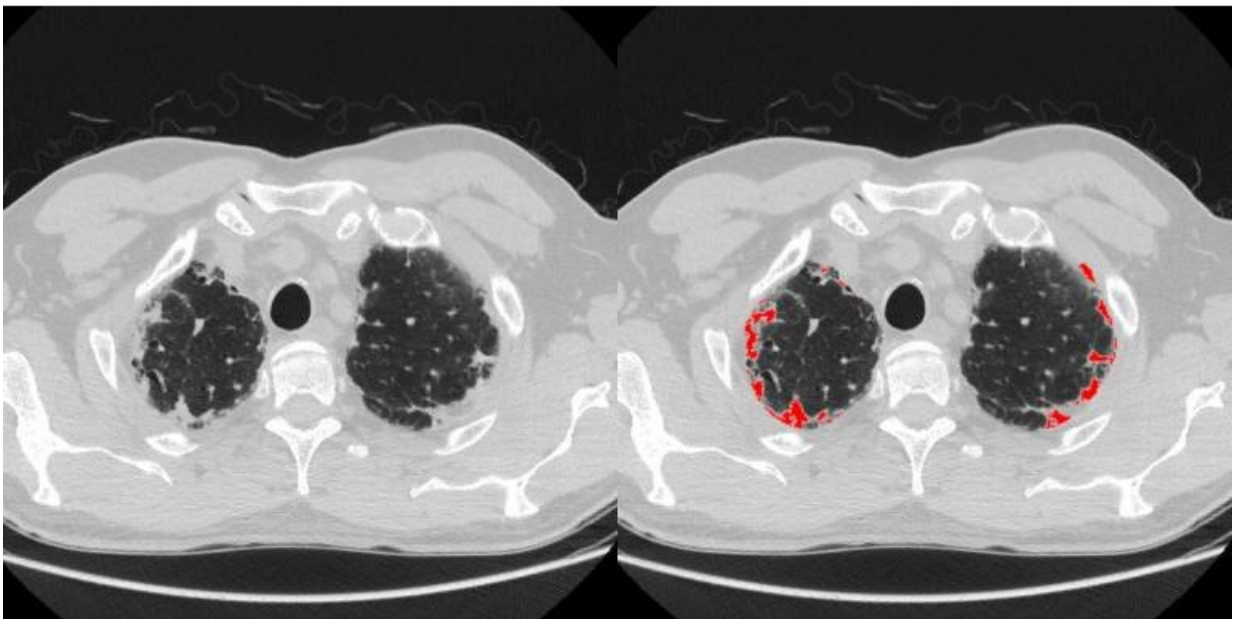


Computerised image analysis identifies new subtype of debilitating lung disease

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Computed tomography images of the upper lungs. The image on the left shows areas of thickening in the periphery of the lungs. The same regions are highlighted in red on the right image following computer analysis of the computed tomography scan. Areas in red correspond to pleuroparenchymal fibroelastosis, a previously under-reported pattern of lung damage seen in patients with lung fibrosis. Credit: University College London

Scientists at UCL have used computer-based imaging analysis to identify new patterns of lung damage in patients with Idiopathic pulmonary fibrosis (IPF), providing the first evidence of a previously unknown and life-limiting lung disease subtype.

IPF is a condition where [scar tissue](#) or fibrosis builds up in the lungs, making them thick and hard. It affects around 3 million people globally (around 32,500 people in the UK), with those affected typically aged between 50 and 70, and has a life expectancy of three to five years.

In this study, published in *EClinicalMedicine*, researchers at the Centre for Medical Image Computing (CMIC) at UCL, developed a computerized image-analysis algorithm, partially based on [deep learning](#), to see if lungs affected by IPF may also have scarring patterns characteristic of another serious, though less well-documented, [lung disease](#) called pleuroparenchymal fibroelastosis (PPFE). It is not known for certain what causes either IPF or PPFE.

Explaining the research, lead author Dr. Joe Jacob (UCL Division of Medicine) said: "Idiopathic pulmonary fibrosis is a [disease](#) that shrinks the lungs and is associated with a life expectancy comparable to many cancers.

"Doctors have suspected for some time that there are distinct subgroups or endotypes of IPF that have unique functional or pathological disease mechanisms which make the condition even more deadly.

"Identifying IPF endotypes could allow a degree of personalized patient management."

In this study, researchers used IPF CT scans from two populations of approximately 145 patients. A radiologist was asked to identify those which also had PPFE and identified extensive PPFE in 31 out of 287 (or

11%) cases.

The research team then applied a computerized imaging tool, developed by Dr. Eyjolfur Gudmundsson at CMIC, UCL to identify PPFE on a subset of scans from one of the populations. The algorithm was subsequently evaluated on scans separately in the two patient populations and concluded that 87 of the 287 scans (or 30%) had clinically important PPFE, a threefold increase on the radiologist's assessment.

The authors say the significant numbers of IPF patient scans, which also had PPFE, shows this is a feature, characteristic of a IPF disease subgroup/endotype.

Dr. Jacob said: "Using this computerized tool we have identified the first endotype in IPF occurring in about 30% of IPF patients.

"The PPFE identified adversely affects survival and appears to occur and progress separate to the fibrosis in the patients' lungs."

The research team also retrospectively matched the IPF images which had PPFE, against patient records, and concluded the presence of PPFE is associated with faster clinical deterioration and earlier death in IPF patients.

Dr. Jacob added: "Our research suggests that PPFE is a previously under-reported disease process that occurs separate to [lung](#) fibrosis but which is associated with an increased risk of death in IPF.

"Our computer algorithm identifies patients with the PPFE endotype of IPF which may inform patient selection in IPF drug trials and clinical management of disease. Our algorithm should be able to measure PPFE worsening over time in patients with IPF."

More information: Eyjolfur Gudmundsson et al, Pleuroparenchymal fibroelastosis in idiopathic pulmonary fibrosis: Survival analysis using visual and computer-based computed tomography assessment, *EClinicalMedicine* (2021). DOI: [10.1016/j.eclinm.2021.101009](https://doi.org/10.1016/j.eclinm.2021.101009)

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