Essential Features of an Interstitial Lung Disease Multidisciplinary Meeting

An International Delphi Survey

Alan K. Y. Teoh^{1,2,3}, Anne E. Holland^{3,4}, Julie Morisset⁵, Kevin R. Flaherty⁶, Athol U. Wells^{7,8}, Simon L. F. Walsh⁸, Ian Glaspole^{3,4}, Wim A. Wuyts⁹, Tamera J. Corte^{1,2,3}, and the ILD MDM Dephi Collaborators

¹Royal Prince Alfred Hospital, Camperdown, New South Wales, Australia; ²University of Sydney, Camperdown, New South Wales, Australia; ³Centre of Research Excellence in Pulmonary Fibrosis, Sydney, New South Wales, Australia; ⁴The Alfred Hospital, Melbourne, Victoria, Australia; ⁵Centre Hospitalier de l'Université de Montréal, Montreal, Quebec, Canada; ⁶University of Michigan, Ann Arbor, Michigan; ⁷National Institute for Health Research Biomedical Research Unit, Royal Brompton and Harefield, National Health Service Foundation Trust, London, United Kingdom; ⁸National Heart and Lung Institute, Imperial College London, London, United Kingdom; and ⁹Unit for Interstitial Lung Diseases, Department of Pulmonary Medicine, University Hospitals Leuven, Belgium

ORCID ID: 0000-0001-7051-112X (A.K.Y.T.).

Abstract

Rationale: The interstitial lung disease (ILD) multidisciplinary meetings (MDM), composed of pulmonologists, radiologists, and pathologists, is integral to the rendering of an accurate ILD diagnosis. However, there is significant heterogeneity in the conduct of ILD MDMs, and questions regarding their best practices remain unanswered.

Objectives: To achieve consensus among ILD experts on essential components of an ILD MDM.

Methods: Using a Delphi methodology, semi-structured interviews with ILD experts were used to identify key themes and features of ILD MDMs. These items informed two subsequent rounds of online questionnaires that were used to achieve consensus among a broader, international panel of ILD experts. Experts were asked to rate their level of agreement on a five-point Likert scale. An *a priori* threshold for consensus was set at a median score 4 or 5 with an interquartile range of 0.

Results: We interviewed 15 ILD experts, and 102 ILD experts participated in the online questionnaires. Five items and two exploratory statements achieved consensus on being essential for an ILD MDM following two questionnaire rounds. There was consensus

that the presence of at least one radiologist, a quiet setting with a visual projection system, a high-quality chest high-resolution computed tomography, and a standardized template summarizing collated patient data are essential components of an ILD MDM. Experts also agreed that it would be useful for ILD MDMs to undergo an annual benchmarking process and a validation process by fulfilling a minimum number of cases annually. Twenty-seven additional features were considered to be either highly desirable or desirable features based on the degree of consensus. Although our findings on desirable features are similar to the current literature, several of these remain controversial and warrant further research. The study also showed an agreement among participants on several future concepts to improve the ILD MDM, such as performing regular self-assessments and conducting research into shared practices to develop an international expert guideline statement on ILD MDMs.

Conclusions: This Delphi study showed consensus among international ILD experts on essential and desirable features of an ILD MDM. Our data represent an important step toward potential collaborative research into future standardization of ILD MDMs.

Keywords: interstitial lung disease; multidisciplinary meeting; Delphi; Essential features

(Received in original form November 16, 2020; accepted in final form June 30, 2021)

A complete list of ILD MDM Dephi Collaborators may be found before the beginning of the REFERENCES.

Ann Am Thorac Soc Vol 19, No 1, pp 66–73, Jan 2022 Copyright © 2022 by the American Thoracic Society DOI: 10.1513/AnnalsATS.202011-14210C Internet address: www.atsjournals.org

Supported by the National Health and Medical Research Council Centre of Research Excellence in Pulmonary Fibrosis (GNT1116371), and by foundation partner, Boehringer Ingelheim, and program partners, Roche and Galapagos.

Author Contributions: Conception and design of study: A.K.Y.T., A.E.H., J.M., I.G., and T.J.C. Data acquisition: A.K.Y.T. Data analysis: A.K.Y.T., A.E.H., and T.J.C. Data interpretation, manuscript drafting, and critical revision: all authors. Final manuscript approval: A.K.Y.T., A.E.H., J.M., K.R.F., A.U.W., S.L.F.W., I.G., W.A.W., and T.J.C. Agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved: all authors.

Interstitial lung disease (ILD) is a heterogeneous group of disorders that cause varying degrees of lung parenchymal inflammation and fibrosis. Although there is an array of specific ILD diagnoses, the clinical presentations of ILD are protean, and the precise recognition of specific ILD groups remains difficult (1). Over the last two decades, there have been increasing efforts to improve the accuracy of the diagnostic pathway for these diseases. In 2002, the American Thoracic Society (ATS) and European Respiratory Society (ERS) released a joint statement on the classification of idiopathic interstitial pneumonia (2) recommending the use of a multidisciplinary meeting (MDM) that included a respiratory clinician, a radiologist, and a histopathologist at its core. These recommendations have been echoed in several subsequent guidelines and position statements (3-5).

The ILD MDM allows for integration of available clinical, radiological and pathological data with the aim of rendering an accurate ILD diagnosis. Flaherty and colleagues reported significantly increased interobserver diagnostic agreement and confidence when relevant clinical, radiological, and pathological data were dynamically exchanged in an MDM setting (6). This approach has also been validated in several other studies (7–9). Furthermore, the diagnostic performances of physicians regularly attending ILD MDMs, irrespective of experience level, were greater than those without access to these meetings (10), suggesting an educational benefit of ILD MDMs. Although these findings demonstrate the integral role of ILD MDMs as the "gold standard" role for ILD diagnosis, many questions regarding the best practice for ILD MDMs remain unanswered (10). Although current international guidelines are able to recommend the basic membership of an ILD MDM, little beyond that is known (11). The member composition, level of expertise required, the amount of patient data required, and whether all ILD cases should be discussed at the MDM are just

few of the many ambiguities left unanswered. Several studies have described significant heterogeneity in the manner in which ILD MDMs are conducted, making it difficult to recommend a "minimum standard" (12, 13). Given these inconsistencies, an international standardized approach or guide for conducting ILD MDMs is required to ensure high-quality discussions on the diagnosis and management of ILD.

We conducted a Delphi survey among international ILD clinicians to explore the necessary components of the ILD MDM. The aim of the study was to achieve consensus among ILD clinicians regarding the key components of an ideal ILD MDM. The Delphi model is a well-described approach frequently used to establish consensus among various health professionals on topics where an established evidence base does not exist (14, 15). This approach provides participant anonymity, ensuring that each individual input carries equal weight.

Methods

To inform development of the Delphi survey, recognized experts in the field of ILD were invited by e-mail to participate in individual interviews to identify potential key features of ILD MDMs. These experts were identified from the professional contacts of T.J., J.M. and S.W. Invited experts also had contributed to the field of ILD by research publications, participation in thoracic societies research working groups, or clinical contribution to the care of patients with ILD in their respective centers of practice. We aimed to have a wide geographical representation (North America: C.R., D.L., H.C., and M.S.; South America: J.E. and M.M.; Europe: J.B., K.A., T.M., and V.C.; Asia: Y.K. and Y.I.; and Australia and New Zealand: D.C., N.G., and M.W.) to ensure diversity of each participant's local ILD MDM experience. The interviews were conducted either in person or over the

telephone in a semi-structured format. A.T. conducted all interviews, which were guided by a list of open-ended questions (Table 1). The interviews were digitally recorded and transcribed verbatim. Two reviewers analyzed the transcripts independently. The transcripts were analyzed using a qualitative approach to content analysis, allowing for common themes to be extracted and be used to identify key items or features of an ILD MDM (16). These items were used to form a list of statements for the first round of the modified Delphi survey. These statements were organized into major domains, reflecting a variety of aspects involved in the efficient running of an ILD MDM. We invited a broader international panel of ILD experts with a range of clinical and research experience in the field of ILD to participate in the Delphi surveys. Participants were identified to be attending an ILD MDM on at least a monthly basis, having previously consented to further research involvement from a previous study (10). ILD experts who participated in the semi-structured interviews were excluded from subsequent survey rounds. The ILD experts were invited by e-mail to participate in each round of the Delphi survey.

We conducted a two-round webbased survey between July 2019 and February 2020 in accordance with defined standards of the Delphi methodology (17). The surveys were published in English on a secure online survey platform (Qualtrics, LLC). An online platform was chosen for the ease of disseminating surveys to an international group of participants and allowing responses to be collected within a short period of time. Consent to the study was implied if the participants completed the questionnaires. Participants completed a short baseline demographic section regarding their medical practice and experience prior to the surveys. In the first round, participants were asked to rate their level of agreement on a list of statements detailing key features of an ILD MDM using a five-point Likert scale. Each item was scored as 1 (strongly disagree), 2 (disagree), 3 (neutral), 4

Correspondence and requests for reprints should be addressed to Alan K. Y. Teoh, M.B. B.S., Department of Respiratory Medicine, Royal Prince Alfred Hospital, 50 Missenden Road, Camperdown 2050 NSW, Australia. E-mail: Alan.Teoh@health.nsw.gov.au.

This article has a related editorial.

This article has an online supplement, which is accessible from this issue's table of contents at www.atsjournals.org.

(agree), and 5 (strongly agree). Experts were allowed to provide feedback following each statement and could add statements they considered relevant if they were not included in the original list. An a priori threshold of consensus was defined for the study as a median score of 4 or 5 with an interquartile range (IQR) of 0 for an essential feature of an ILD MDM. An interquartile range of 0 ensures that the distribution of responses truly reflects agreement and excludes a bimodal distribution where a smaller but important proportion of respondents may disagree. A median score of 4 or 5 with an IQR of 1 was considered to indicate desirable features. Other statements that did not meet these criteria were considered as "disagreement". In round 2, participants were given the distribution of group answers for statements that did not reach consensus. Participant comments and feedback for each statement were also provided. Participants were then asked to once again rate their level of agreement on a five-point Likert scale for statements that did not reach consensus and any additional statements identified in round 1

We reported the results of this study according to the proposed methodological standards for Delphi studies (17). Participant responses remained anonymous during result analysis. STATA version 15.1 (StataCorp) was used for all statistical analyses. The study was approved by the human research ethics committee of Royal Prince Alfred Hospital, Sydney (Protocol No X18–0354 and LNR/18/RPAH/497).

Results

Expert Qualitative Interviews

Fifteen of 17 (88%) invited ILD experts agreed to participate in the semistructured individual interviews. All experts were based in ILD referral centers across nine countries with 13 (87%) being involved in weekly ILD MDMs. Common themes were identified from the interview transcripts and informed development of the Delphi surveys (Table E1). These were subsequently organized into five major domains, namely "MDM team structure", "MDM infrastructure", "MDM organization and administration", "MDM clinical decision-making process", and "Future concept and directions". Table 1. Questions for initial expert interview guide

Questions

- 1. Could you describe your experience in the field of ILD and your involvement in ILD MDM(s)?
- 2. What do you think the role of the ILD MDM is?
- 3. Who do you think should be involved in the ILD MDM?
- 4. What do you think are the key elements in an ILD MDM?
- 5. How much preparation goes into the ILD MDM?
- 6. How do you come to a consensus for each case discussed at the MDM?
- 7. What do you think are challenges that ILD MDM commonly face?
 - 8. How do you think an ILD MDM could improve itself?

Definition of abbreviations: ILD = interstitial lung disease; MDM = multidisciplinary meeting.

Delphi Survey Results

A total of 134 ILD experts were invited by e-mail to participate in the Delphi survey, of whom 102 (76%) from 29 countries completed the first Delphi round. Subsequently, 94 out of 102 (92%) responded in the second round. In total, 91% of the experts actively participated in an MDM at an ILD referral center and 97% reported that their MDMs were attached to a university or academic hospital. Their characteristics are presented in Table 2.

First Round

Out of the 50 statements that were included in the first Delphi round, consensus was reached for a total of 5 statements (median 4 or 5 and IQR 0) (Table 3). Experts agreed that the presence of at least one radiologist is essential for an ILD MDM to function. Experts also agreed that it is essential that the ILD MDM has access to a visual projection system allowing real-time viewing of radiological images and that a high-quality chest high-resolution computed tomography (HRCT) scan is required for the discussion of cases. The two remaining statements that met consensus were considered to be explorative statements rather than current features of ILD MDMs (Table 6). A total of 36 statements met the threshold for being desirable features of an ILD MDM (median 4 or 5 and IQR 1), and 9 statements that did not meet either threshold were labeled as "disagreement".

Second Round

Out of the 45 statements that did not reach consensus in the first Delphi round, 39 were taken forward into round 2 without alterations. Five statements were rephrased to improve clarity based on study participants' feedback, and one statement was divided into two questions to enable a more accurate rating. Participants' feedback in round 1 generated four additional statements. The final list for round 2 contained 50 statements.

Two further statements achieved the *a* priori threshold for consensus in the second round (Table 3). Experts agreed that it was essential for an ILD MDM to be conducted in a quiet setting, allowing for easy interaction among its members. There was also consensus that it was essential for the ILD MDM to summarize collated patient data and information onto a standardized template. A total of 27 statements met the *a* priori threshold for being desirable features of an ILD MDM with a median score of either 4 or 5 and an IQR of 1. These statements were further subcategorized based on the level of agreement among experts. A total of 10 statements with a median score of 5 were considered "highly desirable" features of an ILD MDM (Table 4), whereas 13 statements with a median score of 4 were listed as "desirable" features (Table 5). Similar to round 1, the remaining four statements describing future concepts of ILD MDMs were listed separately.

Concepts and Future Direction

A total of six statements from both Delphi rounds fulfilled criteria for either being essential or desirable, were conceptual, and explored possible methods of improving the quality of ILD MDMs (Table 6). There was consensus among the experts that MDMs undergoing an annual benchmarking process once an international minimum standard has been established would be a useful approach to improve ILD MDMs. Experts also agreed

Table 2. Expert characteristics

Characteristics	Delphi Survey	
Sex, F, <i>n</i> /total (%)	39/102 (38%)	
Specialty, n (%)	Pulmonologist Radiologist Research/academic Other	95 (93%) 2 (2%) 2 (2%) 3 (3%)
Years of experience, <i>n</i> (%)	<5 yr 5–10 yr 11–15 yr 16–20 yr 21–30 yr	9 (8.8%) 27 (26.5%) 25 (24.5%) 16 (15.7%) 19 (18.6%)
Referral center MDM, <i>n</i> /total (%) MDM attached to university/academic hospital, <i>n</i> /total (%)	>30 yr 93/102 (91%) 99/102 (97%)	6 (5.9%)
Frequency of MDM, n (%)	More than once/weekly Weekly Fortnightly Monthly Other	8 (7.8%) 55 (53.9%) 26 (25.5%) 10 (9.8%) 2 (2.0%)
Duration of MDM, n (%)	30 min 31–60 min 61–90 min 91–120 min >120 min	3 (3.0%) 6 (5.9%) 48 (47.0%) 30 (29.4%) 16 (15.7%)
Number of cases discussed per meeting, n (%)	1-5 cases 6-10 cases 11-15 cases 16-20 cases >20 cases	2 (2.0%) 38 (37.3%) 19 (18.6%) 4 (3.9%) 39 (38.2%) 2 (2.0%)

Definition of abbreviation: MDM = multidisciplinary meeting.

that ILD MDMs should undergo a validation process by fulfilling a minimum number of case discussions annually. Experts felt that it would be highly desirable for ILD MDMs to occur on a regular basis to maintain its members' expertise, to perform self-assessments using an internationally collated case database, and for further research to be conducted into shared practices among ILD MDMs to develop an internationally agreed minimum standard. Lastly, experts agreed that the development of an international expert statement or guideline would be useful to provide guidance on running an optimal ILD MDM.

Discussion

In this study, we conducted semistructured interviews with a panel of ILD experts and identified items representing various aspects of an ideal ILD MDM, ranging from its member composition to the clinical decision-making process involved. Following a Delphi process involving an international panel of ILD experts, we identified seven items that achieved consensus as essential features of an ILD MDM. We also further identified a total of 27 items which experts considered to be desirable features that could be incorporated into an ILD MDM, allowing a standard set of criteria for an ideal ILD MDM to be constructed for standardization purposes.

The strong agreement among experts on the presence of at least one radiologist at an ILD MDM is in accordance with multiple iterations of international guidelines supporting radiologists as core members. Unsurprisingly, experts also strongly agreed that a high-quality HRCT scan was essential for all cases being discussed at the ILD MDM. This finding attests to the impact of having radiological data in rendering a consensus ILD diagnosis, with a previous study demonstrating that incorporation of HRCT

Table 3. Statements meeting consensus for essential features of an ILD MDM (median 4/5; IQR 0)

Consensus–Essential Items

It is essential to have at least one radiologist present at an ILD MDM.

It is essential for the ILD MDM to have access to a visual projection system allowing real time viewing of CT scan images.

A good quality high resolution CT scan is required for every case being discussed at the ILD MDM.

It is essential for the ILD MDM to be conducted in a quiet setting which allows for easy interaction among members.* It is essential for the ILD MDM to summarize collated patient information onto a standardized template.*

Definition of abbreviations: CT = computed tomography; ILD = interstitial lung disease; IQR = interquartile range; MDM = multidisciplinary meeting. *Statements that met consensus after second Delphi round.

Table 4. Items meeting criteria for highly desirable features of an ILD MDM (median 5; IQR 1)

Highly Desirable Items

More than one pulmonologist present at an ILD MDM. At least one pathologist present at an ILD MDM when there are histopathological data available. At least one member of the ILD MDM has at least five years ILD-focused experience/training. ILD MDM to serve as an education platform for specialist trainees and fellows. Pathologists should review biopsy specimens prior to the ILD MDM. Clinical history, a high-resolution CT scan and autoimmune serology should be available as a minimum dataset before a case can be presented at the ILD MDM. Pulmonary function testing comprising of at least spirometry and DL_{CO} is required for every case being discussed at the ILD MDM. ILD MDM should report a consensus diagnosis. ILD MDM should discuss initial treatment and management recommendations. An ILD MDM diagnosis should be a provisional diagnosis that may require a representation at an MDM at a later date when new

information is available.

Definition of abbreviations: CT = computed tomography; $DL_{CO} =$ diffusing capacity for carbon monoxide; ILD = interstitial lung disease; IQR = interquartile range; MDM = multidisciplinary meeting.

data led to a change in more than 50% of clinicians' first-choice diagnoses and an improvement in diagnostic confidence (18). The technical requirements of an HRCT for the diagnosis of ILD have also been described by the Fleischner Society (19). The importance placed by experts on having a visual projection system for realtime review of HRCT images is also paramount, where the ability for radiologists to convey specific information of an image and the educational benefit obtained from other clinicians have been described in other medical MDMs (20). Similarly, the strong agreement on having the ILD MDM in a quiet setting likely stems from experts' recognition of the negative impacts of background noise on member interactions and discussions during an MDM (21). Although not

specific to ILD MDMs, these have yet to be mandated in official guidelines or statements. Experts agreed on the importance of using a standardized template to be used to collate patient data for an ILD MDM. The Thoracic Society of Australia and New Zealand recently published a position statement advocating for a standardized format for data presentation, an approach that unfortunately remains variable (5).

Somewhat surprisingly, there was expert agreement on the merit of ILD MDMs undergoing a validation process by fulfilling a minimum number of case discussions annually and to undergo a benchmarking process against an international minimum standard. In the absence of a standardized validation process, an argument can be made that the expertise of an ILD MDM will increase over time with increasing numbers of case discussions. However, participant feedback from the survey highlighted concerns that validation of an ILD MDM merely by annual case numbers could deter smaller and newer ILD MDMs. Furthermore, case quantity as a sole criterion to validate ILD MDMs is surely inadequate. The ability of an ILD MDM to improve diagnostic agreement stems from a linked evidence approach, by which its efficacy is defined by its ability to change a clinical diagnosis and subsequent management, rather than direct evidence of its impact on patient health outcomes (22). However, Walsh and colleagues previously showed poor agreement between expert MDMs for hypersensitivity pneumonitis and nonspecific interstitial pneumonia (8). Quality assurance approaches such as using peer observers to

Table 5. Items meeting criteria for desirable features of an ILD MDM (median 4; IQR 1)

Desirable Items

Having more than one member from each discipline attending the ILD MDM to generate a more dynamic discussion.

ILD MDM should have a chair to moderate and guide its discussions.

The ILD MDM should allow the attendance of external physicians, either in person or via videoconferencing to present their cases. The ILD MDM should be a stand-alone meeting dedicated to the discussion of ILD cases only.

A meeting coordinator should be present to collate essential information required for every case prior to the meeting.

Histopathology images are required for cases that are being discussed at the ILD MDM in which lung biopsies have been performed. There should be processes in place for communicating MDM outputs to relevant stakeholders (i.e., referring physicians, other clinical service providers).

The ILD MDM should report on the degree of confidence of the diagnosis.

The ILD MDM should report a list of differential diagnoses if a confident diagnosis was not achieved.

The ILD MDM should adhere to current and available standardized diagnostic guidelines.

The ILD MDM should have a documented strategy on prioritizing urgent cases for the meeting."

Research terminologies (e.g., IPAF) can be used as consensus final or provisional diagnoses.

The ILD MDM should review its policies and protocols at least annually.*

Definition of abbreviations: ILD = interstitial lung disease; IPAF = idiopathic pneumonia with autoimmune features; IQR = interquartile range; MDM = multidisciplinary meeting.

*Items with interquartile range of 3-4.

 Table 6.
 Future concepts that met criteria for either being essential or desirable features to be incorporated into ILD MDMs

Future Concepts and Direction	Median (IQR) Score
It would be useful for every MDM to undergo an annual benchmarking process once an international minimum standard has been established.	4 (0)
It would be useful for every ILD MDM to undergo a validation process by fulfilling a minimum number of case discussions annually.	4 (0)
It would be useful for every ILD MDM to occur on a regular basis to maintain its expertise.	5 (1)
It would be beneficial to conduct further research into shared practices in ILD MDM to develop an internationally agreed minimum standard.*	4 (1)
It would be useful for every ILD MDM to regularly perform a self- assessment using an internationally collated database of cases *	4 (1)
It would be useful for an international expert statement/guideline to be developed to provide guidance on running an optimal ILD MDM.*	4 (1)

Definition of abbreviations: ILD = interstitial lung disease; IQR = interquartile range; MDM = multidisciplinary meeting.

*Items identified following round 2.

review the quality of MDMs have been explored in the field of oncology and could be explored in ILD MDMs (23, 24). However, the diagnostic accuracy of ILD MDMs has never been validated, and consequently, the best approach to benchmark an ILD MDM remains elusive. Nonetheless, the consensus among participants in our study highlights the strong international collective desire for the development of a validated minimum standard of an ILD MDM and a framework for a benchmarking process of ILD MDMs.

The degree of agreement seen with several highly desirable statements warrants further discussion. In our study, the presence of a pathologist at an ILD MDM was considered to be a highly desirable feature, whereas current guidelines recommend them as core ILD MDM members. A possible explanation is that experts viewed that clinicians and radiologists are more wellplaced to contribute in discussions of a greater case spectrum in the ILD MDM, whereas pathologists contributed only in cases with available histopathological data. However, when the statement was rephrased in round 2 to address this, the degree of agreement did not change. A more likely explanation is the potential challenges encountered in smaller MDMs or geographically remote centers, where access to pathologists may be limited. Certainly, our findings should not imply a lack of importance of the crucial role of an expert

ILD pathologist in the discussion of a patient with ILD for whom histopathology is available. Experts also agreed that it would be highly desirable to have at least one member with five years of ILD-focused experience attending the MDM. The proposed threshold of five years was derived from the expert interviews and has not been studied. Furthermore, the exact definition of experience and which ideal member requiring the additional experience have not been established.

Additionally, the finding that it was highly desirable for ILD MDM to deliver management recommendations is somewhat controversial. The role of MDMs in the management of oncological patients is clear where the diagnosis has already been established and evidence-based treatment options are available. In contrast, ILD MDMs have historically focused on disease characterization and diagnosis formulation (25). Major recent advances in therapeutic options may account for the expanded role of the ILD MDM, where clinicians may need more guidance in managing their patients. However, treatment often depends on individual patient factors such as frailty, comorbidities, and personal wishes, many of which cannot be addressed by an MDM panel that has not met the patient. However, our findings do suggest that there is a broad desire for cooperative assistance and advice in the development of therapeutic management plans for the complex ILD

patient, although the potential role of the ILD MDM in this is yet to be established.

Experts also agreed that it was desirable for ILD MDMs to adhere to available standardized ILD clinical practice guidelines. These guidelines provide a framework that clinicians can use to evaluate patients presenting with ILD. However, it is important to recognize that the fundamental hallmark of the ILD MDM is the integration of multidisciplinary data to render a diagnosis, one that is particularly important in ILD cases that do not fit into clinical practice guidelines. Surprisingly, there was also agreement among experts that research terminologies such as idiopathic pneumonia with autoimmune features (IPAF), could be used as consensus diagnoses. IPAF was proposed as a standardized research term to define patients with overlapping features that do not fit established diagnostic criteria for a connective tissue disease-associated ILD (26). Our findings suggest that there is an international recognition of this IPAF cohort in clinical practice, and that clinicians are keen to use the term in their clinical practice. However, it remains uncertain whether IPAF does indeed represent a separate clinical entity, with implications on prognosis and management remaining unknown.

Nonetheless, the majority of desirable features are already present at ILD MDMs in varying degrees with similar findings reported in a recent systematic review on ILD MDMs (27). Despite the agreement seen in our study, standardization and incorporation of these features are very often constrained by geographical distances and local resource availability. This can potentially account for the small number of statements reaching consensus in our study. However, this could also reflect the collective notion among study participants that only a small number of features are truly essential to run a high-quality ILD MDM without hindering the applicability of these features to smaller and newer ILD MDMs. The burgeoning number and frequency of MDMs are likely to increase workloads of radiologists and pathologists involved, potentially impacting negatively on the MDMs' ability to function effectively (28). The recent coronavirus disease (COVID-19) pandemic has resulted in the rapid uptake of videoconferencing technologies in ILD MDMs (29). Our study findings on essential and desirable features are likely to remain relevant in such hybrid or virtual MDMs. Despite these recent advances in data sharing

and videoconferencing technologies providing viable platforms for MDMs, the ideal format remains unknown, and further research is needed to evaluate the benefits of virtual discussions (30, 31). Research into shared ILD MDM practices, collaborative data sharing, and self-assessments are suggested approaches that warrant further consideration. The UK's National Health Service is one example where an established program of national clinical audits informed a range of policy changes leading to improved patient health outcomes (32). However, successful audits require robust evidencebased guidelines, clinical leadership, and buyin by professional bodies and stakeholders. Perhaps an initial approach is to explore the practicalities of performing self-assessments at an individual and local level, where resources may be more readily available.

Our study has several limitations. Participants recruited were predominantly pulmonologists (93%) with the remainder from other medical specialties (radiology, academic research, immunology, and rheumatology). Of note, no pathologists were recruited onto the study, and their absence as a recommended core member of an ILD MDM could potentially bias our results. Furthermore, we did not have demographic data from participants who did not respond to the invitation, hence will not be able to determine if potential bias was solely from the recruitment strategy or from a lack of response from other specialties. A majority of participants were from ILD referral centers and expert MDMs attached to academic hospitals. Hence, our study was unable to capture sentiments or opinions of clinicians participating in smaller or newer MDMs. However, we sought to establish consensus in an area with limited evidence, and we believe that our participant cohort

enabled us to achieve this aim. We used a rigorous definition of consensus, which resulted in a small number of essential features and a larger number of desirable features. A more lenient definition of consensus would have resulted in more features classified as "essential"; however. this could deter establishment of MDMs in less resource-rich settings. Some potentially important aspects of an MDM were not included in the Delphi, such as the number of participants, level of expertise required, frequency of meetings, and number of case discussions per meeting. However, the experts did not identify these issues for inclusion in round 2. Lastly, opinions captured in a Delphi process are not equivalent to evidence-based facts. Expert opinions from the Delphi process should be subjected to further studies to validate any proposed future ILD MDM models, ideally by demonstrating impact on patient outcomes.

Conclusions

In conclusion, using a Delphi model in surveying an international panel of ILD experts, our study was able to identify the essential and desirable features of an ILD MDM. Our study is an important step toward standardization of ILD MDMs. Our data may guide future collaborative research into development of international guideline reco-mmendations for ILD MDMs, based on high-quality evidence.

<u>Author disclosures</u> are available with the text of this article at www.atsjournals.org.

Acknowledgment: The authors thank all the ILD MDM Delphi collaborators who have generously agreed to participate in this study.

ILD MDM Delphi Collaborators: Huzaifa Adamali, J. Shirine Allam, Sofia Antillon, Katherina M. Antoniou, Rodrigo Athanazio, Sergey Avdeev, Alexander Averyanov, Arata Azuma, Bruno Baldi, Elisabetta Balestro, Rebecca Bascom, Shalini Bastiampillai, Lutz Beckert, Jüergen Behr, Paul Beirne, David Bennett, Raphael Borie, Demosthenes Bouros, Ben Brockway, Kevin Brown, Francisco Javier Callejas González, Diego Castillo, Ronald Chacon Chaves, Daniel Chambers, Sally Chapman, Nazia Chaudhuri, Harold Collard, Vincent Cottin, Bruno Crestani, Jesper Rømhild Davidsen, Devesh J. Dhasmana, Sahajal Dhooria, Juan Ignacio Enghelmayer, Alexandre Todorovic Fabro, Puneet Garcha, Nicole Goh, Alejandro Gomez, Christopher Grainge, Tomohiro Handa, Tristan Huie, Gary Hunninghake, Yoshikazu Inoue, Helen Jo, Kerri Johannson, Rene Jonkers, Eoin Judge, Yasemin Kabasakal, Leticia Kawano Dourado, Gregory Keir, Nasreen Khalil, Yet Hong Khor, Melissa King Biggs, Maria Kokosi, Yasuhiro Kondoh, Vasillis Kouranos, Michael Kreuter, David Lederer, Su Ying Low, Joachim Müller Quernheim, Toby Maher, Eliane Mancuzo, George Margaritopoulos, Carol Mason, Mariano Mazeini, Nesrin Mogulkoc, Maria Molina, Yuben Moodley, António Morais, Anoop Nambiar, Imre Noth, Hilario Nunes, Takashi Ogura, Oguzhan Okutan, Nina Patel, Carlos Pereira, Wojciech Piotrowski, Venerino Poletti, Silvia Quadrelli, Elzbieta Radzikowska, Pilar Rivera Ortega, Christopher J. Ryerson, Mauricio Salinas, Rafaela Sanchez, Recep Savas, Moises Selman, Adrian Shifren, Maria Raquel Soares, Eman Sobh, Jin Woo Song, Paolo Spagnolo, Martina Sterclova, Irina Strambu, Mary E. Strek, Takafumi Suda, Gabriela Tabaj, Jasna Tekavec Trkanjec, Fatma Tokgoz Akyil, Claudia Toma, Rade Tomic, Hiromi Tomioka, Daniel Traila, Lauren Troy, Sergio Trukillo, Argyrios Tzouvelekis, Carlo Vancheri, Brenda Elena Varela, Francesco Varone, Martina Vasakova, Elizabeth Veitch, Vanesa Vicens Zygmunt, Thomas Wessendorf, Glen Westall, Marlies Wijsenbeek, Margaret L. Wilsher, Jeremy Wrobel, Cesar Yoshito Fukuda, and Chris Zappala

References

- Kaner RJ, Brown KK, Martinez FJ. AJRCCM: 100-Year Anniversary. Progress in Interstitial Lung Disease. Am J Respir Crit Care Med 2017; 195(9):1104–1107.
- 2 American Thoracic Society; European Respiratory Society. American Thoracic Society/European Respiratory society international multidisciplinary consensus classification of the idiopathic interstitial pneumonias. This joint statement of the American Thoracic Society (ATS), and the European Respiratory Society (ERS) was adopted by the ATS board of directors, June 2001 and by the ERS Executive Committee, June 2001. Am J Respir Crit Care Med 2002;165:277–304.
- 3 Bradley B, Branley HM, Egan JJ, Greaves MS, Hansell DM, Harrison NK, et al.; British Thoracic Society Interstitial Lung Disease Guideline Group, British Thoracic Society Standards of Care Committee; Thoracic

Society of Australia; New Zealand Thoracic Society; Irish Thoracic Society. Interstitial lung disease guideline: the British Thoracic Society in collaboration with the Thoracic Society of Australia and New Zealand and the Irish Thoracic Society. *Thorax* 2008;63:v1–v58. [Published erratum appears in *Thorax* 63:1029.]

- 4 Travis WD, Costabel U, Hansell DM, King TE, Jr., Lynch DA, Nicholson AG, et al. An official American Thoracic Society/European Respiratory Society statement: update of the international multidisciplinary classification of the idiopathic interstitial pneumonias. *Am J Respir Crit Care Med* 2013;188:733–748.
- 5 Prasad JD, Mahar A, Bleasel J, Ellis SJ, Chambers DC, Lake F, et al. The interstitial lung disease multidisciplinary meeting: A position statement from the Thoracic Society of Australia and New Zealand and the Lung Foundation Australia. *Respirology* 2017;22:1459–1472.
- 6 Flaherty KR, King TE Jr, Raghu G, Lynch JP III, Colby TV, Travis WD, et al. Idiopathic interstitial pneumonia: what is the effect of a

multidisciplinary approach to diagnosis? *Am J Respir Crit Care Med* 2004;170:904–910.

- 7 Thomeer M, Demedts M, Behr J, Buhl R, Costabel U, Flower CD, et al.; Idiopathic Pulmonary Fibrosis International Group Exploring N-Acetylcysteine I Annual (IFIGENIA) study group. Multidisciplinary interobserver agreement in the diagnosis of idiopathic pulmonary fibrosis. *Eur Respir J* 2008;31:585–591.
- 8 Walsh SLF, Wells AU, Desai SR, Poletti V, Piciucchi S, Dubini A, et al. Multicentre evaluation of multidisciplinary team meeting agreement on diagnosis in diffuse parenchymal lung disease: a case-cohort study. Lancet Respir Med 2016;4:557–565.
- 9 De Sadeleer LJ, Meert C, Yserbyt J, Slabbynck H, Verschakelen JA, Verbeken EK, et al. Diagnostic ability of a dynamic multidisciplinary discussion in interstitial lung diseases: a retrospective observational study of 938 cases. Chest 2018;153:1416–1423.
- 10 Walsh SLF, Maher TM, Kolb M, Poletti V, Nusser R, Richeldi L, et al. Diagnostic accuracy of a clinical diagnosis of idiopathic pulmonary fibrosis: an international case-cohort study. *Eur Respir J* 2017;50: 1700936.
- 11 Castillo D, Walsh S, Hansell DM, Vasakova M, Cottin V, Altinisik G, et al.; ERICE ILD working group. Validation of multidisciplinary diagnosis in IPF. Lancet Respir Med 2018;6:88–89.
- 12 Jo HE, Corte TJ, Moodley Y, Levin K, Westall G, Hopkins P, et al. Evaluating the interstitial lung disease multidisciplinary meeting: a survey of expert centres. BMC Pulm Med 2016;16:22.
- 13 Richeldi L, Launders N, Martinez F, Walsh SLF, Myers J, Wang B, *et al.* The characterisation of interstitial lung disease multidisciplinary team meetings: a global study. *ERJ Open Res* 2019;5:00209-2018.
- 14 Graham B, Regehr G, Wright JG. Delphi as a method to establish consensus for diagnostic criteria. J Clin Epidemiol 2003;56:1150–1156.
- 15 Murphy MK, Black NA, Lamping DL, McKee CM, Sanderson CF, Askham J, et al. Consensus development methods, and their use in clinical guideline development. *Health Technol Assess* 1998;2:i–iv, 1–88.
- 16 Cavanagh S. Content analysis: concepts, methods and applications. *Nurse Res* 1997;4:5–16.
- 17 Diamond IR, Grant RC, Feldman BM, Pencharz PB, Ling SC, Moore AM, et al. Defining consensus: a systematic review recommends methodologic criteria for reporting of Delphi studies. J Clin Epidemiol 2014;67:401–409.
- 18 Aziz ZA, Wells AU, Hansell DM, Bain GA, Copley SJ, Desai SR, et al. HRCT diagnosis of diffuse parenchymal lung disease: inter-observer variation. *Thorax* 2004;59:506–511.
- 19 Lynch DA, Sverzellati N, Travis WD, Brown KK, Colby TV, Galvin JR, et al. Diagnostic criteria for idiopathic pulmonary fibrosis: a Fleischner Society White Paper. Lancet Respir Med 2018;6:138–153 10.1016/ S2213–2600(17)30433–2.

- 20 Li J, Robertson T, Hansen S, Mansfield T, Kjeldskov J. Multidisciplinary medical team meetings: a field study of collaboration in health care. Presented at the OZCHI 2008 Proceedings. December 8, 2008, Cairns, Australia. p. 73–80.
- 21 Oeppen RS, Davidson M, Scrimgeour DS, Rahimi S, Brennan PA. Human factors awareness and recognition during multidisciplinary team meetings. *J Oral Pathol Med* 2019;48:656–661.
- 22 Merlin T, Lehman S, Hiller JE, Ryan P. The "linked evidence approach" to assess medical tests: a critical analysis. Int J Technol Assess Health Care 2013;29:343–350.
- 23 Harris J, Green JSA, Sevdalis N, Taylor C. Using peer observers to assess the quality of cancer multidisciplinary team meetings: a qualitative proof of concept study. J Multidiscip Healthc 2014;7:355–363.
- 24 Johnson CE, Slavova-Azmanova N, Saunders C. Development of a peerreview framework for cancer multidisciplinary meetings. *Intern Med J* 2017;47:529–535.
- 25 Walsh SLF. Multidisciplinary evaluation of interstitial lung diseases: current insights: number 1 in the series "Radiology" edited by Nicola Sverzellati and Sujal Desai. *Eur Respir Rev* 2017;26: 170002.
- 26 Fischer A, Antoniou KM, Brown KK, Cadranel J, Corte TJ, du Bois RM, et al.; ERS/ATS Task Force on Undifferentiated Forms of CTD-ILD. An official European Respiratory Society/American Thoracic Society research statement: interstitial pneumonia with autoimmune features. Eur Respir J 2015;46:976–987.
- 27 Furini F, Carnevale A, Casoni GL, Guerrini G, Cavagna L, Govoni M, et al. The role of the multidisciplinary evaluation of interstitial lung diseases: systematic literature review of the current evidence and future perspectives. *Front Med (Lausanne)*. 2019; 6:246.
- 28 Kane B, Luz S, O'Briain DS, McDermott R. Multidisciplinary team meetings and their impact on workflow in radiology and pathology departments. *BMC Med* 2007;5:15.
- 29 Mackintosh JA, Glenn L, Barnes H, Dunn E, Bancroft S, Reddy T, et al. Benefits of a virtual interstitial lung disease multidisciplinary meeting in the face of COVID-19. *Respirology* 2021;26:612–615 10.1111/resp. 14062.
- 30 Grewal JS, Morisset J, Fisher JH, Churg AM, Bilawich AM, Ellis J, et al. Role of a regional multidisciplinary conference in the diagnosis of interstitial lung disease. Ann Am Thorac Soc 2019;16:455–462.
- 31 Fujisawa T, Mori K, Mikamo M, Ohno T, Kataoka K, Sugimoto C, et al. Nationwide cloud-based integrated database of idiopathic interstitial pneumonias for multidisciplinary discussion. Eur Respir J 2019;53: 1802243.
- 32 Stewart K, Bray B, Buckingham R. Improving quality of care through national clinical audit. *Future Hosp J* 2016;3:203–206.