

Please take a minute to understand more about *pulmonary* hypertension



Rare, serious, misunderstood

Pulmonary hypertension (PH for short) causes high blood pressure in the blood vessels connecting the heart and lungs (the pulmonary arteries). In a *healthy cardiovascular system*, the right-side to the heart pumps blood into the lungs to pick up oxygen and this oxygen-rich blood is then pumped around the body by the left-side of the heart.

When a person develops PH, the walls of the pulmonary arteries become stiff and thickened, or blocked by blood clots. This makes it difficult for them to expand; and trying to pump blood through these tightened, narrowed, scarred or blocked arteries puts increasing strain on the right side of the heart as it tries to do its job. The essential task of pumping blood into the lungs to pick up oxygen which can then be circulated to every cell in the body becomes much harder.

Living with the chronic condition can also have a significant impact on emotional wellbeing and self-esteem as people deal with the frustrations caused by PH and their worries about the future.

Pulmonary hypertension is rare.

Around **8,000 people** are diagnosed with PH in the UK. It can affect anyone, regardless of age or ethnic background. It is more common in women than men.

PH affects people's abilities

to carry out basic tasks and get around. People with this condition often look well at rest and it's only through a simple activity such as climbing the stairs that they may experience symptoms:



PH can be *associated with another medical condition* such as congenital heart disease, connective tissue disease, HIV infection or sickle cell anaemia.

A small number of people with PH develop it without having another medical condition – this is known as idiopathic pulmonary arterial hypertension (IPAH).

How is PH treated?

PH responds to a range of highly advanced treatments which can relax the arteries to help increase blood flow and reduce pressure, improve symptoms, slow disease progression and reverse damage to the heart and lungs. However, there is no cure.

Some people with pulmonary arterial hypertension require lung and /or heart transplants.

The type of PH caused by a build-up of blood clots, called chronic thromboembolic pulmonary hypertension (CTEPH), may be cured by undergoing major surgery to remove the clots.



Scan here to learn more about PH



The Pulmonary
Hypertension
Association (PHA UK)
is the only charity
in the UK dedicated
to supporting those
affected by PH.

It provides information, support and advice; and funds research, educational programmes and activities to promote better understanding, diagnosis and treatment of the disease. It is free to become a member.

Find out more or join online at www.phauk.org

Registered Charity Number: 1120756



