Indirect Impact of the COVID-19 Pandemic on Activity and Outcomes of Transcatheter and Surgical Treatment of Aortic Stenosis in England

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BACKGROUND: Aortic stenosis requires timely treatment with either surgical aortic valve replacement (SAVR) or transcatheter aortic valve replacement (TAVR). This study aimed to investigate the indirect impact of coronavirus disease 2019 (COVID-19) on national SAVR and TAVR activity and outcomes.

METHODS: The UK TAVR Registry and the National Adult Cardiac Surgery Audit were used to identify all TAVR and SAVR procedures in England, between January 2017 and November 2020. The number of isolated aortic valve replacement (AVR), AVR+coronary artery bypass graft surgery, AVR+other surgery, and TAVR procedures per month was calculated. Separate negative binomial regression models were fit to monthly procedural counts, with functions of time as covariates, to estimate the expected change in activity during COVID-19.

RESULTS: We included 15,142 TAVR cases, 13,357 isolated AVR cases, 8,550 AVR+coronary artery bypass graft cases, and 6,773 AVR+other cases. Before March 2020 (UK lockdown), monthly TAVR activity was rising, with a slight decrease in the SAVR activity during 2019. We observed a rapid and significant drop in TAVR and SAVR activity during the COVID-19 pandemic, especially for elective cases. Cumulatively, over the period March to November 2020, we estimated an expected 4,989 (95% CI, 4,020–5,959) cases of aortic stenosis who have not received treatment.

CONCLUSIONS: This study has demonstrated a significant decrease in TAVR and SAVR activity in England following the COVID-19 outbreak. This situation should be monitored closely, to ensure that monthly activity rapidly returns to expected levels. There is potential for significant backlog in the near-to-medium term and potential for increased mortality in this population.

GRAPHIC ABSTRACT: A graphic abstract is available for this article.

Key Words: aortic valve • aortic valve stenosis • disease outbreaks • registries • survival

The coronavirus disease 2019 (COVID-19) presents a global health crisis and has resulted in significant excess mortality. Many countries have imposed restrictions based on social distancing and movement (ie, lockdowns), with the aim of mitigating and managing the spread of COVID-19. A UK-wide lockdown was initiated on March 23, 2020, with a second national lockdown imposed in England at beginning of November 2020. The lockdown restrictions, and the pandemic, have resulted in widespread changes in operational activity of health services. Many health care systems have faced significant pressure on services, particularly within critical
WHAT IS KNOWN

- The coronavirus disease 2019 (COVID-19) pandemic has resulted in widespread changes in operational activity of health services.
- The impact, from a national perspective, of COVID-19 on activity and outcomes of surgical and transcatheter aortic valve replacement is unknown.

WHAT THE STUDY ADDS

- We show that there has been a significant decrease in transcatheter aortic valve replacement and surgical aortic valve replacement activity during COVID-19.
- Cumulatively, over the period March to November 2020, we estimated an expected 4989 (95% CI, 4020–5959) cases of severe aortic stenosis who have not received treatment.
- There is potential for significant backlog in the near-to-medium term and potential for increased mortality in this population.

Nonstandard Abbreviations and Acronyms

| Abbreviation | Description |
|--------------|-------------|
| AVR | aortic valve replacement |
| CABG | coronary artery bypass graft |
| COVID-19 | coronavirus disease 2019 |
| LOS | length of stay |
| NICOR | National Institute for Cardiovascular Outcomes Research |
| SAVR | surgical aortic valve replacement |
| TAVR | transcatheter aortic valve replacement |

The aim of this study was to investigate the activity and postprocedural outcomes of all AVRs in contemporary practice, from a national perspective, and to investigate the indirect impact of COVID-19 on activity and outcomes. The intention is to estimate the effect of reduced activity on projected backlog of cases.

METHODS

Because of the sensitive nature of the data collected for this study, requests to access the dataset from qualified researchers trained in human subject confidentiality protocols may be sent to the National Institute for Cardiovascular Outcomes Research (NICOR). The analytical code used for this study is available upon reasonable request, for the purposes of reproducibility.

UK TAVR Registry

The UK TAVR registry collects data for every TAVR procedure undertaken within the UK. Data collection occurs prospectively at each contributing center and is submitted to the NICOR. Data collection is mandated for all centers licensed to undertake TAVR procedures. We extracted data from NICOR on all TAVR procedures undertaken in England between January 2017 and November 2020.

UK National Adult Cardiac Surgery Audit

The NICOR National Adult Cardiac Surgery Audit contains data on all major heart operations undertaken in the United Kingdom. Each center is responsible for prospective data collection and submission of this to NICOR. We extracted all SAVRs undertaken in England between January 2017 and November 2020. We defined SAVR to be any procedure recorded as being a valve replacement, where the aortic valve implant type was recorded as being mechanical, biological, homograft, or autograft replacement. We further categorized SAVRs into (1) isolated AVR, (2) AVR with coronary artery bypass graft (CABG), or (3) AVR with other surgery (AVR+other). Here, other surgery was any mitral valve procedure, tricuspid valve procedure, pulmonary valve procedure, major aortic surgery, or other cardiothoracic procedures.

Data Flows During COVID-19

The British Cardiovascular Intervention Society and the Society for Cardiothoracic Surgery have made significant efforts to maintain data flows during COVID-19. NICOR has provided weekly uploads of data throughout the pandemic. Thus, at the time of analysis, we had updated data until the end of November.
2020. Nonetheless, to reflect the possibility that some individual centers might have stopped submitting data to NICOR during the pandemic or have delays in submitting data, we define a set of rapid-data-submitting centers to be those that submitted at least 1 case (of either TAVR or SAVR) in November 2020 (latest month of our analysis). We perform sensitivity analysis (of the analyses described below) on this subset of centers. Similarly, we also considered sensitivity analyses focusing on the subset of centers that submitted at least 1 case (of either TAVR or SAVR) across every month in 2020.

Outcomes

The primary outcome was presentation and treatment of aortic stenosis with AVR. As secondary outcomes, we considered 30-day mortality and postprocedural length of stay (LOS). Mortality information was provided by linking the UK TAVR registry and the National Adult Cardiac Surgery Audit with the office for national statistics civil registration of deaths dataset. Linkage was made based on each patient’s NHS number. We defined postprocedural LOS to be the number of days between the TAVR/SAVR procedure and hospital discharge.

Statistical Analysis

We excluded any cases in which the age of the patient at the time of the procedure was under 18 years. Additionally, we excluded cases where the NHS number was missing or with missing procedure urgency. Finally, we removed any duplicate cases in either datasets, identified using NHS number, age, sex, admission date, and date/time of the procedure.

In all analyses, we stratified by procedure type (ie, isolated AVR, AVR+CABG, AVR+other, or TAVR). We made no formal comparisons between these procedural types, since there are several confounding factors surrounding the decision-making between TAVR/SAVR (many of which are not captured in the dataset).

Cases performed between February 1 and November 30, 2020, were defined into a during–COVID-19 group, with any case performed in these same months across the preceding years being defined into a pre–COVID-19 group. The February 1, 2020, was chosen since the first COVID-19 case reported in England was on January 28, 2020.

We report patient baseline characteristics for each procedure type, as whole cohorts and across the during COVID-19 and pre–COVID-19 groups. Continuous variables were reported using the mean with SDs. Categorical variables were presented as frequencies of occurrence with relative percentages. Comparisons between continuous variables were made using t tests/ANOVA, while comparisons between categorical variables were made using the $\chi^2$ test. Predicted procedural risk was quantified using the UK TAVR clinical prediction model for all TAVR procedures and the Logistic EuroSCORE clinical prediction model for all SAVR procedures. For the purposes of calculating the risk predictions, missing values in any predictor variables were set to risk factor absent, representing a plausible missingness process in the registries.

The number of procedures per month was calculated across the full study period, separately for each procedure type. Percentage increase or decrease in monthly activity was calculated for each during–COVID-19 month, against the respective pre–COVID-19 months. We fitted negative binomial models to the number of TAVR/SAVR procedures performed per month between January 2017 and December 2019, using time as covariate, which was modeled continuously to capture trends in activity and as a factor variable of month to capture seasonality. This model was used to estimate the expected number of TAVR/SAVR procedures per month in 2020, to compare with the observed activity level.

For each of the 4 procedural types, we compared mortality up to 30 days, across the during–COVID-19 and pre–COVID-19 groups by fitting a Cox proportional hazards model, with the COVID-19 group as a covariate, adjusting for the linear predictor of either the UK TAVR prediction model or the Logistic EuroSCORE, as appropriate. Differences in postprocedural LOS between the during–COVID-19 and pre–COVID-19 groups were also investigated by fitting a Cox proportional hazards model. The proportional hazards assumption was checked by examining the Schoenfeld residuals.

All analyses were performed using R, version 4.0.0, along with the tidyverse suite of packages, and the survival package.

Ethics Approval

In the efforts to understand the impact of the COVID-19 pandemic on cardiology services, extraordinary government permission was obtained to evaluate anonymized records from these databases through an agreement with NHS Digital. This work was endorsed by (1) Scientific Advisory Group for Emergencies (a body responsible for ensuring timely and coordinated scientific advice is made available to decision makers to support UK cross-government decisions in the Cabinet Office Briefing Room), (2) NHS England—a public body of the Department of Health and Social Care, and (3) NHS Improvement, responsible for overseeing NHS trusts. NICOR, which houses the British Cardiovascular Intervention Society registry, has support under section 251 of the NHS Act 2006 to use patient information for approved medical research without informed consent. For this rapid NHS evaluation, health data analysis was enabled under Section 254 of the Health and Social Care Act 2012.

RESULTS

The UK TAVR Registry included 15,741 procedures across the study period, of which we included 15,142. The National Adult Cardiac Surgery Audit registry included 108,881 cases during the study period, of which we included 28,680 SAVR procedures, comprised of 13,357 (46.6%) isolated AVR, 8550 (29.8%) AVR+CABG, and 6773 (23.6%) AVR+other cases.

Table 1 shows the baseline characteristics of each procedural group. The mean age of isolated AVR, AVR+CABG, AVR+other, and TAVR was 67.7, 72.2, 62.9, and 81.3, respectively. Across all surgical groups, the majority of cases were male. The mean Logistic EuroSCORE was 7.50%, 10.7%, and 14.1% for isolated AVR, AVR+CABG, and AVR+other, respectively, while the mean UK TAVR prediction model was 3.11% (Table 1).
TAVR and SAVR Activity

There has been an increase in the number of TAVR procedures performed per month between January 2017 and December 2019, with the majority of procedures being elective (Figure 1). While the number of monthly AVR+CABG and AVR+other procedures remained relatively stable pre-2020, there was a slight decrease in the number of elective isolated AVR cases per month in 2019. The average number of elective isolated AVR cases per month was 250 in 2017 and 252 in 2018, while the monthly activity in 2019 decreased from 226 cases in January to 173 cases by December (Figure 1). After March 1, 2020, there was a significant drop in activity across all AVR procedures compared with historic levels (Figure 2). There was a slight recovery in AVR activity in May to August 2020.

Importantly, similar findings were found in the subgroup of rapid-data-submitting centers (Figure I in the Data Supplement). In particular, in this subset of centers, the observed rapid drop in cases during March and April 2020 persisted. Interestingly, activity in these rapid-data-submitting centers has actually returned (at

**Table 1. Baseline Characteristics of the Surgical Aortic Valve Replacement and TAVR Cases Included in This Analysis**

|                  | Isolated AVR | AVR+CABG | AVR+other | TAVR |
|------------------|--------------|----------|-----------|------|
| n                | 13357        | 8550     | 6773      | 15142|
| Age, y; mean (SD)| 67.66 (11.64)| 72.23 (8.26) | 62.89 (14.06) | 81.25 (7.24) |
| Women (%)        | 5073 (38.0)  | 1905 (22.3) | 2269 (33.5) | 6727 (44.5) |
| CCS angina status, % |                |          |           |      |
| No angina        | 7648 (57.4)  | 2548 (29.9) | 4497 (66.6) | 10624 (75.2) |
| Class I          | 1628 (12.2)  | 881 (10.3)  | 730 (10.8) | 1045 (7.4) |
| Class II         | 2318 (21.2)  | 2978 (35.0) | 973 (14.4) | 1732 (12.3) |
| Class III or IV  | 1222 (9.2)   | 2109 (24.8) | 557 (8.2) | 726 (5.1) |
| NYHA, %          |              |          |           |      |
| Class I          | 1389 (10.5)  | 885 (10.5)  | 1085 (16.3) | 1032 (7.4) |
| Class II         | 5226 (39.6)  | 3490 (41.4) | 2272 (34.1) | 3246 (23.1) |
| Class III or IV  | 6598 (49.9)  | 4052 (48.1) | 3298 (49.6) | 9756 (69.5) |
| Previous MI, %   | 785 (5.9)    | 2182 (25.6) | 332 (4.9) | 2099 (14.6) |
| Previous PCI, %  | 674 (5.1)    | 969 (11.5)  | 231 (3.5) | 2389 (16.8) |
| Previous cardiac surgery, % | 929 (7.5) | 211 (2.7)  | 949 (15.2) | 2404 (17.5) |
| Diabetic, %      | 2554 (19.2)  | 2477 (29.1) | 799 (11.8) | 3464 (24.3) |
| Current/ex-smoker, % | 6995 (52.8) | 5233 (61.6) | 3352 (49.9) | 6466 (49.8) |
| Creatinine, umol/L; mean (SD) | 89.58 (45.91) | 95.90 (54.14) | 96.07 (58.07) | 104.97 (62.54) |
| History of neurological disease, % | 1142 (8.6) | 872 (10.3)  | 866 (10.2) | 2229 (15.5) |
| Extracardiac arteriopathy, % | 675 (5.1) | 1132 (13.3) | 484 (7.2) | 1867 (13.2) |
| Atrial fibrillation or flutter, % | 1175 (8.9) | 1147 (13.5) | 1602 (23.9) | 3850 (28.6) |
| ≥1 vessel with >50% diameter stenosis, % | 620 (5.3) | 7464 (96.8) | 265 (5.0) | 4082 (31.8) |
| PA systolic, mean (SD) | 27.09 (19.81) | 20.09 (22.61) | 38.59 (22.15) | 37.98 (15.13) |
| LV function, %   |              |          |           |      |
| Good (LVEF, >50%) | 10554 (79.7) | 6018 (70.9) | 4750 (70.7) | 10116 (72.5) |
| Moderate (LVEF, 31%–50%) | 2080 (15.7) | 1968 (23.2) | 1568 (23.4) | 2559 (18.3) |
| Poor (LVEF, <30%) | 614 (4.6) | 504 (5.9)  | 397 (5.9) | 1282 (9.2) |
| Height, m; mean (SD) | 1.68 (0.10) | 1.70 (0.10) | 1.71 (0.11) | 1.65 (0.10) |
| Weight, kg; mean (SD) | 82.85 (18.38) | 83.65 (16.80) | 82.45 (19.08) | 76.00 (17.32) |
| Nonelective, %   | 3038 (22.7)  | 2735 (32.0) | 2307 (34.1) | 2977 (19.7) |
| Logistic EuroSCORE, mean (SD) | 750 (5.84) | 10.68 (11.08) | 14.05 (14.47) | NA |
| UK TAVR CPM, mean (SD) | NA | NA | NA | 3.11 (2.39) |

The numbers in some categories might not sum to the total due to missing data. AVR indicates aortic valve replacement; CABG, coronary artery bypass graft; CCS, Canadian Cardiovascular Society; CPM, clinical prediction model; LV, left ventricle; LVEF, left ventricular ejection fraction; MI, myocardial infarction; NA, not available; NYHA, New York Heart Association; PA, pulmonary artery; PCI, percutaneous coronary intervention; and TAVR, transcatheter aortic valve replacement.
least approximately) to expected levels in September to November 2020 (Figure I in the Data Supplement). This indicates that levels of AVR activity have started to recover following the initial rapid drop caused by the first national lockdown.

The number of elective SAVR procedures was below 150 cases per month between March to June 2020 for each of isolated AVR, AVR+CABG, and AVR+other (Figure 1). In contrast, there remained >100 elective TAVR cases per month after March 2020. The percentage change in monthly activity between 2020 and historic levels was lower for TAVR than SAVR with a maximum percentage difference of 86.5%, 80.7%, 83.4%, and 55.7%, for isolated AVR, AVR+CABG, AVR+other, and TAVR, respectively (Figure 2B).

Figure 3 shows the expected (from the negative binomial model) and actual monthly AVR activity during 2020. For the first few months after lockdown, the estimated
difference (95% CI) in the number of TAVR cases per month compared with those expected based on historic trends was −2 (−40 to 35) in March 2020, −229 (−264 to −193) in April 2020, −191 (−229 to −154) in May 2020, and −129 (−166 to −92) in June 2020 (Figure 3B). The estimated decrease in isolated AVR activity was −171 (−201 to −140), −231 (−257 to −205), −177 (−205 to −148) and −96 (−124 to −69), across March to June 2020, respectively. Similar observations were made for AVR+CABG and AVR+other cases (Figure 3B). Cumulatively, over the period March to November 2020, this amounts to an estimated expected drop of 4989 (95% CI, 4020–5959) cases of severe aortic stenosis in England, of which 1683 (95% CI, 1429–1937) were for isolated AVR, 1038 (95% CI, 848–1229) were for AVR+CABG, 703 (95% CI, 519–887) were for AVR+other, and 1565 (95% CI, 1223–1906) were for TAVR.

Evolution of Patient Demographics and Procedural Risk

Table 2 shows patient baseline characteristics of isolated AVR cases across the pre–COVID-19 and during–COVID-19 groups. For isolated AVR, the mean age was significantly lower in the during–COVID-19 group than the pre–COVID-19 group (P<0.001), and there was a significantly higher Canadian Cardiovascular Society (CCS) angina status (P=0.002), NYHA class (P<0.001) and mean pulmonary artery (PA) systolic pressure (P<0.001). For TAVR cases, the mean age, proportion of current/ex-smokers, and mean creatinine were significantly lower in the during–COVID-19 group compared with the pre–COVID-19 group (Table 3). There was a lower proportion of TAVR cases with previous myocardial infarction (P<0.001), previous cardiac surgery (P<0.001), and extracardiac

Figure 2. Number of transcatheter aortic valve replacement (TAVR) and surgical aortic valve replacement (AVR) procedures per month compared to historic levels. A, Temporal plot of the number of TAVR and surgical aortic valve replacement procedures per month during 2020, compared with monthly averages (minimum and maximum shown by shaded region) across all other years in the dataset. B, Percentage change between the mean monthly activity in 2017 to 2019 and the monthly activity in 2020; negative percentage change denotes increase in 2020 over historic levels. AVR indicates aortic valve replacement; and CABG, coronary artery bypass graft.
arteriopathy ($P=0.001$) in the during–COVID-19 group. Similar findings were found for AVR+CABG (Table I in the Data Supplement) and AVR+other (Table II in the Data Supplement). We also explored differences in baseline characteristics between pre–COVID-19 and during–COVID-19 groups, restricting to just 2019 and 2020 data (to look at changes in most contemporary practice); the findings were quantitatively similar (Tables III through VI in the Data Supplement).

Overall surgical AVR procedural risk, as estimated by the Logistic EuroSCORE, has remained relatively stable over time (Figure II in the Data Supplement). While the mean UK TAVR prediction model was significantly lower in the during–COVID-19 group compared with the pre–COVID-19 group ($P<0.001$; Table 3), this was largely driven by 2017 cases (Figure II in the Data Supplement). Indeed, upon comparing cases in February to November 2019 with corresponding months in 2020, we found that there was no significant difference in the UK TAVR clinical prediction model between February and November 2020, compared with corresponding months in 2019 (Table VI in the Data Supplement).

Between 2017 and December 2019, there has been a steady increase in the monthly TAVR activity in the lowest quantiles of risk strata, while the monthly activity in the highest quantiles of risk strata has remained relatively stable (Figure III in the Data Supplement). In contrast, the monthly activity of surgical AVR has been gradually decreasing through time across all quantiles of risk strata (Figures IV through VI in the Data Supplement).

**Outcomes**

The overall Kaplan-Meier estimates of 30-day survival were 98.5% for isolated AVR, 95.8% for AVR+CABG,
94.8%, for AVR+other, and 97.5% for TAVR. For isolated AVR, AVR+other, and TAVR, we found no significant difference in mortality hazards up to 30 days post-procedure between the pre–COVID-19 group and the during–COVID-19 group (Table 4). In contrast, mortality hazards up to 30 days post-procedure were significantly higher in patients undergoing AVR+CABG during–COVID-19 compared with the pre–COVID-19 group (hazard ratio, 1.41 [95% CI, 1.05–1.89]).

The median LOS following TAVR was 3 days (interquartile range, 2–5 days) in the pre–COVID-19 group and 2 days (interquartile range, 1–3 days) in the during–COVID-19 group. The median (interquartile range) LOS in the pre–COVID-19 group for isolated AVR,

### Table 2. Baseline Characteristics of the Isolated Aortic Valve Replacement Cases Included in the Pre–COVID-19 and During–COVID-19 Groups, as Defined in the Methods Section

| n | Pre–COVID-19 | During–COVID-19 | P value |
|---|--------------|----------------|--------|
| n | 9803 | 1549 | <0.001 |
| Age, y; mean (SD) | 67.95 (11.74) | 66.05 (11.22) | <0.001 |
| Women, % | 3791 (38.7) | 539 (34.8) | 0.004 |
| CCS angina status, % | 0.002 |
| No angina | 5616 (57.5) | 895 (57.9) | |
| Class I | 1198 (12.3) | 175 (11.3) | |
| Class II | 2095 (21.4) | 299 (19.3) | |
| Class III or IV | 860 (8.8) | 178 (11.5) | |
| NYHA, % | <0.001 |
| Class I | 1016 (10.5) | 138 (9.0) | |
| Class II | 3908 (40.3) | 512 (33.5) | |
| Class III or IV | 4782 (49.3) | 878 (57.5) | |
| Previous MI, % | 572 (5.9) | 77 (5.0) | 0.190 |
| Previous PCI, % | 479 (4.9) | 80 (5.2) | 0.721 |
| Previous cardiac surgery, % | 676 (7.4) | 116 (8.1) | 0.401 |
| Diabetic, % | 1835 (18.8) | 334 (21.6) | 0.010 |
| Current/ex-smoker, % | 5146 (52.9) | 800 (52.2) | 0.625 |
| Creatinine, umol/L; mean (SD) | 89.81 (46.49) | 91.12 (47.34) | 0.597 |
| History of neurological disease, % | 848 (8.7) | 123 (8.0) | 0.380 |
| Extracardiac arteriopathy, % | 510 (5.2) | 72 (4.7) | 0.391 |
| Atrial fibrillation or flutter, % | 899 (9.2) | 100 (6.6) | 0.001 |
| ≥1 vessel with >50% diameter stenosis, % | 448 (5.2) | 78 (5.9) | 0.296 |
| PA systolic, mean (SD) | 26.79 (18.67) | 32.75 (26.93) | <0.001 |
| LV function, % | 0.398 |
| Good (LVEF, >50%) | 7760 (79.8) | 1206 (78.3) | |
| Moderate (LVEF, 31%–50%) | 1524 (15.7) | 259 (16.8) | |
| Poor (LVEF, <30%) | 439 (4.5) | 75 (4.9) | |
| Height, m; mean (SD) | 1.68 (0.10) | 1.70 (0.11) | <0.001 |
| Weight, kg; mean (SD) | 82.80 (18.08) | 84.53 (19.60) | <0.001 |
| Nonelective, % | 2109 (21.5) | 462 (29.8) | <0.001 |
| Logistic EuroSCORE, mean (SD) | 7.55 (8.35) | 6.98 (8.73) | 0.013 |

This only included cases in February to November each year, and the numbers in some categories might not sum to the total due to missing data. CCS indicates Canadian Cardiovascular Society; COVID-19, coronavirus disease 2019; L V, left ventricle; L VEF, left ventricular ejection fraction; MI, myocardial infarction; NYHA, New York Heart Association; PA, pulmonary artery; and PCI, percutaneous coronary intervention.
AVR+CABG, and AVR+other was 7 (5–9) days, 8 (6–12) days and 9 (6–15) days, respectively, with these being 6 (5–9) days, 7 (6–11) days, and 8 (6–14) days in the during–COVID-19 group. For AVR+CABG procedures performed in the during–COVID-19 period, the adjusted hazard ratio (95% CI) for early discharge was 1.09 (1.02–1.17), showing significantly shorter LOS (Table 5). TAVR patients in the during–COVID-19 group were also significantly more likely for early discharge, up to 2 days post-TAVR (Table 5).

**DISCUSSION**

This study is the first to investigate activity and outcomes following all AVR procedures in contemporary practice, including the potential indirect impacts of the COVID-19 pandemic. We observed a rapid decrease in TAVR and SAVR activity during the COVID-19 pandemic. Over the period March to November 2020, this decline in activity accounts for an estimated 4989 patients with aortic valve replacement. For AVR+CABG procedures performed in the during–COVID-19 period, the adjusted hazard ratio (95% CI) for early discharge was 1.09 (1.02–1.17), showing significantly shorter LOS (Table 5). TAVR patients in the during–COVID-19 group were also significantly more likely for early discharge, up to 2 days post-TAVR (Table 5).

The treatment of severe aortic stenosis has evolved from SAVR being the default treatment modality to TAVR now being an evidence-based option at all surgical risk categories.14–16 We observed changes in TAVR and SAVR activity, with steadily increasing TAVR activity and corresponding slight decreases in elective isolated AVR cases, up to 2019. This supports previous findings in this area.19 Although TAVR is currently only approved for inoperable or high-risk cases in the United Kingdom, the evidence of equivalence in low-risk cases is accumulating.18,31 This may partially explain our finding of a steadily increasing proportion of TAVR cases within the lowest quantile of risk, before 2020. The observed decrease in SAVR activity before COVID-19 also provides evidence that the clinical envelope for TAVR has expanded into lower risk cases within real-world contemporary practice.

Inevitably, the initiation of national lockdown in the United Kingdom was associated with a significant reduction in the monthly number of AVR procedures being performed. We found a relatively smaller fall in TAVR than SAVR. One potential explanation is that TAVR has a low probability of requiring stay in an intensive treatment unit, which is important given constraints during the pandemic. Indeed, during the pandemic, patients with severe symptomatic aortic stenosis were recommended to be treated by TAVR where appropriate.32 However, given that the UK TAVR registry and the National Adult Cardiac Surgery Audit dataset do not contain information on the decision-making behind the SAVR or TAVR choice, we are not able to investigate this directly. Importantly, we observed evidence that monthly activity levels across SAVR and TAVR are returning to expected levels toward the end of the study period (September to November 2020), particularly in centers that rapidly submit data to NICOR. Nonetheless, there will inevitably remain a backlog of cases incurred by the first national lockdown in the United Kingdom.

The observed reduction in AVR activity was largely driven by elective cases. A possible explanation is that the UK government's response to the pandemic was to recommend cancelation of elective procedures.33 This was made to allow a restructuring of hospital services, thereby allowing more staff and resource to deal with the increased admissions due to COVID-19. Another hypothesis for this observed reduction in elective cases could be secondary to patients being less active during COVID-19 and hence less AS-related symptoms resulting in nondiagnosis of AS or lack of urgent need for intervention. Interestingly, we observed that monthly activity for TAVR and SAVR had a slight recovery in May and June 2020. We were unable to investigate the reasons for this, but previous studies have made similar observations.6 Again, one could speculate that this relates to decreasing demands on health care systems as the pandemic evolved, with an aim to resume elective activity once the peak of the pandemic had passed. Given that England has entered a second national
lockdown in November 2020, it will remain to be seen if there is another drop in elective AVR activity. Elective cases were being encouraged to continue through the second lockdown.

Nevertheless, despite restructure of health care services nationally during COVID-19, overall procedural risk has remained relatively constant through time. In many ways, some individual baseline characteristics of those undergoing SAVR during COVID-19 were lower risk than pre-COVID patients. There was inevitably an element of careful selection in patients eligible for SAVR during the initial lockdown, particularly for elective cases that were advised to be canceled. This might partially explain these findings, due to the complex (multivariable) interactions between procedural risk and individual baseline characteristics. Nonetheless, it is important to note that overall procedural risk for SAVR (quantified by the EuroScore) and TAVR (quantified by the UK TAVR clinical prediction model) was not significantly different between the pre–COVID-19 (2019 months) and during–COVID-19 groups.

While the observed temporal changes in activity are perhaps unsurprising, these findings raise important implications for health care resource planning in the near-to-medium term. Namely, the results suggest that there will likely be significant increased future demand for TAVR and SAVR. This will lead to an inevitable increase in waiting times and associated adverse impacts on outcomes. Recommendations for how to manage this challenge are emerging. It was not possible for us to forecast future demand for AVR since we do not have information on patients who are candidates for AVR but who have not currently undergone the procedure. However, based on the available data, we estimated that, cumulatively, between March and November 2020, there were 4989 (95% CI, 4020–5959) cases of severe aortic stenosis who have not received treatment in England. Previous studies have shown that, under normal circumstances, the median wait time for TAVR is 80 days. Thus, assuming these figures apply to AVR normal circumstances, the median wait time for TAVR is 3742 if procedures are made at 2495 patients will remain untreated by 80 days (3742 if procedures are made at 2495 patients will remain untreated by 80 days). Such figures can give further indications of the expected deaths while waiting for intervention.

Several limitations should be noted. First, we make no statistical comparisons between isolated AVR, AVR+CABG, AVR+other, or TAVR groups. Any such comparisons would be subject to confounding by indication. This means that we were not able to investigate changes in patient-level propensity to undergo SAVR versus TAVR through the COVID-19 period. Second, while we used the Logistic EuroSCORE to summarize overall SAVR procedural risk, this model is known to overpredict mortality risk. However, this model is commonly used for benchmarking in national cardiovascular registries, and we use the model in the same capacity here. Third, this analysis is limited to procedures in England; however, given that COVID-19 has caused changes in health care utilization globally, one might expect similar findings in other health care settings. Finally, some delays in data reporting during the pandemic might contribute to some of the results; however, significant efforts have been made to maintain data flows with weekly uploads of data. Additionally, we undertook a sensitivity analysis of rapid-data-submitting centers, which indicated quantitatively similar results to the main analysis, particularly regarding the drastic decrease in activity following the first UK lockdown. Further work should explore whether activity is returning in later months, as suggested by this sensitivity analysis.

In conclusion, this study has demonstrated a significant drop in TAVR and SAVR activity following the COVID-19 outbreak in the United Kingdom. The case mix of patients who have undergone AVR during the COVID-19 period was similar to the case mix seen in the pre–COVID-19 period. There was evidence that activity is starting to return to expected levels by the end of study follow-up. Nonetheless, there will be a backlog of cases caused by the initial lockdown period, suggesting that there will be a sharp rise in demand for AVR intervention in the near-to-medium term, with the potential for an upturn in mortality in patients waiting to be treated.
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