

Documentation and Coding

OCTOBER 2025

Pulmonary Fibrosis

At Healthfirst, we are committed to helping providers accurately document and code their patients' health records. This tip sheet is intended to assist providers and coding staff with the documentation and ICD-10-CM selection on services submitted to Healthfirst specifically for common types of **Pulmonary Fibrosis**. It provides information from industry sources about proper coding practice. However, this document does not represent or guarantee that Healthfirst will cover and/or pay for services outlined. Coverage decisions are based on the terms of the applicable subscriber contract/ evidence of coverage and the provider's participation agreement. This includes the determination of any amounts that Healthfirst or the member owes the provider.

ICD-10-CM Codes and Descriptions

Code	Description			
J84.10 [†]	Pulmonary fibrosis, unspecified			
J84.112	Idiopathic pulmonary fibrosis			
J84.178	Other interstitial pulmonary diseases with fibrosis in diseases classified elsewhere Note: Code first underlying disease, such as: • progressive systemic sclerosis (M34.0) • rheumatoid arthritis (M05.00-M06.9) • systemic lupus erythematosis (M32.0-M32.9)			

[†]Use only in the event no other code describes the condition.

Treatment

Treatment for Pulmonary Fibrosis					
Supplemental Oxygen	Pulmonary Rehabilitation	Symptom Management	Medication	Lung Transplantation	

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Diagnostic Testing

Imaging Test Lur		ing Function Test		ssue Sample iopsy)	ВІ	ood Test	
•	Chest X-ray CT Scan Echocardiogram Bronchoscopy	•	Pulmonary Function Test Pulse Oximetry Exercise Stress Test Arterial Blood Gas Test	•	Surgical Biopsy	•	Providers may test to evaluate patient's liver and kidney function

Clinical Documentation

Clinical Documentation Should Include:				
Status of Condition	Stable, Improved, Worsening			
Risk Factors	Personal History, Family History, Environmental, Medications, Radiation, Occupational			
Link Associated Conditions with Terms	Due to, Secondary to, or Associated with			
Treatment Plan	Family and/or individual counseling Diagnostic testing Specify root cause of pulmonary fibrosis			

Coding Tips

The conditions listed below are **inclusive** of the following terms when coding:

Codes	Conditions	
J84.10	 Capillary fibrosis of lung Cirrhosis of lung (chronic) NOS Fibrosis of lung (atrophic) (chronic) (confluent) (massive) (perialveolar) (peribronchial) NOS Induration of lung (chronic) NOS Post-inflammatory pulmonary fibrosis 	
J84.112	Cryptogenic fibrosing alveolitis Idiopathic fibrosing alveolitis	
J84.114	Hamman-Rich syndrome	
J68.1	Pulmonary edema due to chemicals, gases, fumes, and vapors	

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Pulmonary fibrosis due to

- Inhalation of chemicals, gases, fumes, or vapors (J68.4)
- Pulmonary fibrosis following radiation (J70.1) should never be coded with J84.10 or J84.112
- **J84.1*** Other interstitial pulmonary diseases with fibrosis can be coded with pulmonary fibrosis (chronic) due to inhalation of chemicals, gases, fumes or vapors (**J68.4**).

Coding Example

Case 1	65-year-old man with idiopathic pulmonary fibrosis returns to the office for a follow-up visit. Last visit he reported having dyspnea when walking two blocks, which is now resolved. He is currently taking pirfenidone and states he is compliant. He has no other complaints. Patient instructed to return for follow-up in one month.
Diagnosis Reported J84.112	Idiopathic pulmonary fibrosis
Rationale	Provider documented patient has idiopathic pulmonary fibrosis and is actively being treated with pirfenidone.

References

- ICD-10-CM Official Coding Guidelines FY2026
- Coding Clinic Advisor
- Understanding All Types of Pulmonary Fibrosis
- Pulmonary Fibrosis Treatment Options

Questions?

Contact us at #Risk Adjustments and clinical Documentation@healthfirst.org.

For additional documentation and coding guidance, please visit the coding section on HFproviders.org

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