Supplement

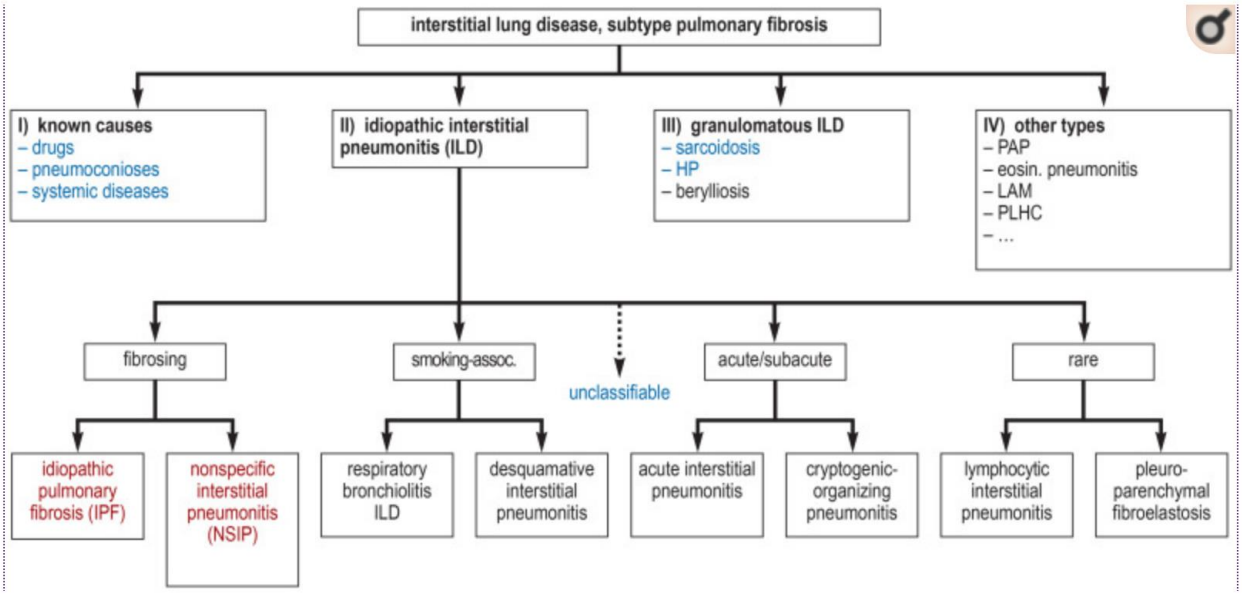
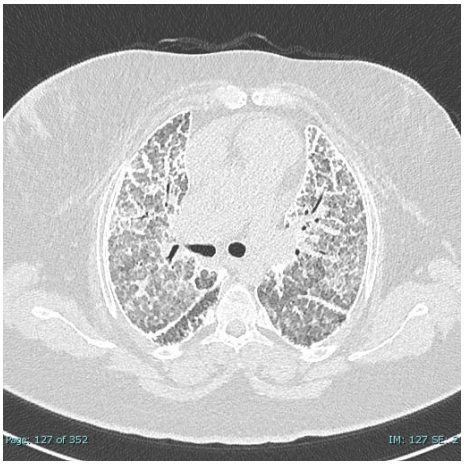
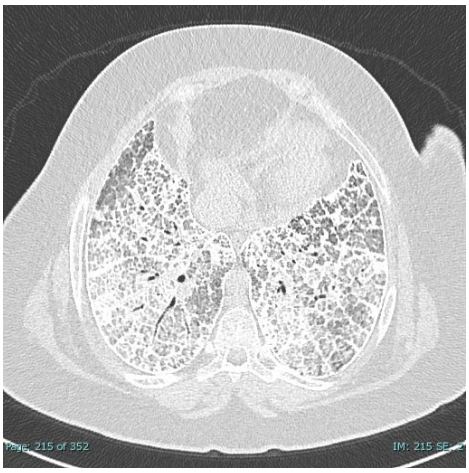


Figure S1 Categorisation of interstitial lung disease (ILD):
I) known causes, II) idiopathic types, III) granulomatous types, IV) further types.
HP: hypersensitivity pneumonitis; ILD: interstitial lung disease; LAM: lymphangioleiomyomatosis; PAP: pulmonary alveolar proteinosis; PLHC: pulmonary Langerhans-cell histiocytosis [2].



S2a



S2b

Figure S2 Chest CT scan of a 31-year-old female with the diagnosis of pulmonary alveolar microlithiasis. Extensive, basal accentuated calcifications (S2b) as well as ground glass opacities (S2a).