

Supporting Information

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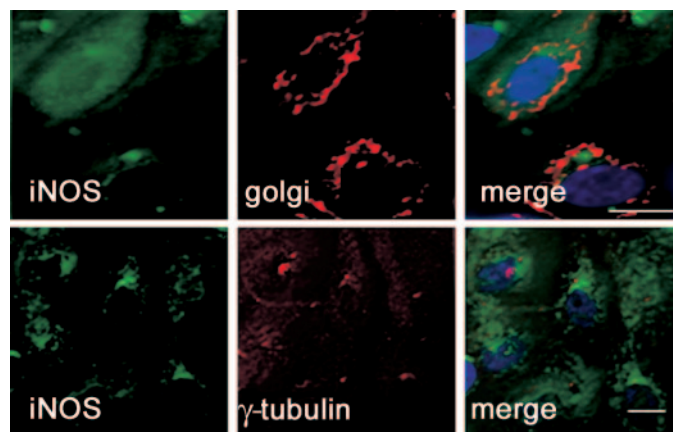


Fig. S1. Characterization of iNOS aggregates in primary normal human bronchial epithelial (NHBE) cells. Primary NHBE cells were grown to full differentiation at the air/liquid interface and stimulated for 30 h with a cytokine mixture similar to that used in Figs. 4 and 5. Cells were then fixed, stained with DAPI to visualize nuclei (blue), and immunolabeled using antibodies against iNOS (green), resident Golgi protein giantin (red), or centrosome marker γ -tubulin (red). (Scale bar, 10 μ m.)