

Resting heart rate and the risk of death and cardiovascular complications in patients with type 2 diabetes mellitus

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Abstract

Aims/hypothesis An association between resting heart rate and mortality has been described in the general population and in patients with cardiovascular disease. There are, however, few data exploring this relationship in patients with type 2 diabetes mellitus. The current study addresses this issue.

Methods The relationship between baseline resting heart rate and all-cause mortality, cardiovascular death and major cardiovascular events (cardiovascular death, non-fatal myocardial infarction or non-fatal stroke) was examined in

11,140 patients who participated in the Action in Diabetes and Vascular Disease: Preterax and Diamicron Modified Release Controlled Evaluation (ADVANCE) Study.

Results A higher resting heart rate was associated with a significantly increased risk of all-cause mortality (fully adjusted HR 1.15 per 10 bpm [95% CI 1.08, 1.21], $p < 0.001$), cardiovascular death and major cardiovascular outcomes without adjustment and after adjusting for age and sex and multiple covariates. The increased risk associated with a higher baseline resting heart rate was most obvious in patients with previous macrovascular complications (fully adjusted HR for death 1.79 for upper [mean 91 bpm] vs lowest [mean 58 bpm] fifth of resting heart rate in this subgroup [95% CI 1.28, 2.50], $p = 0.001$).

Conclusions/interpretation Among patients with type 2 diabetes, a higher resting heart rate is associated with an increased risk of death and cardiovascular complications. It remains unclear whether a higher heart rate directly mediates the increased risk or is a marker for other factors that determine a poor outcome.

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Keywords Heart rate · Outcome · Type 2 diabetes mellitus

Abbreviations

ADVANCE Action in Diabetes and Vascular Disease:
Preterax and Diamicron Modified Release
Controlled Evaluation

bpm Beats per minute

Several studies have demonstrated an association between resting heart rate and all-cause and cardiovascular mortality [1, 2]. This relationship has been described both in the general population and in patients with cardiac disease. In particular, an increased heart rate predicts an unfavourable

outcome in patients with heart failure and following an acute coronary syndrome [1]. There are, however, few data specifically exploring the relationship between resting heart rate and cardiovascular outcome in patients with diabetes mellitus. This is surprising given the accessibility of the measure, the strength of the observed relationship in other populations, the high and increasing prevalence of diabetes and the association between heart rate and glucose and insulin levels. In addition, diabetes mellitus is commonly associated with abnormal function of the autonomic nervous system, the main regulator of resting heart rate. The aim of the current study is, therefore, to explore the relationship between resting heart rate at baseline and cardiovascular events and all-cause mortality in a large cohort of patients with type 2 diabetes mellitus.

Methods

The Action in Diabetes and Vascular Disease: Preterax and Diamicon Modified Release Controlled Evaluation (ADVANCE) Study (ClinicalTrials.gov number NCT00145925) was a factorial trial that recruited 11,140 patients with type 2 diabetes mellitus from 215 centres in 20 countries between June 2001 and March 2003 [3, 4]. The study made two randomised comparisons: a double-blind assessment of the efficacy of a fixed combination of perindopril–indapamide (2 mg/0.625 mg for 3 months, increasing if tolerated to 4 mg/1.25 mg thereafter) vs placebo and an open-label evaluation of an intensive glucose-lowering regimen based on gliclazide modified release targeting an HbA_{1c} of $\leq 6.5\%$ (48 mmol/mol) vs standard guideline-based glycaemic control. The study was approved by the local ethics committee for each participating centre and all participants provided written informed consent. Detailed study methods [5] and the main results [3, 4] of the ADVANCE Study have been previously reported. Briefly, participants in the ADVANCE Study were all at least 55 years of age and had been diagnosed with type 2 diabetes after the age of 30 years. In addition, they were required to have had at least one major macro- or microvascular complication of diabetes or one or more additional cardiovascular risk factors.

Baseline data included estimates of the duration of diabetes, indices of diabetic control and detailed information regarding lifestyle, including self-reported estimates of average weekly exercise. Baseline resting heart rate was measured at the same time as the blood pressure using a digital monitor (Omron HEM-705CP; Omron Healthcare, Lake Forest, IL, USA). Recordings were made after the patient had been resting for a minimum of 5 min in the seated position. Three readings were taken and the mean of the latter two was used. Weight, height, urinary albumin/creatinine ratio, serum creatinine, fasting lipid levels and HbA_{1c} were also measured at baseline.

The current study assesses the relationship between baseline resting heart rate and all-cause mortality, cardiovascular

death and major cardiovascular events (a composite of cardiovascular death, non-fatal myocardial infarction or non-fatal stroke; a primary outcome of the ADVANCE Study) over the entire length of the trial. All outcomes were validated by an independent adjudication committee who were unaware of treatment allocation.

Statistical analyses Baseline resting heart rate was considered both as a continuous variable and after grouping into approximately equal fifths according to its quintiles. Categorical data were summarised using percentages and compared using χ^2 tests. Normally distributed continuous data were summarised as mean (SD) and comparisons made using the *t* test or a general linear model to determine any difference across fifths. Skewed data are presented as median (inter quartile range) and compared using the Kruskal–Wallis test.

Cox regression analyses were used to calculate the HR with 95% CIs. These were performed without adjustment and in models adjusted for age and sex and for multiple covariates (age, sex, ADVANCE Study blood pressure treatment arm, ADVANCE Study glycaemic control arm, body mass index, duration of diabetes, HbA_{1c}, urinary albumin/creatinine ratio, estimated glomerular filtration rate, systolic blood pressure, diastolic blood pressure, history of hospitalisation for heart failure, participation in moderate and/or vigorous exercise for >15 min at least once weekly, total cholesterol, triacylglycerol level, atrial fibrillation, treatment with calcium channel blockers and treatment with β -blockers). Subgroup analyses were performed assessing outcome in patients who had and had not experienced a macrovascular complication prior to randomisation in the ADVANCE Study, according to treatment with β -blockers at the time the heart rate was recorded and stratified by the presence or absence of atrial fibrillation (on an ECG performed at baseline or previously [6]). The linearity of the relationship between resting heart rate and outcome was assessed through likelihood tests by fitting the fifths of resting heart rate as an ordinal variable and residual non-linearity was tested by comparison with fitting the fifths as a categorical variable [7]. Competing risk analyses were performed to determine whether a higher heart rate conferred an increased risk of all-cause mortality, compared with other outcomes [8]. Analyses were performed using SPSS v18 (SPSS, Chicago, IL, USA) and Stata 11 (StataCorp LP, College Station, TX, USA) and all significance levels are two-sided.

Results

Of the 11,140 patients randomised into the ADVANCE Study, two individuals did not have their baseline resting

heart rate recorded. The remaining 11,138 patients are included in the current analyses. Their mean heart rate was 74 (SD 12) beats per minute (bpm), with a range of 33 to 140 bpm. Baseline average resting heart rates were slightly lower in men (73 [SD 12] bpm) than women (75 [SD 11] bpm, $p < 0.001$) and in older patients (73 [SD 12] bpm in patients ≥ 65 years old vs 75 [SD 12] bpm in those < 65 years of age, $p < 0.001$). Table 1 shows the relationship between heart rate fifth and a variety of clinical variables.

During a median follow-up of 4.4 years, 879 (8%) patients died, 468 (4%) from cardiovascular causes. One thousand patients (9%) suffered a major cardiovascular event during

follow-up. A higher resting heart rate was associated with a greater hazard of death (all-cause and cardiovascular) and major cardiovascular events (Fig. 1). Adjustment for use of other cardiovascular medications (statins, angiotensin-converting enzyme inhibitors or angiotensin receptor blockers), alternate measures of renal function (creatinine or creatinine clearance), lipid fractions (LDL-cholesterol and HDL-cholesterol levels) or other measures of average weekly exercise (participation in vigorous exercise > 15 min per week or number of exercise sessions > 15 min per week) made minimal difference to the observed hazard associated with a higher heart rate (data not shown). The hazard associated with a faster resting heart rate was

Table 1 Characteristics of the study population by baseline resting heart rate fifths

Characteristic	All patients	Fifth 1 (lowest)	Fifth 2	Fifth 3	Fifth 4	Fifth 5 (highest)	<i>p</i> value for trend
<i>n</i>	11,138	2,066	2,393	2,178	2,195	2,306	–
Mean heart rate (bpm)	74	57	67	74	80	91	–
Heart beat range (bpm)	33–140	33–63	64–70	71–76	77–83	84–140	–
Age (years)	65.8±6.4	67.1±6.3	65.9±6.4	65.5±6.5	65.5±6.4	65.1±6.2	<0.001
Female sex, <i>n</i> (%)	4,733 (42)	664 (32)	989 (41)	973 (45)	1,050 (48)	1,057 (46)	<0.001
Active blood pressure treatment arm, <i>n</i> (%)	5,568 (50)	1,024 (50)	1,195 (50)	1,072 (49)	1,127 (51)	1,150 (50)	0.56
Active glycaemic control treatment arm, <i>n</i> (%)	5,570 (50)	1,007 (49)	1,216 (51)	1,050 (48)	1,098 (50)	1,199 (52)	0.09
Systolic blood pressure (mmHg)	145±22	147±22	144±21	143±21	145±21	146±21	0.11
Diastolic blood pressure (mmHg)	81±11	79±11	80±11	81±11	81±11	82±11	<0.001
Body mass index (kg/m ²)	28.3±5.2	28.8±5.0	28.5±5.0	28.2±5.3	28.2±5.2	28.2±5.3	<0.001
Smoker, <i>n</i> (%)	1,550 (14)	234 (11)	314 (13)	333 (15)	324 (15)	345 (15)	<0.001
Regular exercise ^a , <i>n</i> (%)	5,112 (46)	1,029 (50)	1,149 (48)	994 (46)	970 (44)	970 (42)	<0.001
History of MI, <i>n</i> (%)	1,334 (12)	392 (19)	316 (13)	251 (12)	205 (9)	170 (7)	<0.001
History of stroke, <i>n</i> (%)	1,022 (9)	168 (8)	197 (8)	198 (9)	217 (10)	242 (10)	0.001
History of heart failure ^b , <i>n</i> (%)	356 (3)	66 (3)	71 (3)	66 (3)	71 (3)	82 (4)	0.38
Atrial fibrillation, <i>n</i> (%)	846 (8)	199 (10)	154 (6)	128 (6)	173 (8)	192 (8)	0.68
Serum total cholesterol (mmol/l)	5.20±1.19	4.94±1.08	5.14±1.17	5.21±1.20	5.30±1.18	5.37±1.27	<0.001
Serum triacylglycerols (mmol/l)	1.96±1.29	1.84±1.14	1.88±1.15	1.91±1.21	2.04±1.35	2.11±1.52	<0.001
Duration of diabetes (years)	7.9±6.4	7.7±6.6	7.6±6.3	8.0±6.5	8.2±6.2	8.2±6.2	0.001
HbA _{1c} (%)	7.5±1.6	7.2±1.3	7.3±1.4	7.5±1.6	7.7±1.6	7.8±1.8	<0.001
HbA _{1c} (mmol/mol)	59±17	55±15	57±16	59±17	60±17	62±19	<0.001
Creatinine (μmol/l)	87±25	91±23	87±24	85±24	84±27	86±28	<0.001
Estimated GFR (ml min ⁻¹ [1.73 m] ⁻²)	78±25	75±20	78±22	79±27	80±28	79±26	<0.001
Urinary ACR (mg/mmol) ^c	1.70 (0.80, 4.50)	1.42 (0.68, 3.61)	1.60 (0.74, 4.10)	1.58 (0.79, 3.90)	1.90 (0.90, 5.17)	2.00 (0.90, 5.50)	<0.001
β-Blocker, <i>n</i> (%)	2,729 (25)	938 (45)	682 (28)	439 (20)	365 (17)	305 (13)	<0.001
Calcium channel antagonist, <i>n</i> (%)	3,426 (31)	666 (32)	706 (30)	638 (29)	647 (29)	769 (33)	0.38

Data are mean±SD, except where indicated otherwise. GFR was estimated using the Modification of Diet in Renal Disease formula ACR, albumin/creatinine ratio; MI, myocardial infarction

^a Defined as moderate and/or vigorous exercise for > 15 min at least once per week (self-reported)

^b Requiring hospitalisation

^c Median (interquartile range)

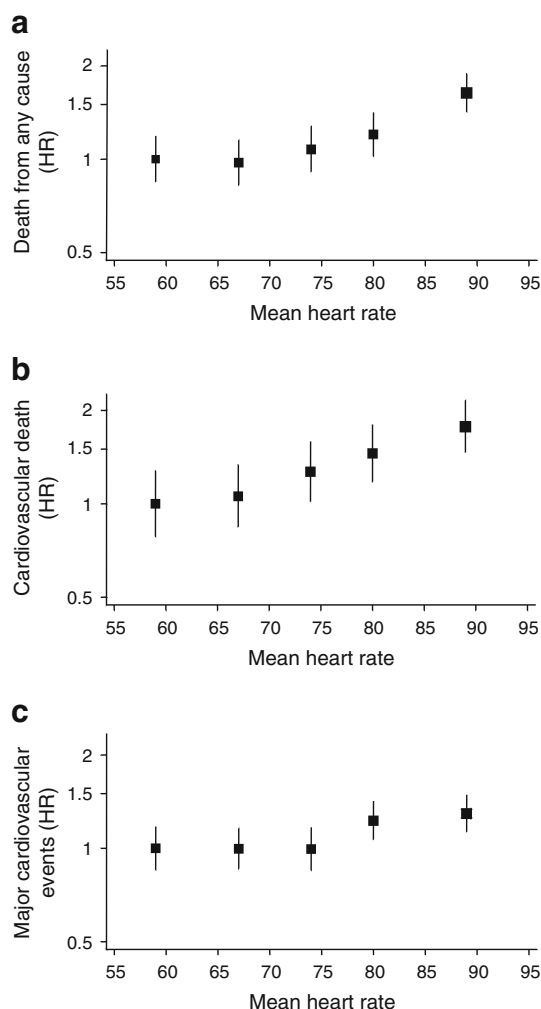


Fig. 1 Baseline resting heart rate (by fifths) and the risk of death from any cause, cardiovascular death and major cardiovascular events. Data were hazard adjusted for age, sex, ADVANCE Study blood pressure treatment arm, ADVANCE Study glycaemic control arm, body mass index, duration of diabetes, HbA_{1c}, urinary albumin/creatinine ratio, estimated glomerular filtration rate, systolic blood pressure, diastolic blood pressure, history of hospitalisation for heart failure, participation in moderate and/or vigorous exercise for >15 min at least once weekly, total cholesterol, triacylglycerol level, atrial fibrillation, treatment with calcium channel blockers and treatment with β -blockers. **a** Death from any cause ($p<0.001$ for trend); **b** cardiovascular death ($p<0.001$ for trend); **c** major cardiovascular events ($p=0.002$ for trend)

also similar in unadjusted and age- and sex-adjusted models (see electronic supplementary material Figs 1–3).

The relationship between baseline heart rate and each outcome was linear ($p<0.001$ for death, $p=0.003$ for cardiovascular death and $p=0.05$ for major cardiovascular events) with no evidence of non-linearity ($p=0.14$ for death, $p=0.96$ for cardiovascular death and $p=0.48$ for major cardiovascular events). After adjustment for multiple covariates the hazard of death increased by 15% (95% CI 8%, 21%; $p<0.001$) for every 10 bpm increase in resting heart rate. Likewise, the hazard of cardiovascular death increased by 16% (95% CI

7%, 25%; $p<0.001$) and for all cardiovascular events by 8% (95% CI 2%, 14%; $p=0.009$). This increased hazard was evident in both male and female patients (fully adjusted HR for death 1.11 per 10 bpm [95% CI 1.04, 1.19], $p=0.002$ in men and fully adjusted HR for death 1.22 per 10 bpm, [95% CI 1.10, 1.36], $p<0.001$ in women). In competing risk models (using all-cause mortality as the competing event) the sub-distribution hazard of cardiovascular death increased by 15% (95% CI 7%, 25%; $p<0.001$) and for all major cardiovascular events by 7% (95% CI 1%, 13%; $p=0.01$).

The adverse prognosis associated with a higher heart rate was most evident in diabetic patients who had had a previous macrovascular event (myocardial infarction, stroke, hospital admission for a transient ischaemic attack, hospital admission for unstable angina, coronary revascularisation, peripheral revascularisation or amputation secondary to peripheral vascular disease; Fig. 2). In this high-risk subgroup the fully adjusted HR for death was 1.79 (95% CI 1.28, 2.50; $p=0.001$) for patients in the upper fifth of resting heart rate (mean 91 bpm) when compared with those in the lowest fifth (mean 57 bpm). Likewise, patients with a history of coronary heart disease (myocardial infarction, hospital admission for unstable angina or coronary revascularisation) who had a faster resting heart rate were at particularly high risk (fully adjusted HR for death 1.23 per 10 bpm [95% CI 1.10, 1.37], $p<0.001$ in patients with prior coronary heart disease and fully adjusted HR 1.13 per 10 bpm [95% CI 1.06, 1.21], $p<0.001$ in patients without any known coronary heart disease).

The relationship between baseline heart rate and death was similar regardless of therapy with β -blockers (p for interaction between β -blocker use and heart rate=0.57; Fig. 3). Eight hundred and forty-six patients had current or previous, ECG-confirmed, atrial fibrillation. Of these patients, 431 were in atrial fibrillation on the ECG performed at their baseline visit. The relatively small number of patients with current or prior atrial fibrillation limits the power to detect any association between heart rate and outcome in this subgroup. Therefore, although no interaction between atrial fibrillation and heart rate was detected statistically (p for interaction=0.62; Fig. 3), no definitive conclusions can be drawn in this population.

Discussion

The current data confirm that in patients with type 2 diabetes mellitus a higher resting heart rate is a predictor of all-cause mortality, cardiovascular death and major cardiovascular events. This association remains after adjustment for a variety of other factors that may influence heart rate and impact on outcome in this setting. Indeed, the strength of the relationship was stronger in these adjusted models. It is more evident among diabetic patients who have

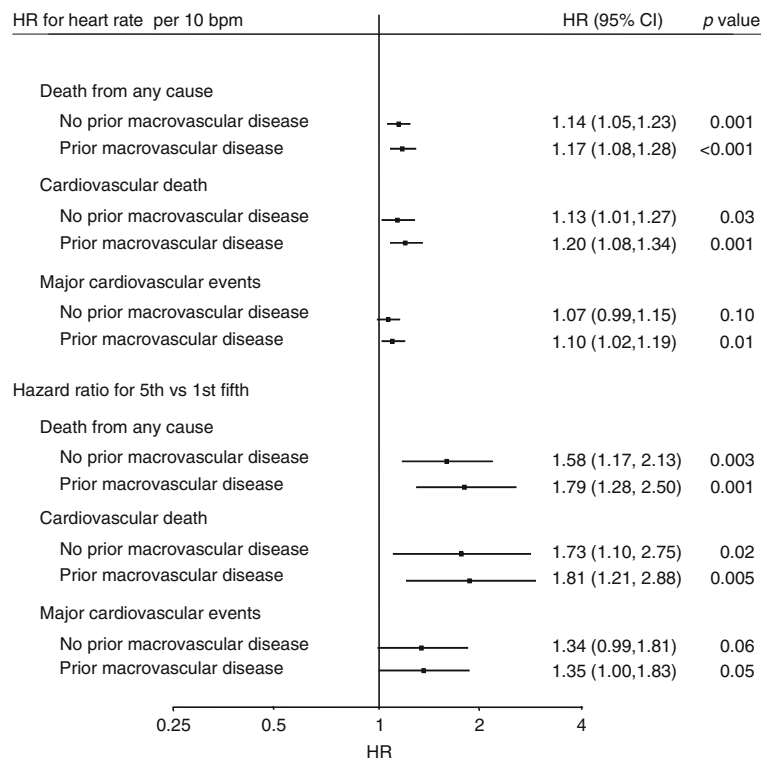


Fig. 2 Baseline resting heart rate and the risk of death from any cause, cardiovascular death and major cardiovascular events stratified by a history of prior macrovascular complications. Macrovascular disease: myocardial infarction, stroke, hospital admission for a transient ischaemic attack, hospital admission for unstable angina, coronary revascularisation, peripheral revascularisation or amputation secondary to peripheral vascular disease. Data were hazard adjusted for age, sex, ADVANCE Study blood pressure treatment arm, ADVANCE Study

glycaemic control arm, body mass index, duration of diabetes, HbA_{1c}, urinary albumin/creatinine ratio, estimated glomerular filtration rate, systolic blood pressure, diastolic blood pressure, history of hospitalisation for heart failure, participation in moderate and/or vigorous exercise for >15 min at least once weekly, total cholesterol, triacylglycerol level, atrial fibrillation, treatment with calcium channel blockers and treatment with β -blockers

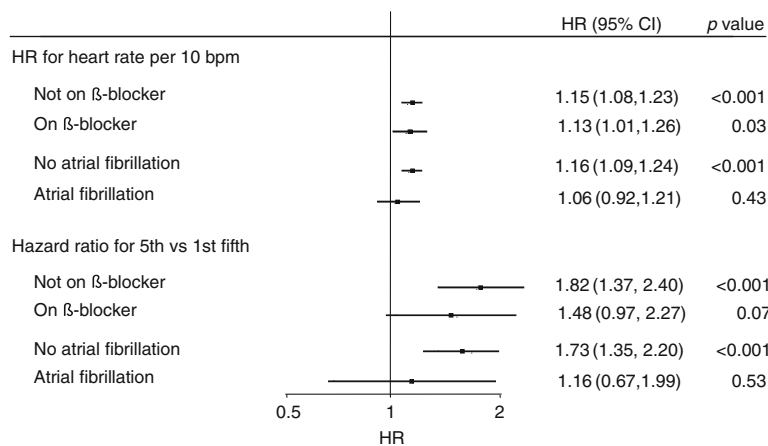


Fig. 3 Baseline resting heart rate and the risk of death from any cause stratified by treatment with β -blockers and by atrial fibrillation. Data were hazard adjusted for age, sex, ADVANCE Study blood pressure treatment arm, ADVANCE Study glycaemic control arm, body mass index, duration of diabetes, HbA_{1c}, urinary albumin/creatinine ratio, estimated glomerular filtration rate, systolic blood pressure, diastolic

blood pressure, history of hospitalisation for heart failure, participation in moderate and/or vigorous exercise for >15 min at least once weekly, total cholesterol, triacylglycerol level, atrial fibrillation (only in analysis stratified by treatment with β -blockers), treatment with calcium channel blockers and treatment with β -blockers (only in analysis stratified by history of atrial fibrillation)

had a prior vascular event, is present in both men and women, and is consistent regardless of treatment with β -blockers.

Comparison with prior studies There are few data assessing the prognostic importance of resting heart rate in patients with type 2 diabetes mellitus. Long-term follow-up of 302 such patients recruited into the Swiss cohort of the World Health Organization Multinational Study of Vascular Disease in Diabetes did, however, report a relationship between resting heart rate and both all-cause and cardiovascular mortality [9]. This relationship was maintained after accounting for a variety of potentially confounding factors, although in that population no adjustments were made for the use of β -blockers, participation in regular exercise or HbA_{1c}. Interestingly, a similar relationship was not observed in patients with type 1 diabetes mellitus [9].

Recently, the Euro Heart Survey Investigators have assessed the relationship between heart rate and 1 year outcome in 751 diabetic patients and 1,756 non-diabetic individuals, all with stable coronary heart disease [10]. The main finding of this study was that in the diabetic patients a higher resting heart rate was associated with an increased risk of all-cause mortality. This relationship persisted after adjustment for a range of clinical factors although there were only 37 deaths in the diabetic cohort, limiting the ability to accurately correct for potential confounders. Likewise, that study did not account for the type or duration of diabetes, nor any measure of diabetic control or microvascular disease.

In the Euro Heart Survey cohort, there was no significant relationship between resting heart rate and a composite outcome of death, myocardial infarction and stroke [10]. The absence of an association when non-fatal cardiovascular events were considered in addition to all-cause death might imply that the link between higher heart rate and death was driven primarily by non-cardiovascular events. Our data, which address this issue in a much larger cohort followed for considerably longer, suggest that this is not the case. This is consistent with the data in healthy men [11] and in patients with coronary heart disease [12]. It also concurs with the data from Switzerland [9] and with two other small studies that, although focussing on ECG-derived QT interval variables, also reported an association between resting heart rate and cardiovascular outcome in patients with diabetes [13, 14]. An alternate explanation for the Euro Heart Survey findings is that resting heart rate is more closely associated with fatal than non-fatal cardiovascular events. This has been reported in other cohorts [15] but is not evident in the current study.

The nature of any relationship between resting heart rate and cardiovascular events remains to be fully defined. Several studies have reported a linear relationship [16, 17] but a

variety of other patterns have also been described, including J-shaped [18, 19], U-shaped [20] and sigmoid [21] associations. In the current cohort the relationship appears to be predominantly linear but with a suggestion of attenuation at lower heart rates.

Potential mechanisms The key question is whether a high resting heart rate is partly causative of an adverse outcome (and if so to what extent) or whether it is merely a marker of other factors that determine prognosis, in particular concomitant disease and/or autonomic dysfunction. Several mechanisms have been proposed whereby a more rapid heart rate might be directly deleterious, all of which apply at least as much to patients with diabetes as to the general population. These include the detrimental effects of a high heart rate on myocardial oxygen demand and supply [1], long-term adverse effects on the vasculature [22–24] and myocardial function [25] and an increased risk of ventricular dysrhythmia [1, 2] and sudden cardiac death [11]. In addition to direct effects, an increased heart rate is associated with a variety of factors that may themselves adversely affect prognosis, such as poorer general fitness, obesity, high blood pressure and a less favourable lipid profile [26, 27], all of which are strongly associated with type 2 diabetes. As far as possible, the current analyses have adjusted for these potential confounders.

Diabetes may also directly influence resting heart rate. Hyperinsulinaemia and elevated blood glucose levels have both been associated with a higher heart rate [28–30]. Furthermore, dysfunction of the autonomic nervous system is a common complication of diabetes, even in the absence of other overt neuropathic complications [31–33]. Damage to cardiovascular autonomic nerve fibres results in sympathetic dominance, with higher resting heart rate, and has been associated with a greatly increased risk of death [31, 34]. Evidence of cardiac autonomic neuropathy can be detected early after diagnosis in many patients with diabetes [31, 35] and autonomic dysfunction may even pre-date the diagnosis of diabetes itself [36, 37]. In healthy adults recruited in the Atherosclerosis Risk in Communities Study a resting heart rate in the upper quarter was associated with a 60% increased risk of developing diabetes compared with those with heart rates in the lowest quarter [37]. This may just reflect an association between heart rate and physical fitness but it underscores the complex inter-relationship between diabetes and heart rate.

Randomised studies provide some limited evidence that heart rate plays a direct role in outcome. β -Blockers reduce mortality after acute myocardial infarction and in patients with heart failure, and at least some of the benefits appear to be secondary to a reduction in heart rate [38, 39]. It is, however, difficult to exclude other effects of β -blockade.

In contrast, ivabradine selectively reduces heart rate without other direct effects. In the BEAUTIFUL (morBidity–mortality EvAIUaTion of the If inhibitor ivabradine in patients with CAD and left ventricULar dysfunction) study ivabradine was compared with placebo in patients with coronary heart disease and left ventricular systolic dysfunction and no overall benefit was demonstrated, although, in the pre-specified subgroup with a baseline heart rate ≥ 70 bpm, a significant reduction in the secondary endpoint of myocardial infarction or coronary revascularisation was observed [40]. In the SHIFT (Systolic Heart Failure with If Inhibitor) study of patients with stable chronic heart failure, ivabradine reduced the primary endpoint, a composite endpoint of cardiovascular death and hospitalisation owing to heart failure, by 18% (95% CI 10%, 25%) [41], with changes in heart rate appearing to explain most of the observed benefit [42]. Overall, therefore, despite a strong pathophysiological basis, the evidence that heart rate exerts a direct causative role remains limited. Likewise, there is no definitive evidence to support the use of specific rate-limiting medication to improve outcome, except perhaps in patients with stable heart failure.

Strengths and limitations The current study is by far the largest to date to explore the relationship between heart rate and outcome in patients with diabetes. The patient population was drawn from many different countries and is ethnically diverse. It is well characterised and has been carefully followed. Importantly, the number of independently adjudicated outcomes ensures that the study has considerable power and allows statistical adjustment for multiple potentially confounding factors, including several that have not been addressed by prior studies. Nonetheless, it is impossible to fully correct for all potential confounders and, in particular, no specific measures of cardiac autonomic neuropathy were available. Likewise, although we corrected for a prior history of heart failure requiring hospital admission, we were unable to adjust for the presence or severity of any heart failure symptoms at baseline.

Conclusion Among patients with diabetes, a higher heart rate is associated with an increased risk of death and cardiovascular complications. It remains unclear, however, whether a higher heart rate directly mediates the increased risk or whether it is a marker for other characteristics that determine a poor outcome.

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Duality of interest M. Woodward, S. Zoungas, A. Patel, G. Mancina, N.R. Poulter and J. Chalmers have received lecture fees and/or travel expenses from Servier. M. Woodward, S. Zoungas, A. Patel, N.R. Poulter and J. Chalmers have also received grant support from Servier. Otherwise, the authors declare that there is no duality of interest associated with this manuscript.

Contribution statement GSH, MW, AR, GDB, TN and JC were responsible for the conception and design of the study. GSH, MW and QL were involved in the analysis of data and all authors were involved in the interpretation of data. GSH, MW and JC drafted the article and all authors revised the article and gave final approval of the version to be published.

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