Elevated APACHE score, hypoxaemia and mechanical ventilation are risk factors for mortality in ILD. No increased mortality was found with steroid use. Evidence on advanced age is inconclusive. Survival may be improving over time, but still remains high.

ABSTRACT Data on interstitial lung disease (ILD) outcomes in the intensive care unit (ICU) is of limited value due to population heterogeneity. The aim of this study was to examine risk factors for mortality and ILD mortality rates in the ICU.

We performed a systematic review using five databases. 50 studies were identified and 34 were included: 17 studies on various aetiologies of ILD (mixed-ILD) and 17 on idiopathic pulmonary fibrosis (IPF). In mixed-ILD, elevated APACHE score, hypoxaemia and mechanical ventilation are risk factors for mortality. No increased mortality was found with steroid use. Evidence is inconclusive on advanced age. In IPF, evidence is inconclusive for all factors except mechanical ventilation and hypoxaemia. The overall in-hospital mortality was available in 15 studies on mixed-ILD (62% in 2001–2009 and 48% in 2010–2017) and 15 studies on IPF (79% in 1993–2004 and 65% in 2005–2017). Follow-up mortality rate at 1 year ranged between 53% and 100%.

Irrespective of ILD aetiology, mechanical ventilation is associated with increased mortality. For mixed-ILD, hypoxaemia and APACHE scores are also associated with increased mortality. IPF has the highest mortality rate among ILDs, but since 1993 the rate appears to be declining. Despite improving in-hospital survival, overall mortality remains high.

Introduction

The interstitial lung diseases (ILD) encompass over 200 different entities, sharing common clinical features of progressive inflammation with variable scarring of the interstitium and alveoli [1]. The associated deterioration in lung function leads to both substantial morbidity and mortality. Acute exacerbation of ILD is defined as an acute respiratory deterioration characterised by evidence of new widespread alveolar
infiltrates [2]. While best characterised in idiopathic pulmonary fibrosis (IPF), acute exacerbations are relevant in other forms of ILD and often lead to intensive care unit (ICU) admission [3]. Notably, progression of disease and infections are the most common causes of ICU admissions in patients with IPF [3]. However, there is a paucity of data on ICU outcomes in patients with ILD. Research on ICU mortality has been complicated by evolving diagnostic guidelines [4–6], which has led to significant variation in inclusion criteria from one study to the next. Ultimately, this makes it difficult for clinicians to provide relevant information on prognosis and outcomes to patients based on findings in a given study.

IPF is the most common form of idiopathic interstitial pneumonia (IIP) [5]. Mallick et al. [7] reported an overall ICU mortality of 87% in ventilated patients from studies published before 2006. Many studies in the ILD literature focus only on ICU outcomes in IPF because this diagnosis generally carries the worst prognosis in ILD [8]. However, it is not clear that other forms of ILD confer a similar prognosis. Making this distinction is difficult, but clinically important as articles considering ICU outcomes in mixed-ILD populations often include patients with IPF.

Given the heterogeneity of disease and diagnosis, various stages of disease and the myriad of triggers for ICU admission [9], our objective was to review current knowledge on variables associated with increased risk of mortality in the ICU, evaluate the factors that affect clinical outcomes in the ICU, determine mortality rates and identify important areas for future research.

Methods

Search strategy and selection criteria
We conducted a systematic review with relevant papers identified via PubMed, Scopus, Cochrane, Embase and Scielo searches for English and Spanish articles with the keywords "ILD", "interstitial pneumonia", "interstitial pneumonitis", "IIP", "IPF", "intensive care unit" and "outcomes", among other terms (supplementary material e-table 1). Our target population was adult patients with ILD admitted to the ICU for whom mortality data were available. Patients who presented in acute respiratory failure and then underwent a lung biopsy during their ICU stay were included. Patients admitted to the ICU due to complications from elective lung biopsy were excluded from this review. Studies on patients with acute interstitial pneumonia were also excluded as prior review articles on this topic have been previously published [10, 11].

The complete search strategy was performed on November 21, 2017. All titles and abstracts were evaluated for further review independently by two authors (figure 1). A total of 348 unique articles were reviewed. Of these, 298 were excluded because: ICU outcomes were not evaluated; different target populations were included (acute respiratory distress syndrome, lung transplant, children, etc.); they consisted of case reports of less than three patients; or were letter to the editor and review articles. Conference abstracts were not reviewed if they had not been published in the aforementioned databases. Of 50 articles reviewed in full text, 16 articles were excluded (lung biopsy for ILD: n=4; unknown lung infiltrates in the ICU: n=4; polymyxin and extracorporeal membrane oxygenation: n=2; acute interstitial pneumonia: n=2; coal worker pneumoconiosis: n=1; acute respiratory distress syndrome: n=1; myositis: n=1; and use of bronchoscopy: n=1). In total, 34 studies were identified for inclusion in this review (table 1). Studies of patients with IPF were separated from those with heterogeneous patient population (mixed-ILD). 17 papers included only IPF patients; the remaining 17 included mixed-ILD.

Data extraction and quality assessment
A data extraction form was developed for the purpose of the review and included research design, study sample, source of bias, data collection and follow-up rates. We did not use other tools available in the literature given that all the studies, except for one, presented retrospective information based on series of cases without a comparator group. Two authors (J.A. Huapaya and E.M. Wilfong) assessed the relevance of articles independently. Queries were resolved through discussion with the senior author (S.K. Danoff). No meta-analysis was considered because of the heterogeneity of the population included in the studies, lack of consistent evaluation of all the risk factors for mortality (the majority of studies only reported information on a few risk factors and did not mention others), lack of consistent analysis (only a few studies performed multivariate analysis) and a wide range of years included in the analysis of each study.

Mortality rates estimation
Mortality rates were extracted from each study at different time-points: ICU, in-hospital and follow-up mortality (3, 6 and 12 months). For simplicity of analysis, overall mortality included the available in-hospital and ICU mortality. Mean percentage of mortality rates was not used for the calculation of overall mortality as this could have represented a source of bias given different sample sizes. Therefore, the
total number of deaths was calculated for each study and was then divided by the total number of patients at risk.

Results

One prospective and 33 retrospective studies (one that also reports 10 patients who were prospectively evaluated) [12] were included in the analysis (table 1). 26 studies included less than 50 patients, six studies included between 50 and 100 patients, and two studies included more than 100 patients. The largest cohort had 209 patients and the smallest one had only three patients. IPF specific data from the studies with mixed population was not available to be added to the IPF specific studies. Mixed-ILD patients included IPF, connective tissue disease (CTD)-ILD, drug-induced ILD, hypersensitivity pneumonitis, and

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other aetiologies. Overall in-hospital mortality according to geographic regions is presented in supplementary material e-table 5.

**Mortality in mixed-ILD studies**

Our review identified 780 patients with mixed causes of ILD in 17 studies [12–28]. 15 studies had information on ICU or in-hospital mortality, reporting a total of 685 patients. The remaining two studies had mortality at post-hospitalisation follow-up (2 and 3 months, respectively) [24, 25]. ICU admission occurred due to acute respiratory failure secondary to infections and disease progression in 30–75% of cases. 5–10% of admissions occurred due to pneumothorax, pulmonary embolism, congestive heart failure, were drug induced or due to other causes. The overall in-hospital mortality (15 studies) was 52%. Notably, half of these patients had IPF (table 2). Stratifying the studies according to their year of publication showed the overall in-hospital calculated mortality was 62% during the period 2001–2009 and 48% during 2010–2017 (supplementary material e-table 3). Cut-offs for different years were used to evaluate if this trend was consistent throughout different periods (supplementary material e-table 4). Notably, mortality increases with time post hospitalisation (table 2). Some studies reported 1-year mortality of 54–59% [14, 28], whereas others reported 1-year mortality of 85–96% [12, 16].

**Mortality in IPF studies**

17 studies of IPF included 554 patients (table 2) [3, 29–44]. 15 of these studies had information on ICU or in-hospital mortality accounting for 526 patients; the remaining two studies had mortality at post-discharge follow-up (2 and 3 months, respectively) [29, 44]. The overall in-hospital mortality of the 15 studies was 68%, which is lower than previously reported [7]. Notably, studies from the 1990s reported mortality rates around 80–100% [30, 32, 36], whereas recent studies in large populations reported
mortality rates of 50–66% [37, 41]. Reasons for ICU admissions were similar as for mixed-ILD. Stratifying the 15 studies according to their year of publication, the overall in-hospital mortality was 79% during the period 1993–2004 and 65% during 2005–2017 (supplementary material e-table 3). Similarly to mixed-ILD studies, different year cut-offs were used to repeat the analysis (supplementary material e-table 4). Despite improving overall survival, mortality increased over time post hospitalisation as two studies reported 100% mortality at 1-year of follow-up (table 2) [30, 43]. Notably, the cause of death post hospitalisation is not available and, therefore, it is not possible to assign these deaths directly to IPF.

### Variables associated with increased mortality

The current evidence on different factors and their association with mortality is presented in the following sections (table 3).

| First author [ref.] | Inclusion years | Total included | IPF | Overall deaths | ICU mortality % | In-hospital mortality % | Overall in-hospital mortality % |
|---------------------|-----------------|----------------|-----|----------------|---------------------|-----------------------------|---------------------------------|
| **Mixed-ILD**        |                 |                |     |                |                     |                             |                                 |
| CHURG [13]          | Not provided    | 12             | 12  | 9              | 2                   | 13                          | 36 (11 months)                |
| FERNANDEZ-PEREZ [14]| 02/2002–07/2006 | 94             | 94  | 30             | 50                  | 53                          | 59 (1 year)                   |
| FUMEAUX [15]        | 12/1996–03/2001 | 14             | 14  | 11             | 14                  | 100                         | 100                            |
| GAUDRY [16]         | 01/2002–04/2009 | 27             | 27  | 22             | 23                  | 77                          | 85                            |
| GONZALEZ [17]       | 01/2004–05/2015 | 37             | 37  | 35             | 23                  | 62                          | 62                            |
| GUNDO [18]          | 01/2008–12/2010 | 120            | 120 | 96             | 60                  | 60                          | 60                            |
| HUÉ [12]            | 06/2006–04/2009 | 27             | 27  | 13             | 17                  | 63                          | 63                            |
| MARTINEZ [19]       | 12/2007–12/2013 | 36             | 36  | 25             | 25                  | 55                          | 69                            |
| MOLINA/MOLINA [20]  | 01/1986–06/2002 | 20             | 20  | 14             | 20                  | 95                          | 100                            |
| OLSON [21]          | 01/2005–01/2007 | 4              | 4   | 4              | 4                   | 100                         | 100                            |
| OSAMA [22]          | 01/2013–07/2015 | 91             | 91  | 68             | 9                   | 10                          | 10                            |
| PARK [23]           | 1993–04/2006    | 10             | 10  | 0              | 0                   | 60                          | 60                            |
| TACHIBANA [24]      | 1/2003–12/2009  | 62             | 62  | NI             | NI                  | 90                          | 90 (3 months)                 |
| TONG [25]           | 10/2001–09/2006 | 33             | 33  | NI             | NI                  | 44                          | 44 (2 months)                 |
| VAL-DUPUY [26]      | 7/1993–12/2009  | 72             | 72  | 16             | 33                  | 46                          | 46                            |
| YOKOYAMA [27]       | 01/2005–12/2010 | 38             | 38  | 16             | 21                  | 54                          | 54                            |
| ZAFRANI [28]        | 01/2002–03/2013 | 83             | 83  | 9              | 34                  | 41                          | 41                            |
| **Total**           |                 | 780            | 685 | 334            | 353                 |                             |                                 |
| **IPF**             |                 |                |     |                |                     |                             |                                 |
| AKIRA [29]          | Not provided    | 17             | 17  | NI             | NI                  | 48                          | 48 (3 months)                 |
| AL-HAMEED [30]      | 09/1988–12/2001 | 25             | 25  | 25             | 24                  | 84                          | 96                            |
| AMBERGON [31]       | 05/2001–06/2002 | 5              | 5   | 5              | 4                   | 80                          | 80 (1.5 years)                |
| BUKET [32]          | 01/1989–06/1998 | 15             | 15  | 15             | 13                  | 73                          | 86                            |
| KIM [33]            | 01/1990–09/2003 | 11             | 11  | 11             | 7                   | 70                          | 63                            |
| KONDOR [34]         | Not provided    | 3              | 3   | 3              | 0                   | 0                            | 0                             |
| MOLICA [35]         | 01/2000–01/2007 | 34             | 34  | 34             | 29                  | 85                          | 85 (6 months)                 |
| NANA [36]           | Not provided    | 7              | 7   | 7              | 6                   | 85                          | 85                            |
| ODA [37]            | 04/2010–03/2013 | 209            | 209 | 209            | 138                 | 66                          | 66                            |
| PAPRII [38]         | 2007–06/2013    | 17             | 17  | 17             | 11                  | 65                          | 65                            |
| PARAM [39]          | 01/1996–12/2006 | 7              | 7   | 7              | 6                   | 86                          | 86 (8 months)                 |
| RANGAPPA [3]        | 01/1996–12/2006 | 24             | 24  | 24             | 22                  | 67                          | 92                            |
| SATYAN [40]         | 01/1995–07/2000 | 38             | 38  | 38             | 23                  | 45                          | 61                            |
| SONG [41]           | 1990–2009       | 90             | 90  | 90             | 45                  | 50                          | 50                            |
| STERN [42]          | 09/1990–10/1999 | 23             | 23  | 23             | 22                  | 96                          | 96                            |
| VIANELLO [43]       | 01/2005–04/2013 | 18             | 18  | 18             | 10                  | 55                          | 55 (3 months)                 |
| YOKOYAMA [44]       | 04/1998–06/2004 | 11             | 11  | NI             | NI                  | 55                          | 55 (3 months)                 |
| **Total**           |                 | 554            | 526 | 526            | 360                 |                             |                                 |

Data are presented as n, unless otherwise stated. NI: not included in the overall mortality analysis. *, patients with radiological diagnosis of usual interstitial pneumonia who were not labelled as IPF in the original study but were considered IPF for our analysis; #: these studies provided information on follow-up mortality only; +: all the patients of 17 studies included in the systematic review; §: number of patients included in the ICU and in-hospital mortality analysis, [24, 25, 29, 44] were excluded.
| Risk factor                              | Studies in mixed-ILD patients | Mortality association | Studies in IPF patients | Mortality association |
|-----------------------------------------|------------------------------|-----------------------|-------------------------|-----------------------|
| Advanced age                            | 2 positive studies [14, 27], 5 negative studies [12, 18, 22, 26, 28⁺] | Inconclusive: 1 large study in each side | 1 positive study [37], 4 negative studies [3, 41, 43⁺, 44] | Inconclusive: 1 large study found an association; another large study and four small studies did not |
| Elevated APACHE score                   | 3 positive studies [12, 14, 18], 3 negative studies [17, 26, 27⁺] | Large studies report a significant association | 1 positive study [3], 2 negative studies [43⁺, 44] | Inconclusive: studies with small sample size |
| Histology showing diffuse alveolar damage* | 3 studies [12, 13, 21] | Small studies report high mortality in the univariate analysis; not enough data to assess for an association | 3 studies [29, 31, 34] | Small studies report high mortality in the univariate analysis; not enough data to assess for an association |
| Hypoxaemia                              | 3 positive studies [14, 22⁺⁺, 27⁺⁺], 1 negative study [18⁺⁺] | Large studies report a significant association | 3 negative studies [41⁺, 43⁺, 44] | No association was found; 2 studies with <20 patients each |
| Mechanical ventilation⁺                 | 6 positive studies [12, 14⁺⁺, 17, 18, 26, 27], 2 negative studies [22, 28] | Multiple studies report high mortality in the univariate analysis; 6 studies report a significant association | Multiple studies report high mortality in the univariate analysis; not enough data to assess for an association |
| Steroids                                | 3 negative studies [14, 26⁺⁺, 28⁺⁺] | No association was found in 3 studies; 1 reported use of steroids was a protective factor | 1 negative study [41] | Inconclusive: only 1 study with a small sample size is available |

APACHE: Acute Physiology And Chronic Health Evaluation; NIMV: noninvasive mechanical ventilation.⁺⁺: no bivariate analysis is provided due to the small sample sizes and high mortality reported in these studies.⁺⁺⁺: high rates of mortality associated with mechanical ventilation (70–100% for IPF and 27–100% in mixed-ILD) precluded bivariate or multivariate analysis in more studies other than those reported in this table.⁺⁺⁺: an association was found in the bivariate analysis but not in the multivariate.⁺⁺⁺: no direct comparison between survivors and nonsurvivors. The study primary compared NIMV success group versus NIMV failure group. However, no patient died in the former group and all 10 died in the latter group.⁺⁺⁺: a trend towards significance.⁺⁺⁺: this study reported significantly lower arterial oxygen saturation but comparable arterial oxygen tension in nonsurvivors versus survivors [bivariate analysis only].⁺⁺⁺: significantly higher arterial oxygen tension/inspired oxygen fraction ratio in survivors versus nonsurvivors at the start of NIMV but not on admission.⁺⁺⁺⁺: higher positive end-expiratory pressure was associated with increased mortality.⁺⁺⁺⁺: this study showed that the use of steroids was a protective factor in the multivariate analysis.
Age
Increased age has been evaluated as a risk factor of ICU mortality, especially in large studies [14, 18, 22, 26, 28, 37, 41]. In mixed-ILD, using multivariate analysis, Fernandez-Perez and co-workers [14] showed increasing age was a risk factor for in-hospital mortality (OR 1.07, 95% CI 1.01–1.14) and 1-year mortality (HR 1.03, 95% CI 1.00–1.05) in patients who required invasive mechanical ventilation (IMV). Yokoyama and co-workers [27] demonstrated a significant association between younger age and hospital survival when evaluating patients with rapidly progressing interstitial pneumonia regardless of the cause of the underlying ILD. However, five other studies failed to replicate those results [12, 18, 22, 26, 28].

Data on age as a risk factor for mortality in IPF is also mixed. Three small studies of less than 25 patients each did not find an association between age and mortality [3, 43, 44]. However, in a study of 209 patients, Oda and co-workers [37] reported that patients aged >80 years had significantly increased ICU mortality compared to those aged <60 years (OR 2.94, 95% CI 1.04–8.303). The study by Song and co-workers [41] does not support this finding when analysing age during acute exacerbation in 209 patients (OR 1.022, 95% CI 0.969–1.078), but demonstrates that age was a prognostic factor on overall IPF survival after diagnosis (HR 1.017, 95% CI 1.001–1.033).

Hypoxaemia
In mixed-ILD, three studies [14, 22, 27] found an association between hypoxaemia and mortality and one study had a trend towards association [18]. Fernandez-Perez and co-workers [14] reported a significant difference (p<0.001) in arterial oxygen tension/inspiratory oxygen fraction (\(\text{PaO}_2/\text{FiO}_2\)) ratio between survivors (median [interquartile range, IQR] 252 (158–388)) and nonsurvivors (median [IQR] 117 (71–182)). In the multivariate analysis, each 10-unit increase in the \(\text{PaO}_2/\text{FiO}_2\) ratio was protective for in-hospital mortality (OR 0.96, 95% CI 0.92–0.99) and 1-year survival (HR 0.99, 95% CI 0.97–1.00).

In IPF studies, three studies failed to show an association between hypoxaemia and mortality [41, 43, 44]. The size of two studies was very small (<20 patients) and the third study (n=90) found an association in the bivariate but not the multivariate analysis.

Acute physiology and chronic health evaluation (APACHE) scores
APACHE scores predicts ICU mortality based on the first ICU day [45]. Six studies in mixed-ILD patients reported APACHE scores and hospital mortality: three studies found a positive association [12, 14, 18] and one study showed a trend towards association [27]. Hue and co-workers [12] report that an increase of 1 point in APACHE II score was associated with an OR of 1.05 (95% CI 1.00–1.11, p=0.040) in 27 patients with fibrotic lung disease. Gungor and co-workers [18] report a median APACHE score of 19 (IQR 16–23) in survivors and 29 (IQR 23–36) in nonsurvivors (p=0.001). Similarly, Fernandez-Perez and co-workers [14] evaluated mortality risk using the APACHE III scores and found a HR of 1.33 (95% CI 1.18–1.50) with regards to hospital mortality (median of 57 in survivors and 78 in nonsurvivors). While not reaching statistical significance, Yokoyama and co-workers [27] showed a trend in association between APACHE II score and mortality with a mean score of 9.5 for survivors and 11.4 for nonsurvivors (p=0.056). It is important to note that age and \(\text{PaO}_2\), identified above as independent risk factors, are both components of the APACHE score and, thus, there may be overlap in their predictive roles. Two small studies found no association between APACHE and mortality [17, 26].

In IPF studies, only Rangappa and Moran [3] found significantly higher APACHE III scores in nonsurvivors (HR 1.02, 95% CI 1.004–1.031). They reported a 2% increase in hazard per unit increase in APACHE III score. However, two small studies including 29 patients did not find an association [43, 44].

Histology
Older studies report a high number of biopsies (50–100%) with diffuse alveolar damage and suggest it was a poor prognostic factor in patients with ILD in the ICU. Biopsies were mainly done for work-up for new unknown infiltrates [12, 13, 21, 29, 31, 33, 34, 39]. However, no bivariate analysis was provided due to the small sample sizes and high mortality reported in these studies [13, 21, 29, 31]. Nowadays, biopsies are not routinely performed because lung biopsy in acute ILD is associated with increased in-hospital mortality (16%) compared to elective biopsies (1.7%) [46]. This higher mortality may be attributable to the severity of illness rather than to the biopsy itself.
**Immunosuppression**

The use of immunosuppressant medications including steroids and cyclophosphamide, among others, is controversial in the treatment of ILD patients in the ICU, and the benefit likely depends on underlying ILD aetiology. Two large mixed-ILD studies did not find an association between steroid use and increased mortality [14, 28]. In fact, a retrospective study found that use of high-dose steroids during ICU had a protective effect for in-hospital mortality (OR 0.19, 95% CI 0.04–0.99) [26]. These authors suggest that steroids could be used once infection has been ruled out. However, it is important to highlight the number of patients who received steroids was higher in the groups that included conditions typically responsive to steroids such as CTD-ILD, hypersensitivity pneumonitis, vasculitis and drug-induced ILD. In light of these results, certain patients will likely benefit from this treatment, especially young patients with features suggestive of CTD-ILD or patients with ILD who do not have a significant amount of fibrosis [28].

In IPF patients, Song and co-workers [41] found that steroid use did not predict mortality in 90 patients suggestive of CTD-ILD or patients with ILD who do not have a significant amount of fibrosis [28]. Results, certain patients will likely benefit from this treatment, especially young patients with features of patients who received steroids was higher in the groups that included conditions typically responsive to steroids such as CTD-ILD, hypersensitivity pneumonitis, vasculitis and drug-induced ILD. In light of these results, certain patients will likely benefit from this treatment, especially young patients with features suggestive of CTD-ILD or patients with ILD who do not have a significant amount of fibrosis [28].

Furthermore, ZAFRANI and co-workers [28] showed that IMV was associated with 1-year mortality in a multivariate analysis (OR 5.18, 95% CI 1.18–22.75). In IPF, data precluded calculation of ORs due to high rates of mortality associated with IMV, ranging from 70% to 100% (supplementary material e-table 2). Pooled analysis on mortality rates associated with IMV in mixed-ILD and IPF can be found in the supplementary material.

Two studies have specifically investigated the impact of IMV settings in ILD [3, 14]. High positive end-expiratory pressure (PEEP) defined as >10 cmH₂O was associated with a decreased survival both during hospitalisation (OR 17.26, p<0.001) and at 12 month (OR 4.72, p<0.001) [14]. The benefits of low tidal volumes have not been assessed in prospective studies of ILD; however, some retrospective data does exist. GAUDRY and co-workers [16] raise the question of the positive effect of low tidal volumes (5.9±1.3 mL) for ILD as eight out of 27 patients were successfully weaned off the ventilator. However, FERNANDEZ-PEREZ and co-workers [14] reported lower tidal volumes in nonsurvivors (median (IQR) 7 (6–9) mL·kg⁻¹) compared to survivors (median (IQR) 6 (7–10) mL·kg⁻¹) although the authors acknowledge low tidal volumes were used inconsistently.

**Mode of ventilation**

**Invasive mechanical ventilation**

Poor outcomes for ILD patients have been reported when respiratory failure requires IMV. In mixed-ILD, six studies showed increased mortality in IMV in the bivariate or multivariate analysis [12, 14, 17, 18, 26, 27]; whereas only two studies did not find an association [22, 28]. Multiple studies report high mortality rates associated with IMV, ranging from 27% to 100% in the univariate analysis (supplementary material e-table 2). Furthermore, ZAFRANI and co-workers [28] showed that IMV was associated with 1-year mortality in a multivariate analysis (OR 5.18, 95% CI 1.18–22.75). In IPF, data precluded calculation of ORs due to high rates of mortality associated with IMV, ranging from 70% to 100% (supplementary material e-table 2). Pooled analysis on mortality rates associated with IMV in mixed-ILD and IPF can be found in the supplementary material.

Noninvasive mechanical ventilation

NIMV has also been a subject of study in ILD. Some studies suggest that NIMV can prevent intubation and reduce mortality in some IPF patients [18, 25, 35, 44]. YOKOHAMA and co-workers [27] found that early initiation of NIMV in hypoxaemic patients suspected of having an acute exacerbation of ILD and initiation of NIMV on day one were associated with survival (76.9% survivors versus 8.3% nonsurvivors, p<0.0001). MOLINCA and co-workers [35] found that NIMV significantly reduced the respiratory rate in comparison to admission (mean of 26 versus 36) and prevented intubation in five out of 19 patients but did not improve survival. A major limitation of this study, however, was a significant variation in disease severity between the patients who initially received NIMV (APACHE II 19.5±5.9) and IMV (APACHE II 24.2±6) [35]. Nonetheless, the authors concluded that NIMV might be beneficial for early intervention in less severe cases, as well as a means of palliation by decreasing respiratory rate. The use of NIMV in the setting of less severe respiratory failure was further supported by GUNOR and co-workers [18] Individuals treated with non-continuous NIMV alone had a mortality rate of only 10.7% (out of 28 patients). Survival was particularly high among patients with an APACHE II score <20 supporting the use of NIMV in less severe presentations. A study of 11 acute exacerbation IPF patients showed that NIMV prevented intubation in five patients with severe hypoxaemia (PaO₂/FIO₂ 156.4±43.9). However, similar to the
previous studies, survivors had lower APACHE II scores (14.5±2.6) [44]. Pooled analysis on mortality rates associated with NIMV in mixed-ILD and IPF can be found in the supplementary material.

Other risk factors
Multiple risk factors have been evaluated in individual studies that have not been re-evaluated or have failed to replicate in others. Here, we present a brief summary of those that reached significance in multivariate analysis and were performed in large studies. In mixed-ILD patients, ZAFRAN and co-workers [28] identified that ILD-related pulmonary hypertension and computed tomography fibrosis were associated with a higher 12-month mortality (HR 3.21 (95% CI 1.45–7.11) and 2.36 (95% CI 1.05–5.32), respectively). Also, acute kidney injury (OR 10.60, 95% CI 2.25–49.97) was found to be associated with in-hospital mortality, which is consistent with other studies in the critical care literature [48]. In IPF studies, ODA and co-workers [37] reported that cotrimoxazole (OR 0.28, 95% CI 0.13–0.60) and macrolides (OR 0.37, 95% CI 0.15–0.87) were independently associated with decreased mortality, although this finding has not been evaluated in other studies. Finally, SONG and co-workers [41] found C-reactive protein and lymphocytes in the bronchoalveolar lavage fluid were associated with mortality risk after correcting for several factors (age, sex and PaO2/FIO2 ratio among others).

Discussion
Given the increasing availability of ICU care, evaluating which risk factors are associated with increased mortality becomes important to counsel patients and their families on likely outcomes. To our knowledge, this is the first systematic review that presents a thorough analysis on risk factors for mortality in patients with ILD in the ICU. While the small sample size and heterogeneity of many studies precluded the identification of statistically significant predictors of mortality or cause–effect relationships, some studies identified risk factors that may also indicate severity of disease. In patients with mixed-ILD, elevated APACHE score, hypoxaemia and mechanical ventilation were associated with increased mortality in the majority of the studies. In patients with IPF, only IMV was associated with increased mortality. Notably, while IMV is associated with overall poor prognosis, patient outcomes may be improved with lung protective strategies such as low PEEP and low tidal volumes, especially when there is a reversible condition. FERNANDEZ-PEREZ and co-workers [14] hypothesised that ILD patients may have limited recruitable lung. Thus, at higher PEEP, they were possibly experiencing over-distention injury, which was supported by elevated peak and plateau pressures [14]. Similarly, RANGAPPA and MORAN [3] suggest that high PEEP might be detrimental in IPF for the aforementioned reasons. However, high airway pressure may not necessarily cause injury if the pressure is from distending chest wall or collagen in the lung. Alternatively, as patients with severe hypoxaemia are given high PEEP, this might also represent an indicator of disease severity. This is why some authors suggest that IMV should be restricted to patients who can be eligible for lung transplant, which is an intervention that has been shown to improve mortality [49].

The use of steroids is still controversial. Our study did not find increased mortality associated with steroid use in patients with mixed-ILD. Their benefit likely depends on the underlying aetiology of ILD. Patients with steroid responsive diseases, such as hypersensitivity pneumonitis and CTD-ILD, probably derive benefit; this is likely untrue for IPF. Carefully designed clinical trials in various types of ILD are critically needed to assess the role of immunosuppression. The evidence on advance age is inconclusive, as positive and negative results have been reported in different studies. Regarding other factors in patients with IPF, evidence is also inconclusive for advanced age, elevated APACHE score and steroids use. The explanation of this phenomenon might rely not only on the size of the studies but also on the high mortality of IPF, which precludes a bivariate analysis. It is not clear why no association between hypoxaemia and mortality was found in three studies in IPF but the size of two studies was very small (<20 patients) and the third study (n=90) found an association in the bivariate but not the multivariate analysis. A potential explanation for this observation can be that the degree of hypoxaemia is severe in both survivors and nonsurvivors to find a significant difference. Therefore, this observation needs to be evaluated in large studies, as it is unexpected. Research in high flow nasal cannula has been proposed to be an alternative to conventional oxygen to correct hypoxaemia in patients with IPF [50] but the evidence is limited to series of cases on patients with interstitial pneumonias [51] and patients with do-not-intubate orders [52].

Our study also reports a down-trending mortality rate from 1993 to 2017 in patients with ILD (supplementary material e-table 3) which confirms the results from a recent large cohort study that reported decreased mortality from 2006 to 2012 in patients with IPF [53]. This observation could be explained by the role of new ventilation parameters such as low tidal volumes described in the acute respiratory distress syndrome network trial [54] and could explain its potential benefit in ILD. An alternative explanation could be secular trends in ICU care such as better haemodynamic management and aggressive antibiotic coverage. However, our calculation has limitations including the wide range of
years included in the studies, different available interventions over time, and the fact that mortality in the mixed-ILD group does not represent non-IPF ILD population alone as it includes some patients with IPF. Notably, mortality increases with time post hospitalisation with rates of 54–96% in mixed-ILD and 100% in IPF [12, 14, 16, 28]. Thus, ICU admission with ILD appears to carry prognostic implications beyond the actual hospital stay. Our study also presents exploratory data on mortality and failure rates associated with NIMV and IMV over time. In mixed-ILD, the mortality rate associated with NIMV was 49% compared to 71% with IMV, and the failure rate of NIMV was 52% (supplementary material). This observation can represent an indicator of less severity as patients who were offered NIMV could have had only mild or moderate disease. Esquinas Rodriguez and Ferrari [55] postulate that NIMV can reduce alveolar stress during tidal ventilation by improving lung compliance and increasing functional residual capacity. Therefore, NIMV needs further research as our exploratory data and some studies suggest it can prevent intubation and reduce mortality in some patients [18, 25, 35, 44]. Further discussion of this topic in patients with IPF can be found in the supplementary material.

This review has several limitations including the inclusion of retrospective studies of limited sample size, variable statistical analyses (including lack of multivariate modelling and lack of time-to-event analysis), heterogeneous subsets of ILD patients, and diverse evaluation of risk factors. We were not able to evaluate differences in populations over time given patient heterogeneity (in terms of underlying ILD, severity and treatments), different definitions for IPF used before the American Thoracic Society/European Respiratory Society guidelines were published [4], the inclusion of exacerbations caused by numerous aetiologies with different prognosis, and a wide overlap in the years included in each study. These limitations might have affected our results as some studies have conclusions in opposite directions, which made the evidence for some factors inconclusive. We should also highlight that mixed-ILD studies have the problem of combining patients with potentially worse prognosis such as IPF and others with conditions that might have a better prognosis such as CTD-ILD or drug-induced ILD [26]. Therefore, the findings presented in our manuscript should be interpreted with caution. However, we have to acknowledge that ILD is not a common disease and the current literature is only based, mainly, on series of cases. While some lessons are clear, this review highlights many remaining questions. Multi-centre cohort studies would be invaluable to enrol more patients over a short period of time, better characterise different types of ILD, detect geographical variations and help evaluate if emerging management therapies are affecting survival. The latter point is especially important in light of recent IPF and IIP guidelines [5, 6]. Prospective trials are also needed, particularly to investigate ventilation strategies.

Conclusions
Irrespective of ILD aetiology, mechanical ventilation is associated with increased mortality. For mixed aetiology ILD, hypoxaemia and APACHE scores are also associated with increased mortality. No increased mortality was found with steroids use. IPF has the highest mortality rate among ILDs, but since 1993 the rate appears to be declining. Despite improving in-hospital survival, overall mortality remains high.

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