



Interstitial pneumonia with autoimmune features: challenges and controversies

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Since the proposal of the research entity interstitial pneumonia with autoimmune features (IPAF) in 2015, many questions remain unanswered and challenges remain in its definition

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Abstract

The presence of clinical, serological and/or radiological features suggestive, but not confirmatory, of a defined connective tissue disease in patients with interstitial lung disease is a relatively frequent occurrence. In 2015, the European Respiratory Society and the American Thoracic Society proposed classification criteria for the interstitial pneumonia with autoimmune features (IPAF) research entity to capture such patients in a standardised manner, with the intention of nurturing clinical research. This initiative resulted in the publication of several series of IPAF patients, with significant variation between cohorts in clinical characteristics, outcome and the application of IPAF criteria in patient selection. From this increasing body of published work, it has become apparent that revision of IPAF criteria is now required in order to justify the eventual designation of IPAF as a standalone diagnostic term, as opposed to a provisional entity put forward as a basis for clinical research. This review covers the current state of IPAF, conclusions that can and cannot be drawn from the IPAF evidence base, and ongoing uncertainties that require further expert group consideration.

Introduction

The diffuse parenchymal lung diseases are differentiated from the outset by the presence of an established cause or association. Connective tissue diseases (CTDs) represent one of the more common causes of non-idiopathic interstitial lung disease (ILD). Almost all CTDs may be associated with ILD, but the individual CTDs vary in the prevalence and most frequent morphological pattern of ILD. CTDs are defined by strict diagnostic criteria that, outside of systemic sclerosis (SSc), exclude ILD as a diagnostic factor. When ILD complicates a previously defined CTD, the diagnosis is generally straightforward, amounting to CTD-ILD (with the exception of drug-induced disease). However, a reasonable proportion of patients present with ILD in the presence of features suggestive of, but not confirmatory for, a defined CTD. Whether such patients behave and respond similarly to CTD-ILD, or more akin to their idiopathic counterparts, has been the focus of much literature over the last decade. Despite defined classification criteria for research under the title interstitial pneumonia with autoimmune features (IPAF), challenges and controversies continue to surround this group of patients.

Why was IPAF proposed?

A sizable proportion of patients with ILD present with clinical, serological and/or radiological features suggestive of CTD, but lacking features to meet the established diagnostic criteria of defined CTDs [1–7].



Up to 2015, this overlap between idiopathic interstitial pneumonia (IIP), especially idiopathic pulmonary fibrosis (IPF) and CTD-ILDs (figure 1) was described using a variety of terminologies, including “undifferentiated CTD-ILD” [8], “undifferentiated connective tissue syndromes” [9, 10], “lung-dominant CTD” [11], “early CTD”, “*formes frustes*” of CTD [1], “occult CTD”, “overlap CTD”, “lung-limited CTD” [2] and “autoimmune-featured ILD” [12]. These terms largely overlap yet do not exactly match. Corresponding studies provided accumulating evidence that CTD features were relevant in the setting of ILD even when falling short of fulfilling existing classification criteria for a specific CTD. However, research was hampered by heterogeneity in terminology and definition [13].

In 2015, the European Respiratory Society (ERS) and American Thoracic Society (ATS) Task Force on Undifferentiated Forms of Connective Tissue Disease-Associated Interstitial Lung Disease proposed classification criteria for a so-called research category IPAF [14], with the aim of providing a consensus terminology and definition, and to foster research. The Task Force included respiratory physicians, rheumatologists and immunologists. The IPAF classification criteria are summarised in table 1.

Advantages of the IPAF concept and definition

The concept of IPAF emphasises that there can be an overlap between IIPs and CTD-ILD and provides a framework to designate and study this condition. Such overlap between entities is common in medicine, and can be due to overlapping manifestations, as seen in overlap CTD, in which features of separate CTDs are present in the same patient, or be related to uncertain aetiology, as in patients with a condition strongly suggesting IPF but exposed to inhaled antigens known to cause chronic fibrotic hypersensitivity pneumonitis [15]. IPAF deals with the overlap between IIPs and CTD-ILD.

The concept of IPAF has highlighted the need for multidisciplinary discussions in the field of CTD-ILD, especially the input of rheumatologists in the ILD clinic [16]. It has already shed light on the importance of a thorough evaluation of patients with apparently idiopathic ILD and on the value of the interaction between medical specialties [17]. Although its management is not standardised, the concept of IPAF facilitates making decisions in patients presenting with ILD and mild CTD features before a longer follow-up clarifies whether CTD is present or not [8], as opposed to the concept of “undifferentiated CTD”, therefore addressing an unmet clinical need when seeing a patient with new-onset ILD.

Outstanding issues in the IPAF concept

A question that remains open is whether IPAF represents a clinical diagnosis, a distinct entity, or just an overlap of existing conditions. A small proportion (10–20%) of patients initially classified as IPAF

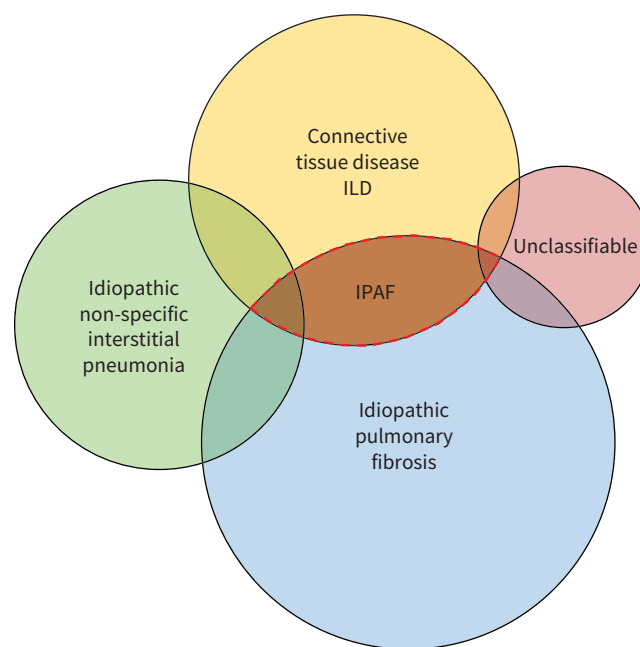


FIGURE 1 Interstitial pneumonia with autoimmune features (IPAF) in the schema of clinical interstitial lung disease (ILD).

TABLE 1 Classification criteria for interstitial pneumonia with autoimmune features

Presence of an interstitial pneumonia (by HRCT or surgical lung biopsy) *and*
 Exclusion of alternative aetiologies *and*
 Does not meet criteria of a defined connective tissue disease *and*
 At least one feature from at least two of these domains:

- a) clinical domain
 - Distal digital fissuring (*i.e.* “mechanic hands”)
 - Distal digital tip ulceration
 - Inflammatory arthritis or polyarticular morning joint stiffness ≥ 60 min
 - Palmar telangiectasia
 - Raynaud’s phenomenon
 - Unexplained digital oedema
 - Unexplained fixed rash on the digital extensor surfaces (Gottron’s sign)
- b) Serological domain

ANA $\geq 1:320$ titre, diffuse, speckled, homogeneous patterns *or*

 - 1) ANA nucleolar pattern (any titre) *or*
 - 2) ANA centromere pattern (any titre)

Rheumatoid factor $\geq 2\times$ upper limit of normal

 - Anti-CCP
 - Anti-dsDNA
 - Anti-Ro (SS-A)
 - Anti-La (SS-B)
 - Anti-ribonucleoprotein
 - Anti-Smith
 - Anti-topoisomerase (Scl-70)
 - Anti-tRNA synthetase (*e.g.* Jo-1, PL-7, PL-12; others are: EJ, OJ, KS, Zo, tRS)
 - Anti-PM-Scl
 - Anti-MDA-5
- c) Morphological domain
 - 1) Suggestive radiology patterns by HRCT (see text for descriptions):
 - i) NSIP
 - ii) OP
 - iii) NSIP with OP overlap
 - iv) LIP
 - 2) Histopathology patterns or features by surgical lung biopsy (figure 2):
 - i) NSIP
 - ii) OP
 - iii) NSIP with OP overlap
 - iv) LIP
 - v) Interstitial lymphoid aggregates with germinal centres
 - vi) Diffuse lymphoplasmacytic infiltration (with or without lymphoid follicles)
 - 3) Multi-compartment involvement (in addition to interstitial pneumonia):
 - i) Unexplained pleural effusion or thickening
 - ii) Unexplained pericardial effusion or thickening
 - iii) Unexplained intrinsic airways disease[#] (by PFT, imaging or pathology)
 - iv) Unexplained pulmonary vasculopathy

ANA: antinuclear antibody; HRCT: high-resolution computed tomography; LIP: lymphoid interstitial pneumonia; NSIP: non-specific interstitial pneumonia; OP: organising pneumonia; PFT: pulmonary function testing.
[#]: includes airflow obstruction, bronchiolitis or bronchiectasis. Reproduced and modified from [14] with permission.

progress to a diagnosis of CTD-ILD [18–22], suggesting that in some cases IPAF does indeed correspond to CTD-ILD early in the course of disease. Also, significant heterogeneity persists within the group designated by IPAF. The respective contribution of the different domains to the classification as IPAF varies between series [23], suggesting variations in referral patterns, with some IPAF series being “close to IPF cases” and others “closer to CTD-ILD cases”. The outstanding issues surrounding the IPAF criteria are detailed below and summarised in table 2.

Inclusion of rheumatological review

Routine involvement of a rheumatologist in the multi-disciplinary evaluation of patients with ILD has been proposed but is not a current prerequisite for the adjudication of IPAF criteria. CHARTRAND *et al.* [24] and

TABLE 2 Unresolved issues of current interstitial pneumonia with autoimmune features (IPAF) criteria

Clinical domain

- When to involve rheumatology and/or immunology in the evaluation of patients meeting IPAF criteria.
- When and how to re-evaluate for a connective tissue disease.
- Inclusion of sicca symptoms, oesophageal dysmotility, proximal muscle weakness and myalgia.
- Clustering in time of autoimmune features.
- Family history of autoimmune disease.
- Role of additional investigations, including nailfold capillaroscopy, salivary gland biopsy, articular radiology and electromyography.

Serological domain

- Inclusion/exclusion of myositis-specific antibodies.
- Inclusion of anti-neutrophil cytoplasmic antibodies.

Morphological domain

- Separation of usual interstitial pneumonia (UIP) from non-UIP – should IPAF criteria be applied differently?
- Specific definitions for the application of multi-compartment involvement.
- Role and indications of lung biopsy in patients with IPAF features.

JEE *et al.* [25], whose clinics routinely included rheumatologists, noted a higher prevalence of features from the clinical domain. Additionally, the IPAF cohort of AHMAD *et al.* [26] was notable for a number of patients with abnormal nailfold capillaroscopy and salivary gland biopsy, investigations not typical of a pulmonologist. In fact, abnormal nailfold capillaroscopy, together with the presence of ILD, may be sufficient to satisfy SSc diagnostic criteria (table 3), removing such cases from the IPAF entity [27]. LEVI *et al.* [28] observed that, following rheumatological review, 40% of selected ILD multi-disciplinary meeting cases received a rheumatology-related ILD diagnosis, including IPAF. One-fifth of IPF diagnoses were reclassified into a range of CTD-ILDs. However, the data supplement would suggest that pulmonologists had access to a narrow range of auto-antibodies, and it is unclear whether defined CTD criteria were enforced. DE LORENZIS *et al.* [29] evaluated the routine involvement of rheumatological expertise in their cases with a suggestion of an underlying rheumatic disorder. Interestingly, a large number of patients underwent additional investigations, including articular radiology, electromyography, nailfold capillaroscopy and salivary gland biopsy. Agreement between pulmonologists and rheumatologists across various CTD clinical and serological features was only fair to moderate, with rheumatologists identifying a greater number of extrapulmonary features of CTD. To a large extent, identifying IPAF criteria depends on the effort made to look for CTD features. Nailfold capillaroscopy [26], biopsy of accessory salivary glands [26, 30, 31] or consultation with a rheumatologist or a dermatologist often identify CTD features that may alter the eventual diagnosis, yet they cannot be easily recommended in all patients with ILD. The decision on how far to search for evidence of an underlying CTD often impacts the eventual diagnosis [32]. The current IPAF criteria do not specify a need for specialist rheumatologist evaluation, but the data summarised above suggest that rheumatological input might help to improve the sensitivity and specificity of the IPAF criteria [33].

Controversies surrounding inclusion/exclusion of items/characteristics

The heterogeneity in IPAF series has prompted some authors to suggest alternative definitions of IPAF [34, 35], to remove some of the individual items from the IPAF group, especially anti-synthetase antibodies [36, 37] (see below), to consider the inclusion of selected additional items (*e.g.* sicca symptoms, oesophageal dysmotility, lymphocytic bronchiolitis on biopsy, proximal muscle weakness with myalgia [38] or antineutrophil cytoplasmic antibodies [39]), or to separate subjects with a usual interstitial pneumonia (UIP) pattern from those with a non-UIP pattern (figures 2 and 3) [35].

The issue of sicca symptom inclusion is an example of some of the uncertainties surrounding the inclusion of specific items in the clinical domain. In reports prior to the IPAF nomenclature, sicca symptoms had been invariably included as an autoimmune clinical feature. However, sicca symptoms are absent from IPAF criteria, as they are considered to lack specificity. While CORTE *et al.* [10] and KINDER *et al.* [9] showed no difference in the prevalence of sicca symptoms between those with and without autoimmune features, VIJ *et al.* [12] reported a prevalence of 57.1% compared to 15.5% in those with autoimmune features and IPF respectively. SEBASTIANI *et al.* [22] observed a similar prevalence of sicca symptoms in their IPAF cohort, suggesting that their inclusion may need to be considered.

Myositis-specific antibodies (MSAs), including anti-synthetase antibodies, are an example of the uncertainties surrounding the serological domain (figure 4a and b). MSAs are currently included in the IPAF serological domain (table 1). GRAHAM *et al.* [37] compared survival within IPAF patients, subdivided

TABLE 3 Various diagnostic criteria proposed for interstitial lung disease (ILD) in patients with features of connective tissue disease (CTD), and the diagnostic criteria for defined CTDs adapted to suit the domains of the interstitial pneumonia with autoimmune features (IPAF) criteria

Domain	Undifferentiated CTD KINDER <i>et al.</i> [9]	Undifferentiated CTD CORTE <i>et al.</i> [10]	Autoimmune-featured ILD VI <i>et al.</i> [12]	Lung-dominant CTD FISCHER <i>et al.</i> [11]	IPAF ERS/ATS [14]	RA ACR/EULAR [7]	SSc ACR/EULAR [6]	Idiopathic inflammatory myopathy EULAR/ACR [4]	Systemic lupus erythematosus EULAR/ACR [3]	Sjögren syndrome ACR/EULAR [5]
Entry requirements	Presence of an interstitial pneumonia. Exclusion of alternative aetiologies. Does not meet criteria for a defined CTD.	Presence of an interstitial pneumonia. Exclusion of alternative aetiologies. Does not meet criteria for a defined CTD.	Presence of an interstitial pneumonia. Exclusion of alternative aetiologies. Does not meet criteria for a defined CTD.	Presence of an interstitial pneumonia. Exclusion of alternative aetiologies. Does not meet criteria for a defined CTD.	Presence of an interstitial pneumonia. Exclusion of alternative aetiologies. Does not meet criteria for a defined CTD.	At least one joint with definite clinical synovitis. Synovitis not better explained by another disease.	Excludes skin thickening sparing the fingers or better explained by another condition.	No better explanation for the symptoms and signs.	ANA \geq 1:80 on HEp-2 cells or an equivalent positive test (ever).	Ocular or oral dryness. Exclusion of other causes.
Clinical	Raynaud's phenomenon. Arthralgias/multiple joint swelling. Morning stiffness. Dry mouth or dry eyes. Proximal muscle weakness. Photosensitivity. Unintentional weight loss. Dysphagia. Recurrent unexplained fever. GORD. Skin changes (rash). Oral ulceration. Non-androgenic alopecia.	Raynaud's phenomenon. Arthralgias/multiple joint swelling. Morning stiffness. Dry mouth or dry eyes. Proximal muscle weakness.	Raynaud's phenomenon. Arthralgias/multiple joint swelling. Morning stiffness. Dry mouth or dry eyes. Proximal muscle weakness. Photosensitivity. Unintentional weight loss. Dysphagia. GORD. Oral ulceration. Leg/foot swelling. Hand ulcers.	Raynaud's phenomenon. Inflammatory arthritis or polyarticular morning joint stiffness \geq 60 min. Distal digital tip ulceration. Palmar telangiectasia. Unexplained digital oedema. Distal digital fissuring (<i>i.e.</i> mechanic hands). Unexplained fixed rash on the digital extensor surfaces (Gottron's sign).	Raynaud's phenomenon. Clinical synovitis (increasing score with increasing small joint involvement). Duration of joint symptoms ($</\geq$ weeks).	Raynaud's phenomenon. Skin thickening. Digital ulcers/pitting scars. Telangiectasia. Abnormal nailfold capillaries.	Proximal muscle weakness. Heliotrope rash. Gottron's papules. Gottron's sign. Dysphagia or oesophageal dysmotility.	Joint involvement. Oral ulcers. Non-scarring alopecia. Subacute cutaneous or discoid lupus. Acute cutaneous lupus. Unexplained fever. Neuropsychiatric (delirium, psychosis, seizure).	Ocular or oral dryness.	
Serological	ANA Anti-Scl70 SSA or SSB Jo-1 RF ESR $>2\times$ normal	ANA (high titre) Anti-centromere ENA Anti-Scl70 Anti-RNP SSA or SSB Jo-1 RF (high titre)	ANA \geq 1:160 Anti-Scl70 Anti-RNP SSA or SSB Jo-1 Anti-Sm RF Anti-CCP Anti-dsDNA ANCA CK Aldolase	ANA $>$ 1:320 or nucleolar or anticentromere Anti-Scl70 Anti-RNP SSA or SSB Anti-tRNA synthetase Anti-Sm Anti-PM-Scl RF $>$ 60 IU·mL ⁻¹ Anti-CCP Anti-dsDNA	ANA \geq 1:320 or nucleolar or anti-centromere Anti-Scl70 Anti-RNP SSA or SSB Anti-tRNA synthetase Anti-Sm Anti-PM-Scl Anti-MDA-5 RF \geq 2 \times ULN Anti-CCP Anti-dsDNA	RF Anti-CCP Elevated CRP or ESR	Anti-centromere Anti-Scl70 Anti-RNA polymerase III	Jo-1 Elevated CK/LDH/AST/ALT	ANA \geq 1:80 Anti-Sm Antiphospholipid antibody Anti-dsDNA Low C3/C4 Leukopenia Thrombocytopenia Autoimmune haemolysis Proteinuria	SSA

Continued

TABLE 3 Continued

Domain	Undifferentiated CTD KINDER <i>et al.</i> [9]	Undifferentiated CTD CORTE <i>et al.</i> [10]	Autoimmune-featured ILD VU <i>et al.</i> [12]	Lung-dominant CTD FISCHER <i>et al.</i> [11]	IPAF ERS/ATS [14]	RA ACR/EULAR [7]	SSc ACR/EULAR [6]	Idiopathic inflammatory myopathy EULAR/ACR [4]	Systemic lupus erythematosus EULAR/ACR [3]	Sjögren syndrome ACR/EULAR [5]
Morphological	Not specified	Not specified	Not specified	Lung histopathology: lymphoid aggregates with germinal centres. Prominent plasmacytic infiltration. Dense perivascular collagen. Extensive pleuritis.	Radiology: NSIP; OP; NSIP+OP; LIP. Lung histopathology: NSIP; OP; NSIP+OP; LIP; interstitial lymphoid aggregates with germinal centres; diffuse lymphoplasmacytic infiltration (with or without lymphoid follicles). Multi-compartment involvement: unexplained pleural effusion or thickening; unexplained pericardial effusion or thickening; unexplained intrinsic airways disease (by PFT, radiology or histopathology); unexplained pulmonary vasculopathy.		Pulmonary arterial hypertension. ILD.	Muscle histopathology: endomysial infiltration of mononuclear cells surrounding, but not invading, myofibres; perimysial and/or perivascular infiltration of mononuclear cells; perifascicular atrophy; rimmed vacuoles.	Lupus nephritis on renal biopsy. Pleural or pericardial effusion. Acute pericarditis.	Labial salivary gland with focal lymphocytic sialadenitis and focus score of ≥ 1 . Ocular staining score. Schirmer's test. Unstimulated whole saliva flow.
Required to meet definition	At least one clinical and one serological feature.	At least one clinical and one serological feature.	At least one clinical and one serological feature.	At least one serological feature or at least two histopathology features.	At least one feature from at least two of the domains.	Summed weighted scores of ≥ 6 for classification as definite RA.	Summed of weighted scores of ≥ 9 for definite SSc.	Probability score calculated by sum of weighted responses to above features.	At least one clinical criterion and ≥ 10 points.	Total score ≥ 4 .

ACR: American College of Rheumatology; ALT: alanine aminotransferase; ANA: anti-nuclear antibody; ANCA: anti-neutrophil cytoplasmic antibody; AST: aspartate aminotransferase; ATS: American Thoracic Society; CCP: cyclic citrullinated peptide; CK: creatinine kinase; CRP: C-reactive protein; dsDNA: double-stranded DNA; ENA: extractable nuclear antibody; ERS: European Respiratory Society; ESR: erythrocyte sedimentation rate; EULAR: European Alliance of Associations for Rheumatology; GORD: gastro-oesophageal reflux disease; LDH: lactate dehydrogenase; LIP: lymphoid interstitial pneumonia; NSIP: non-specific interstitial pneumonia; OP: organising pneumonia; PFT: pulmonary function test; RA: rheumatoid arthritis; RF: rheumatoid factor; RNP: ribonucleoprotein; SSA: Sjögren syndrome-related antibody A; SSB: Sjögren syndrome-related antibody B; SSc: systemic sclerosis; UIP: usual interstitial pneumonia; ULN: upper limit of normal.

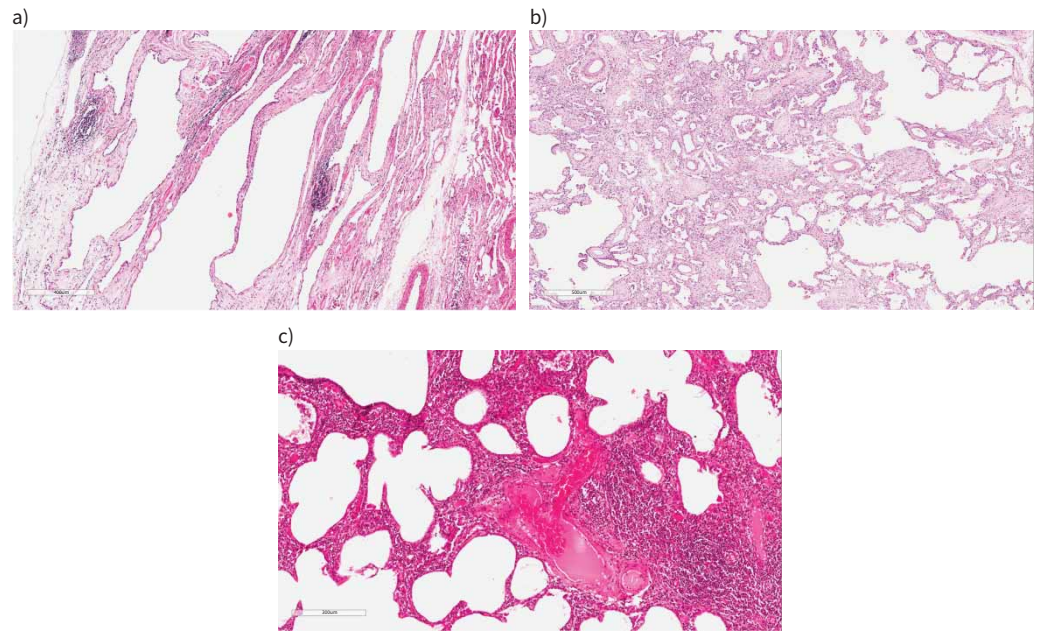


FIGURE 2 Histological patterns of interstitial pneumonia seen in interstitial pneumonia with autoimmune features. **a)** Fibrotic non-specific pneumonia (NSIP): there is diffuse uniform interstitial fibrosis associated with a mild patchy non-specific chronic inflammatory cell infiltrate. **b)** Organising pneumonia (right side) with progression to interstitial fibrosis (left side): buds of granulation tissue merge with established fibrosis. **c)** Lymphoid interstitial pneumonia (LIP) overlapping with cellular NSIP: in this field, the right-hand part of the image shows a density of interstitial inflammation characteristic of LIP.

according to whether patients were 1) positive for MSAs; 2) positive for myositis-associated antibodies (MAAs) but negative for MSAs; or 3) negative for both. The study also included cohorts of idiopathic inflammatory myositis (IIM), CTD-ILD and IPF patients. Interestingly, IPAF patients with MSA (n=35), but not MAAs (n=65), were more frequently characterised by a computed tomography (CT) or pathology pattern of non-specific interstitial pneumonia (NSIP), and had very good outcomes on immunosuppressive treatment, similarly to patients with IIM, and significantly better than IPAF with MAA, or IPAF without MSA/MAA. While a UIP pattern was associated with a worse survival in IPAF overall, and in the IPAF-MAA group, this was not the case in IPAF-MSA, although there were only six cases with IPAF-MSA UIP. Whether a UIP pattern associated with MSA has a better prognosis than the other IPAF

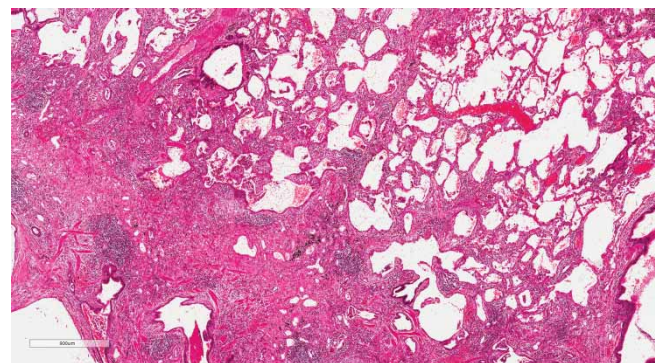


FIGURE 3 Usual interstitial pneumonia (UIP) with features arguing against idiopathic pulmonary fibrosis (IPF). There are areas with established patchy interstitial fibrosis with an occasional fibroblastic focus characteristic of UIP, but also areas where interstitial chronic inflammation predominates, making this case indeterminate for UIP/IPF. Based on pathology alone, either fibrotic hypersensitivity pneumonitis or a connective tissue disease-related interstitial lung disease (interstitial pneumonia with autoimmune features) were favoured over IPF, and multidisciplinary review was recommended.

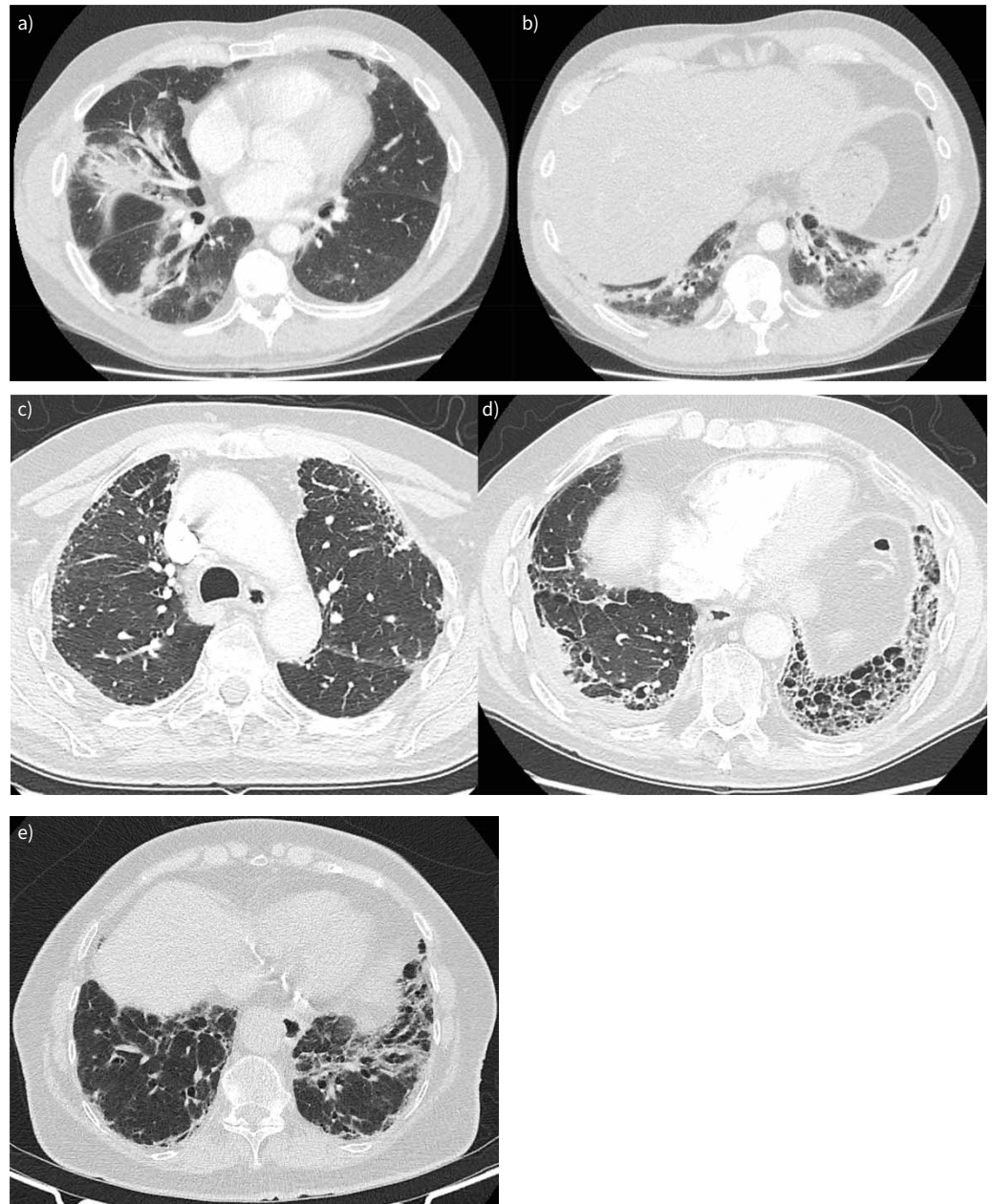


FIGURE 4 a and b) Interstitial pneumonia with autoimmune features (IPAF) in a 60-year-old male presenting with acute onset type 1 respiratory failure. Computed tomography (CT) demonstrates patchy consolidation with a bronchocentric distribution, and some lower lobe volume loss, suggestive of a pattern of fibrotic organising pneumonia. Positive serology for anti-OJ was identified. No clinical features of an idiopathic inflammatory myopathy were evident at presentation. The patient had an excellent response to prednisolone and tacrolimus. c and d) IPAF in an 80-year-old male. The CT demonstrates a usual interstitial pneumonia (UIP) pattern with right-sided pleural thickening/effusion suggestive of multi-compartment involvement. No known history of asbestos exposure was reported and no pleural plaques were evident on CT. The patient's anti-nuclear antibody (ANA) was low titre but in a nucleolar pattern and rheumatoid factor was elevated. The patient was treated with immunomodulation, but demonstrated an idiopathic pulmonary fibrosis-like disease trajectory and died within a few years of diagnosis. e) 84-year-old female presenting with gradually progressive dyspnoea with CT demonstrating a UIP pattern. ANA at presentation was speckled with titre of 1:320. The patient subsequently developed inflammatory arthritis and a positive anti-cyclic citrullinated peptide, resulting in an eventual diagnosis of rheumatoid arthritis and associated interstitial lung disease. Her rheumatoid arthritis was treated with sulfasalazine and the lung disease has remained under observation after the patient declined clinical trial participation. Over the first 12 months there was a 7% relative decline in forced vital capacity and a 26% relative decline in carbon monoxide diffusing capacity.

UIP patients will require confirmation in further studies. HUAPAYA *et al.* [40] reported progressive improvement and/or stabilisation with azathioprine or mycophenolate in patients with IIM. GRAHAM *et al.* [37] suggested that IPAF with MSA should be clinically managed as IIM ILD, and that MSA should potentially be removed from the IPAF criteria.

Clustering in time of the manifestations (*e.g.* concomitant apparition of arthralgia and gastro-oesophageal reflux in a patient with new-onset ILD) may be meaningful clinically, but it is not captured in the IPAF criteria. Identification of a family history of autoimmune disease by CHARTRAND *et al.* [24] in 25% of their IPAF cohort suggests that this factor might also require consideration.

The evaluation of multi-compartment involvement

The IPAF morphological domain requires either the presence of a non-UIP histological pattern or “multi-compartment involvement”, *i.e.* co-existing airways disease, pulmonary vascular disease or pleural disease (figure 4c and d). However, there is a lack of specificity and reproducibility of the multi-compartment involvement subdomain as an independent criterion. There are ongoing difficulties with the identification of co-existent disease processes relevant to the entity of IPAF due to the high prevalence of comorbidities in patients with ILD.

In this regard, the definition of “disproportionate” pulmonary vascular disease, a well-recognised autoimmune phenotype, is especially problematic. In the index IPAF series of OLDHAM *et al.* [41], pulmonary vascular involvement was identified based on a reduction in carbon monoxide diffusing capacity (D_{LCO}) that was disproportionate to forced vital capacity (FVC), captured using an FVC/D_{LCO} threshold. However, whilst standardisation of this sort has advantages, an FVC/D_{LCO} threshold is not, in itself, indicative of disproportionate pulmonary vasculopathy. Pulmonary hypertension is a frequent complication in advanced fibrotic lung disease and, when severe, is associated with major increases in the FVC/D_{LCO} ratio and major reductions in the gas transfer coefficient (K_{CO}). However, pulmonary hypertension is not a disproportionate finding in this scenario and is not an “autoimmune feature”. Moreover, both K_{CO} and the FVC/D_{LCO} ratio are non-specific: influenced alike by pulmonary vasculopathy and the presence of concurrent emphysema. In the “idiopathic” (or tobacco-related) combined pulmonary fibrosis and emphysema syndrome, and in patients with combined emphysema and CTD-ILD, D_{LCO} levels are disproportionately reduced whilst lung volumes, including FVC, tend to be spuriously preserved. Thus, the use of standardised FVC/D_{LCO} or K_{CO} thresholds to define multi-compartment disease may have led to false positive satisfaction of IPAF criteria in some patients. Indeed, this problem may lead to the selective designation of IPAF in patients with advanced ILD or concurrent emphysema, confounding outcome comparisons between patients meeting IPAF criteria and those with IPF or idiopathic NSIP.

The identification of airway disease also poses difficulties, whether identified by ventilatory patterns or CT findings. It is often difficult to decipher whether the presence of airway disease is in favour of an underlying CTD, what exactly constitutes airway disease in subjects with fibrotic ILD, and how the confounding effect of tobacco smoking can be dealt with rigorously. An obstructive or mixed ventilatory defect may result from concurrent asthma or smoking-related chronic obstructive pulmonary disease. Furthermore, no pulmonary function threshold has been validated in the identification of airway disease when there is concurrent ILD. The presence of “standalone bronchiectasis” on CT in lung regions distant from interstitial fibrosis (as seen in rheumatoid arthritis (RA) or Sjögren disease, but quite rarely in IPF) is likely to be a robust multi-compartment criterion. However, the distinction between standalone bronchiectasis and traction bronchiectasis is not always straightforward, especially when ILD is extensive.

Thus, multi-compartment disease, when obviously present, is an important autoimmune feature but its definition will need to be carefully considered when IPAF criteria are eventually revised. We favour a robust approach in which pulmonary vasculopathy is considered to be disproportionate only when ILD is mild in severity (whether defined by CT or pulmonary function variables) and emphysema is absent or limited in extent. There is no current consensus on pulmonary function or CT extent thresholds that might be harnessed for this purpose. This will require careful expert group consideration of whether current pulmonary hypertension criteria should be discarded from the designation of multi-compartment disease in IPAF or modified to specify disproportionate pulmonary vasculopathy. Similarly, it can be argued that airway disease should be designated only when there is standalone bronchiectasis on CT or an unexplained obstructive or mixed ventilatory defect (*i.e.* in the absence of asthma or smoking-related emphysema).

Prognostic implications of the IPAF entity

Ultimately, the introduction of a new entity is useful if it provides prognostic and/or management implications. In the article by OLDHAM *et al.* [41] that quickly followed the IPAF proposal, approximately a

third of patients with an IIP or “undifferentiated” CTD met IPAF criteria, including 18% of their IPF cohort. There have since been numerous studies detailing the features and outcomes for various IPAF cohorts [20, 24–26, 42–48]. A summary and comparison of published IPAF cohorts can be found in table 4 [25]. However, despite the ERS/ATS IPAF research statement, significant cohort heterogeneity remains, resulting at least in part from the retrospective nature of most series and of varying recruitment patterns by centre. The selection of subjects to whom IPAF criteria are applied across studies has differed, as has the application of the multi-compartment morphological domain criterion. Some of the inconsistency in reporting relates to the presence or absence of certain autoantibodies in the systematic screening of patients with IIP. As a result, the various IPAF studies report on slightly different patients and their associations with outcomes.

In the cohort of OLDHAM *et al.* [41], subjects most commonly met IPAF criteria on the basis of combined serological and morphological features. Supporting the validity of the IPAF criteria was the observation that survival trended approximately halfway between that of true CTD-ILD and IPF, an observation that has been noted variably in other studies. In the study by OLDHAM *et al.* [41], once the presence of a UIP pattern was taken into account, survival of UIP-IPAF was identical to IPF, while non-UIP IPAF had outcomes closer to those of CTD-ILD, suggesting that IPAF features in the context of a UIP pattern may be of limited prognostic relevance. However, a subsequent study from the same centre, while identifying honeycombing as being independently associated with adverse survival, did not identify a clear survival difference between definite and inconsistent for UIP radiological patterns in IPAF subjects [49]. A recent systematic review and meta-analysis confirmed the poor outcome associated with a UIP pattern in the context of IPAF [50]. The review acknowledged additional recognised risk factors for IIP mortality in IPAF subjects, including age, male gender, smoking history and lung function parameters. However, on multivariable analysis, older age was the only parameter significantly associated with mortality [50].

Should a UIP pattern be excluded from the IPAF entity?

In IPAF, a UIP pattern is excluded from the morphological domain as an IPAF criterion. However, if additional features suggestive of autoimmune or multi-compartment involvement are observed, including diffuse lymphoplasmacytic infiltration, interstitial lymphoid aggregates, histological involvement of the airways, unexplained pleural disease and unexplained pulmonary vasculopathy, the IPAF morphological domain can be met even with a UIP pattern (table 1) (figure 4c and d). Furthermore, a patient with a UIP pattern and no additional morphological features can be included within IPAF if at least one item in each of the clinical and serological domains is present (table 1). In CTD-ILD, a UIP pattern is generally associated with a better survival than UIP/IPF [51–53]. RA may be an exception, as RA-ILD patients with a definite and extensive UIP pattern on CT and/or a UIP pattern on biopsy have an IPF-like survival [54–58]. The fact that, in most studies, a UIP pattern in the context of CTD-ILD is not associated with an IPF-like behaviour (with the exception of RA-ILD), supported not excluding patients with UIP (providing they met IPAF criteria) from the IPAF entity. However, in light of the poor prognostic impact of a UIP pattern within IPAF, there could be an argument for its removal. Nevertheless, there are a number of issues that will need to be considered before deciding whether IPAF-UIP needs to be considered separately [35]. Not all IPAF-UIPs are equal and demographic/clinical characteristics will impact on prognostic and management implications. Furthermore, it is possible that UIP-IPAF would not have the same outcome as IPF if the IPAF criteria were revised, with a particular focus on allowing satisfaction of the multi-compartment criterion through vasculopathy, just because an IPF patient has PH. In addition, the exact definition of increased inflammation in the context of a UIP biopsy, and whether more intense/numerous diffuse lymphoplasmacytic infiltration and/or interstitial lymphoid aggregates are associated with a better prognosis, remains unclear. Figure 3 is an example of histological UIP with areas of interstitial chronic inflammation.

How does the presence of autoimmune features influence initial evaluation?

At present, no consensus exists on best practice with regard to the investigation and initial management of patients meeting IPAF criteria. We stress that the views expressed below are the combined views of the authors.

According to studies published so far, a UIP pattern in the context of IPAF, whether on CT or at biopsy, is associated with a similar mortality to IPF [41, 44]. In the appropriate clinical context for an IPF diagnosis (age greater than 60 years, male sex and a smoking history), we believe that a probable UIP pattern on CT, even in patients meeting IPAF criteria, may not require histological confirmation of UIP, as management would not necessarily change. In contrast, the threshold for biopsy in the context of a probable UIP pattern on CT is reduced in the younger IPAF patient, particularly if female (see treatment section below). In most studies, although not in all [26], patients with non-UIP IPAF have a similar survival to CTD-ILD, and

TABLE 4 Comparison of interstitial pneumonia with autoimmune features (IPAF) cohorts

Cohort	Inclusion	Clinical	Serological	Morphological	Positive in all three domains	Treatment	Outcomes (IPAF versus other entities)
JEE <i>et al.</i> [25]	IIP, CTD-ILD, unclassifiable ILD n=228 IPAF 36 IIP 113 CTD-ILD 49 Unclassifiable 30	61.1% Arthritis 38.9% Raynaud's 25% Mechanic's hands 5.6%	83.3% ANA 16.7% RF/CCP 16.7% SSA 0% MA 44.1% ASA 35.3% ANCA 20%	86.1% Radiology NSIP 44% OP 19.4% NSIP/OP 16.7% UIP 16.7% Histology (available in 19.4%) UIP 73.5% NSIP 22.9%	30.6%	Any 83.3% IS 80.6% Antifibrotic 2.8%	TFS IIP worse versus CTD IPAF no different versus IPF or CTD-ILD PFS No difference between groups
OLDHAM <i>et al.</i> [41]	IIP or UCTD n=422 IPAF 144	49.3% Raynaud's 27.8% Arthritis 17.4% Mechanic's hands 10.4%	91.7% ANA 77.6% RF 13% CCP 4.7% SSA 16.6% ASA 0.7%	85% Radiology NSIP 31.9% OP 16.9% NSIP/OP 3.6% UIP 54.6% Histology (available in 57.6%) UIP 73.5% NSIP 22.9%	49.3%	IS 32.2%	IPAF survival worse than CTD-ILD; trend towards better than IPF Non-UIP IPAF similar to CTD-ILD; UIP-IPAF similar to IPF
CHARTRAND <i>et al.</i> [24]	Autoimmune ILD without CTD n=56 IPAF 56	62.5% Raynaud's 39% Arthritis 16.1% Mechanic's hands 28.6%	91.1% ANA 48.2% CCP 10.7% RF 10.7% SSA 42.9% ASA 36%	98% Radiology NSIP 57% OP 7.1% NSIP/OP 14.3% LIP 1.8% UIP 8.9% Histology (available in 64.3%) NSIP 33% UIP 22%	52%	IS 98.2% CS 81.8% CS+IS 76.4%	No deaths reported
AHMAD <i>et al.</i> [26]	Hospitalised with IIP or CTD-ILD n=380 IPF 156 CTD-ILD 167 IPAF 57	47.3% Raynaud's 75% Arthritis 48.1% Mechanic's hands 7.4%	93% ANA 82% RF 7.5% CCP 9.4% SSA 9.4% ASA 17%	78.9% Radiology NSIP 53% OP 3.5% NSIP/OP 15.8% LIP 1.8% UIP 28% Histology (available in 28.1%) NSIP 31% UIP 19%	Not reported	IS 28.6% CS 67.9% Antifibrotic 5.4%	IPAF survival not different to IPF

Continued

TABLE 4 Continued

Cohort	Inclusion	Clinical	Serological	Morphological	Positive in all three domains	Treatment	Outcomes (IPAF versus other entities)
Ito <i>et al.</i> [20]	IIP, serological and morphological domains met n=98 IPAF 98	Not reported	100% ANA 28.6% RF 28.5% CCP 15.3% SSA 18.4% ASA 13.3%	100% Radiology NSIP 64.3% OP 20.4% NSIP/OP 15.3% Histology (available in 17.3%) NSIP 20.6% OP 5% NSIP/OP 20% UIP 3.1%	Not applicable	CS 27.6% CS+IS 44.9%	Not applicable
Dai <i>et al.</i> [46]	All ILD n=1429 IPAF 177 Other ILDs 252	20.3% Raynaud's 12.9% Arthritis 4.5% Mechanic's hands 1.7%	92.1% ANA 49.2% RF 12.4% CCP 5.6% SSA 36.1% ASA 3.9%	95.5% Radiology NSIP 61.6% OP 22% NSIP/OP 11.9% UIP 4.5%	Not reported	CS 72.3%	IPAF worse survival than non-IPAF
YOSHIMURA <i>et al.</i> [47]	Chronic fibrotic ILD n=194 IPF 163 NSIP 31 IPAF 32	53.1% Arthritis 28.1% Raynaud's 18.8% Mechanic's hands 12.5%	71.9% ANA 28.1% RF 21.9% CCP 9.4% SSA 9.4% ASA 6.3%	96.9% Radiology NSIP 40.6% NSIP/OP 12.5% Histology NSIP 59.4%	21.9%	CS 59.4% IS 34.4% Antifibrotic 25%	IPAF survival better than non-IPAF IPAF-NSIP better survival than non-IPAF-NSIP IPAF-NSIP and IPAF-IPF similar outcomes
KELLY and MOUA [44]	Defined UCTD-ILD n=151 IPAF 101 IPF 50	Not reported Raynaud's 54.5% Arthritis 29.7% Mechanic's hands 10.9%	Not reported ANA 18.8% RF 18% CCP 3.8% SSA 35.6% ASA 1%	Not reported Radiology NSIP 64.4% OP 3.9% NSIP/OP 4% LIP 2% UIP 11.9% Histology (available in 50.5%) NSIP 13.7% UIP 23.5% OP 23.5% Non-diagnostic 37.2%	Not reported	Not reported	IPAF survival better than IPF IPAF-UIP worse survival than IPAF-non-UIP IPAF-UIP similar to IPF
LIM <i>et al.</i> [48]	All ILD n=305 IPAF 54 CTD-ILD 76 IPF 175	31.5% Raynaud's 17.6% Arthritis 76.5% Mechanic's hands 5.9%	90.7% ANA 63.3% RF 28.6% CCP 14.3% SSA 8.2%	81.5% Radiology NSIP 63% OP 7.7% NSIP/OP 5.1% UIP 25.9%	Not reported	Not reported	IPAF survival better than IPF, worse than CTD-ILD

ANA: anti-nuclear antibody; ANCA: anti-neutrophil cytoplasmic antibody; ASA: anti-synthetase autoantibodies; CCP: cyclic citrullinated peptide; CS: corticosteroid; CTD: connective tissue disease; CTD-ILD: connective tissue disease-associated interstitial lung disease; IIP: idiopathic interstitial pneumonia; ILD: interstitial lung disease; IPF: idiopathic pulmonary fibrosis; IS: immunosuppression; LIP: lymphocytic interstitial pneumonia; MA: myositis autoantibodies; NSIP: non-specific interstitial pneumonia; OP: organising pneumonia; PFS: progression-free survival; RF: rheumatoid factor; SSA: Sjögren syndrome-related antibody A; TFS: transplant-free survival; UCTD: undifferentiated connective tissue disease; UIP: usual interstitial pneumonia. Reproduced and modified from JEE *et al.* [25] with permission.

most ILD specialists would probably treat in a similar manner, at least initially. In our opinion, while identification of a histological NSIP pattern would support an immunosuppressive approach, histological confirmation of a UIP pattern would lead to a more cautious approach to immunosuppression, with frequent monitoring and earlier consideration of antifibrotic therapy, should there be progression. Finally, the finding of associated morphological features of IPAF on a biopsy otherwise characterised by a UIP pattern, would lead to further interrogation and monitoring over time to detect additional features suggestive of an underlying CTD. Whether cryobiopsy samples, rather than those obtained through a surgical biopsy, are sufficient to detect these additional features in the context of IPAF has not been established, and the decision as to which biopsy procedure to undertake and whether to proceed to surgical biopsy if less invasive techniques are inconclusive needs to be made on a case-by-case basis. In light of the many complexities in defining IPAF criteria, these cases will always benefit from discussion by a multidisciplinary team, with input from pulmonologists, radiologists, pathologists, rheumatologists and, in some instances, immunologists.

Monitoring of autoimmune features over time

ILD may be the initial manifestation of a CTD, and it is therefore possible for patients with ILD, including IPAF, to manifest a defined CTD at a later timepoint (figure 4e) [59]. SEBASTIANI *et al.* [22] observed IPAF evolution to definite CTD in 13.5% of cases. A definite CTD diagnosis of Sjögren disease, RA and polymyositis was possible after a mean of 31 months (range 7–71 months). In two cases, these diagnoses were made after the development of relevant antibody positivity which was absent at baseline. In a cohort of patients with an initial diagnosis of idiopathic NSIP, 10% developed clinical manifestations of a defined CTD at a median of almost 2 years after their ILD diagnosis [60]. It is important to recognise that IPAF is not a prerequisite for the subsequent development of a CTD.

All patients with an ILD should be routinely evaluated for the presence of CTD symptoms and signs, with re-evaluation occurring at each visit and for the duration of follow-up, even in the case of a highly confident diagnosis of IPF. In cases of IPAF, the treating clinician should continually search for those features which will add to the CTD diagnostic puzzle. Cases which satisfy IPAF criteria should be considered for formal rheumatological evaluation, as a defined CTD, following established criteria (table 3), will better inform treatment decisions. The goals are to achieve a highly confident diagnosis and to inform a patient of the aetiology, natural history and treatment of their disease. However, lower degrees of diagnostic confidence are often the best that can be achieved at presentation, with the need for an iterative diagnostic approach, and repeated multi-disciplinary discussions as needed, taking into account response to initial treatment, disease behaviour and development of additional features over time. Re-evaluation with repeat anti-nuclear antibody (ANA) testing is unlikely to be informative [61], noting that a positive ANA on its own, even when in an anti-centromere pattern, is not sufficient for systemic lupus erythematosus or SSc. However, repeating extended antibody testing should be considered, particularly in those with radiological/histological phenotypes of NSIP, organising pneumonia (OP) or a combination of both, and/or clinical features suggestive of, but not confirmatory for, SSc or IIM. Involvement of an immunologist may be necessary in such a scenario, in order not to miss the rare autoantibodies that are not systematically assessed. The development of an inflammatory arthralgia in a patient with ILD should herald the potential development of RA, noting that positive serology is not necessary for a diagnosis (figure 4e). Just as important as the search for definite CTD features are the efforts to exclude a CTD, particularly in the case of UIP radiology. Where the IPAF criteria are satisfied only for low titre ANA and perhaps borderline multi-compartment involvement (figure 4c and d), a diagnosis of IPF should be considered, although perhaps with a degree of confidence.

Treatment of IPAF

As IPAF is a research entity and not a diagnosis, studies specifically addressing the management of IPAF are lacking. There have been no randomised controlled trials specifically conducted in IPAF. Management strategies must therefore be based on expert opinion and are largely extrapolated from CTD-ILD studies [17, 62]. It is uncertain whether a specific management distinct from that of IPF and from CTD-ILD is needed; however, research is required to address this question. In the absence of a general approach that could be applied to any patient with lung disease fitting the definition of IPAF, management is individualised.

Similarly to other ILDs, especially IPF [63], pulmonary rehabilitation, long-term oxygen supplementation therapy if appropriate, and treatment of gastro-oesophageal reflux if present, are indicated, as well as prevention of infection and bone health assessment. In advanced disease, lung transplantation should be considered, extrapolating satisfactory outcomes in appropriately selected patients with CTD, including SSc [64].

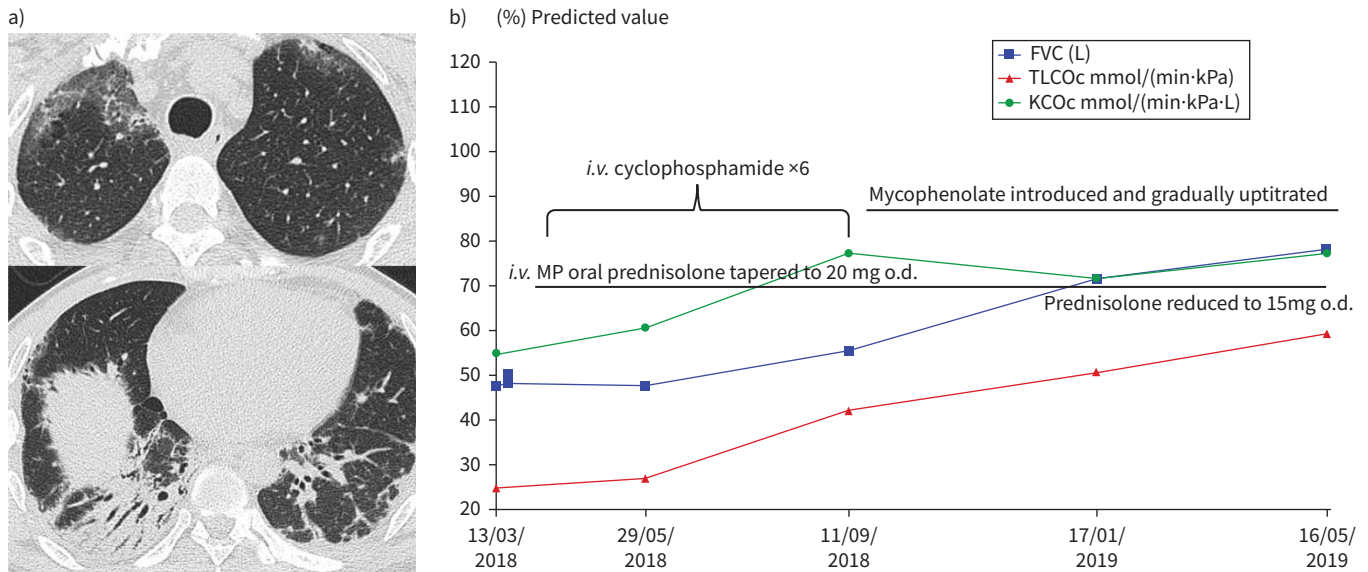


FIGURE 5 Interstitial pneumonia with autoimmune features (IPAF) in a 32-year-old male presenting with subacute cough and dyspnoea over 2 months. On presentation, he was in type 1 respiratory failure requiring supplemental oxygen. **a)** His computed tomography (CT) demonstrated upper zone ground glass opacity and fibrotic organising pneumonia in the lung bases. Anti-nuclear antibody was positive at 1:1280, although extended serology was negative. There were no clinical features of an autoimmune condition. IPAF criteria were met by morphological and serological domain. **b)** A significant response was observed with immunomodulation, with ongoing lung function improvement over a protracted period. FVC: forced vital capacity; *i.v.*: intravenous; KCOc: gas transfer coefficient, corrected for haemoglobin; MP: methylprednisolone; o.d.: once daily; TLCoc: transfer factor of the lung for carbon monoxide, corrected for haemoglobin.

In the series of IPAF published to date where treatment is reported [23, 25], a majority of patients have received glucocorticoids and/or immunosuppressive therapy, including mycophenolate, azathioprine, cyclophosphamide, calcineurin inhibitors (cyclosporine and tacrolimus) and occasionally rituximab, illustrating that many clinicians would treat subjects with IPAF as they would treat CTD-ILD [65–68]. Figure 5 depicts a case meeting IPAF criteria treated similarly to CTD-ILD with pulsed methylprednisolone and cyclophosphamide, followed by maintenance mycophenolate and prednisolone. A small minority of patients in published series have received antifibrotic agents [25], which may reflect the fact that until recently these drugs were only approved and available for patients with IPF. In these series, however, efficacy of management is virtually impossible to assess in the absence of a control group and given the high clinical heterogeneity.

There are data stemming from studies of patients with unclassifiable ILD, some of which include patients with IPAF. In one study of patients with unclassifiable ILD, intravenous pulse cyclophosphamide was suggested to stabilise lung function [69]; a subset of patients in this study had IPAF and these seemed to benefit more from the treatment regimen, although none of them had a UIP pattern. This suggests that patients with IPAF and a non-UIP pattern may benefit from immunomodulation, although this needs confirmation.

In a retrospective cohort of 52 patients who met criteria for IPAF, 28 received mycophenolate after a median time of 22 months [70]. Although changes in FVC% and D_{LCO} % were not significantly different between the mycophenolate-treated and untreated groups, there was a trend toward more rapid baseline decline of both FVC% and D_{LCO} % in the mycophenolate-treated cohort before *versus* after mycophenolate therapy. This suggests that patients with IPAF might benefit from mycophenolate therapy; however, this view is again subject to many possible biases.

In a multicentre, double-blind, randomised, placebo-controlled phase 2 trial, 253 patients who had progressive fibrosing unclassifiable ILD were randomised to receive pirfenidone *versus* placebo [71]. Randomisation was stratified by concomitant mycophenolate mofetil use and presence or absence of IPAF. Analysis of the primary endpoint, the mean predicted change in FVC from baseline over 24 weeks measured by daily home spirometry, was affected by intraindividual variability, which prevented application of the prespecified statistical model. Over 24 weeks, predicted mean change in FVC measured

by site spirometry was lower in patients given pirfenidone than placebo (treatment difference 95.3 mL, 95% CI 35.9–154.6; $p=0.002$). Compared with the placebo group, patients in the pirfenidone group were less likely to have a decline in FVC of more than 5% or more than 10%. At week 24, mean change in D_{LCO} from baseline was -0.7% (standard deviation 7.1) for the pirfenidone group and -2.5% (8.8) for the placebo group, and mean change in 6-min walk distance from baseline was -2.0 m (68.1) for the pirfenidone group and -26.7 m (79.3) for the placebo group. Changes from baseline in University of California San Diego–Shortness of Breath Questionnaire, Leicester Cough Questionnaire score, cough visual analogue scale and St George’s Respiratory Questionnaire scores were similar between the pirfenidone and placebo groups at week 24. The safety and tolerability profile was acceptable. This study suggests that patients with progressive fibrosing unclassifiable ILD could benefit from pirfenidone treatment. Only 33 patients with IPAF were included in this study, precluding any conclusion regarding this subgroup.

Nintedanib, a tyrosine kinase inhibitor with antifibrotic properties, has demonstrated efficacy in IPF (the INPULSIS studies) [72], in SSc-associated ILD (the SENSICIS study) [73] and in patients with ILD and a progressive fibrosing phenotype (the INBUILD study) [74]. In the SENSICIS study, approximately half of the patients were also receiving mycophenolate, demonstrating that nintedanib can be combined with mycophenolate with acceptable tolerance. In the INBUILD study, 114 participants (17%) had unclassifiable ILD; however, the proportion of them who met IPAF criteria is unknown. Sensitivity analysis of the annual rate of decline in FVC ($\text{mL}\cdot\text{year}^{-1}$), excluding aetiological subgroups of ILD diagnosis, demonstrated homogeneity of results across subgroups [75]. Although the INBUILD trial was not designed or powered to provide evidence for a benefit of nintedanib in any specific diagnostic subgroup, the results suggest that nintedanib reduces FVC decline in patients who have a chronic fibrosing ILD and progressive phenotype, irrespective of the underlying ILD diagnosis. It can be inferred that subjects with IPAF who have a progressive phenotype can benefit from nintedanib treatment.

Overall, possible treatment decisions in patients classified as with IPAF include observation without drug therapy, immunomodulation with glucocorticoids and/or immunosuppressants, antifibrotic drugs, and a combination (upfront or sequential) of immunomodulation and antifibrotics. Currently, treatment decisions must be based on a careful evaluation of benefit to risk ratio in the individual subject and should be discussed in a multidisciplinary setting. Eventually, the choice of first-line therapy is based on the global assessment of the patient, including the age and gender of the patient, the type of extrapulmonary manifestations present, the underlying CTD for which there is a “flavour”, the chest CT pattern and the histopathological pattern if available, the severity of functional impairment, the existence of disease progression, the patient’s expectations, the comorbidities and co-medications, and the time elapsed since the diagnosis of ILD (figure 6). Of those, the demographics and the imaging phenotype are often the main elements to take into consideration, separating a predominantly fibrosing from a predominantly inflammatory phenotype.

As an example, a 65-year-old male patient with a chest CT pattern of NSIP and a UIP pattern on biopsy might not benefit from corticosteroids and/or immunosuppressive drugs even if lymphocytic follicles are present on biopsy and high titre of antinuclear antibodies with nucleolar pattern are found; a sensible decision in such case may be to observe and monitor disease progression, and later to treat using antifibrotics, in case the clinical behaviour is that of a progressive phenotype. Conversely, a 50-year-old female with a chest CT suggesting NSIP and a high titre of antinuclear antibodies with nucleolar pattern, who declined lung biopsy, will likely be treated similarly to SSc-associated ILD even in the absence of any other manifestation of CTD. Similarly, a chest CT pattern of OP associated with NSIP is highly suggestive of autoimmune myositis and is likely to be managed as such by most ILD experts, even in the absence of autoantibodies identified precluding the classification of such cases as autoimmune myositis.

Next to these extreme examples, many cases fall into a spectrum that encompasses both inflammatory/autoimmune and fibrotic features, hence presenting very challenging management decisions. In the future, it is likely that a combination of approaches [76] may be used in many patients with IPAF. One key element to keep in mind is the detrimental effect of high-dose glucocorticoids and azathioprine therapy in patients with IPF, potentially applicable also to IPAF patients with UIP. Furthermore, as IPAF is not a validated as a standalone diagnosis, our view is that in most cases we should, for management purposes, view IPAF-UIP as synonymous with IPF. The designation of IPAF as a diagnosis deprives these patients of access to antifibrotic therapy at presentation. This is a particular problem if pulmonary vasculopathy is viewed as a separate process in IPF and is the basis for the designation of multi-compartment disease. As discussed earlier, OLDHAM *et al.* [41] classified UIP patients as having IPAF based on an inflexible FVC/ D_{LCO} ratio and one other criterion. In the majority of cases, the existing validated diagnosis in patients



FIGURE 6 Factors to consider in the management of patients fulfilling interstitial pneumonia with autoimmune features criteria. CTD: connective tissue disease; CT: computed tomography.

with IPAF-UIP is IPF, with the crucial consequence that these patients are eligible for antifibrotic therapy at presentation.

Progress in management is critically dependent on randomised clinical trials and of prospective registry studies dedicated specifically to patients with IPAF, with the aim of guiding individualised medicine based on better phenotyping and more precise characterisation of all features present.

Conclusion

The authors who collaborated on the ERS/ATS IPAF research statement indicated upfront that this represented a “first draft”, to be considered as a starting point for future work. It is not surprising, therefore, that numerous suggestions have been proposed to improve and refine the IPAF criteria. There remains an important ongoing need for the IPAF entity, serving a valuable purpose in identifying those cases with autoimmune features that do not meet criteria for defined CTDs. Much more is known about these cases than in 2015 when IPAF was proposed, and it might now be time to work on a “second draft” of the ERS/ATS IPAF statement.

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