme, National Institute for Health Research, London, UK; 5Research Fellow, Department of Medical Statistics, Faculty of Epidemiology and Public Health, London School of Hygiene and Tropical Medicine, London WC1E 7HT, UK; 6Lay representative, Society of Cardiothoracic Surgery of Great Britain and Ireland, London WC2A 3PE, UK; 7Chair, Aortic Dissection Awareness UK, Ireland, UK; 8Patient representative, The CardiacCovid study, St Bartholomew’s Hospital, Barts Health NHS Trust, London EC1A 7BE, UK; 9Professor in Applied Medical Statistics. Department of Medical Statistics, Faculty of Epidemiology and Public Health, London School of Hygiene and Tropical Medicine, London WC1E 7HT, UK; 10Consultant Cardiothoracic Surgeon, St Bartholomew’s Hospital, Barts Health NHS Trust, London EC1A 7BE, UK

Aims
The outbreak of COVID-19 was potentially stressful for everyone and possibly heightened in those having surgery. We sought to explore the impact of the pandemic on recovery from cardiac surgery.

Methods and results
A prospective observational study of 196 patients who were ≥18years old undergoing cardiac surgery between March 23 and July 4, 2020 (UK lockdown) was conducted. Those too unwell or unable to give consent/complete the questionnaires were excluded. Participants completed (on paper or electronically) the impact of event [Impact of Events Scale-revised (IES-R)] (distress related to COVID-19), depression [Centre for Epidemiological Studies Depression Scale (CES-D)], and EQ-5D-5L [(quality of life, health-related quality of life (HRQoL)] questionnaires at baseline, 1 week after hospital discharge, and 6 weeks, 6 months and 1 year post-surgery. Questionnaire completion was >75.0% at all timepoints, except at 1 week (67.3%). Most participants were male [147 (75.0%)], white British [156 (79.6%)] with an average age 63.4years. No patients had COVID-19. IES-R and CES-D were above average at baseline (indicating higher levels of anxiety and depression) decreasing over time. HRQoL pre-surgery was high, reducing at 1 week but increasing to almost pre-operative levels at 6 weeks and exceeding pre-operative levels at 6 months and 1 year. IES-R and CES-D scores were consistently higher in women and younger patients with women also having poorer HRQoL up to 1-year after surgery.

Conclusions
High levels of distress were observed in patients undergoing cardiac surgery during the COVID-19 pandemic with women and younger participants particularly affected. Psychological support pre- and post-operatively in further crises or traumatic times should be considered to aid recovery.

Registration
Clinicaltrials.gov ID:NCT04366167.
Introduction

The World Health Organization declared the outbreak of SARS-CoV-2 (COVID-19), a public health emergency of international concern on January 30, 2020. By the end of March 2020, there had been almost 34,000 deaths worldwide, with patients experiencing underlying cardiovascular disease having a particularly poor prognosis. The impact on those undergoing surgery for cardiovascular disease was unknown, although reports from the H1N1 pandemic indicated that an ‘unexpected and dramatically extraordinary hospital course’ could be expected. Early surgical studies from COVID-19, albeit small samples \((n<35)\), concluded that surgery in those with COVID-19 had a high risk of death and that surgery might accelerate disease progression in those incubating COVID-19.

At the beginning of the COVID-19 pandemic there was much uncertainty, and very little was known about COVID-19. The UK government ordered the stopping all essential contact and travel on March 16, 2020 and entered a full ‘stay at home’ lockdown on March 23, 2020. While the need to suspend UK elective surgery was recognized, urgent and emergency cases continued to require proper management while protecting resources for the response to COVID-19. Inevitably, cardiothoracic surgical services were severely affected. There was a >50% reduction in dedicated cardiac theatres and ICU beds, and in some countries services were restructured to form regional cardiac surgery hubs. The UK experienced a 52% reduction in dedicated cardiac theatres and ICU beds, and in some countries services were restructured to form regional cardiac surgery hubs.

The impact of the COVID-19 pandemic on recovery from cardiac surgery

Cardiac surgery patients \((n=196)\)

- High baseline distress & depression levels decreased over time
- HRQoL returned to pre-op levels at 6 wks and exceeded pre-op at 6 mths and 1 year
are at increased risk of depression and anxiety and those with depressive or anxiety symptoms tend to have poorer post-operative outcomes and health-related quality of life (HRQoL). Therefore, it was anticipated that the COVID-19 outbreak would be additional stressful for people, especially in those with underlying chronic diseases who are at higher risk for COVID-19 or in those having urgent or emergency cardiac surgery. Equally, symptoms of a traumatic event occur after 1 month but can also be delayed by several months, with older adults particularly at risk. Thus, we sought to explore and describe the impact of the COVID-19 pandemic on recovery up to 1 year from heart surgery.

Methods

Study design and setting

The CardiacCovid study is a prospective single-centre (UK) observational cohort study conducted between March 2020 (UK lockdown started for the COVID-19 pandemic) and August 2021 in one of the largest cardiovascular centres in Europe. Participants gave written informed consent to participate and were given questionnaires relating to the impact of the event (the pandemic), depression and HRQoL to complete for baseline (pre-surgery), 1 week after hospital discharge and at 6 weeks, 6 months, and 12 months after surgery.

The study complies with the Declaration of Helsinki and received National Health Service Health Research Authority Yorkshire and The Humber Sheffield Research Ethics Committee approval (reference 20/YH/0132, 15.04.2020). The study is registered with clinicaltrials.gov (NCT04366167) and is reported in accordance with the Strengthening the Reporting of Observational Studies in Epidemiology guidelines.

Study population: inclusion and exclusion criteria

Adult patients (≥18 years of age) undergoing cardiac surgery between March 23, 2020 and July 4, 2020 (UK COVID-19 national lockdown, when elective surgery was mainly suspended) were eligible for inclusion. Potential participants were identified from in-patient and theatre lists by a member of the study team and were approached prior to surgery where possible, or prior to hospital discharge. As research ethics approval was received on April 15, 2020, patients who had surgery prior to this date, and who were not in-patients at study commencement, were also contacted retrospectively. If recruited to the study after surgery, retrospective completion of the baseline questionnaires reflecting on pre-surgery state was undertaken. Those unable or unwilling to give written informed consent and/or to complete the questionnaires were excluded.

Data collection and measurement

Clinical data

Demographic information, pre-operative risk factors (including EuroSCORE (pre-operative risk assessment score)), medical history, pre-operative details and immediate post-operative outcome (for example, hospital length of stay, mortality) were obtained from the local National Institute for Cardiovascular Outcomes Research adult cardiac surgery database (https://www.nicor.org.uk/national-cardiac-audit-programme/datasets/). Other clinical data, for example COVID-19 status on admission (defined as a positive COVID-19 polymerase chain reaction test prior to surgery) and depression history, were obtained from local electronic sources and the participant, recorded on a standardized proforma and entered onto a bespoke database to link with the other clinical data and questionnaire data. Whether participants had COVID-19 during follow-up, COVID-19 vaccinations or participated in cardiac rehabilitation was obtained via the telephone call at 1 year.

Questionnaires

Participants completed the following questionnaires for baseline (pre-surgery status collected either contemporaneously or retrospectively), 1 week after hospital discharge and at 6 weeks, 6 months and 1 year after surgery:

(a) IES-R (Impact of Events Scale-revised) Impact of event scale: The Impact of Events Scale (IES) is a psychometrically robust questionnaire and one of the most widely used measures of event-specific distress and measures distress experienced by serious life changes/events. The IES-R (revised version) is a 22-item self-report scale where each item is reported on a five point Likert scale from 0 (not at all) to 4 (extremely) with respect to how distressing each item has been during the past week. Scale scores are formed for the three subscales, which reflect intrusion (for example, intrusive thoughts/feelings, nightmares; 8 items) avoidance (for example numbing of responses, avoidance of feelings; 8 items), and hyperarousal (for example anger, irritability, difficulty concentrating; 6 items). The total mean IES-R score is and a score of ≥33 has been suggested as the best cut-off for a probable diagnosis of post-traumatic stress disorder (PTSD).

(b) EQ-SD-5L (health-related quality of life): The EQ-SD-5L is a standardized, simple generic measure of health-related quality of life, which exhibits excellent psychometric properties across a broad range of populations, conditions and setting and is well received by patients. It has been the National Institute for Health and Care Excellence preferred adult HRQoL measure since 2008 and is the most widely used general HRQoL measure in UK and Europe. It consists of five dimensions (mobility, self-care, usual activities, pain/discomfort, anxiety/depression) each with five levels of health response reflecting escalating problems in each domain. These dimensions can be converted to a single index value, which reflects how good or bad a health state is according to the preferences of the general population of a country/region with a higher value indicating better HRQoL. Additionally, the EQ-SD-5L contains a visual analogue scale (VAS) ranging from 0 to 100 (0 being the worst possible health imaginable and 100 being the best possible health imaginable).

(c) CES-D (Centre for Epidemiological Studies Depression Scale). The CES-D is a well-known, freely available, widely used tool for depressive symptoms in a broad range of populations. It is psychometrically robust in cardiovascular patients and has previously been used in cardiac surgery patients. The CES-D is a 20-item self-report adult instrument designed to measure common symptoms (behavioural, cognitive and affective) of depression that have occurred over the past week, such as poor appetite, hopelessness, pessimism, and fatigue. All questions are answered on a scale of 0–3, with 0 indicating no symptom presence and 3 representing symptoms ‘most or all of the time’. CES-D scores range from 0 to 60 with higher scores indicating more severe depressive symptoms. A score of 16 or higher identifies subjects with potentially clinically meaningful depression.

Collectively, the questionnaires take approximately 20 min to complete. Both the CES-D and IES-R were available to use without permission or cost. Trust-level approval was already in place for electronic EQ-SD-5L use, and separate approval was sought and obtained from the EuroQol Research Foundation for EQ-SD-5L paper use, noting its deviation from protocol use (some patients completed the baseline measure retrospectively). Questionnaires were completed using the Amplitude™ patient reported outcomes platform, or on paper (with return stamped self-addressed envelope provided), depending on participant preference. If paper copies were received, responses were entered onto the Amplitude™ system by a member of the research team. Efforts to minimize loss to follow-up included using up to three automated (email) reminders and at least one telephone call from a member of the research team at each time-point. Furthermore, a member of the research team was available to assist the participant in completing the questionnaires, if needed.

Patient and public involvement

This study has had active patient and public involvement (PPI) through concept, delivery, analysis, interpretation, and dissemination. A specific PPI panel of 11 patient members was established, with two members representing patients on the interdisciplinary study steering group.
Statistical methods
Baseline characteristics and scores were summarized using means and standard deviations (SD) or medians and interquartile ranges (IQR) for continuous variables and counts and percentages for categorical variables. The characteristics were compared to those of patients undergoing surgery at the same centre in the corresponding time-period in 2019 (pre-pandemic). The baseline scores for IES-R, EQ-5D-5L, and CES-D were compared descriptively for those where questionnaires were completed retrospectively following surgery vs. those where questionnaires were completed prospectively prior to surgery.

Given the unknown impact, spread and duration of the COVID-19 pandemic at the time the study was designed, it was not possible to predict the number of patients likely to be enrolled in the study, nor the proportion who would be diagnosed with COVID-19. Therefore, a formal sample size calculation was not performed.

Separate mixed effects models for repeated measures (MMRM) were used to assess changes in IES-R, EQ-5D-5L, and CES-D scores over time by including the baseline measurements as an additional outcome and an indicator variable for follow-up visit only. This model accounts for the baseline values of each measure, while also incorporating all follow-up measurements, and enables the inclusion of participants with missing measurements. Unstructured variance-covariance matrices were used to allow for the anticipated correlations between repeated measurements over time. In order to assess the adjusted effects of age (years), sex (male or female), ethnicity (Asian, black or white), diabetes (yes or no), and surgery urgency (elective or emergency/urgent) on each of the outcome scores, MMRMs including an indicator variable for follow-up visit, each of the aforementioned covariates (with age modelled as a linear effect), and interactions between each covariate and follow-up visit were used. Finally, the unadjusted effect of the EuroSCORE-II on each of the outcome scores was assessed using MMRMs containing an indicator variable for follow-up visit, a linear effect for the EuroSCORE-II, and their interaction. The impact of the EuroSCORE-II was assessed separately from other pre-specified covariates of interest because calculation of this risk score involves age, sex, diabetes, and surgery urgency. For each of the models described effects at each follow-up visit were estimated and corresponding 95% confidence intervals and P-values calculated. The MMRMs used assume that any missing outcome data is missing at random. All analyses were conducted using Stata IC version 16.0 by authors EB, MD, and TC.

Results
Participants
During the study period 325 patients had surgery of whom 298 (91.7%) were screened for participation and 203 (68.1%) provided written informed consent (Figure 1). A further seven participants were excluded resulting in analysis being conducted on n = 196 (Figure 1). Retrospective completion of baseline questionnaires occurred 73.0% and overall questionnaire completion was >75.0% at all timepoints, except 1 week after surgery (67.3%) (Figure 1). The cohort characteristics are detailed in Table 1. Overall, the majority of patients were male (75.0%), of white background (79.6%) undergoing urgent or emergency surgery (59.7%) (as elective surgery was effectively postponed) with a median EuroSCORE of 1.6. No patients had COVID-19 at time of surgery. The characteristics of the cohort were similar to a comparative sample during non-COVID times (2019) at this centre (Table 1), although it is noted that 21 patients who died after surgery but before enrolment into this study were excluded who were likely to be of higher surgical risk (Figure 1). Particular differences in the study population included a smaller number of patients undergoing surgery (325 vs. 581), a smaller proportion undergoing elective surgery (40.3 vs. 57.8%) and having a longer length of hospital stay (9.0 days vs. 7.0 days), but lower mortality (0.5 vs. 2.8%). During the year of follow-up there were 7 (3.6%) deaths, 23 (11.7%) tested positive for COVID-19, 161 (82.1%) received COVID-19 vaccinations and 109 (55.6%) completed cardiac rehabilitation.

The impact of event, health-related quality of life and depression scores
Table 2, Figure 2 and Supplementary material online, Table S1 show the results from each questionnaire at each time-point. No differences were observed in baseline scores comparing those who completed the questionnaires retrospectively or prospectively (data not shown).

The impact of the pandemic
The impact of the pandemic was high at baseline and decreased over the time following surgery. The observed mean IES-R score was higher than expected at baseline (17.4) remaining high at 1 week (15.6), 6 weeks (15.4) and 6 months (15.4) and decreasing to 10.8 at 1-year post surgery. There is a clear trend over 1 year in a decreasing impact with a reduction of 5.65 points (95% CI 2.92 to 8.39) from the MMRM model and closed to the expected levels (Table 2, Figure 2 and Supplementary material online, Table S2).

Overall, 34/196 (17.3%) scored >33 (suggestive of PTSD) at baseline reducing to 11/149 (7.5%) at 1-year. All subscale scores were higher at baseline and decreasing over time, although hyperarousal responses were lower at all time-points than intrusion or avoidance responses, where similar levels were observed.

Health-related quality of life
HRQoL was high pre-surgery (mean 0.73), reducing at 1 week (mean 0.58) and then increasing to almost pre-operative levels at 6 weeks (mean 0.72). The mean score at 6 months and 1-year exceeded pre-surgery levels (mean 0.78 and 0.80, respectively) with the increase from baseline at 1-year estimated to be 0.06 (95% CI 0.01–0.10). Over a third of participants still had issues in all categories except self-care at 1-year. In respect of the VAS scores (out of 100) there was a progressive improvement in the overall HRQoL after surgery at each time-point.

Depression
Mean CES-D score was 12.3 pre-surgery, which increased at 1 week [an estimated increase from baseline of 2.53 (95% CI 0.91–4.15)] and then decreasing over time such that at 1-year post-surgery there was a change decrease from baseline of 1.96 (95% CI 0.05–1.96). At baseline, 28.9% scored >16 (indicative of depression) with 22.2% continuing to score highly at 1-year after surgery.

The association of age, sex, ethnicity, diabetes, surgical risk and surgery urgency on outcome
The impact of the pandemic
Older participants were affected less by the pandemic at all timepoints (Table 3) although female sex was associated with higher IES-R scores, compared to men, with the point estimate consistently higher at all timepoints (Figure 3A). Similarly, greater pre-operative risk (higher EuroSCORE) was associated with lower IES-R score, but at baseline only (Table 3). No evidence of associations with ethnicity, surgery urgency or diabetes status was observed, although the numbers in some categories were small (see Supplementary material online, Table S2).

Health-related quality of life
Although no association with age (Table 3), diabetes, surgery urgency or ethnicity was observed (see Supplementary material online, Table S2), female sex was associated with lower HRQoL at 1 week, 6 weeks and 1-year after surgery (with the direction of point estimate also
indicating lower HRQoL at 6 months) (Figure 3B). Similarly, greater pre-operative risk (higher EuroSCORE) was associated with lower EQ-5D-5L score at 6 weeks (Table 3).

**Depression**
Female gender has higher observed CES-D scores, indicating higher levels of depression at all time-points with the largest difference seen at 1 week [4.29 (95% CI 0.40–8.18] (Figure 3C). Similarly, older age was associated with lower CES-D scores at baseline, 6 weeks and 6 months after surgery with an observed difference also seen at 1 week. No association was observed with ethnicity, surgery urgency, diabetes, or EuroSCORE (Table 3 and Supplementary material online, Table S2).

**Discussion**
We undertook a prospective observational cohort study at one of the largest cardiovascular centres in Europe, to explore and describe the impact of the COVID-19 pandemic on recovery up to 1 year from heart surgery. Overall, we observed high levels of pandemic-related distress, and found that anxiety and depression were higher at baseline and at 1-week after surgery. These levels then declined over time, with stabilized levels at 6 weeks and 6 months and the lowest levels observed at 1-year. Similarly, HRQoL was high pre-surgery and had returned to pre-operative levels by 6 weeks after surgery and continued improve exceeding pre-operative levels at 6 months and 1-year, which is the expected trajectory in non-COVID times. Approximately 17.3% had
Table 1  Characteristics of study participants undergoing cardiac surgery during the COVID-19 pandemic (n = 196) and a comparative pre-COVID population from 2019

| Variable | Pre-COVID data [March 23 to July 4, 2019] n(%) or mean (SD) (unless otherwise stated) (n=581) | Pre-COVID data [March 23 to July 4, 2019] n(%) or mean (SD) (unless otherwise stated) (n=581) |
|----------|------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------|
| Demographics |                                                                                                   |                                                                                                   |
| Age [years/median (IQR)] | 65.0 [57.0–72.0] | 65.0 [56.0–73.0] |
| Sex (Female) | 49 (25.0) | 140 (24.1) |
| Ethnicity |                                                                                                   |                                                                                                   |
| White | 156 (79.6) | 196 (33.7) |
| Asian | 29 (14.8) | 93 (16.0) |
| Black | 11 (5.6) | 26 (4.5) |
| Other | 0 (0.0) | 29 (5.0) |
| Not stated/not known | 0 (0.0) | 237 (40.8) |
| Medical history |                                                                                                   |                                                                                                   |
| Previous MI (yes) | 72 (36.7) | 186 (32.6) |
| Previous cardiac surgery (yes) | 7 (3.6) | 25 (4.3) |
| Renal function/dialysis | 1 (0.5) | 7 (1.2) |
| History of pulmonary disease | 17 (8.7) | 44 (7.6) |
| History of neurological disease | 12 (6.1) | 38 (6.5) |
| History of neurological dysfunction | 5 (2.6) | 14 (2.4) |
| History of depression | 27 (13.8) | Not routinely recorded |
| Symptoms |                                                                                                   |                                                                                                   |
| NYHA class |                                                                                                   |                                                                                                   |
| I/II | 131 (66.8) | 354 (60.9) |
| III/IV | 65 (33.2) | 227 (39.1) |
| CCSC classification |                                                                                                   |                                                                                                   |
| 0/I/II | 119 (60.7) | 382 (65.7) |
| III/IV | 77 (39.3) | 199 (34.3) |
| Cardiac risk factors |                                                                                                   |                                                                                                   |
| Current smoker | 22 (11.2) | 71 (12.5) |
| Hypertension | 146 (74.5) | 439 (77.8) |
| Diabetes | 50 (25.5) | 170 (29.3) |
| BMI (kg/m²/median [IQR]) | 27.6 [24.7–30.5] | 27.4 [24.6–31.1] |
| Examination and investigation |                                                                                                   |                                                                                                   |
| LVEF: Good | 138 (70.4) | 438 (75.4) |
| Number of diseased vessels |                                                                                                   |                                                                                                   |
| 0 | 59 (30.1) | 229 (39.4) |
| 1 | 20 (10.2) | 25 (4.3) |
| 2 | 27 (13.8) | 72 (12.4) |
| 3 | 58 (29.6) | 171 (29.4) |
| 4+ | 32 (16.3) | 84 (14.5) |
| Pre-operative risk assessment |                                                                                                   |                                                                                                   |
| EuroSCORE [median (IQR)] | 1.6 [1.1–3.0] | 1.7 [1.1–3.3] |
| Intra-operative details |                                                                                                   |                                                                                                   |
| Operative priority |                                                                                                   |                                                                                                   |
| Elective | 79 (40.3) | 336 (57.8) |
| Urgent | 104 (53.1) | 213 (36.7) |
| Emergency | 13 (6.6) | 31 (5.3) |
| Salvage | 0 (0.0) | 1 (0.2) |
scores suggestive of PTSD at baseline. This is between the wide range (7.6–53.8%) reported in general population studies during the same time-period.31,32,33 Similarly, post-surgery IES-R scores suggestive of PTSD at all other time-points were within reported ranges (4–24%) in a variety of cardiovascular populations (not including cardiac surgery) in non-COVID times.34 As stated previously, we found scores indicative of clinical depression mirrored the trajectory of pandemic-related distress. However, those scoring ≥16 at all post-operative time-points (excluding 1-week after surgery) were also within ranges observed in pre-pandemic times (pre-CABG depression: 19 to 37%; post-CABG depression: 15–33%),35 although evidence is limited.

The stabilization of pandemic-related distress and depression scores at 6 weeks and 6 months, and the increase in anxiety/depression as measured on the EQ-5D-5L between these time-points could potentially reflect the unfolding COVID-19 situation in the UK. Lockdown 2 occurred between 2nd November 2020 and 2nd December 2020 while lockdown 3 commenced on 6th January 2021 with decreasing restrictions introduced from 8th March 2021. For many, the 6 week and 6 month time-points would have fallen within lockdown 1 and lockdown 2. Furthermore, the first COVID-19 vaccination was administered in the UK on 8th December 2020 and by the end of the 1-year follow-up for the study over 68 million vaccinations had been given.36 This may have had a positive influence on some 6 month responses and the 1-year responses, considering that 81.2% of our participants had their vaccination by the time they completed their 1-year questionnaires. Despite this, just under half of participants were denied the opportunity to access to cardiac rehabilitation which may be why high levels of pain and discomfort (83.8%) and impairment of undertaking usual activities (73.1%) were still observed at 6 weeks post-surgery.

We observed that age, pre-surgical risk and sex had an impact on outcome. Older participants, who are particularly at risk of traumatic events,20 and those with higher pre-operative surgical risk, actually reported less pandemic-associated distress compared with younger participants and those with lower risk, respectively. Similarly, increased age was associated with less depression, particularly at 6 weeks and 1-year. A potential reason for these findings, voiced by some participants and our PPI group, was that in these circumstances participants felt that the need for surgery outweighed the risks of the pandemic and they had ‘less to lose’. Considering sex, women were disproportionately affected by the pandemic as they tend to work in economically vulnerable positions, undertake more unpaid care work and are at increased risk of abuse during isolation periods37—all which are likely to have had an impact on their psychological and physical health.38 Although our results reflect pre-pandemic findings in that women, compared to men, are more likely to experience depression at time of cardiac surgery,39 suffer greater post-operative morbidity burden40 and have worse quality of life,41 the women in our study also reported higher levels of pandemic-related distress up to 1-year after surgery. Since anxiety is associated with poorer recovery after cardiac surgery42 this also could have contributed to the poorer HRQoL we observed.

Another key finding to note were the results 1 week after discharge. Although pandemic-related distress was reduced from baseline, a decrease in HRQoL particularly related to pain and increased depression were observed. Poorer outcomes at 1 week were also reflected in the reduced response rate (see limitations) as many indicated they felt too unwell to respond to the questionnaires. Traditionally, although not evidence-based, patients receive a follow-up appointment approximately 6 weeks after surgery. Certainly, it has recently been found that patients suffer highest morbidity 1 week post-surgery and that 44% of patients would like an earlier review, including perhaps a telephone call which could help reassure, alleviate anxiety or assist in detecting early signs of complications.43 In addition to this, some patients will have experienced prolonged periods of isolation, both in hospital and on discharge home, due to the pandemic lockdown rules, which can impact on psychological and physical health.44 Therefore,
patients are likely to benefit from earlier review routines, but particularly during challenging times.

It is also interesting to note that despite conducting this study at the beginning of the pandemic and during the first UK lockdown, no patients in our study had COVID-19 at the time of enrolment. This offers the advantage that COVID-19 does not confound the results at time of enrolment, although 23 (11.7%) subsequently reported having COVID-19 during the 1-year follow-up study period.

Overall, this study addresses the significant gaps in the current evidence in this area of cardiovascular care. It is a methodologically robust study using a prospective design and is one of the largest cardiac surgery studies exploring recovery during the COVID-19 pandemic. This study also greatly benefited from the input of a PPI group to ensure the study was feasible and inclusive to potential participants. This was particularly important due to the possibly stressful circumstances patients were being approached under, and in retaining them over the course of a challenging year which included additional COVID-19 waves and national lockdowns. Despite these strengths, this study has several limitations. Firstly, due to the organizational restrictions imposed in this early phase of the pandemic, the majority of patients (73%) were recruited after surgery and completed the baseline questionnaires retrospectively. While prospective completion is preferred, as prospective application has been shown to lead to recall bias and lower HRQoL scores in trauma45 and intensive care46 patients, the difference is not thought to be clinically relevant.46 Despite this potential bias retrospective evaluation of health status is still considered more appropriate than applying population norms.46 Equally, on analysis we did not observe any differences in baseline scores comparing those who completed the questionnaires prospectively or retrospectively. Thus, the impact of this pragmatic approach is likely to have little impact on the results. Secondly, although questionnaire response rates were good at all timepoints, a lower response rate was achieved at 1 week after discharge, despite using a hybrid method of delivery which is considered best practice.48 Mainly, this related to the retrospective nature of recruitment for those who had surgery prior to ethics approval being obtained and missing this time-point. However, as stated previously, we also observed poorer outcomes at this time-point which likely contributed to the willingness and ability of participants to complete them. Thirdly, this is an observational study and is limited by factors associated with this design, mainly selection bias and confounders, but also the type of data collected. However, considerable effort was made to address these limitations. Selection bias was minimized by attempting to recruit all eligible consecutive patients and by our hybrid approach to questionnaire delivery to limit barriers and encourage questionnaire completion to reduce loss to follow-up. We also adjusted for known confounders but we did not have detailed data on all health and psychosocial circumstances in the follow-up period which may have influenced questionnaire responses. We are also limited by only having data

|                     | Baseline n = 196 | 1 week n = 132 | 6 weeks n = 159 | 6 months n = 159 | 1 year n = 149 |
|---------------------|------------------|---------------|-----------------|-----------------|--------------|
| **IES-R**           |                  |               |                 |                 |              |
| IES-R, mean (SD)    | 17.4 (17.2)      | 15.6 (15.6)   | 13.3 (15.4)     | 12.6 (15.4)     | 10.8 (12.6)  |
| IES-R, median (IQR) | 11.5 (4.0–26.0)  | 10.5 (4.0–21.5)| 7.0 (2.0–20.0)  | 7.0 (2.0–18.0)  | 5.0 (1.0–17.0)|
| IES-R change from baseline, mean (95% CI) | — | 0.43 (−2.98, 2.13) | −2.56 (−5.29, 0.16) | −4.34 (−7.08, −1.61) | −5.65 (−8.39, −2.92) |
| IES-R score > 33 (PTSD), n (%) | 34 (17.3) | 17 (12.9) | 19 (12.0) | 14 (8.8) | 11 (7.5) |
| Subscale: void, mean (SD) | 6.2 (6.8) | 5.2 (5.9) | 4.6 (6.0) | 4.5 (5.8) | 4.1 (5.4) |
| Subscale: hyperarousal, mean (SD) | 4.5 (5.1) | 4.2 (4.4) | 3.7 (4.5) | 3.3 (4.5) | 2.6 (3.5) |
| Subscale: intrusion, mean (SD) | 6.7 (6.8) | 6.0 (6.2) | 5.0 (6.0) | 4.8 (6.0) | 4.1 (4.9) |

| **EQ-SD-5L**       |                  |               |                 |                 |              |
| EQ-SD-5L index, mean (SD): score range 0–100 | 0.73 (0.24) | 0.58 (0.24) | 0.72 (0.18) | 0.78 (0.22) | 0.80 (0.20) |
| EQ-SD-5L index, median (IQR) | 0.75 (0.63–0.91) | 0.63 (0.44–0.75) | 0.76 (0.65–0.81) | 0.83 (0.68, 1.0) | 0.84 (0.72, 1.0) |
| EQ-SD-5L change from baseline, mean (95% CI) | — | −0.16 (−0.21, −0.12) | −0.03 (−0.07, 0.01) | 0.04 (−0.002, 0.08) | 0.06 (0.01, 0.10) |
| Dimension: mobility, n (%)a | 75 (38.3%) | 71 (53.8%) | 55 (34.4%) | 58 (36.3%) | 55 (36.9%) |
| Dimension: self-care, n (%)a | 30 (15.3%) | 71 (53.8%) | 33 (20.6%) | 32 (20.0%) | 19 (12.8%) |
| Dimension: usual activities, n (%)a | 97 (49.5%) | 117 (88.6%) | 117 (73.1%) | 61 (38.1%) | 59 (39.6%) |
| Dimension: pain/discomfort, n (%)a | 119 (60.7%) | 121 (91.7%) | 134 (83.8%) | 93 (58.1%) | 68 (45.6%) |
| Dimension: anxiety/depression, n (%)a | 73 (37.2%) | 59 (44.7%) | 59 (36.9%) | 68 (42.5%) | 52 (34.9%) |
| EQ VAS, mean (SD): score range 0–100 | 61.8 (23.8) | 62.0 (18.5) | 72.6 (16.7) | 76.6 (17.0) | 78.7 (15.3) |

| **CES-D**          |                  |               |                 |                 |              |
| CES-D, mean (SD): score range 0–60 | 12.3 (10.9) | 13.9 (9.9) | 10.8 (10.1) | 10.4 (10.5) | 9.6 (9.7) |
| CES-D, median (IQR) | 10.0 (4.0–17.5) | 12.0 (6.0–20.0) | 8.0 (3.0–16.0) | 7.0 (3.0–15.0) | 7.0 (2.0–13.0) |
| CES-D Change from baseline, mean (95% CI) | — | 2.53 (0.91, 4.15) | −0.38 (−1.95, 1.19) | −1.26 (−3.09, 0.58) | −1.96 (−3.86, −0.05) |
| CES-D score ≥16 (depressed), n (%) | 57 (28.9%) | 46 (34.9%) | 42 (26.3%) | 36 (22.6%) | 33 (22.2%) |

Mean (SD) were calculated using the observed data. The change from baseline results are estimated from the mixed effects models for repeated measures. Mean IES-R, EQ-SD-5L, and CES-D scores estimated using mixed effects models for repeated measures are presented in Supplementary material online, Table Sf. CES-D, Centre for Epidemiological Studies Depression Scale; EQ-SD-5L, EuroQol Health Related Quality of Life instrument; EQ VAS, EuroQol Visual Analogue Scale; IES-R, The Impact of Events Scale (Revised); PTSD, post-traumatic stress disorder; SD, standard deviation. Number (and proportion) experiencing any limitation.
**Figure 2** Mean (95% CI) for (A) The Impact of Events Scale (Revised), (B) EuroQoL Health Related Quality of Life instrument and (C) Centre for Epidemiological Studies Depression Scale questionnaires at baseline, 1 week, 6 weeks, 6 months and 1 year after surgery for participants undergoing cardiac surgery during the COVID-19 pandemic.
Table 3  Association between outcome measures and age, sex and EuroSCORE in participants undergoing cardiac surgery during the COVID-19 pandemic at baseline, 1 week, 6 weeks, 6 months and 1 year after surgery

|                  | Baseline |          |          |          |          |          |          |          |          |
|------------------|----------|----------|----------|----------|----------|----------|----------|----------|----------|
|                  |          | mean difference (95% CI) | P-value | mean difference (95% CI) | P-value | mean difference (95% CI) | P-value | mean difference (95% CI) | P-value |
|                  |          | P         | 1 week   | 6 weeks  | 6 months | 1 year   | 6 months | 1 year   | 1 year   |
| IES-R             |          |          |          |          |          |          |          |          |          |
| Sex              |          |          |          |          |          |          |          |          |          |
| Male             | 0        | 0.06     | 0        | 0.56     | 0        | 0.05     | 0        | 0.11     | 0        |
| Female           | 5.35     | (−0.27, 10.96) | 1.80     | (−4.22, 7.82) | 5.95     | (0.12, 11.78) | 4.43     | (−1.04, 9.91) | 4.09     | (−0.90, 9.07) |
| Age (years)     |          |          |          |          |          |          |          |          |          |
| Male             | −0.23    | (−0.46, 0.005) | 0.06     | (−0.24, −0.49, 0.003) | 0.05     | (−0.23, −0.46, −0.007) | 0.04     | (−0.35, −0.56, −0.14) | 0.001    | (−0.20, −0.39, −0.01) |
| EuroSCORE        | −0.51    | (−0.96, −0.06) | 0.03     | (−0.18, −0.77, 0.41) | 0.56     | (0.26, −0.28, 0.80) | 0.35     | (0.008, −0.40, 0.42) | 0.97     | (0.06, −0.29, 0.41) |
| EQ-5D-SL         |          |          |          |          |          |          |          |          |          |
| Sex              |          |          |          |          |          |          |          |          |          |
| Male             | 0        | 0.09     | 0        | 0.003    | 0        | 0.06     | 0        | 0.39     | 0        |
| Female           | −0.07    | (−0.15, 0.01) | −0.14    | (−0.23, −0.05) | −0.07    | (−0.13, 0.004) | −0.04    | (−0.12, 0.05) | −0.11    | (−0.19, 0.03) |
| Age (years)     | 0.0008   | (−0.003, 0.004) | 0.6      | (−0.001, 0.005, 0.003) | 0.54     | (0.001, −0.002, 0.004) | 0.46     | (0.001, −0.002, 0.004) | 0.53     | (0.0002, −0.003, 0.003) |
| EuroSCORE        | 0.006    | (−0.0007, 0.01) | 0.08     | (−0.006, −0.02, 0.004) | 0.23     | (−0.01, −0.02, −0.004) | 0.002    | (−0.001, −0.007, 0.005) | 0.64     | (−0.0005, −0.006, 0.005) |
| CES-D            |          |          |          |          |          |          |          |          |          |
| Sex              |          |          |          |          |          |          |          |          |          |
| Male             | 3.12     | (−0.37, 6.62) | 4.29     | (0.40, 8.18) | 3.30     | (−0.48, 7.07) | 3.04     | (−0.84, 6.93) | 2.92     | (−0.96, 6.81) |
| Female           | −0.20    | (−0.34, −0.05) | 0.01     | (−0.12, −0.28, 0.04) | 0.14     | (−0.21, −0.36, −0.07) | 0.004    | (−0.21, −0.36, −0.06) | 0.01     | (−0.13, −0.27, 0.02) |

n = 196.

* Estimates are adjusted for age, ethnicity, surgery urgency and diabetes.

** Estimates are adjusted for sex, ethnicity, surgery urgency and diabetes.

*** Estimates are unadjusted.
Figure 3 Questionnaire results (mean [95%CI] by sex in participants undergoing cardiac surgery during the COVID-19 pandemic at baseline, 1 week, 6 weeks, 6 months and 1 year after surgery: (A) The Impact of Events Scale (Revised), (B) EuroQoL Health Related Quality of Life instrument, (C) Centre for Epidemiological Studies Depression Scale.
relating to event-related distress, depression and HROoL. This was a pragmatic decision based on the complexities of the situation, considering participant burden at a difficult time, and also study resources at a time when staff were being redeployed to respond to the acute pandemic phase. However, a questionnaire-only study potentially underrepresents the actual impact and experiences of patients. Therefore, we are conducting a complementary qualitative study to further understand the lived experience and impact of having cardiac surgery during a global pandemic. Finally, we equally did not have the resources to include those who had their surgery cancelled or postponed during this period, due to the higher number of patients that were affected by this. At the end of the first wave it was predicted that if countries increased their normal surgical capacity by 20% after the pandemic it would take a median of 45 weeks to undertake these missed operations. The impact of the pandemic on these patients is likely to be considerable.

Globally, the COVID-19 pandemic is far from over. In March 2022 a large proportion of the world had been infected with the omicron variant and new variants are expected, where some may be more severe than omicron. Differing country-level policies for managing the pandemic as well as vaccine inequality, particularly in low-and middle-income countries poses a significant barrier to a global end to the pandemic. Therefore, our study is likely to be informative internationally for future variants and pandemic waves not only for patients undergoing cardiac surgery, but potentially other complex surgeries as well as interventional cardiology procedures. Of course, pandemics are not the only current global challenge—any scenario impacting on ‘normal’ healthcare delivery, for example conflict, major incidents, and climate emergencies, may impact on pre-intervention distress and subsequent recovery. Our key messages are that high levels of distress were observed relating to the COVID-19 pandemic in patients undergoing cardiac surgery with women and younger participants particularly affected. Equally, pre- and post-surgery psychological support, earlier (remote) follow-up approximately 1-week after hospital discharge and adapting the delivery of cardiac rehabilitation should be considered to aid recovery.

Supplementary material

Supplementary material is available at European Journal of Cardiovascular Nursing online.

Acknowledgements

We gratefully acknowledge all the patients who kindly participated in this study. We also gratefully acknowledge the valuable contributions of Mr Amir Sepehrpour and Mr Jorge Mascaro, who supported participant identification, screening and recruitment, Ms Samantha Cook who supported participant follow-up and The CardiacCovid Patient and Public Involvement (PPI) group who have been involved throughout the study (Mrs Sarah Murray, Mr Gareth Owens, Mr Paul Whitehouse, Mr Steve Stevenson, Mr Michael Connor, Mr Alan Berry, Mr Edward Hyde, Mr Prashant Navaratnarajah), Mr Adrian Bendall, Mr Gilbert Wheeler and Mr Phil Blakelow). Teofila Bueser is a National Institute for Health Research (NIHR) Senior Nurse and Midwife Research Leader. The views expressed in this article are those of the author and not necessarily those of the NIHR, or the Department of Health and Social Care. ©EuroQol Research Foundation. EQ-SDTM is a trade mark of the EuroQol Research Foundation. The EQ-SD was reproduced by permission of EuroQol Research Foundation. Reproduction of this version of the EQ-SD as detailed in this paper is not allowed. For reproduction, use or modification of the EQ-SD (any version), please register your study by using the online EQ registration page: www.euroqol.org.

Funding

None declared.

Conflict of interest: None declared.

Data availability

The data underlying this article cannot be shared publicly due to the ethical restrictions on data sharing for this study. The data will be shared on reasonable request to the corresponding author, if appropriate sponsor and/or ethics committee approval is given.

References

1. Zheng YY, Ma YT, Zhang JY, Xie X. COVID-19 and the cardiovascular system. Nat Rev Cardiol 2020;17:259–260.
2. Person B, Bahouth H, Brauner E, Ben-Ishay O, Bickel A, Kluger YS. Surgical emergencies confounded by H1N1 influenza infection—a plea for concern. World J Emerg Surg 2010; 5:2–5.
3. Galbraith JG, Butler JS, Pead M, Twomey A. H1N1 infection in emergency surgery: a cautionary tale. Int J Surg Case Rep 2010;1:4–6.
4. Lei S, Jiang F, Su W, Chen C, Chen J, Mei W, Liying Z, Yifan J, Liangzeng Z, Dayong L, Zhong-Yuan X, Zhing-Yuan X. Clinical characteristics and outcomes of patients undergoing surgeries during the incubation period of COVID-19 infection. EClinicalMedicine 2020;21.
5. Cai Y, Hao Z, Gao Y, Ping W, Wang Q. Coronavirus disease 2019 in the periparative period of lung resection: a brief report from a single thoracic surgery department in Wuhan, People’s Republic of China. J Thoar Oncole 2020;15:1063–1072.
6. Iacobucci G. COVID-19: all non-urgent elective surgery is suspended for at least three months in England. BMJ 2020;368:m1106.
7. NHS. Clinical guide for the management of cardiothoracic patients during the coronavirus pandemic. 2020.
8. Gauthino M, Chikwe J, Hameed I, Robinson NB, Fremen S, Ruel M. Response of cardiac surgery units to COVID-19: an internationally-based quantitative survey. Circulation 2020;142:300–302.
9. Bonalumi G, di Mauro M, Garata A, Barill F, Gerosa G, Parolari A. The COVID-19 outbreak and its impact on hospitals in Italy: the model of cardiac surgery. Eur J Cardiothor Surg 2020;57:1023–1028.
10. Hussain A, Balmforth D, Yates M, Lopez-Moreno A, Rathwell C, Lambourne J, Roberts N, Lall K, Edmundson S. The pan London emergency cardiac surgery service: coordinating a response to the COVID-19 pandemic. J Card Surg 2020;35:1561–1569.
11. Sanders J, Akowuah E, Cooper J, Kirmani BH, Kanan M, Acharya M, Jegathan R, Krasopoulou G, Ngage D, Deiglurk I, Yiu P, Kendall S, Oo AY. Cardiac surgery outcome during the COVID-19 pandemic: a retrospective review of the early experience in nine UK centres. J Cardiothor Surg 2021;6:43.
12. COVIDSurg Collaborators. Elective surgery cancellations due to the COVID-19 pandemic: global predictive modelling to inform surgical recovery plans. Br J Surg 2020.
13. Tully PJ. Psychological depression and cardiac surgery: a comprehensive review. J Extra Corp Technol 2012;44:224–232.
14. Cromhout PF, Moons P, Thygesen LC, Nashef S, Lambourne J, Roberts N, Lall K, Edmundson S. The pan London emergency cardiac surgery service: coordinating a response to the COVID-19 pandemic. J Card Surg 2020;35:1561–1569.
15. Sanders J, Akowuah E, Cooper J, Kirmani BH, Kanan M, Acharya M, Jegathan R, Krasopoulou G, Ngage D, Deiglurk I, Yiu P, Kendall S, Oo AY. Cardiac surgery outcome during the COVID-19 pandemic: a retrospective review of the early experience in nine UK centres. J Cardiothor Surg 2021;6:43.
16. COVIDSurg Collaborators. Elective surgery cancellations due to the COVID-19 pandemic: global predictive modelling to inform surgical recovery plans. Br J Surg 2020.
17. Tully PJ. Psychological depression and cardiac surgery: a comprehensive review. J Extra Corp Technol 2012;44:224–232.
23. Creaser M, Bell R, Failla S. Psychometric properties of the impact of event scale—revised. Behav Res Ther 2003;41:1489–1496.
24. EuroQol Research Foundation. EQ-SD-5L user guide [Internet]. 2019. Available from: https://euroqol.org/publications/user-guides

25. Feng YS, Kohlmann T, Janssen MF, Buchholz I. Psychometric properties of the EQ-SD-5L: a systematic review of the literature. Qual Life Res 2021;30:647–673.
26. Kim J, Henderson RA, Pocock SJ, Clayton T, Sculptor MJ, Foss KAA, RITA-3 Investigators. Health-related quality of life after interventional or conservative strategy in patients with unstable angina or non-ST-segment elevation myocardial infarction. J Am Coll Cardiol 2005;45:221–228.

27. Brooks R. Euroqol: the current state of play. Health Policy 1996;37:53–72.
28. Radloff LS. The CES-D scale: a self-report depression scale for research in the general population. Appl Psychol Meas 1977;1:385–401.
29. Moon JR, Huh J, Song J, Kang IS, Park SW, Chang SA, Yang JH, Jun TG. The center for epidemiologic studies depression scale is an adequate screening instrument for depression and anxiety disorder in adults with congenital heart disease. Health Qual Life Outcomes 2017;15:176.

30. Mckenzie L, Simpson J, Stewart M. A systematic review of pre-operative predictors of post-operative depression and anxiety in individuals who have undergone coronary artery bypass surgery. Psychol Heal Med 2010;15:74–93.
31. Zhang Y, Ma ZF. Impact of the COVID-19 pandemic on mental health and quality of life among local residents in Liaoning province, China: a cross-sectional study. Int J Environ Res Public Health 2020;17:2381.

32. Forte G, Favieri F, Tambelli R, Casagrande M. COVID-19 Pandemic in the Italian population: validation of a post-traumatic stress disorder questionnaire and prevalence of PTSD symptomatology. Int J Environ Res Public Health 2020;17:1451.
33. Wang C, Pan R, Wan X, Tan Y, Xu L, Ho CS, Ho RC. Immediate psychological responses and associated factors during the initial stage of the 2019 coronavirus disease (COVID-19) epidemic in the general population in China. Int J Environ Res Public Health 2020;17:1729.

34. Tulloch H, Greenman PS, Tasse V. Post-traumatic stress disorder among cardiac patients: prevalence, risk factors, and considerations for assessment and treatment. Behav Sci 2014;5:27–40.
35. Correa-Rodríguez M, Abu Elhusseh M, Suleiman-Martos N, Membrive-Jiménez MJ, Velando-Sartono A, Schmidt-RioValle J, Gómez-Urquiola JL. Prevalence of depression in coronary artery bypass surgery: a systematic review and meta-analysis. J Clin Med 2020;9:909.

36. NHS England. COVID-19 Vaccination Statistics: Week ending Sunday 18th July 2021 [Internet]. 2021. Available from: https://www.england.nhs.uk/statistics/wp-content/uploads/sites/2/2021/07/COVID-19-weekly-announced-vaccinations-22-July-2021.pdf
37. The Organisation for Economic Co-operation and Development (OECD). EQ-5D-5L: a systematic review of the literature. https://euroqol.org/publications/user-guides

38. Fisher AN, Ryan MK. Gender inequalities during COVID-19. Gr Process Integr Relations 2021;24:237–245.
39. Steenman M, Sartipy U. Depression screening in cardiac surgery patients. Heart Lung Circ 2019;28:953–958.

40. Matyal R, Qureshi NQ, Mufarrih SH, Sharkey A, Bose R, Chu LM, Liu DC, Senthilnathan V, Mahmood F, Khabaz KR. Update: gender differences in CABG outcomes–have we bridged the gap? PLoS One 2021;16:e0255170.

41. Peric V, Borzanovic M, Stolic R, Jovanovic A, Sovtic S, Djikic D, Marcetic Z, Dimkovic S. Quality of life in patients related to gender differences before and after coronary artery bypass surgery. Interact Cardiovasc Thorac Surg 2010;10:232–238.

42. Tully PJ, Pedersen SS, Winefield HR, Baker RA, Turnbull DA, Denollet J. Cardiac morbidity risk and depression and anxiety: a disorder, symptom and trait analysis among cardiac surgery patients. Psychol Health Med 2011;16:333–345.

43. Ngase DL, Gooseman MR, Bullment KL, Jarvis MA, Chaudhry MA, Cale AR, Cowen ME. Is six weeks too long for the first outpatient review after cardiac surgery? FORCAST6. Br J Cardiol 2019;26:34.

44. Pietrabissa G, Simpson SG. Psychological consequences of social isolation during COVID-19 outbreak. Front Psychol 2020;11:2201.

45. Spiroki I, Geraerts AJM, Bosel GJ, de Jongh MAC, Polinder S, Haagmans JA. Correspondence of directly reported and recalled health-related quality of life in a large homogeneous sample of trauma patients. Qual Life Res 2019;28:3005–3013.

46. Dinglas VD, Gifford JM, Husain N, Colauttouzi E, Needham DM. Quality of life before intensive care using EQ-5D: patient versus proxy responses. Crit Care Med 2013;41:9–14.

47. Wilson R, Derrett S, Hansen P, Langley J. Retrospective evaluation versus population norms for the measurement of baseline health status. Health Qual Life Outcomes 2012;10:68.

48. Neve OM, van Benthem PPG, Stiggelbout AM, Hensen EF. Response rate of patient reported outcomes: the delivery method matters. BMC Med Res Methodol 2021;21:220.

49. Kluge HH. Statement—Two years on, we could be entering a new phase in the pandemic with plausible hope for stabilization, yet too early to drop our guard [Internet]. World Health Organization; 2022 [cited 25 Mar 2022]. Available from: https://www.euro.who.int/en/media-centre/sections/statements/2022/statement-two-years-on,-we-could-be-entering-a-new-phase-in-the-pandemic-with-plausible-hope-for-stabilization,-yet-too-early-to-drop-our-guard

50. Murray CJL. COVID-19 will continue but the end of the pandemic is near. Lancet 2022;399:417–419.

51. The PLOS Medicine Editors. Vaccine equity: a fundamental imperative in the fight against COVID-19. PLOS Med 2022;19:e1003948.