

Table 2**Current real world survival of pulmonary hypertension patients referred to designated pulmonary hypertension centres in the UK³**

Cause of pulmonary hypertension	Median age at diagnosis (years)	Median survival from diagnosis (years)
Idiopathic, heritable and drug-induced pulmonary arterial hypertension	60	5.8
Congenital heart disease	45	> 8*
Connective tissue disease associated pulmonary arterial hypertension with scleroderma	67	3.5
Connective tissue disease associated pulmonary arterial hypertension without scleroderma	61	4.1
Operated chronic thromboembolic pulmonary hypertension	61	> 9*
Unoperated chronic thromboembolic pulmonary hypertension	69	5.7
Pulmonary hypertension due to left heart disease	74	4.5
Pulmonary hypertension due to lung disease and/or hypoxia	68	1.8

*50% mortality of the cohort has not been reached in 2019