



## Mild pulmonary hypertension and premature mortality among 154956 men and women undergoing routine echocardiography

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Mild pulmonary hypertension (as indicated by estimated right ventricular systolic pressure 30.0–39.9 mmHg) is associated with increased risk of all-cause mortality and a substantial component of premature mortality https://bit.ly/3ytwlEP

**Cite this article as:** Stewart S, Chan Y-K, Playford D, *et al.* Mild pulmonary hypertension and premature mortality among 154956 men and women undergoing routine echocardiography. *Eur Respir J* 2022; 59: 2100832 [DOI: 10.1183/13993003.00832-2021].

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This article has supplementary material available from erj.ersjournals.com

This article has an editorial commentary: https://doi.org/10.1183/13993003.02064-2021

Received: 21 March 2021 Accepted: 16 May 2021

## Abstract

*Background* Although mild pulmonary hypertension is known to be associated with increased mortality, its impact on premature mortality is largely unknown.

*Methods* We studied the distribution of estimated right ventricular systolic pressure (eRVSP) among a total of 154956 adults with no evidence of left heart disease investigated with echocardiography. We then examined individually linked mortality, premature mortality and associated life-years lost (LYL) according to eRVSP levels.

Results The cohort comprised 70826 men and 84130 women (aged 61.3±17.7 and 61.4±18.4 years, respectively). Overall, 85173 (55.0%), 49276 (31.8%), 13060 (8.4%) and 7447 (4.8%) cases had eRVSP levels indicative of no (<30.0 mmHg), mild (30.0−39.9 mmHg), moderate (40.0−49.9 mmHg) or severe (≥50.0 mmHg) pulmonary hypertension, respectively. During a median (interquartile range) 5.7 (3.2−8.9) years of follow-up, 38456/154986 (24.8%) individuals died. Compared with eRVSP <30.0 mmHg, age and sex-adjusted hazard ratios for all-cause and cardiovascular-related mortality were 1.90 (95% CI 1.84−1.96) and 1.85 (95% CI 1.74−1.97), respectively, for eRVSP 35.0−39.9 mmHg. Overall, 6256 (54%) men and 7524 (55%) women died prematurely. As a proportion of all deaths, premature mortality rose from 46.7% to 79.2% among those with eRVSP <30.0 versus ≥60.0 mmHg with a mean of 5.1−11.4 LYL each time. However, due to more individuals affected overall, eRVSP 30.0−39.9 mmHg accounted for 58% and 53% of total LYL among men (40 606/70019 LYL) and women (47 333/88 568 LYL), respectively. Conclusions These data confirm that elevated eRVSP levels indicative of mild pulmonary hypertension are associated with increased risk of death. Moreover, this results in a substantive component of premature mortality/LYL that requires more proactive clinical surveillance and management.