Heartbeat: Social isolation is associated with increased mortality after acute myocardial infarction or stroke

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Loneliness and social isolation are associated with an increased risk of premature mortality, yet the specific diseases contributing to these adverse outcomes have not been fully elucidated. In 2016, *Heart* published a meta-analysis that showed 'Poor social relationships were associated with a 29% increase in risk of incident coronary heart disease (pooled relative risk: 1.29, 95% CI 1.04 to 1.59) and a 32% increase in risk of stroke (pooled relative risk: 1.32, 95% CI 1.04 to 1.68).'¹ This paper and the linked editorial² received some of the highest Altmetric scores for all papers published in *Heart*, reflecting the importance of this topic to the general public and healthcare systems. However, while a meta-analysis is a powerful approach to combining data, published studies do not always provide enough detail about other factors that might affect these findings and thus is difficult to translate into patient care or public policy.

In this issue, the association of loneliness and social isolation with cardiovascular disease (CVD) risk is examined in further detail in a UK Biobank cohort of over 479 000 men and women.³ Although unadjusted data suggested that social isolation and loneliness were associated with a significant higher risk of acute myocardial infarction (AMI) and stroke, the detailed information available in the data set allowed adjustment for other known cardiovascular risk factors. In fact, after correction for other risk factors, these associations were significantly attenuated and were no longer statistically significant for the overall cohort. The relative importance of adjustment for health behaviours, depressive symptoms, socioeconomic and other risk factors is shown in figure 1. However, in the subgroup of patients who suffered an AMI or stroke, the fully adjusted model did show that social isolation, but not loneliness, was associated with increased mortality with HR 1.25 (95% CI 1.03 to 1.51) for those with an AMI and 1.32

Figure 1 Entire Biobank cohort (n=479 054). Proportions of the social isolation—AMI and stroke excess risk mediated by biological, behavioural, socioeconomic and health-related factors. AMI, acute myocardial infarction; PERM, percentage of excess risk mediated.

(95% CI 1.08 to 1.61) for those with a stroke (figure 2).

This important study demonstrates the additive CVD risk associated with social isolation after AMI or stroke and thus identifies a group of patients where efforts to reduce social isolation would be likely to have the most effect. The attenuation of risk with adjustment for other known risk factors also emphasises the key role of these linked factors,

particularly patient health behaviours, in reducing CVD risk.

Detection of asymptomatic atrial fibrillation (AF) is an increasing public health concern given the increased prevalence of AF in our ageing population and the stroke risk associated with this cardiac rhythm. Various screening approaches have been proposed in papers previously published in *Heart*, including use of smartphone ECGs

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Figure 2 Participants with incident AMI or stroke (n=9202). Proportions of the social isolation—mortality after AMI or stroke event excess risk mediated by biological, behavioural, socioeconomic and health-related factors. AMI, acute myocardial infarction; PERM, percentage of excess risk mediated.

Figure 3 Contribution of risk factors to total Screening for Asymptomatic Atrial Fibrillation Events (SAAFE) risk score. The variables are age, CHF, CAD, H×W, recent hospitalisation for cardiopulmonary conditions (Hosp), cardiac arrest (SCD), chronic obstructive pulmonary disease (COPD), height and KT×. Note that the first three variables have the greatest contribution to the total risk score. CAD, coronary artery disease; CHF, congestive heart failure; COPD, chronic obstructive pulmonary disease; H×W, height times weight; KT×, kidney transplant; SCD, sudden cardiac death.

in the general population⁴ and handheld ECG devices in hospitalised geriatric patients,⁵ to name just a few.⁶ Linker and colleagues⁷ propose a new risk score based on clinical variables, without an ECG, to identify patients at risk of AF development. The clinical risk factors included in the Screening for Asymptomatic Atrial Fibrillation Events risk score and the contribution of each to the overall model are shown in figure 3. The risk of AF increased from a low of 2% in those with a score of 2.0 or less to a high of 66% in those with a score >4.5. Overall, the model identified 81% of patients with incident AF over 3 years.

In an editorial, Bisson and colleagues⁸ agree that lower cost screening methods for AF are needed but also suggest, 'Another strategy might be to propose early oral anticoagulation in patients with very high risk of AF and of embolic ischaemic stroke, before AF itself is documented. Further knowledge on implication of multiple risk factors in patients without AF is thus warranted and might influence future strategies in stroke prevention (figure 4). The major challenge will be to find a cost-effective prophylaxis with a favourable risk–benefit ratio, which will require randomised trials in such high-risk patients.'

The *Education in Heart* article⁹ summarises the investigation and management of adults with hypertension. Hypertension treatment might seem mundane but effective blood pressure control is essential for prevention of CVD and stroke. Identification of patients with secondary hypertension allows intervention to correct the underlying cause but diagnostic testing is indicated only when clinical characteristics suggest this diagnosis. Recommendations for initiation of therapy and long-term management have been developed by several professional organisation and are summarised in a concise readable format in this article (figure 5).

The *Image Challenge* asks you to interpret the ECG in a patient with a recently implanted pacer.¹² The ECG seems straightforward at first glance but there is more than meets the eye.

Competing interests None declared.

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Figure 4 Possible strategies to be proposed or evaluated in the future when using screening tools identifying patients at risk for AF in clinical practice. AF, atrial fibrillation; CHA₂DS₂-VASc, congestive heart failure, hypertension, age (≥75 years; 2 points), diabetes, stroke/transient ischaemic attack (2 points), vascular disease, age (65–74 years), sex category (female); OAC, oral anticoagulation; RCT, randomised controlled trial.

Figure 5 Flow diagram for selection of class of antihypertensive pharmacotherapy (Adapted from BHS/NICE guidelines [10] and incorporating additional evidence from Williams *et al* [11]). However, see online supplementary table 2 for the rationale against an all-purpose ranking of drugs. ACEi, ACE inhibitor; ARB, angiotensin receptor blocker; BHS, British Hypertension Society; CCB, calcium channel blocker; NICE, National Institute for Health and Care Excellence.

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