

Impact of COVID-19 in patients with heart failure with mildly reduced or preserved ejection fraction enrolled in the DELIVER trial

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Aim

COVID-19 may affect clinical risk in patients with heart failure. DELIVER began before and was conducted during the COVID-19 pandemic. This study aimed to evaluate the association between COVID-19 and clinical outcomes among DELIVER participants.

Methods and results

Participants with chronic heart failure with mildly reduced or preserved ejection fraction (HFmrEF/HFpEF) were randomized to dapagliflozin or placebo across 350 sites in 20 countries. COVID-19 was investigator-reported and the contribution of COVID-19 to death was centrally adjudicated. We assessed (i) the incidence of COVID-19, (ii) event rates before/during the pandemic, and (iii) risks of death after COVID-19 diagnosis compared to risks of death in participants without COVID-19. Further, we performed a sensitivity analysis assessing treatment effects of dapagliflozin vs. placebo censored at pandemic onset. Of 6263 participants, 589 (9.4%) developed COVID-19, of whom 307 (52%) required/prolonged hospitalization. A total of 155 deaths (15% of all deaths) were adjudicated as definitely/possibly COVID-19-related. COVID-19 cases and deaths did not differ by randomized assignment. Death rate in the 12 months following diagnosis was 56.1 (95% confidence interval [CI] 48.0–65.6) versus 6.4 (95% CI 6.0–6.8)/100 participant-years among trial participants with versus without COVID-19 (adjusted hazard ratio [aHR] 8.60, 95% CI 7.18–10.30). Risk was highest 0–3 months following diagnosis (153.5, 95% CI 130.3–180.8) and remained elevated at 3–6 months (12.6, 95% CI 6.6–24.3/100 participant-years). After excluding investigator-reported fatal COVID-19 events, all-cause death rates

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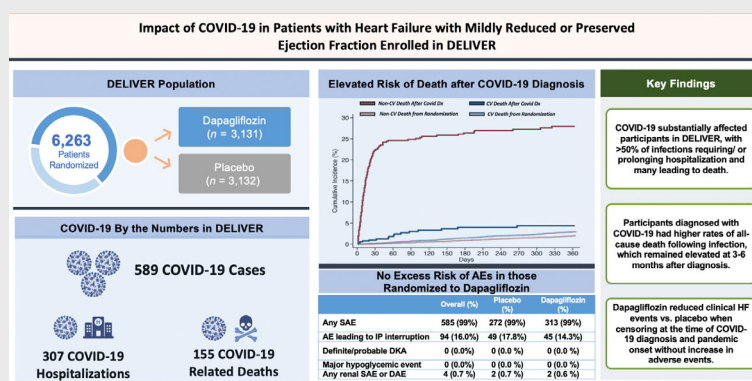
in the 12 months following diagnosis among COVID-19 survivors ($n=458$) remained higher (aHR 2.46, 95% CI 1.83–3.33) than rates for all trial participants from randomization, with censoring of participants who developed COVID-19 at the time of diagnosis. Dapagliflozin reduced cardiovascular death/worsening HF events when censoring participants at COVID-19 diagnosis (HR 0.81, 95% CI 0.72–0.91) and pandemic onset (HR 0.72, 95% CI 0.58–0.89). There were no diabetic ketoacidosis or major hypoglycaemic events within 30 days of COVID-19.

Conclusion

DELIVER is one of the most extensive experiences with COVID-19 of any cardiovascular trial, with >75% of follow-up time occurring during the pandemic. COVID-19 was common, with >50% of cases leading to hospitalization or death. Treatment benefits of dapagliflozin persisted when censoring at COVID-19 diagnosis and pandemic onset. Patients surviving COVID-19 had a high early residual risk.

Clinical Trial Registration: ClinicalTrials.gov Identifier NCT03619213.

Graphical Abstract



Impact of COVID-19 in patients with HF and mildly reduced and preserved ejection fraction. COVID-19, coronavirus disease-2019; HF, heart failure.

Keywords

Heart failure • COVID-19 • Death

Introduction

Patients with cardiovascular (CV) disease, including heart failure (HF), are at elevated risk of complications associated with COVID-19¹; however, few data exist describing the immediate and longer-term risks faced by individuals with HF who develop COVID-19. Large registries and trials may be well-suited to assess such risks; however, CV disease COVID-19 registries have generally limited enrolment to hospitalized COVID-19 patients, have had short follow-up periods, or have lacked central adjudication of clinical events.^{2,3} In addition, CV trials during this period had largely completed enrolment and follow-up prior to pandemic onset,^{4,5} enrolled North American participants only,⁶ and evaluated primary surrogate endpoints.⁶ In addition, the COVID-19 pandemic had a major effect on rates of death, CV outcomes and treatment effects in CV clinical trials.^{5,7} DELIVER (Dapagliflozin Evaluation to Improve the LIVEs of Patients with Preserved Ejection Fraction Heart Failure), the largest trial of HF with mildly

reduced or preserved ejection fraction (HFmrEF/HFpEF), enrolled a population across four continents, was operational before and during the COVID-19 pandemic⁸ and observed a treatment effect favouring dapagliflozin. We sought to examine the incidence and impact of COVID-19 in the DELIVER trial.

Methods

Trial design and changes during the COVID-19 pandemic

DELIVER was an international, prospective, multicentre, double-blind, event-driven clinical trial assessing the safety and efficacy of dapagliflozin 10 mg once daily versus placebo in patients with HFmrEF or HFpEF. The design and primary results of the DELIVER trial have been previously published.⁸ In brief, patients with symptomatic HF, elevated natriuretic peptides (NP), and left ventricular ejection fraction (LVEF) >40% were randomized in a 1:1 fashion to dapagliflozin 10 mg once daily or a matching placebo. Patients were also required to have

elevated N-terminal pro-B-type natriuretic peptide (NT-proBNP) and evidence of structural heart disease. The primary endpoint of DELIVER was a composite of time to first CV death or worsening HF event (including HF hospitalization or an urgent HF visit).

DELIVER randomized patients across 350 centers in 20 countries in North America, Latin America, Europe, and Asia. The first patient was randomized in September 2018 and the last patient completed the trial follow-up in March 2022. In response to the rise in severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infections and the declaration of COVID-19 as a pandemic by the World Health Organization on 11 March 2020, the sponsor's clinical project team, in conjunction with the trial executive committee, took several steps to minimize risk to participants and trial staff. Trial enrolment was temporarily paused in all active sites on 27 March 2020 (except Chinese sites which initiated randomization in March 2020) and re-started at various time points depending on region and site; home delivery of study drug was initiated after the resumption of enrolment. Local sites were instructed to collect and report COVID-19 information as adverse events. Investigators reported COVID-19 diagnoses, COVID-19-related events leading to or prolonging hospitalization, and fatal COVID-19 events as adverse events. Infection status was determined by local protocols and could have included probable and lab-confirmed diagnoses. All deaths regardless of assigned cause were centrally adjudicated by an independent endpoints committee (Brigham and Women's Hospital, Boston, MA, and University of Glasgow, Scotland, UK) as definitely, possibly, or not related to COVID-19. Definite relatedness required a clinical diagnosis with a positive polymerase chain reaction or typical imaging of COVID-19 and possible cases required clinical features and treatment of COVID-19 without definitive test. Guidance to sites from trial leadership included recommendations to consider interruption of the investigational product (IP) in patients with type 2 diabetes and known diabetic ketoacidosis (DKA) risk and to continue IP in non-diabetic patients and those without risk factors or prior DKA.

Statistical analyses

The statistical analysis plan for the DELIVER study was modified prior to unblinding to incorporate a COVID-19 sensitivity analysis, censoring participants at the time of COVID-19 diagnosis, and has been previously reported.⁸ The primary objectives of this pre-specified analysis by amendment to the academic statistical analysis plan (SAP) were (1) to describe the incidence of COVID-19 in a large, international randomized clinical trial, (2) to assess the impact of the pandemic on clinical event accrual and treatment effects in DELIVER, and (3) to assess the health risks faced by participants with HFmrEF/HFpEF after COVID-19 diagnosis.

We described the incidence of COVID-19 associated, hospitalization, and death during the trial based on investigator-reported adverse events. Baseline characteristics at the time of randomization were compared between participants who did versus did not develop COVID-19 and those with severe versus non-severe disease as reported by investigators. Severe disease was defined as requiring or prolonging hospitalization or resulting in an investigator-reported fatal COVID-19 event. Baseline characteristics were summarized as means and standard deviations, medians and interquartile ranges, or frequencies and percentages, as appropriate. ANOVA, Kruskal–Wallis, and chi-square tests were used to test for between-group differences.

To assess the impact of the COVID-19 pandemic on changes in event rates, we compared event rates (per 100 participant-years) between a pre-pandemic and pandemic period. The pre-pandemic period was

defined as before the World Health Organization's declaration of COVID-19 as a pandemic on 11 March 2020. Clinical outcomes assessed included (1) the primary composite outcome, (2) components of the primary composite, and (3) all-cause death. We further compared the proportion of all deaths attributable to CV causes (vs. non-CV causes) and the proportion of primary events due to worsening HF events (vs. CV death) between the two periods. Comparisons were adjusted for time since randomization. The associations between COVID-19 and risks of time-to-event outcomes were estimated by creating time-updated covariates to represent pre- versus post-COVID-19 diagnosis status at the patient level, as well as pre-pandemic and pandemic period (i.e. set to 0 for all patient follow-up prior to 11 March 2020 and set to 1 for all follow-up subsequent to 11 March 2020), such that comparisons were made between participants based on the time they have spent in the study from randomization or from 11 March 2020. We also conducted a dedicated sensitivity analysis evaluating the treatment effects of dapagliflozin versus placebo on clinical outcomes with censoring at the time of pandemic onset.

To assess the health and safety risks faced by participants after COVID-19 diagnosis, we evaluated (1) drug interruptions and withdrawals and (2) adverse events in the 30 days following COVID-19 diagnosis overall and by treatment assignment. In addition, we assessed the risk of clinical outcomes in the 12 months after incident COVID-19 using Cox proportional hazards models. To evaluate excess short-term risk after COVID-19 diagnosis, we compared these risks to those faced by all trial participants from the time of enrolment, censoring participants who developed COVID-19 at the time of diagnosis. Cox models were adjusted for age, sex, race, geographic region, body mass index, HF duration, NT-proBNP, glycated haemoglobin, estimated glomerular filtration rate, baseline use of loop diuretics and angiotensin receptor–neprilysin inhibitor, and history of type 2 diabetes, chronic obstructive pulmonary disease, sleep apnoea, atherosclerotic vascular disease, prior HF hospitalization, smoking, atrial fibrillation/atrial flutter.

All analyses were performed in Stata version 16 (StataCorp, College Station, TX, USA). A nominal p -value of ≤ 0.05 was considered statistically significant.

Results

COVID-19 among participants in DELIVER

At the time of first reported COVID-19 infection in the trial (16 March 2020), 5153 (82.3%) of the total trial population had been randomized, 363 (5.8%) had already experienced a primary endpoint event (out of a total of 1122 primary events in the trial), and 168 (2.7%) had died; >75% of trial follow-up occurred on or after 11 March 2020. Among the full study population of 6263 participants in DELIVER, 589 (9.4%) were diagnosed with COVID-19 during the trial, 314/3131 (10.0%) of which were randomized to dapagliflozin and 275/3132 (8.7%) randomized to placebo (*Graphical Abstract*). Most participants with COVID-19 had symptomatic disease (92%) and the majority ($n = 307$, 52.1%) required hospitalization or had prolongation of an existing hospitalization (*Figure 1*). Reported COVID-19 cases occurred across all four enrolling regions: Europe and Saudi Arabia ($n = 377$), Asia ($n = 18$), Latin America ($n = 128$) and North America ($n = 66$).

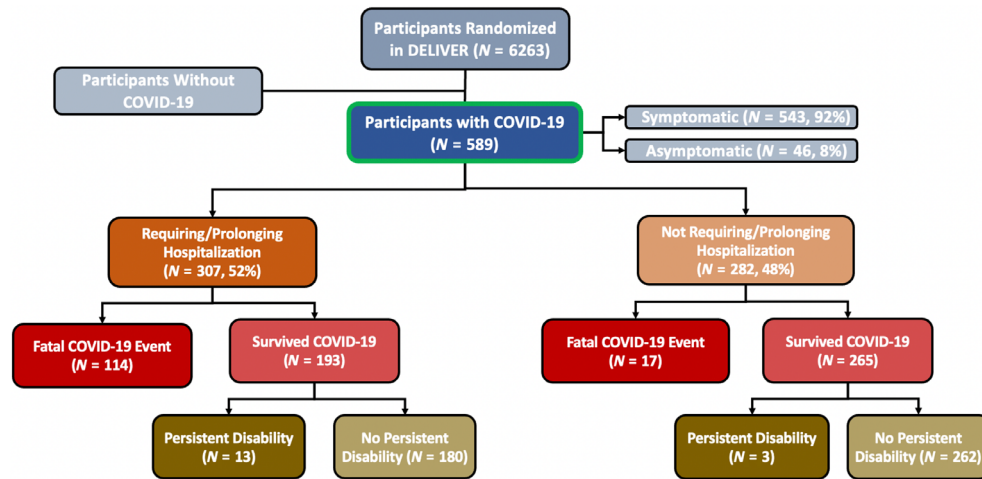


Figure 1 Impact of COVID-19 on participants in DELIVER. Persistent disability was based on investigator report.

Patients who developed COVID-19 during the trial were similar in age and sex but were more likely to be white, enrolled in Europe/Saudi Arabia, and had greater symptomatic impairment by baseline Kansas City Cardiomyopathy Questionnaire total symptom score (KCCQ-TSS) than those who did not develop COVID-19. Mean LVEF was similar between those who did ($54.1 \pm 8.4\%$) and did not ($54.2 \pm 8.8\%$) develop COVID-19. Rates of pulmonary comorbidities, including chronic obstructive pulmonary disease and smoking, were also similar between groups; however, participants with COVID-19 were more likely to have cardiometabolic comorbidities, including diabetes, atherosclerotic cardiovascular disease, and obesity. Older patients and those with higher baseline NT-proBNP levels were more likely to have severe COVID-19 (requiring/prolonging hospitalization or resulting in a fatal COVID-19 event, $n = 324$) versus non-severe COVID-19 (Table 1). Baseline characteristics in the COVID-19 population were similar by treatment assignment (online supplementary Table S1). COVID-19 diagnosis rates over time are summarized in online supplementary Figure S1.

Rates of trial outcomes prior to and during the COVID-19 pandemic

The primary composite endpoint occurred at a rate of 11.0 (95% confidence interval [CI] 9.9–12.2) per 100 participant-years during the pre-pandemic period (prior to 11 March 2020); rates were numerically lower during the pandemic (7.9, 95% CI 7.4–8.5 per 100 participant-years), but not statistically different when adjusting for the time from randomization ($p = 0.34$). However, the proportion of primary composite events attributable to worsening HF events (HF hospitalization or urgent HF visit) was lower during the pandemic period compared with the pre-pandemic period (68.7% vs. 82.8%, $p < 0.01$). All-cause death rates were higher during the pandemic (8.2, 95% CI 7.6–8.7 per 100 participant-years) versus pre-pandemic periods (5.0, 95% CI 4.3–5.8 per

100 participant-years; $p = 0.05$); the proportion of deaths related to non-CV causes also rose during the pandemic period (Table 2).

Health risks after COVID-19 diagnosis

Adverse events

Among DELIVER participants on study drug at the time they developed COVID-19 ($n = 533$), 91 (17.1%) had study drug interruption or withdrawal after reported infection; most ($n = 84$, 92%) were temporary interruptions with subsequent resumption of study drug. Study drug interruption/withdrawal did not differ by treatment assignment to dapagliflozin versus placebo (15.9% vs. 18.4%, $p = 0.44$), but was more common overall in patients with severe COVID-19 (28.6%) versus non-severe COVID-19 (3.7%) (online supplementary Table S2). Adverse events in the 30 days following COVID-19 diagnosis are listed in Table 3. There were four investigator-reported renal adverse events which were serious in nature or resulted in IP discontinuation and two episodes of volume depletion among participants within 30 days of COVID-19 diagnosis. There were no reported hypoglycaemia or DKA events among trial participants within 30 days of COVID-19 diagnosis. Adverse event rates did not differ by treatment assignment (Table 3). Adverse events by COVID-19 severity and by region of enrolment are listed in online supplementary Tables S3 and S4, respectively.

Among patients with COVID-19 during the trial ($n = 589$), 181 (30.7%) died on or after the date of COVID-19 diagnosis (100/314 participants randomized to dapagliflozin and 81/275 participants randomized to placebo, $p = 0.53$). The majority ($n = 131$, 72.4%) of these deaths were linked by investigators to the index COVID-19 diagnosis, of which an independent clinical events committee adjudicated 122 deaths to be definitely or probably related to COVID-19. The clinical events committee determined an additional 33 deaths to be definitely or probably related to COVID-19, for a total of 155 deaths with direct attribution to COVID-19 (15% of total deaths). Deaths within 30 days or at any time following a COVID-19 diagnosis, fatal COVID-19 adverse events, or

Table 1 Baseline characteristics among trial participants by COVID-19 status

Characteristics	Developed COVID-19 (n = 589)	Did not develop COVID-19 (n = 5674)	p-value	Severe COVID-19 (n = 324)	Non-severe COVID-19 (n = 265)	p-value
Age (years)	71.2 ± 9.6	71.7 ± 9.5	0.17	72.1 ± 9.2	70.0 ± 10.1	0.009
Age group (years)			0.31			0.021
≤65	154 (26.1%)	1350 (23.8%)		70 (21.6%)	84 (31.7%)	
>65–75	229 (38.9%)	2183 (38.5%)		135 (41.7%)	94 (35.5%)	
>75	206 (35.0%)	2141 (37.7%)		119 (36.7%)	87 (32.8%)	
Men	319 (54.2%)	3197 (56.3%)	0.31	186 (57.4%)	133 (50.2%)	0.08
Race			< 0.001			0.11
White	508 (86.2%)	3931 (69.3%)		281 (86.7%)	227 (85.7%)	
Asian	19 (3.2%)	1255 (22.1%)		10 (3.1%)	9 (3.4%)	
Black or African American	21 (3.6%)	138 (2.4%)		10 (3.1%)	11 (4.2%)	
American Indian or Alaska Native	20 (3.4%)	169 (3.0%)		7 (2.2%)	13 (4.9%)	
Other	21 (3.6%)	181 (3.2%)		16 (4.9%)	5 (1.9%)	
Geographic region			< 0.001			0.26
Europe and Saudi Arabia	377 (64.0%)	2628 (46.3%)		219 (67.6%)	158 (59.6%)	
Asia	18 (3.1%)	1208 (21.3%)		9 (2.8%)	9 (3.4%)	
Latin America	128 (21.7%)	1053 (18.6%)		63 (19.4%)	65 (24.5%)	
North America	66 (11.2%)	785 (13.8%)		33 (10.2%)	33 (12.5%)	
Comorbidities						
History of:						
AFF	321 (54.5%)	3231 (56.9%)	0.25	180 (55.6%)	141 (53.2%)	0.57
Stroke	56 (9.5%)	541 (9.5%)	0.98	33 (10.2%)	23 (8.7%)	0.54
Dyslipidemia	412 (69.9%)	3578 (63.1%)	< 0.001	218 (67.3%)	194 (73.2%)	0.12
Type 2 diabetes mellitus	304 (51.6%)	2502 (44.1%)	< 0.001	172 (53.1%)	132 (49.8%)	0.43
Chronic obstructive pulmonary disease	73 (12.4%)	619 (10.9%)	0.27	46 (14.2%)	27 (10.2%)	0.15
Peripheral artery disease	42 (7.1%)	337 (5.9%)	0.24	27 (8.3%)	15 (5.7%)	0.21
Sleep apnoea	59 (10.0%)	426 (7.5%)	0.029	28 (8.6%)	31 (11.7%)	0.21
Myocardial infarction	180 (30.6%)	1459 (25.7%)	0.011	102 (31.5%)	78 (29.4%)	0.59
Prior HF hospitalization flag	219 (37.2%)	2320 (40.9%)	0.08	132 (40.7%)	87 (32.8%)	0.048
Any atherosclerotic cardiovascular disease	356 (60.4%)	3196 (56.3%)	0.06	201 (62.0%)	155 (58.5%)	0.38
Smoking status			0.61			0.67
Current	41 (7.0%)	443 (7.8%)		20 (6.2%)	21 (7.9%)	
Former	207 (35.1%)	2054 (36.2%)		113 (34.9%)	94 (35.5%)	
Never	341 (57.9%)	3177 (56.0%)		191 (59.0%)	150 (56.6%)	
Time from diagnosis of HF			0.024			0.88
0–3 months	41 (7.0%)	527 (9.3%)		23 (7.1%)	18 (6.8%)	
>3–6 months	47 (8.0%)	545 (9.6%)		28 (8.6%)	19 (7.2%)	
>6–12 months	66 (11.2%)	776 (13.7%)		40 (12.3%)	26 (9.8%)	
>1–2 years	109 (18.5%)	886 (15.6%)		58 (17.9%)	51 (19.2%)	
>2–5 years	146 (24.8%)	1423 (25.1%)		80 (24.7%)	66 (24.9%)	
>5 years	180 (30.6%)	1512 (26.7%)		95 (29.3%)	85 (32.1%)	
Body mass index (kg/m ²)	31.9 ± 6.1	29.6 ± 6.1	< 0.001	31.8 ± 6.3	32.1 ± 5.8	0.63
Body mass index group (kg/m ²)			< 0.001			0.18
<18.5 (underweight)	2 (0.3%)	52 (0.9%)		0 (0.0%)	2 (0.8%)	
18.5–24.9 (normal weight)	62 (10.5%)	1281 (22.6%)		41 (12.7%)	21 (7.9%)	
25.0–29.9 (overweight)	182 (30.9%)	1891 (33.4%)		100 (30.9%)	82 (30.9%)	
30.0–34.9 (Class I obesity)	179 (30.4%)	1395 (24.6%)		94 (29.0%)	85 (32.1%)	

Table 1 (Continued)

Characteristics	Developed COVID-19 (n = 589)	Did not develop COVID-19 (n = 5674)	p-value	Severe COVID-19 (n = 324)	Non-severe COVID-19 (n = 265)	p-value
35.0–39.9 (Class II obesity)	101 (17.1%)	697 (12.3%)		51 (15.7%)	50 (18.9%)	
≥40 (Class III obesity)	63 (10.7%)	352 (6.2%)		38 (11.7%)	25 (9.4%)	
NYHA class at baseline			0.92			0.06
I	0 (0.0%)	1 (0.0%)		–	–	
II	441 (74.9%)	4272 (75.3%)		232 (71.6%)	209 (78.9%)	
III	147 (25.0%)	1384 (24.4%)		92 (28.4%)	55 (20.8%)	
IV	1 (0.2%)	17 (0.3%)		0 (0.0%)	1 (0.4%)	
KCCQ-TSS	66.2 ± 21.5	70.4 ± 22.2	< 0.001	64.1 ± 20.6	68.7 ± 22.4	0.015
LVEF (%)	54.1 ± 8.4	54.2 ± 8.8	0.93	54.2 ± 8.5	54.1 ± 8.3	0.85
LVEF group (%)			0.52			0.87
≤49	200 (34.0%)	1916 (33.8%)		110 (34.0%)	90 (34.0%)	
50–59	201 (34.1%)	2055 (36.2%)		108 (33.3%)	93 (35.1%)	
≥60	188 (31.9%)	1703 (30.0%)		106 (32.7%)	82 (30.9%)	
NT-proBNP (ng/L)	975 [569–1621]	1017 [626–1768]	0.019	1050 [638–1686]	824 [502–1452]	< 0.001
NT-proBNP in AFF	1351 [951–2309]	1402 [964–2208]	0.61	1399 [1003–2286]	1274 [882–2318]	0.43
NT-proBNP when no AFF	666 [451–1187]	718 [470–1299]	0.14	768 [491–1292]	610 [406–1074]	0.002
ECG AFF	227 (38.5%)	2417 (42.6%)	0.06	135 (41.7%)	92 (34.7%)	0.08
Systolic blood pressure (mmHg)	130.0 ± 15.5	128.1 ± 15.3	0.003	129.4 ± 15.5	130.7 ± 15.5	0.29
Diastolic blood pressure (mmHg)	73.9 ± 10.2	73.9 ± 10.4	0.85	74.1 ± 10.4	73.5 ± 10.0	0.50
HbA1c (%)	6.9 ± 1.7	6.6 ± 1.4	< 0.001	7.0 ± 1.8	6.7 ± 1.5	0.10
Pulse rate (bpm)	70.5 ± 11.4	71.6 ± 11.8	0.028	71.1 ± 10.9	69.7 ± 12.0	0.13
Creatinine (μmol/L)	105.5 ± 33.2	102.2 ± 30.8	0.013	108.6 ± 33.6	101.7 ± 32.3	0.011
eGFR (ml/min/1.73 m ²)	59.3 ± 18.5	61.2 ± 19.2	0.018	57.3 ± 18.3	61.6 ± 18.5	0.005
Medication use						
Loop diuretics	463 (78.6%)	4348 (76.7%)	0.29	268 (82.7%)	195 (73.6%)	0.007
ACEi	230 (39.0%)	2065 (36.4%)	0.21	136 (42.0%)	94 (35.5%)	0.11
ARB	220 (37.4%)	2052 (36.2%)	0.57	107 (33.0%)	113 (42.6%)	0.016
ARNI	17 (2.9%)	284 (5.0%)	0.022	9 (2.8%)	8 (3.0%)	0.86
Beta-blocker	484 (82.2%)	4693 (82.7%)	0.73	274 (84.6%)	210 (79.2%)	0.09
MRA	239 (40.6%)	2428 (42.8%)	0.30	143 (44.1%)	96 (36.2%)	0.05
Pacemaker	75 (12.7%)	587 (10.3%)	0.07	40 (12.3%)	35 (13.2%)	0.75
ICD	15 (2.5%)	98 (1.7%)	0.15	7 (2.2%)	8 (3.0%)	0.51

Values are presented as mean ± standard deviation or median [25th–75th percentile] unless otherwise indicated.

Severe COVID-19 defined as COVID-19 infection requiring hospitalization and/or resulting in a fatal COVID-19 event as determined by investigators. COVID-19 infection was determined by investigator report and may or may not have been confirmed with laboratory testing.

ACEi, angiotensin-converting enzyme inhibitor; AFF, atrial fibrillation/flutter; ARB, angiotensin receptor blocker; ARNI, angiotensin receptor–neprilysin inhibitor; ECG, electrocardiogram; eGFR, estimated glomerular filtration rate; HbA1c, glycated haemoglobin; HF, heart failure; ICD, implantable cardioverter-defibrillator; KCCQ-TSS, Kansas City Cardiomyopathy Questionnaire total symptom score; LVEF, left ventricular ejection fraction; MRA, mineralocorticoid receptor antagonist; NT-proBNP, N-terminal pro-B-type natriuretic peptide; NYHA, New York Heart Association.

deaths definitely or possibly related to COVID-19 as determined by the clinical events committee did not differ significantly by treatment assignment (Table 3).

Treatment effects

A previously reported sensitivity analysis demonstrated consistent treatment benefits of dapagliflozin versus placebo when censoring participants at the time of their COVID-19 diagnosis.⁸ In the overall trial, rates of all-cause death were numerically lower in participants randomized to dapagliflozin versus placebo (hazard

ratio [HR] 0.94, 95% CI 0.83–1.07). This observed trend appeared marginally stronger in the pre-specified analysis with censoring participants at the time of COVID-19 diagnosis (HR 0.89, 95% CI 0.78–1.02). The effects of dapagliflozin versus placebo on the primary endpoint were similar when considering the full trial population (HR 0.82, 95% CI 0.73–0.92) and after censoring at the time of COVID-19 diagnosis (HR 0.81, 95% CI 0.72–0.91). We conducted an additional sensitivity analysis, censoring participants at the time of pandemic onset (11 March 2020). Dapagliflozin consistently reduced the time CV death or worsening HF event (HR 0.72, 95% CI 0.58–0.89) when censoring at pandemic onset.

Table 2 Event rates prior to and during the COVID-19 pandemic

	Before 11 March 2020	On or after 11 March 2020	RR pandemic period to pre-pandemic period ^a (95% CI)	Time-stratified p-value
Event rates, per 100 participant-years (95% CI)				
Primary composite	11.0 (9.9–12.2)	7.9 (7.4–8.5)	0.93 (0.79–1.09)	0.34
All-cause death	5.0 (4.3–5.8)	8.2 (7.6–8.7)	1.24 (1.00–1.54)	0.05
CV death	2.9 (2.4–3.5)	3.8 (3.4–4.2)	1.10 (0.82–1.48)	0.52
Worsening HF event	9.1 (8.1–10.2)	5.5 (5.0–5.9)	0.86 (0.72–1.03)	0.11
Hospitalization for HF	7.9 (7.0–8.9)	5.0 (4.6–5.5)	0.90 (0.75–1.09)	0.30
Urgent HF visit	1.5 (1.1–2.0)	0.8 (0.7–1.0)	0.59 (0.36–0.94)	0.026
Proportion of all-cause death attributable to CV causes	98/164 (59.8%)	394/853 (46.2%)	–	<0.001
Proportion of the primary composite due to a worsening HF event	299/361 (82.8%)	523/761 (68.7%)	–	<0.001
Proportion of all worsening HF events which were due to HF hospitalization	257/299 (86.0%)	461/524 (88.0%)	–	0.40

CI, confidence interval; CV, cardiovascular; HF, heart failure; RR, risk ratio.

^aThe associations between COVID-19 and risks of time-to-event outcomes were estimated by creating time-updated covariates to represent pre- versus post-COVID-19 diagnosis status at the patient level.

Table 3 Adverse events by treatment arm in the 30 days after COVID-19 diagnosis and deaths related to COVID-19

Adverse Events	Overall	Randomized to placebo	Randomized to dapagliflozin
Any SAE	585 (99.3%)	272 (98.9%)	313 (99.7%)
Any AE leading to discontinuation of IP	11 (1.9%)	5 (1.8%)	6 (1.9%)
Any AE leading to interruption of IP	94 (16.0%)	49 (17.8%)	45 (14.3%)
Any amputation	0 (0.0%)	0 (0.0%)	0 (0.0%)
Any definite or probable diabetic ketoacidosis	0 (0.0%)	0 (0.0%)	0 (0.0%)
Any major hypoglycaemic event	0 (0.0%)	0 (0.0%)	0 (0.0%)
Any SAE or DAE suggestive of volume depletion	2 (0.3%)	2 (0.7%)	0 (0.0%)
Any renal SAE or DAE	4 (0.7%)	2 (0.7%)	2 (0.6%)
Deaths following COVID-19 diagnosis			
Any death within 30 days of COVID-19 diagnosis	156 (26.5%)	70 (25.5%)	86 (27.4%)
Any death after COVID-19 diagnosis	181 (30.7%)	81 (29.5%)	100 (31.8%)
Fatal COVID-19 adverse event (investigator-reported)	131 (22.4%)	57 (21.0%)	74 (23.6%)
Death definitely or possibly related to COVID-19 (by CEC) ^a	142 (24.1%)	62 (22.5%)	80 (25.5%)

No statistically significant differences by randomized treatment assignment.

AE, adverse event; CEC, clinical endpoints committee; DAE, adverse events leading to discontinuation of study drug; IP, investigational product; SAE, serious adverse event.

^aAn additional 13 deaths (for a total of 155 deaths) without an investigator-reported COVID-19 diagnosis were adjudicated by the CEC as definitely or possibly related to COVID-19.

Risks of death after COVID-19

All-cause death rate in the 12 months following diagnosis was 56.1 (95% CI 48.0–65.6) per 100 participant-years compared to 6.4 (95% CI 6.0–6.8) per 100 participant-years among all trial participants censored at the time of COVID-19 (adjusted HR [aHR] 8.22, 95% CI 6.86–9.85). All-cause death was highest in the 3 months following COVID-19 diagnosis (153.5, 95% CI 130.3–180.8 per 100 participant-years) but remained elevated even 3–6 months after COVID-19 diagnosis (12.6, 95% CI 6.6–24.3 per 100 participant-years). Death rates declined 6–12 months from COVID-19 diagnosis (5.2, 95% CI 2.3–11.6 per

100 participant-years (Figure 2). Results were primarily driven by excess non-CV mortality (online supplementary Figure S2). After excluding immediately fatal COVID-19 events, all-cause death rates in the 12 months following diagnosis among COVID-19 survivors ($n=458$) remained multifold higher (17.0, 95% CI 12.8–22.6 per 100 participant-years; aHR 2.46, 95% CI 1.83–3.33) than all trial participants from randomization, censoring participants who developed COVID-19 at the time of diagnosis. Results were largely consistent by region (online supplementary Table S5). Rates of the primary endpoint and its components were numerically higher in the year following COVID-19 diagnosis as compared

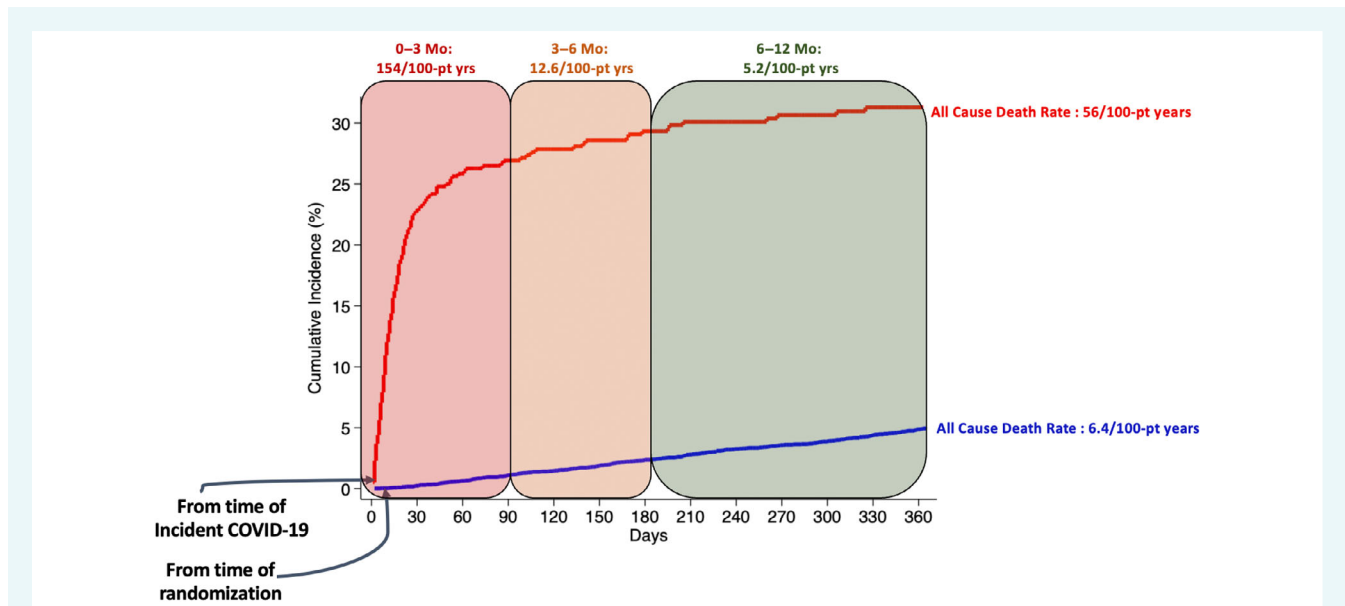


Figure 2 Mortality risk after COVID-19 compared to trial participants without COVID-19. Kaplan–Meier curves in blue indicate event rates among participants from the time of randomization and censored at the time of COVID-19 adverse event reporting.

Table 4 Risk of clinical outcomes following COVID-19 diagnosis

Event rates per 100 participant-years, 95% CI	Full trial population, censored at COVID-19 diagnosis (n = 6263)	All COVID-19 patients in the 12 months after diagnosis (n = 589)	HR, unadjusted	HR, adjusted ^a	All COVID-19 patients in the 12 months after diagnosis, excluding fatal COVID-19 events (n = 458)	HR, unadjusted	HR, adjusted ^a
Primary endpoint	8.7 (8.2–9.2)	8.9 (5.9–13.5)	1.20 (0.78–1.83)	1.30 (0.85–1.99)	8.9 (5.9–13.5)	1.21 (0.79–1.86)	1.32 (0.86–2.03)
All-cause death	6.4 (6.0–6.8)	56.1 (48.0–65.6)	7.97 (6.69–9.51)	8.22 (6.86–9.85)	17.0 (12.8–22.6)	2.40 (1.79–3.22)	2.46 (1.83–3.33)
CV death	3.5 (3.2–3.8)	5.7 (3.5–9.3)	1.48 (0.89–2.44)	1.55 (0.93–2.56)	5.3 (3.2–8.9)	1.39 (0.83–2.33)	1.45 (0.86–2.44)
Worsening HF events	6.4 (6.0–6.9)	5.3 (3.1–9.1)	1.06 (0.61–1.84)	1.20 (0.69–2.08)	5.3 (3.1–9.1)	1.08 (0.62–1.87)	1.23 (0.70–2.13)

Immediately fatal adverse were determined and reported by the investigators.

CI, confidence interval; CV, cardiovascular; HF, heart failure; HR, hazard ratio.

^aAdjusted for age, sex, race, geographic region, body mass index, HF duration, N-terminal pro-B-type natriuretic peptide, glycated haemoglobin, estimated glomerular filtration rate, baseline use of loop diuretics and angiotensin receptor–neprilysin inhibitor, and history of type 2 diabetes, chronic obstructive pulmonary disease, sleep apnoea, atherosclerotic vascular disease, prior HF hospitalization, smoking, atrial fibrillation/atrial flutter.

with all participants from randomization with censoring at the time of COVID-19 in both unadjusted and adjusted models (Table 4).

COVID-19 vaccination

Overall, 2722 trial participants received at least one vaccination for COVID-19, representing 47% of the trial population. There were 6456 vaccinations in 2722 participants during the trial; 277 participants received a single COVID-19 vaccination, 1156 received two vaccine doses, and 1289 received three vaccinations. Among participants who developed COVID-19 during the trial period (n = 589), 222 (38%) received at least one dose of COVID-19 vaccination, and 184 (31%) received at least two doses. Of these patients with COVID-19 who completed a primary vaccination series (≥ 2 doses), 90 (49%) received a third (i.e., booster) vaccine.

Discussion

A substantial proportion of participants with HFmrEF or HFpEF in the DELIVER trial were diagnosed with COVID-19 during the course of the study, with >50% of infections requiring/prolonging hospitalization and many leading to death. The pandemic period was associated with greater all-cause mortality and consequently relatively lower rates of worsening HF events compared to the pre-pandemic period. Participants diagnosed with COVID-19 had higher rates of all-cause death following infection, which remained residually elevated at 3–6 months after diagnosis, after excluding immediately fatal COVID-19 events and after extensive covariate adjustment. Adverse events leading to treatment discontinuation, major hypoglycaemia, DKA, and renal adverse events did not differ by treatment groups, as were all deaths and deaths ascribed to COVID-19. Dapagliflozin reduced clinical HF events as compared

to placebo when censoring at the time of COVID-19 diagnosis and pandemic onset. To our knowledge, DELIVER represents one of the most extensive international experiences with COVID-19 of any CV trial during the pandemic. These data provide a global perspective on the immediate to long-term impact of the COVID-19 pandemic in patients with HFmrEF or HFpEF.

In this large, international trial of patients with HFmrEF or HFpEF, COVID-19 was reported in nearly 10% of trial participants during the study period, with over half of infections requiring or prolonging hospitalization and nearly one in five resulting in a fatal outcome. These data highlight the burden and disease severity associated with COVID-19 in a large and growing segment of the HF population. Trial participants with cardiometabolic (but not pulmonary) comorbidities were more likely to have COVID-19; these data substantiate early reports identifying existent CV conditions as associated with increased susceptibility to viral infection and potential to more severe disease.^{1,9,10} Participants with severe forms of COVID-19 in DELIVER were older and had higher NP levels at baseline compared with those with non-severe disease. Prior observational data from a large, claims database¹¹ and a clinical registry¹² found NP levels at hospital admission to be prognostically important in indicating severity of COVID-19 disease; our data potentially extend the prognostic potential of NP levels to ambulatory settings prior to COVID-19 diagnosis. Existent endothelial, microvascular, and myocardial abnormalities, and reduced immune responses in more severe HF phenotypes characterized by higher NP levels may contribute to the higher incidence of more severe forms of COVID-19.^{13–16} Overall, among patients with HFmrEF or HFpEF, risks of COVID-19 are high and often associated with severe disease phenotypes. Innovations in prevention, rapid detection and early mitigation of COVID-19 should be prioritized in this high-risk patient population.

The pandemic period was associated with a rise in all-cause death and a decline in the proportion of primary outcome events attributable to worsening HF (vs. CV death) compared with the pre-pandemic period in DELIVER. Deaths attributable to or influenced by COVID-19 likely contributed significantly to the observed rise in all-cause death. A smaller trial assessing haemodynamic monitoring of HF patients found no difference in death rates between pre-pandemic and pandemic period, though was limited to North American participants only and was operational for a shorter period of time during the pandemic.⁷ Concurrent observational studies during the early pandemic reported declines in hospitalizations for acute CV conditions, including HF hospitalizations.^{17,18} Potential reasons include the effects of stay-at-home messaging, concerns about the safety and accessibility of hospital care, and the potential effects of environmental factors (e.g. reduced pollution, lower exposure to other viral pathogens [e.g. influenza], and availability of sodium and high caloric foods), though direct evidence for the latter is limited. Concurrently, reports indicated increased out-of-hospital cardiac arrests¹⁹ and lower survival/return of spontaneous circulation rates during the pandemic.²⁰ Implantation of primary and secondary prevention defibrillators also declined.²¹ These trends may contribute to the observed shift between worsening HF events and CV death observed during the pandemic.^{5,6} These findings may have

important implications for the design and conduct of cardiovascular trials, particularly as COVID-19 remains endemic; the use of adaptive clinical trial designs may help to deal with uncertainty in event rate accrual caused by unexpected trial disruptions such as COVID-19. However, COVID-19 had little impact on treatment effect estimates of dapagliflozin versus placebo on clinical outcomes. These data come in contrast to prior trials in which the pandemic period was associated with general attenuation treatment effects.^{5,7} Reasons for these discrepant findings are unclear but may be related to DELIVER remaining active in later timepoints during the pandemic. Importantly, given the post-randomization nature of the exposure (the COVID-19 pandemic), natural evolution of fatal and non-fatal events in a time-to-event design with a composite endpoint may have also contributed, as non-fatal events among high-risk participants are generally accrued early (i.e. before the pandemic) while fatal events generally occur later (i.e. during the pandemic)²²; conclusions about the causes of the changing epidemiology of clinical events during later trial periods should therefore be interpreted with caution.

Study drug interruptions or withdrawals following COVID-19 diagnosis were largely temporary and occurred at similar rates among those randomized to dapagliflozin or placebo. Guidance to sites largely mirrored clinical practice and included consideration of study drug interruption among participants with COVID-19 who were at high risk for or with prior history of DKA. There was no reported major hypoglycaemia or DKA events and adverse renal events were rare in the 30 days following COVID-19 diagnosis; precise reasons for interruption or withdrawal were not systematically available. Poor nutritional intake and volume depletion during acute COVID-19 infection raised concern that sodium–glucose cotransporter 2 inhibitor (SGLT2i) use might further increase risks of euglycaemic DKA.²³ However, a retrospective cohort of patients with diabetes and COVID-19 found no evidence of increased risk of DKA among patients receiving SGLT2i. This study observed potential favourable effects on endothelial function and reductions in oxidative stress with this therapy, which may have implications in attenuating the severity of acute viral illness.²⁴ The Dapagliflozin in Respiratory Failure in Patients With COVID-19 (DARE-19) trial randomized 1250 patients with cardiometabolic risk factors who were hospitalized with COVID-19 to dapagliflozin versus placebo and also found no increase in the risks of acute kidney injury or DKA among participants randomized to dapagliflozin^{25,26}; however, only 90 participants (7.2%) had a history of HF in DARE-19. Data from DELIVER extend the available safety data of dapagliflozin with standard of care guidance to a large cohort of patients with HFmrEF or HFpEF; in fact, similar safety profiles were observed in clinical trials of patients with HFpEF conducted prior to the COVID-19 pandemic. Deaths following a COVID-19 diagnosis did not statistically differ by treatment groups. Collectively, the data from this DELIVER analysis and DARE-19 are reassuring in regard to the tolerability of SGLT2i in patients that develop COVID-19, and support consideration of the continuation of SGLT2i (with usual care guidance) in stable HF with COVID-19. Of note, additional studies including from two large platform trials (Randomised Evaluation of COVID-19 Therapy [RECOVERY] and Accelerating COVID-19 Therapeutic Interventions and Vaccines

4 ACUTE [ACTIV-4A]) have assessed the safety and efficacy of SGLT2i in patients hospitalized with COVID-19, independent of HF status. These data, in aggregate, will further clarify the role of SGLT2i in patients with COVID-19.

Participants in DELIVER who developed COVID-19 during the trial faced increased risks of all-cause death following diagnosis, compared to those who did not. Importantly, elevated all-cause death risk persisted 3–6 months after diagnosis, after adjustment for clinically relevant confounders, and when assessing only the population surviving their COVID-19 event. Rates of the primary endpoint and its components were numerically higher in the year following COVID-19 as compared to participants without COVID-19 in both unadjusted and adjusted models. A prior claims-based analysis found that among patients with HF, COVID-19 hospitalization carried a significantly elevated prognostic risk of death compared to similar patients hospitalized with acute HF; however, follow-up was limited to only the in-hospital setting.¹ Recent evidence from >500 000 registry participants found an increased incidence of new HF diagnoses in patients in the year following COVID-19 diagnosis; however, the registry findings were limited to those without prior known HF.² Relatively small sample sizes and limited post-COVID-19 follow-up in DELIVER affected statistical power to detect excess CV death risk among participants diagnosed with COVID-19. Our data from DELIVER extend the potential downstream risks of COVID-19 to patients with existing HF in ambulatory and hospitalized settings. Similar trends have been observed with influenza infection; a recent secondary analysis of two large HF trials found that trial participants faced >4× greater risk of death following influenza diagnosis compared with participants without influenza infection, though the excess risks after COVID-19 diagnosis in DELIVER were larger.²⁷ Extended surveillance programmes among HF patients with COVID-19 (and other viral disease) are needed to fully understand and attempt to mitigate possible longer-term collateral effects of COVID-19 illness.²⁸

Limitations

The results of our analysis should be considered in the context of several potential limitations. First, COVID-19 diagnosis was determined by adverse event reporting from site investigators; local regulations, access to and availability of testing may have varied over time and by site or region, which may have led to ascertainment bias and undercounting of true COVID-19 cases. The inclusion of possible COVID-19 cases without polymerase chain reaction confirmation may have also affected the results, though all deaths were centrally adjudicated for relatedness to COVID-19. Second, unmeasured confounding and natural changes during a trial may in part explain the observed findings. Differences in trial event rates before and during the pandemic should be interpreted in the context of time-to-first endpoints, in which early accrual of non-fatal events may have affected the observed findings. Third, data on in-hospital resource use (e.g. mechanical ventilation and haemodynamic monitoring) and COVID-19 specific treatment (e.g. corticosteroids) were not available, nor was specific detailed assessment of potential cardiopulmonary sequelae of COVID-19

disease. Fourth, drug interruption and withdrawal rates were obtained from adverse event reporting; the specific reasons for study drug interruption and timing of interruption/resumption were unavailable and underreporting during acute illness (e.g. during COVID-19 hospitalization) may be possible. Fifth, event reporting may have been impacted by operational challenges during the pandemic. Sixth, data on COVID-19 variants were not available; time-dependent variation in the risk of COVID-19 infection influenced by circulating variants may have impacted the observed findings. Sixth, availability, eligibility, and access to COVID-19 vaccination may have varied significantly across sites and regions during the trial period; reasons for not obtaining COVID-19 vaccine were not systematically captured. Finally, most patients who developed COVID-19 did so substantially after randomization, so post-randomization treatment comparisons are only hypothesis generating and should be interpreted with caution.

Conclusion

In DELIVER, COVID-19 was common, with >50% of diagnoses requiring/prolonging hospitalization and a high proportion leading to death. Serious adverse events, adverse events leading to treatment discontinuation, DKA, renal adverse events, hypoglycaemia, and deaths following COVID-19 diagnosis did not differ between dapagliflozin and placebo arms and permanent treatment withdrawals were rare. Dapagliflozin improved clinical outcomes when censoring at the time of COVID-19 diagnosis and at the time of pandemic onset. Patients surviving a COVID-19 diagnosis face residual elevations in clinical risk in the months following this event.

Supplementary Information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Conflict of interest: M.N.K. has received research grant support from AstraZeneca and Boehringer Ingelheim; has served as a consultant or on an advisory board for Alnylam, Amgen, Applied Therapeutics, AstraZeneca, Bayer, Boehringer Ingelheim, Eli Lilly, Esperion Therapeutics, Janssen, Lexicon, Merck (Diabetes and Cardiovascular), Novo Nordisk, Sanofi, Pharmacosmos and Vifor Pharma; has received other research support from AstraZeneca; and has received honoraria from AstraZeneca, Boehringer Ingelheim, and Novo Nordisk. B.L.C. has received consulting fees from Amgen, Cardurion, Corvia, and Novartis. M.V. has received research grant support, served on advisory boards, or had speaker engagements with American Regent, Amgen, AstraZeneca, Bayer AG, Baxter Healthcare, Boehringer Ingelheim, Chiesi, Cytokinetics, Lexicon Pharmaceuticals, Merck, Novartis, Novo Nordisk, Pharmacosmos, Relpsa, Roche Diagnostics, Sanofi, and Tricog Health, and participates in clinical trial committees for studies sponsored by AstraZeneca, Galmed, Novartis, Bayer AG, Occlutech, and Impulse Dynamics. C.S.P.L. is supported by a Clinician Scientist Award from the National Medical Research Council of Singapore; has received research support from Bayer and Roche Diagnostics; has served as consultant or on the Advisory Board/ Steering Committee/Executive Committee for Actelion, Alleviant Medical, Allysta Pharma, Amgen, AnaCardio AB, Applied Therapeutics, AstraZeneca, Bayer, Boehringer Ingelheim, Boston Scientific, Cytokinetics, Darma Inc.,

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References

- Bhatt AS, Jering KS, Vaduganathan M, Claggett BL, Cunningham JW, Rosenthal N, et al. Clinical outcomes in patients with heart failure hospitalized with COVID-19. *JACC Heart Fail* 2021;9:65–73. <https://doi.org/10.1016/j.jchf.2020.11.003>
- Salah HM, Fudim M, O'Neil ST, Manna A, Chute CG, Caughey MC. Post-recovery COVID-19 and incident heart failure in the National COVID Cohort Collaborative (N3C) study. *Nat Commun* 2022;13:4117. <https://doi.org/10.1038/s41467-022-31834-y>
- Alger HM, Rutan C, Williams JH, Walchok JG, Bolles M, Hall JL, et al. American Heart Association COVID-19 CVD Registry powered by Get With The Guidelines. *Circ Cardiovasc Qual Outcomes* 2020;13:e006967. <https://doi.org/10.1161/CIRCOUTCOMES.120.006967>
- Pfeffer MA, Claggett B, Lewis EF, Granger CB, Køber L, Maggioni AP, et al.; PARADISE-MI Investigators and Committees. Angiotensin receptor-neprilysin inhibition in acute myocardial infarction. *N Engl J Med* 2021;385:1845–1855. <https://doi.org/10.1056/NEJMoa2104508>
- Ponikowski P, Kirwan BA, Anker SD, McDonagh T, Dorobantu M, Drozd J, et al.; AFFIRM-AHF Investigators. Ferric carboxymaltose for iron deficiency at discharge after acute heart failure: A multicentre, double-blind, randomised, controlled trial. *Lancet* 2020;396:1895–1904. [https://doi.org/10.1016/S0140-6736\(20\)32339-4](https://doi.org/10.1016/S0140-6736(20)32339-4)
- Lindenfeld J, Zile MR, Desai AS, Bhatt K, Ducharme A, Horstmanshof D, et al. Haemodynamic-guided management of heart failure (GUIDE-HF): A randomised controlled trial. *Lancet* 2021;398:991–1001. [https://doi.org/10.1016/S0140-6736\(21\)01754-2](https://doi.org/10.1016/S0140-6736(21)01754-2)
- Zile MR, Desai AS, Costanzo MR, Ducharme A, Maisel A, Mehra MR, et al. The GUIDE-HF trial of pulmonary artery pressure monitoring in heart failure: Impact of the COVID-19 pandemic. *Eur Heart J* 2022;43:2603–2618. <https://doi.org/10.1093/eurheartj/ehac114>
- Solomon SD, McMurray JJV, Claggett B, de Boer RA, DeMets D, Hernandez AF, et al.; DELIVER Trial Committees and Investigators. Dapagliflozin in heart failure with mildly reduced or preserved ejection fraction. *N Engl J Med* 2022;387:1089–1098. <https://doi.org/10.1056/NEJMoa2206286>
- Targher G, Mantovani A, Wang XB, Yan HD, Sun QF, Pan KH, et al. Patients with diabetes are at higher risk for severe illness from COVID-19. *Diabetes Metab* 2020;46:335–337. <https://doi.org/10.1016/j.diabet.2020.05.001>
- Bhatt AS, Adler ED, Albert NM, Anyanwu A, Bhadelia N, Cooper LT, et al. Coronavirus disease-2019 and heart failure: A scientific statement from the Heart Failure Society of America. *J Card Fail* 2022;28:93–112. <https://doi.org/10.1016/j.cardfail.2021.08.013>
- Cunningham JW, Claggett BL, Jering KS, Vaduganathan M, Bhatt AS, Rosenthal N, et al. Prognostic value of natriuretic peptides and cardiac troponins in COVID-19. *Circulation* 2021;144:177–179. <https://doi.org/10.1161/CIRCULATIONAHA.121.054969>
- Bhatt AS, Daniels LB, de Lemos J, Goodrich E, Bohula EA, Morrow DA; AHA COVID-19 CVD Registry Investigators. Multi-marker risk assessment in patients hospitalized with COVID-19: Results from the American Heart Association COVID-19 Cardiovascular Disease Registry. *Am Heart J* 2023;149–156. <https://doi.org/10.1016/j.ahj.2022.12.014>
- Giustino G, Pinney SP, Lala A, Reddy VY, Johnston-Cox HA, Mechanick JL, et al. Coronavirus and cardiovascular disease, myocardial injury, and arrhythmia: JACC focus seminar. *J Am Coll Cardiol* 2020;76:2011–2023. <https://doi.org/10.1016/j.jacc.2020.08.059>
- Lala A, Johnson KW, Januzzi JL, Russak AJ, Paranjpe I, Richter F, et al.; Mount Sinai COVID Informatics Center. Prevalence and impact of myocardial injury in patients hospitalized with COVID-19 infection. *J Am Coll Cardiol* 2020;76:533–546. <https://doi.org/10.1016/j.jacc.2020.06.007>
- Libby P, Lüscher T. COVID-19 is, in the end, an endothelial disease. *Eur Heart J* 2020;41:3038–3044. <https://doi.org/10.1093/eurheartj/ehaa623>

16. Lowenstein CJ, Solomon SD. Severe COVID-19 is a microvascular disease. *Circulation* 2020;**142**:1609–1611. <https://doi.org/10.1161/CIRCULATIONAHA.120.050354>
17. Solomon MD, McNulty EJ, Rana JS, Leong TK, Lee C, Sung SH, et al. The Covid-19 pandemic and the incidence of acute myocardial infarction. *N Engl J Med* 2020;**383**:691–693. <https://doi.org/10.1056/NEJMc2015630>
18. Bhatt AS, Moscone A, McElrath EE, Varshney AS, Claggett BL, Bhatt DL, et al. Fewer hospitalizations for acute cardiovascular conditions during the COVID-19 pandemic. *J Am Coll Cardiol* 2020;**76**:280–288. <https://doi.org/10.1016/j.jacc.2020.05.038>
19. Baldi E, Sechi GM, Mare C, Canevari F, Brancaglione A, Primi R, et al.; Lombardia CARE Researchers. Out-of-hospital cardiac arrest during the Covid-19 outbreak in Italy. *N Engl J Med* 2020;**383**:496–498. <https://doi.org/10.1056/NEJMc2010418>
20. Chan PS, Girotra S, Tang Y, Al-Araji R, Nallamothu BK, McNally B. Outcomes for out-of-hospital cardiac arrest in the United States during the coronavirus disease 2019 pandemic. *JAMA Cardiol* 2021;**6**:296–303. <https://doi.org/10.1001/jamacardio.2020.6210>
21. Filipecki A, Orszulak M, Tajstra M, Kowalski O, Skrzypek M, Kalarus Z, et al. Cardiac implantable electronic devices procedures and their recipients characteristic during COVID-19 pandemic: 3.8 million population analysis. *Cardiol J* 2022;**29**:27–32. <https://doi.org/10.5603/CJ.a2021.0170>
22. Claggett B, Pocock S, Wei LJ, Pfeffer MA, McMurray JJV, Solomon SD. Comparison of time-to-first event and recurrent-event methods in randomized clinical trials. *Circulation* 2018;**138**:570–577. <https://doi.org/10.1161/CIRCULATIONAHA.117.033065>
23. Koufakis T, Pavlidis AN, Metallidis S, Kotsa K. Sodium-glucose co-transporter 2 inhibitors in COVID-19: Meeting at the crossroads between heart, diabetes and infectious diseases. *Int J Clin Pharm* 2021;**43**:764–767. <https://doi.org/10.1007/s11096-021-01256-9>
24. Khunti K, Ruan Y, Davies J, Field BCT, Harris S, Kosiborod M, et al.; ABCD COVID-19 Diabetes National Audit Investigators. Association between SGLT2 inhibitor treatment and diabetic ketoacidosis and mortality in people with type 2 diabetes admitted to hospital with COVID-19. *Diabetes Care* 2022;**45**:2838–2843. <https://doi.org/10.2337/dc22-0357>
25. Kosiborod MN, Esterline R, Furtado RHM, Oscarsson J, Gasparyan SB, Koch GG, et al. Dapagliflozin in patients with cardiometabolic risk factors hospitalised with COVID-19 (DARE-19): A randomised, double-blind, placebo-controlled, phase 3 trial. *Lancet Diabetes Endocrinol* 2021;**9**:586–594. [https://doi.org/10.1016/S2213-8587\(21\)00180-7](https://doi.org/10.1016/S2213-8587(21)00180-7)
26. Heerspink HJL, Furtado RHM, Berwanger O, Koch GG, Martinez F, Mukhtar O, et al. Dapagliflozin and kidney outcomes in hospitalized patients with COVID-19 infection: An analysis of the DARE-19 randomized controlled trial. *Clin J Am Soc Nephrol* 2022;**17**:643–654. <https://doi.org/10.2215/CJN.14231021>
27. Shen L, Jhund PS, Anand IS, Bhatt AS, Desai AS, Maggioni AP, et al. Incidence and outcomes of pneumonia in patients with heart failure. *J Am Coll Cardiol* 2021;**77**:1961–1973. <https://doi.org/10.1016/j.jacc.2021.03.001>
28. Califf RM. Avoiding the coming Tsunami of common, chronic disease: What the lessons of the COVID-19 pandemic can teach us. *Circulation* 2021;**143**:1831–1834. <https://doi.org/10.1161/CIRCULATIONAHA.121.053461>