Supplemental Figure S3.

(a)

The annual %FVC change rate between ILPs other than IPF and CTD-ILD in the no antifibrotic group for (a) the 1st year after PF-ILD diagnosis and (b) the observation period after PF-ILD diagnosis.

* Differences were assessed using the Wilcoxon rank sum test.

Abbreviations: %FVC; percentage of forced vital capacity; NS, not significant; ILPs, idiopathic interstitial pneumonias; IPF, idiopathic pulmonary fibrosis; CTD-ILD, connective tissue disease-related interstitial lung disease.