To the Editor:

Pulmonary arterial hypertension (PAH) is a rare disease characterised by progressive pulmonary vascular remodelling, resulting in right ventricular failure, which is the principal cause of death [1]. Excessive bodyweight is a well-established independent risk factor for the development of cardiovascular disease and mortality [2]. However, data have emerged for a variety of chronic cardiovascular diseases, particularly heart failure, showing lower long-term mortality in obese patients [3]. This has been termed the “obesity paradox”. It challenges clinical reasoning and the significant healthcare expenditure directed toward weight loss in the general population.

An analysis of the US-based REVEAL (Registry to Evaluate Early and Long-term PAH Disease Management) found obese PAH patients had a reduced risk of death, despite important limitations of this registry [4]; foremost, the inclusion of prevalent patients and disparities in the standard of care available. Additional studies of pulmonary hypertension (PH) have shown obesity carries a survival benefit across a number of patient populations including incident patients [5-10].

Explanations for this paradox include obese patients having greater metabolic reserve to meet the increased catabolic burden, oxidative stress and systemic inflammation associated with heart failure [11]. However, this is a contentious area and many studies supporting the hypothesis are only observational. They may miss the effect of illness-induced weight-loss. Obese patients may also present earlier and be treated more appropriately earlier in time [12]. Indeed, a large recent study conducted in the French reference centre for PH has suggested there is no protective effect from obesity [13]. However, this population included a significant proportion of drug-induced PAH patients, which may have skewed the results.

We investigated the Scottish Pulmonary Vascular Unit (SPVU) incident PAH population in order to evaluate whether obesity carries a survival benefit. All patients with suspected PAH in Scotland (population 5.1 million) are referred to the SPVU prior to right heart catheterisation, so this study represents a true national incident population.

We performed a retrospective longitudinal observational study of all consecutive, newly diagnosed idiopathic and hereditary PAH patients managed by the SPVU between 1997 and 2016. Parameters collected at diagnosis included haemodynamics (right atrial pressure (RAP), pulmonary artery pressure (PAP) and pulmonary capillary wedge pressure (PCWP)), N-terminal pro-brain natriuretic peptide (NT-proBNP), 6-min walk distance (6MWD), functional class (FC), age and body mass index (BMI). All patients were treatment-naïve prior to right heart catheterisation. The treatment commenced was at the discretion of the treating specialist, in accordance with contemporary guidelines. World Health Organization-defined BMI categories were used, with a BMI of 30 kg·m$^{-2}$ representing obesity. A survival analysis including Kaplan–Meier curves and Cox regression analysis was performed.

Analysis of pulmonary arterial hypertension patients in Scotland across 20 years supports a recent French study suggesting there is no protective effect from obesity for this disease, in contrast to emerging evidence from the USA and China [14].

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The analysis included 266 patients, 260 with idiopathic PAH and six with hereditary PAH. The prevalence of obesity was 35.7%. The nonobese group (BMI <30 kg·m\(^{-2}\)) comprised 171 patients, with a mean BMI of 24.7±3.3 kg·m\(^{-2}\). The obese group (BMI ≥30 kg·m\(^{-2}\)) comprised 95 patients, with a mean BMI of 35.9±5.7 kg·m\(^{-2}\). Mean age at diagnosis was 60.9±18.2 years and 59.1±14.3 years for the nonobese and obese groups, respectively. 49% of the nonobese group were female, compared to 77% in the obese group (p<0.001). Cardiac index, pulmonary vascular resistance, NT-proBNP and 6MWD were similar for both groups. The percentage of patients in FC I or II was higher in the nonobese group, 18.7% versus 14.9% for the obese group, although this did not reach statistical significance (p=0.054).

A number of haemodynamic parameters were significantly worse in the obese group. Mean RAP was 9.8±5.5 mmHg for the obese group and 7.5±5.1 mmHg for the nonobese group (p<0.001). Mean PAP was 51.7±10.7 mmHg for the obese group and 48.3±13.5 mmHg for the nonobese group (p=0.001). Mean PCWP was 8.6±3.9 mmHg for the obese group and 7.5±3.1 mmHg for the nonobese group (p=0.029).

Using Kaplan–Meier survival analysis (figure 1), we found no significant difference in survival between the nonobese and obese groups (p=0.358). Cox regression survival analysis using a multivariable model found neither BMI nor obesity were significant prognostic indicators of survival, only age, NT-proBNP, 6MWD and female sex were. BMI significantly correlated only with RAP (r=0.242, p<0.001) and PAP (r=0.221, p<0.001) using Spearman rank correlation.

Further variations to our model were made in order to test whether there was any obesity–mortality interaction. However, no significant difference in survival was observed when comparing overweight to nonoverweight individuals (BMI ≥25 versus <25 kg·m\(^{-2}\)), excluding severely obese individuals (BMI ≥40 kg·m\(^{-2}\)), or separating the cohorts based on age, using age separations at 50 and 65 years.

We acknowledge there are limitations in this retrospective, single-centre observational study. Recruitment of patients over a 20-year period may ignore significant changes in the demographics and treatment of PAH patients. Comorbid conditions including diabetes were not available for analysis. BMI was also the only available measure of adiposity; this does not account for adipose distribution and the effects of oedema at the time of diagnosis. BMI measurements were only recorded once at baseline but decreases in BMI over time may have occurred. The results may not be generalisable to patients with other group 1 PAH and PH aetiologies, and further studies are needed here. The strengths of this study are that the cohort is a large population of incident, treatment-naïve idiopathic and hereditary PAH patients only, assessed by specialists in a multidisciplinary team, with a standardised, high quality of treatment available as part of the UK’s National Health Service.

In contrast to this study, some earlier studies supporting the obesity hypothesis have focused on prevalent patients [4] and patients with several PAH [8, 9] or PH [7] aetiologies. AGARWAL et al. [10] studied 18,450 US-based in-hospital PAH patients and found obese patients had lower adjusted odds of mortality. However, this relied on a database using diagnostic codes rather than haemodynamics, with limited BMI information. The access to healthcare and specialised treatment may also be variable in some US studies [4]. A Chinese study of incident, treatment-naïve idiopathic PAH patients observed the obesity paradox.
was present, but included only three obese patients and had very limited availability to PAH-targeted treatments, due to cost [5].

Moreover, there is no established mechanism for this obesity effect. Some authors have recognised the possibility of it being a statistical artefact or confounding factors being responsible [4, 14]. Indeed, WEATHERALD et al. [13] recently reported no overall association of obesity or BMI with mortality amongst a population of 1255 idiopathic, heritable and drug-induced PAH patients from the French Pulmonary Hypertension Network registry.

Our present study supports the conclusion of this larger, multicentre French study. This holds true despite important differences between the cohorts of both studies. The French study included many patients with drug- and toxin-induced PAH, representing as much as 29% of the obese group. Rates of obesity were lower in the French cohort (30% versus 36% with BMI $\geq 30$ kg·m$^{-2}$). Baseline parameters including FC, 6MWD and haemodynamics were worse in the SPVU cohort. However, both cohorts are similar in terms of age, BMI and gender splits. WEATHERALD et al. [13] also reported significantly worse haemodynamic parameters in their obese group compared to their nonobese group, as have we.

In conclusion, this study of idiopathic and heritable PAH patients demonstrates no survival benefit from obesity, and mortality was not significantly associated with BMI or obesity. We hope this gives further support to recent evidence from WEATHERALD et al. [13] suggesting the obesity paradox is not present in PAH.

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