Diagnosis	☐Idiopathic Pulmonary Fibrosis		
Basis for IPF Diagnosis	☐Chest HRCT is "consistent with UIP" and Atypical features are ABSENT ☐Surgical biopsy consistent with "definite UIP" or "probable UIP" And HRCT is without definite HRCT features of a specific alternate etiology		
Chest HRTC Features	☐Subpleural predelection ☐Basilar dominant ☐Honeycomb changes present ☐Paucity of ground glass		
Pathological Pattern	☐UIP	☐UIP/NSIP	☐Other
Family history of pulmonary fibrosis	☐Yes	☐No	