Heartbeat: interaction of renin–angiotensin–aldosterone blocking drugs with COVID-19 disease susceptibility and severity

doi:10.1136/heartjnl-2020-318164

Catherine M Otto

Angiotensin converting enzyme 2 (ACE2) is the primary receptor for the SARS-CoV-2 virus raising concern that patients with cardiovascular disease being treated with an ACE inhibitor or angiotensin receptor blocker (ARB) drug might have altered ACE2 expression leading to increased risk of COVID-19 infection or a more severe disease course. In this issue of *Heart*, Hippisley-Cox and colleagues¹ addressed this question using routinely collected data in a prospective cohort study in England of 8.28 million participants aged 20–99 years. COVID-19 disease was diagnosed in 19 486 patients and 1286 required treatment in the intensive care unit (ICU). Even after adjustment for confounders, ACE inhibitors were associated with a significantly reduced risk of COVID-19 disease (adjusted HR 0.71, 95% CI 0.67 to 0.74) and no increased risk of ICU care (adjusted HR 0.89, 95% CI 0.75 to 1.06). Similarly, adjusted HRs for ARBs were 0.63 (95% CI 0.59 to 0.67) for COVID-19 disease and 1.02 (95% CI 0.83 to 1.25) for ICU care. However, there were significant interactions with ethnicity for ACE inhibitors and ARB (both $p < 0.001$) for COVID-19 RT-PCR diagnosed disease outcome (table 1).

In the accompanying editorial, Straw and Witte² point out that ‘Observational data are challenging to work with, requiring care to avoid the pitfalls of observed and hidden interactions and biases while balancing the need to make the results accessible to non-statisticians. The reliability of conclusions drawn from observational data depends on their interpretation and on the data themselves.’ However, given the rigorous approach in this study, they that this data ‘should reassure physicians and patients that these vital medications are not implicated in COVID-19 and should not be stopped.’ This study also underscores the principles ‘that the cardiovascular system is not the same across ethnicities, that good supportive care in the absence of specific therapies is essential and that the independent scientific process based on precise peer review must be protected, even when authors and editors are under immense pressure.’

Correspondence to Professor Catherine M Otto, Division of Cardiology, University of Washington, Seattle, WA 98195, USA; cmotto@uw.edu

Table 1 Adjusted HRs (95% CI) for risk of COVID-19 positive test associated with ACE inhibitor and ARB exposure by ethnic group

	ACE inhibitor		ARB	
	Adjusted HR (95% CI)	P value	Adjusted HR (95% CI)	P value
White	0.66 (0.63 to 0.70)	<0.001	0.56 (0.52 to 0.62)	<0.001
Indian	0.74 (0.61 to 0.90)	0.003	0.66 (0.52 to 0.82)	<0.001
Pakistani	0.83 (0.64 to 1.09)	0.182	0.78 (0.57 to 1.06)	0.114
Bangladeshi	0.97 (0.72 to 1.31)	0.847	0.74 (0.49 to 1.13)	0.164
Other Asian	0.81 (0.64 to 1.03)	0.084	0.96 (0.73 to 1.23)	0.726
Caribbean	1.05 (0.87 to 1.28)	0.480	0.70 (0.53 to 0.92)	0.010
Black African	1.31 (1.08 to 1.59)	0.005	1.24 (0.99 to 1.58)	0.062
Chinese	0.73 (0.30 to 1.79)	0.575	1.53 (0.77 to 3.01)	0.223
Other ethnic group	0.82 (0.67 to 1.05)	0.122	1.09 (0.86 to 1.39)	0.475

HRs are comparing risks of COVID-19 in users versus non-users of ACE inhibitor and ARB, and are adjusted for age, sex, deprivation, geographical region, comorbidities (including hypertension included as a binary variable) and other medications listed in table 1 of Hippisley-Cox and colleagues.¹

ARB, angiotensin receptor blocker.

Similarly, a meta-analysis of 10 studies enrolling 9890 hypertensive subjects found that the risk of severe or lethal COVID-19 was no increased in patients on treatment with an ACE inhibitor or ARB compared with patients not on these medications (for

ACE inhibitors OR: 0.90; 95% CI 0.65 to 1.26; for ARBs OR 0.92; 95% CI 0.75 to 1.12).³

These two studies in *Heart* complement the randomised clinical trial (RCT) recently presented at the European Society

Figure 1 Forest plots of ORs of cardiovascular disease (CVD) (International Classification of Diseases, Tenth Revision definition) in homeless versus housed individuals from (A) all studies in meta-analysis (n=9), (B) North American studies (n=6) and (C) European studies (n=3). (D) Forest plot of ORs of hypertension (a subset of CVD) (n=2).

of Cardiology Annual Scientific Sessions. In the Angiotensin Receptor Blockers and Angiotensin-converting Enzyme Inhibitors and Adverse Outcomes in Patients With COVID-19 (BRACE-CORONA) trial,⁴ 659 patients hospitalized with COVID-19 infection who were chronically on an ACE inhibitor or ARB were randomised to 30-days of suspension vs continuation of these medications. The mean age was relatively young (56 years) and 40% were women, with hypertension in 100% but heart failure in only 1.4%. Overall, the primary outcome of days alive and out of the hospital at 30 days was no different in those suspending or continuing ACE-inhibitor or ARB therapy. The congruence between this RCT and the publications in this issue of *Heart* shows the power of carefully performed and analyzed observational studies.

Homelessness is an often-underappreciated risk factor for cardiovascular disease (CVD). 'Homelessness is a global crisis which has increased at an alarming rate over the last decade; worldwide it is estimated that nearly 1.6 billion people have inadequate shelter, resulting in the United Nations announcing its first-ever resolution on homelessness on 19 February 2020.'⁵ In a meta-analysis of 17 articles addressing both the risk of CVD and interventions for CVD in homeless

Figure 2 Integrating fractional myocardial mass with FFR_{CT} analysis. A conceptual image of how integrating FFR_{CT} with myocardium at risk and myocardial blood flow could be applied in future iterations of CTCA analysis. CTCA, CT coronary angiography; FFR, fractional flow reserve; FFR_{CT} , CT derived FFR; LAD, left anterior descending.

populations, Al-Shakarchi and colleagues⁶ found an almost threefold increased risk of CVD in homeless individuals pooled (OR 2.96; 95% CI 2.80 to 3.13; $p < 0.0001$) and a higher CVD mortality rate than non-homeless individuals (age-standardised mortality ratio range: 2.6–6.4) (figure 1).

As Lane concludes in her editorial,⁵ 'We need to gain a better understanding of the magnitude and types of CVD among the

homeless; the specific barriers to health-care faced by homeless people; and identify effective models of care and adapt/tailor these into current CVD services and existing homeless services.' 'One important tangible difference healthcare professionals can implement immediately is to treat everyone with the same level of professionalism and compassion and not discriminate or stigmatise people who are homeless.'

An interesting imaging study by Keulards and colleagues⁷ demonstrates that CT flow-based measurement of myocardial mass is closely related to invasively measured myocardial perfusion supporting the use of CT-mass for estimating the extent of myocardium-at-risk in patients with proximal coronary artery disease. The novelty of this approach is put into context by Ihdahid and Sellers⁸ who discuss how fractional flow reserve measured by CT (FFR_{CT}) techniques might allow us to 'integrate both physiology and myocardium at risk into a single non-invasive test' (figure 2).

The *Education in Heart* article in this issue⁹ provides an overview of interventional approaches to treatment of chronic thromboembolic pulmonary hypertension (CTEPH), comparing surgical pulmonary endarterectomy (PEA) versus balloon pulmonary angioplasty (BPA) (figure 3). As they point out 'BPA is now sufficiently developed that appropriately powered RCTs comparing PEA versus BPA for segmental and subsegmental (levels 3 and 4) CTEPH could be planned in centres with equivalent experience in both techniques. However, consensus in the interpretation of imaging and standardisation of the surgical and BPA approach between

Figure 3 Chronic thromboembolic pulmonary hypertension (CTEPH) treatment algorithm.¹⁰ #Multidisciplinary: pulmonary endarterectomy surgeon, pulmonary hypertension (PH) expert, BPA interventionist and radiologist. ¶Treatment assessment may differ depending on the level of expertise. +BPA without medical therapy can be considered in selected cases. BPA, balloon pulmonary angioplasty.

centres will be important to deliver meaningful comparisons.’

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not required.

Provenance and peer review Commissioned; internally peer reviewed.

© Author(s) (or their employer(s)) 2020. No commercial re-use. See rights and permissions. Published by BMJ.

To cite Otto CM. *Heart* 2020;**106**:1451–1453.

Heart 2020;**106**:1451–1453.
doi:10.1136/heartjnl-2020-318164

ORCID iD

Catherine M Otto <http://orcid.org/0000-0002-0527-9392>

REFERENCES

- Hippisley-Cox J, Young D, Coupland C, *et al*. Risk of severe COVID-19 disease with ACE inhibitors and angiotensin receptor blockers: cohort study including 8.3 million people. *Heart* 2020;106:1503–11.
- Straw S, Witte KK. Observational data during the COVID-19 pandemic: opportunity with uncertainty. *Heart* 2020;106:1461–2.
- Flacco ME, Acuti Martellucci C, Bravi F, *et al*. Treatment with ACE inhibitors or ARBs and risk of severe/lethal COVID-19: a meta-analysis. *Heart* 2020;106:1519–24.
- Lopes RD on behalf of the BRACE CORONA Investigators. Continuing versus suspending ACE inhibitors and ARBs: Impact of adverse outcomes in hospitalized patients with COVID-19-The BRACE CORONA Trial. Presented at ESC Scientific Sessions; Sept 1, 2020.
- Lane DA. Homelessness: a ‘cause of the causes’ of cardiovascular disease? *Heart* 2020;106:1456–7.
- Al-Shakarchi NJ, Evans H, Luchenski SA, *et al*. Cardiovascular disease in homeless versus housed individuals: a systematic review of observational and interventional studies. *Heart* 2020;106:1483–8.
- Keulards DCJ, Fournier S, van 't Veer M, *et al*. Computed tomographic myocardial mass compared with invasive myocardial perfusion measurement. *Heart* 2020;106:1489–94.
- Ihdayhid AR, Sellers SL. Novel method for assessing myocardium at risk: a new arrow in the diagnostic quiver of coronary CT. *Heart* 2020;106:1458–60.
- Hoole SP, Jenkins DP. Chronic thromboembolic pulmonary hypertension: interventional approaches. *Heart* 2020;106:1525–31.
- Kim NH, Delcroix M, Jais X, *et al*. Chronic thromboembolic pulmonary hypertension. *Eur Respir J* 2019;53:1801915.