### Pathways for Navigating ILD and IPF:

The Journey to Early Recognition, Diagnosis, and Patient-Centered Disease Management

# **Final Report Live Educational Initiative**

A free CME/CNE evening symposium



**Breathing Science is Life**.

### Key Features of Program Design

#### **Background**

The **online enduring case-based simulations** was developed first and then (4) live evening symposia employed structure of the cases developed for the online simulation. Images and video components from the simulation engaged learners in the multimedia live presentations to include Audience Response System (ARS) and provision of an infographic clinical reference aid. Learners will step through decisions in (3) case simulations to test and reinforce their skills in diagnosis, treatment, and management of ILD/IPF as well as effective patient communication strategies.





**Breathing Science is Life**.

### Accreditation, Audience and Outcomes Strategy

Accreditation Details: In support of improving patient care, NJH is accredited by the Accreditation Council for Continuing Medical Education (ACCME) to provide continuing medical education for physicians. NJH is also accredited by the Accreditation Council for Pharmacy Education (ACPE) and the California Board of Registered Nursing (CBRN) to provide continuing education for the healthcare team. *NJH designated the online program for 2.5 AMA PRA Category 1 Credits and the live evening symposia for a maximum of 2.5 AMA PRA Category 1 Credits*<sup>™</sup> and 3.0 nursing contact hours.

**Target Audience:** Pulmonologists, Radiologists, Pathologists along with Primary Care Physicians, Nurse Practitioners, and Physician Assistants who treat patients with Interstitial Lung Disease. Registered Nurses will be targeted for the live activities.

**Educational Outcomes Strategy:** Outcomes will be measured via participation totals, specialty, designation, pre-test, post-test, clinically-based decisions in case simulations, interactive polling questions, and evaluations. The metrics will demonstrate participation, satisfaction, engagement, and change in knowledge, competency, and performance to achieve Moore's Level 5 outcomes.



### Program Faculty





#### Michael P. Mohning, MD

Assistant Professor, Department of Medicine, Division of Pulmonary, Critical Care, Interstitial Lung Disease Program National Jewish Health Denver, CO



**Zulma X. Yunt, MD** Assistant Professor of Medicine, Department of Medicine, Division of Pulmonary, Critical Care & Sleep Medicine National Jewish Health Denver, CO



#### Jesse Roman, MD

CEO, Jane and Leonard Korman Respiratory Institute Jefferson Health – National Jewish Health Ludwig Kind Professor of Medicine and Pharmacology & Experimental Therapeutics Enterprise Division Chief, Pulmonary, Allergy & Critical Care Medicine Jefferson Health Philadelphia, PA



### Final Report: Program Overview



#### **Objectives**

- Describe best practices for diagnosing IPF based on the most recent evidence-based guidelines.
- Differentiate IPF from other interstitial lung diseases through comprehensive assessment, diagnosis, and/or referral to ILD specialty centers.
- Apply recent clinical data and guidelines to the management and treatment selection of IPF.
- Determine appropriate communication strategies for addressing quality of life issues in patients with ILD.

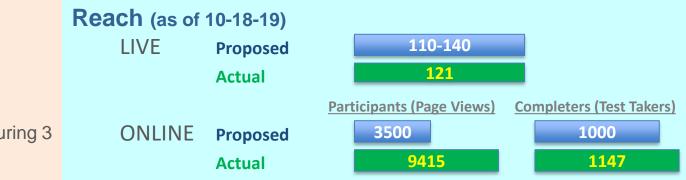


#### **Target Audience**

Pulmonologists, Radiologists, Pathologists, Primary Care Physicians, Nurse Practitioners, Physician Assistants and Nurses who treat patients with Idiopathic Pulmonary Fibrosis

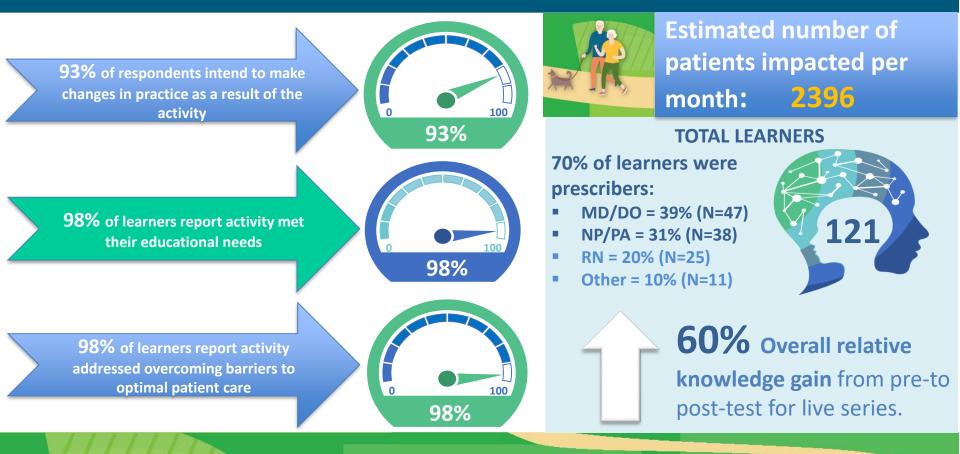
#### Format

- 4 live meetings consisting of interactive, case-based presentations and breakout workshops
- Online enduring activity featuring 3 patient cases



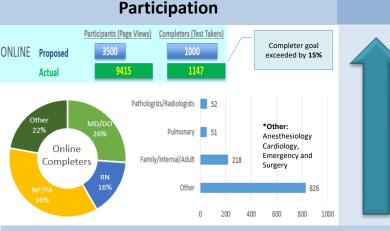
### Final Report: Live Program Dashboard





### Final Report: Online Program Dashboard





#### Satisfaction

I loved this activity! First one of it's kind. Very interactive, very informative, and fun!

I enjoyed the interactive format!

This course covered an important type of subject that is very common in the community.

#### Learner Impact

- **116%** overall relative gain in knowledge from pre to post test for live meetings
- **92%** of learners indicated that they planned to make changes to practice as a result of the education provided.

#### Performance

**71%** of prescribers could identify antifibrotic therapy as an appropriate IPF treatment option immediately after the initial education or upon remediation.



#### Persistent Gaps/Needs

#### Identified by traditional prepost assessment:

 Variable understanding existed related to making a diagnosis based on HRCT images





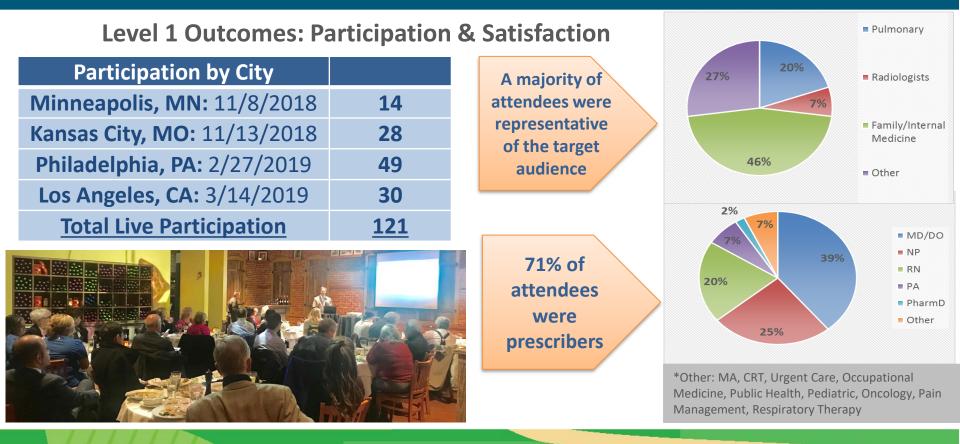
### Identified in case-simulation platform performance:

- Diagnosing IPF based on HRCT images
- Distinguishing IPF from other ILD's

Kn to diagnose Wilma w/ IPF	38	17	45
Kn cannot discern type of IPF	35	23	42

### Level 1 Outcomes (Live)





### Level 2 Outcomes (Live)



**Level 2 Outcomes: Learning & Satisfaction** N=69

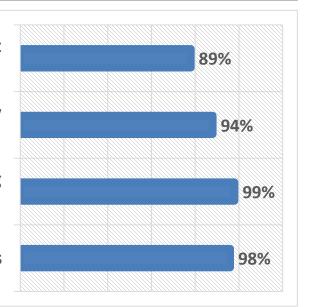
Participants report the activity was "Good" to "Excellent" at:

Improving your ability to treat or manage your patients

Enhancing your ability to apply the LOs to practice

Reinforcing and/or improving your current skills

Meeting your educational needs



"Enjoyed this educational activity. Looking forward to more in the future."

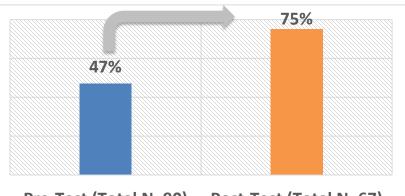
"There is a need for more programs like this to reach healthcare teams (not just physicians) in the community."

"Wonderful speakers. Made information easy to understand."

### Level 3 & 4 Outcomes (Live)

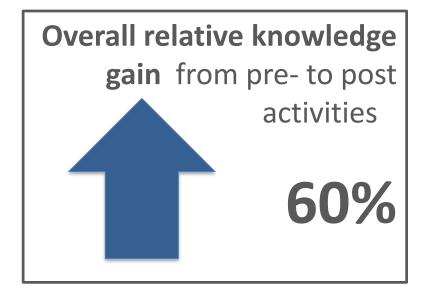


#### Level 3&4 Outcomes: Knowledge/Competence (Pre-Test/Post-Test)



Pre-Test (Total N=90) Post-Test (Total N=67)

Level 3 and 4 outcomes were measured by comparing pre-and post-test answers. Attendees' responses to these questions demonstrated that **participants gained knowledge as a result of the activity.** 



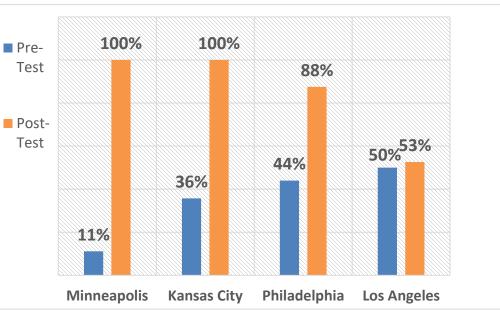


#### Level 3 Outcomes: Knowledge

*Learning Objective: Apply recent clinical data and guidelines to the management and treatment selection of IPF* 

Q1: When should a patient with IPF be referred for lung transplant evaluation?

- A. After first exacerbation
- B. When the patient has shown a significant decline in FVC
- C. Lung transplant is not beneficial in IPF
- D. At time of diagnosis
- E. After they have failed anti-fibrotic therapy



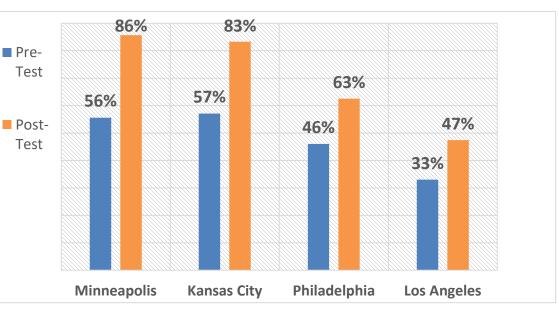


#### Level 3 Outcomes: Knowledge

Learning Objective: Describe best practices for diagnosing IPF based on the most recent evidence-based guidelines

Q4: Which histologic pattern is seen in the setting of idiopathic pulmonary fibrosis?

A. Usual interstitial pneumonia
B. Non-specific interstitial pneumonia
C. Lymphocytic interstitial pneumonia
D. Desquamative interstitial pneumonia
E. Diffuse alveolar damage





#### Level 3 Outcomes: Knowledge

*Learning Objective: Determine appropriate communication strategies for addressing quality of life issues in patients with ILD.* 

Q10: What is the best approach to discussing IPF disease management with patients and caregivers?

- A. Carefully review with patients and caregivers the data that shows IPF prognosis is approximately 2.5 years
- B. Recommend antifibrotics and pulmonary rehabilitation
- C. Ask patients and caregivers to conduct research online
- D. Explain to patients that while IPF is an incurable disease, there are strategies that can be employed to enhance overall QOL

Pre-Test
 Post-Test
 Post-Test
 Minneapolis
 Kansas City
 Philadelphia
 Los Angeles

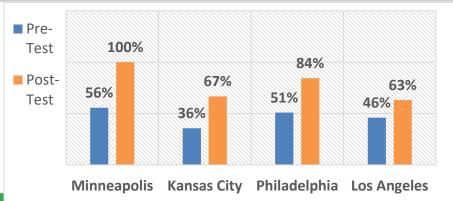


#### Level 4 Outcomes: Competence

Learning Objectives: Apply recent clinical data and guidelines to the management and treatment selection of IPF **and** Differentiate IPF from other interstitial lung diseases through comprehensive assessment, diagnosis, and/or referral to ILD specialty centers.

Q2: A 50 year old female presents with progressive shortness of breath and dry cough. She also notes arthralgias and joint swelling. On exam you note crackles in the bases bilaterally and she has ambulatory oxygen desaturation. A HRCT scan is done which reveals bibasilar reticulation and ground glass opacities. Lab testing reveals a positive anti-CCP antibody and hand x-rays reveal erosive arthritis. What is the next step in evaluation and management?

- A. Start patient on anti-fibrotic medication
- B. Refer for surgical lung biopsy
- C. Perform bronchoscopy
- D. Start immunosuppression for treatment of CTD-ILD
- E. Close monitoring with serial CT imaging



### Evaluation Results (Live)

### **Level 4 Outcomes: Evaluation Data**

- **93%** of respondents report that they intend to make changes in practice as a result of the activity.
- 100% of respondents report that the material was presented in an objective manner and free of commercial bias
- **98%** of respondents report that the activity addressed strategies for overcoming barriers to optimal patient care
- **99%** of respondents report that the content presented was evidence-based and clinically relevant







### Level 4 Outcomes (Live)



### Self-Reported Performance N=576

## 88%

of those in practice report that the activity provided new ideas or information they have used in practice 76%

report one or more of their patients have already benefitted from the information learned 75%

report that the infographic associated with this activity is a helpful guide for diagnosing IPF

"The information was immediately used. The next day at work I spoke to 1 nurse and 3 patients diagnosed with ILD."



The four live meetings featured interactive case-based polling questions using Poll Everywhere. These questions are in addition to the pre/post-test and are asked throughout the lecture in a test and teach format. Patient charts and films were shown to answer the questions. The data allowed the presenter to understand the baseline knowledge, as well as to get more data from participants to help elucidate some of the findings in our preliminary analysis of the online enduring program data.

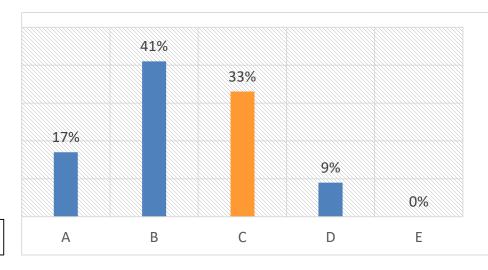
#### Which study would you like to order?

Low Recolution CT Scan MR Cheet High resolution CT (HRCT) Full complement of cheat X-ray PCT scan	Response options	Count	Percentage	
	Low Resolution CT Scan	0	0%	
	MRI Chest	2	9%	
	High resolution CT (HRCT)	20	91%	
-	Full complement of chest X-rays (PA, expiratory, decubitus, lordotic)	0	0%	
	PET scan	0	0%	

- Case Study #1 (JACOB): 49 y.o. Caucasian male, married; Mild pulmonary hypertension (diagnosed 3 years ago); Progressive dyspnea; Raynaud's phenomenon; Hashimoto's thyroiditis; Hypertension; GERD
- Given the HRCT findings (shown in presentation) what is your diagnosis?

N=62

- A. Idiopathic Pulmonary Fibrosis (IPF)
- B. Connective tissue disease (CTD)-ILD
- C. Need more information\*
- D. Non-specific interstitial pneumonia
- E. Cryptogenic organization pneumonia





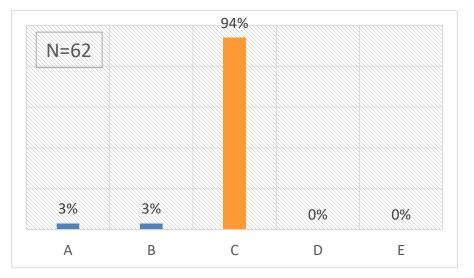




Case Study #1 (JACOB): 49 y.o. Caucasian male, married; Mild pulmonary hypertension (diagnosed 3 years ago); Progressive dyspnea; Raynaud's phenomenon; Hashimoto's thyroiditis; Hypertension; GERD

#### What is your diagnosis?

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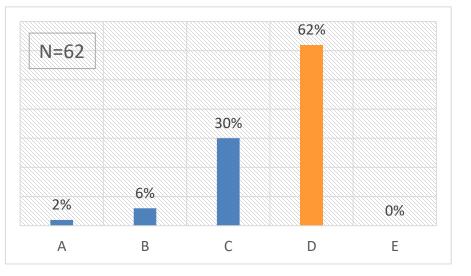


Case Study #1 (JACOB): 49 y.o. Caucasian male, married: Mild n

49 y.o. Caucasian male, married; Mild pulmonary hypertension (diagnosed 3 years ago); Progressive dyspnea; Raynaud's phenomenon; Hashimoto's thyroiditis; Hypertension; GERD

#### What medication will you order?

A. Nintedanib (over pirfenidone)
B. Pirfenidone (over nintedanib)
C. Either nintedanib or pirfenidone
D. Prednisone and mycophenolatemofetil
E. Warfarin



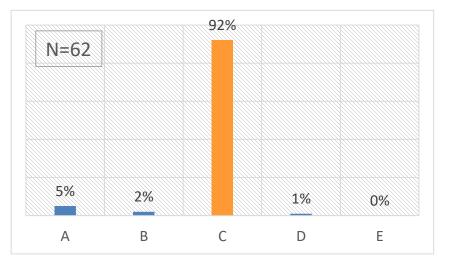




Case Study #2 (RAYMOND): 74 y.o. Caucasian male; coronary artery disease; myocardial infarction 26 years ago; COPD; obstructive sleep apnea; GERD; Surgery (3-vessel CABG 20 years ago)

#### Which of the following studies should be performed next?

- A. Bronchoscopy with bronchoalveolarlavage
- B. Lung biopsy
- C. High Resolution CT Scan (HRCT)
- D. Exhaled Nitric Oxide
- E. Standard CT without contrast



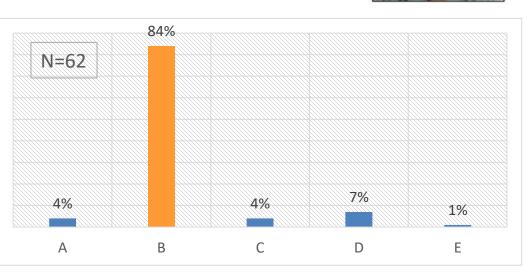




Case Study #2 (RAYMOND): 74 y.o. Caucasian male; coronary artery disease; myocardial infarction 26 years ago; COPD; obstructive sleep apnea; GERD; Surgery (3-vessel CABG 20 years ago)

#### What is Raymond's diagnosis?

- A. Connective tissue disease ILD
- **B. Idiopathic pulmonary fibrosis**
- C. Desquamative interstitial pneumonia
- D. Acute interstitial pneumonia
- E. Cryptogenic organization pneumonia







68 y.o. African American female; chronic cough for past 12 months; post nasal drainage; GERD;

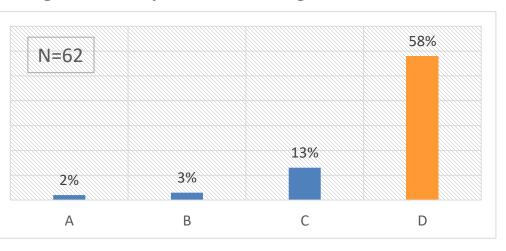
- A. Carefully review with patients and caregivers the data that shows IPF prognosis is approximately 2.5 vears
- B. Recommend antifibrotics and pulmonary rehabilitation

Case Study #3 (WILMA):

Hypertension; Hypothyroidism

- C. Ask patients and caregivers to conduct research online
- D. Explain to patients that while IPF is an incurable disease, there are strategies that can be employed to enhance overall QOL

What is the best approach to discussing IPF disease management with patients and caregivers?

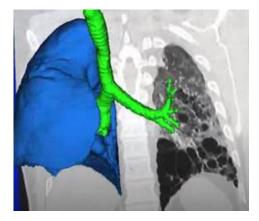






### **Online Activity: Final Status Report**





Launched October 19, 2018: https://learning.freecme.com/a /30328PAgeVqR

### **ProDoctor**

Pathways for Navigating ILD and IPF: The Journey to Early Recognition, Diagnosis, and **Patient-Centered Disease Management** 



### **Online Case Simulation Platform**



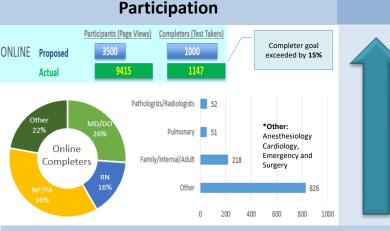
### ProDoctor

The online activity uses ProDoctor's innovative simulation platform to highlight three patient cases (1 CTD-ILD and 2 IPF) each with accompanying HRCT images, HRCT reconstructions, 3D animations and radiology impression. Learners are challenged to make decisions regarding the workup, tests, and differential diagnosis of ILD in all three cases. Key learning points were reinforced with an infographic clinical reference aid developed for the use in both the live and online activity.



### Final Report: Online Program Dashboard





#### Satisfaction

I loved this activity! First one of it's kind. Very interactive, very informative, and fun!

I enjoyed the interactive format!

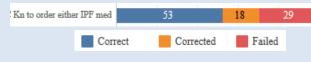
This course covered an important type of subject that is very common in the community.

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#### Persistent Gaps/Needs

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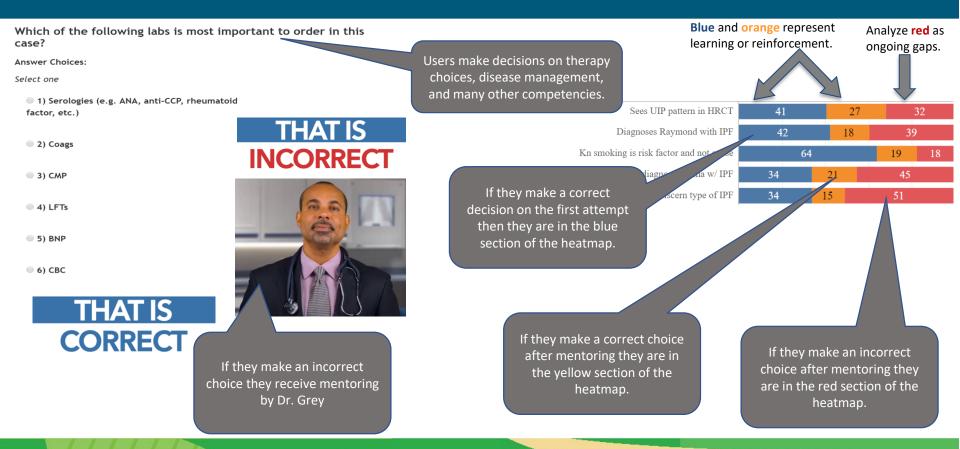
Identified in case-simulation platform performance:

- Diagnosing IPF based on HRCT images
- Distinguishing IPF from other ILD's

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### ProDoctor Heatmaps – Level 5 Outcomes

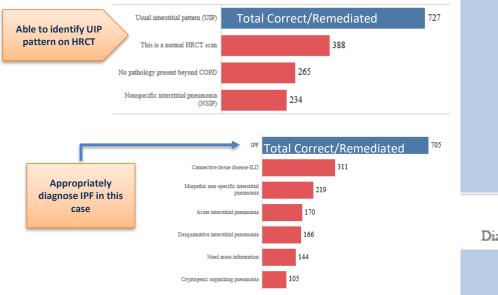




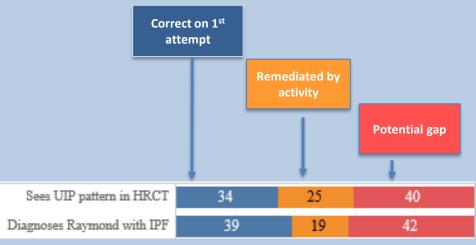


**Learning Objective 1**: Describe best practices for diagnosing IPF based on the most recent evidence-based guidelines.

### **ProDoctor Heatmap Data:**



#### Learner performance after viewing simulation:

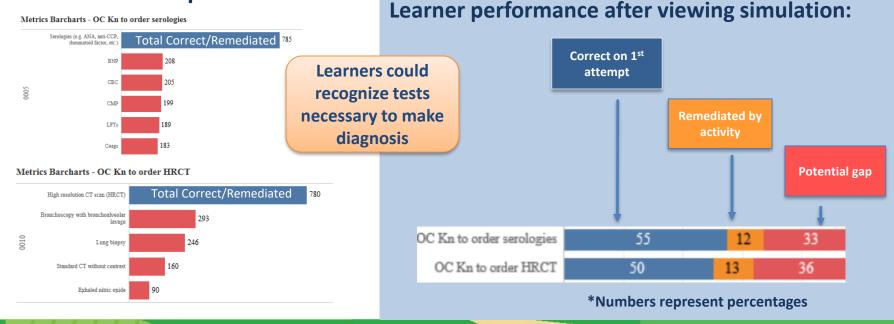


\*Numbers represent percentages



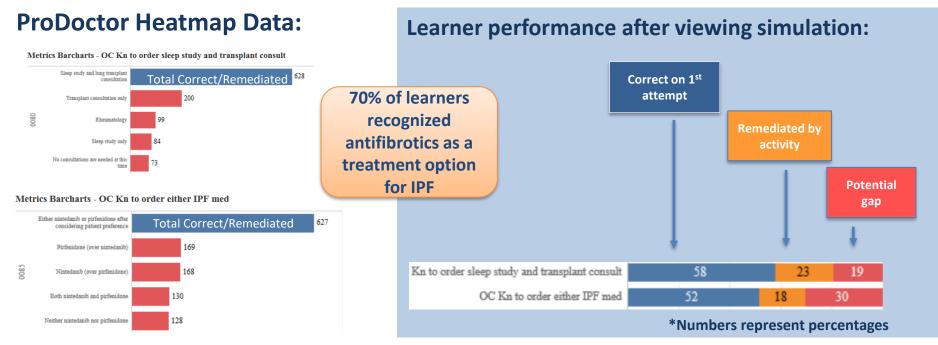
**Learning Objective 2:** Differentiate IPF from other interstitial lung diseases through comprehensive assessment, diagnosis, and/or referral to ILD specialty centers.

#### **ProDoctor Heatmap Data:**





Learning Objective 3: Apply recent clinical data and guidelines to the management and treatment selection of IPF.





Learning Objective 4:Determine appropriate communication strategies for addressing quality of life issues in patients with ILD.

### **ProDoctor Heatmap Data:**

#### Metrics Barcharts - Kn this is Wilmas new normal

#### Unfortunately, this could be your new **Total Correct/Remediated** 634 Correct on 1st normal attempt 070 243 Learners Your lungs will probably get better Remediated by incorporated 175 activity I have no idea appropriate communication Metrics Barcharts - Encourages Wilma to investigate transplant Potential gap strategies. IPF is unpredictable. I recommend we **Total Correct/Remediated** 639 begin to investigate this option and initiate a referral. Okay, it is a little too early for 174 transplant referral. You can wait. 0075 Kn this is Wilmas new normal 22 Yeah, lung transplant is a bad idea 152 anyway Encourages Wilma to investigate transplant 64 10 You should have a lung transplant; I 124 am going to refer you.

#### \*Numbers represent percentages

Learner performance after viewing simulation:

### Performance Comparison



45 47

38

47

58

59

60

60

63

54 41

49

40

21

26

18

28

18

27

27

18

38

10

46

#### Pulmonary Prescribers (N=21)

12% of pulmonary prescribers were correct on first attempt after the education. The remainder were effectively remediated by the education on 86% of topics.

	Sees UIP pattern in HRCT		89		11
	Diagnoses Raymond with IPF	89		11	
	Kn smoking is risk factor and not cause		80		20
	Kn to diagnose Wilma w/ IPF		67 33		3
	Kn cannot discern type of IPF		67	17	17
	Kn this is Wilmas new normal		71		29
	Encourages Wilma to investigate transplant		100		
	Sees NSIP pattern in HRCT image		83		17
	r HRCT for Jacob		100		
	OC Kn to order serologies		67	22	11
	OC Kn to order HRCT		89		11
	OC Kn to order echocardiogram		67 3		3
	OC Kn consults needed		67	22	11
	OC Kn PAH intervention is supplemental O2		86		14
	OC Kn to order pul rehab		83		- 17
	OC Kn Wilma does not want IPF drugs		100		
	$\operatorname{OC}\nolimits\operatorname{Kn}$ to order sleep study and transplant consult		67	17	17
	OC Kn to order either IPF med		67	17	17
	OC Kn to order PFTs and 6 min walk test		86		14
	OC Kn to request rheum and pulm hyp specialist		86		14
	OC Kn Dx is connective tissue disease ILD		100		
	${\rm Kn}{\rm Rx}$ s/b predinsone and mycophenolate mofetil		71	14	14
	OC Kn to add trimethoprim-sulfamethoxazole	29	14	57	
/	OC Kn hing biopsy consult is inappropriate		86		14
	rescribe inhaler for COPD		78	1	1 11

#### Primary Care Providers (N=134)

	Sees UIP pattern in HRCT	
	Diagnoses Raymond with IPF Kn smoking is risk factor and not cause Kn to diagnose Wilma w/ IPF	
	Kn cannot discern type of IPF	
	Kn this is Wilmas new normal	
	Encourages Wilma to investigate transplant	
	Sees NSIP pattern in HRCT image	
	r HRCT for Jacob	
	OC Kn to order serologies	
	OC Kn to order HRCT	
	OC Kn to order echocardiogram	
	OC Kn consults needed	
	OC Kn PAH intervention is supplemental O2	
	OC Kn to order pul rehab	
	OC Kn Wilma does not want IPF drugs	
	OC Kn to order sleep study and transplant consult	
	OC Kn to order either IPF med	
	OC Kn to order PFTs and 6 min walk test	
	OC Kn to request rheum and pulm hyp specialist	
	OC Kn Dx is connective tissue disease ILD	7
	C Kn Rx s/b predinsone and mycophenolate mofetil	
2	OC Kn to add trimethoprim-sulfamethoxazole	
	OC Kn lung biopsy consult is inappropriate	
	rescribe inhaler for COPD	

Failed

Corrected

Performance improved in

all areas. The largest

persistent gaps for

primary care providers

relate to making the

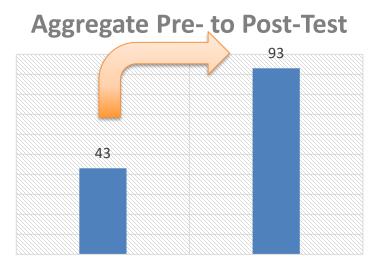
diagnosis of IPF.

Correct

Both groups had difficulty with this question in Jacob's

### **Online Enduring Metrics**





Pre-Test (N=1887)

Post-Test (N=1151)

#### Gain in Knowledge

Participants demonstrated a 116% relative gain in knowledge and competence as a result of this activity.

In addition to questions posed in the online simulations, a set of standard pre/post questions were presented to learners upon entry into the activity via FreeCME and after completion of the ProDoctor simulation.



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Pathways for Navigating ILD and IPF: The Journey to Early Recognition, Diagnosis, and Patient-Centered Disease Management

Released On October 19, 2018 Media Type Internet

Expires On

October 18, 2019

Completion Time Up to 150 minutes

Specialty Pulmonology, Radiology, Pathology, Primary Care Topic(s) Interstitial Lung Disease, Diagnosing IPF, Quality of Life, Communication Strategies

Clicking Start Activity indicates that you have reviewed the CME/CE information for this activity.

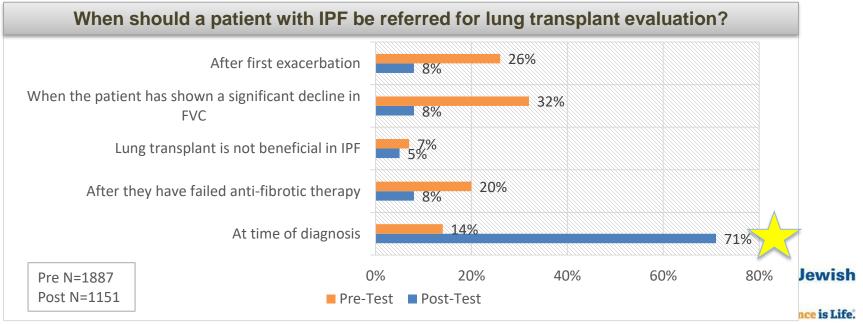


### Online Pre-Post Test Question



#### Level 3: Outcomes: Knowledge

Learning Objective: Determine appropriate communication strategies for addressing quality of life issues in patients with ILD.

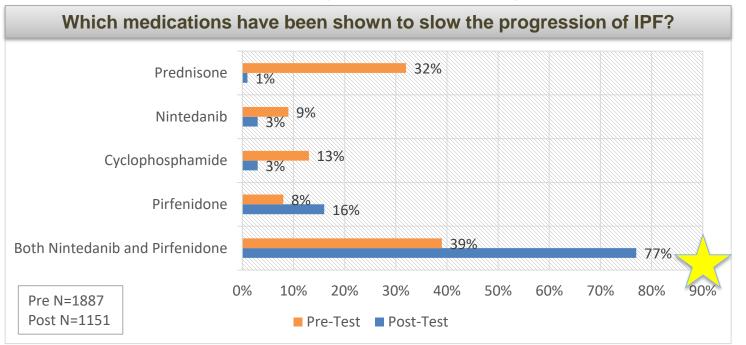


### **Online Pre-Post Test Question**



#### Level 3 Outcomes: Knowledge

Learning Objective: Apply recent clinical data and guidelines to the management and treatment selection of IPF.

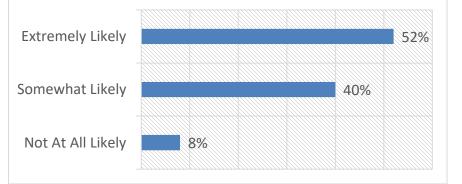


### **Online Enduring Evaluation Results**



### Level 4 Outcomes: Competence (Evaluation Results)

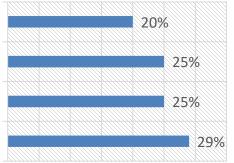
As a result of what I learned, I intend to make changes in my practice:



As a result of what I learned, I intend to make the following changes in my practice:

#### Modify treatment plans

Use alternative communication methodologies with patients... Incorporate different diagnostic strategies into patient evaluation Change my screening/prevention practice



N=1122

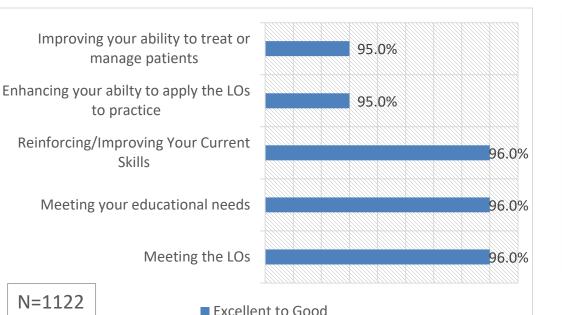
**Note: 92%** of Participants reported that they were somewhat or extremely likely to make a change in their practice

### **Online Enduring Evaluation Results**



### Level 4 Outcomes: Competence (Evaluation Results)

#### Participants report the activity was "Excellent" to "Good" at:



#### **Evaluation**

94% reported the material was presented without commercial bias

**97%** reported the content presented was evidencebased and clinically relevant

N=1122

to practice

Skills

### Key Take-Aways



#### Four overarching themes emerged from live/online data:

- 1) Early intervention and diagnosis
  - "Early intervention is key"
  - "Differentiating IPF from other ILD's"
- 2) Communicating with patients
  - "Discussing long term disease management"
  - "Being upfront with patients"
- 3) Appropriate referrals
  - "Refer to lung transplant early"
  - "Refer to specialty centers"
- 4) Selecting the appropriate treatment
  - "Multidisciplinary approach to treatment"
  - "When to start treatment"



Participants in the live session reflected on key lessons learned

### **Recommendations for Future Education**



- Asthma Management
- Pulmonary Hypertension
- Lung Transplant
- More details on the anti-fibrotic meds and how they low the progression of IDF
- Management of difficult to treat patients
- Targets of investigational agents