

Supplementary Materials

Modeling airway dysfunction in asthma using synthetic mucus biomaterials

Daniel Song, Ethan Iverson, Logan Kaler, Shahed Bader, Margaret A. Scull, Gregg A. Duncan

*Correspondence to: Gregg Duncan; email: gaduncan@umd.edu

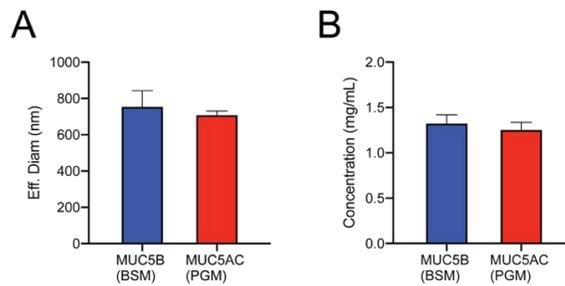


Fig. S1. (A) Effective hydrodynamic diameter of MUC5B (BSM) and MUC5AC (PGM) solutions measured in 10 mM NaCl using dynamic light scattering (NanoBrook Omni; Brookhaven Instruments). (B) Concentration of 1% mucin solutions measured using BCA assay.

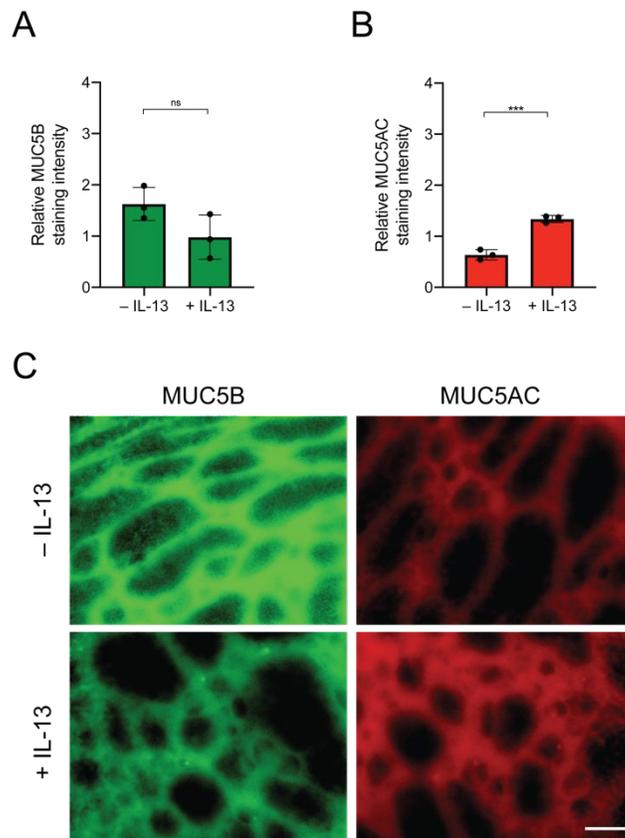


Fig. S2. (A) Relative MUC5B staining intensity from *en face* immunostaining. (B) Relative MUC5AC staining intensity. (C) Additional *en face* immunostaining images of mucus gels produced from unstimulated (-IL-13) or IL-13-stimulated (+IL-13) BCI-NS1.1 HAE cultures. Scale bar = 20 μ m.

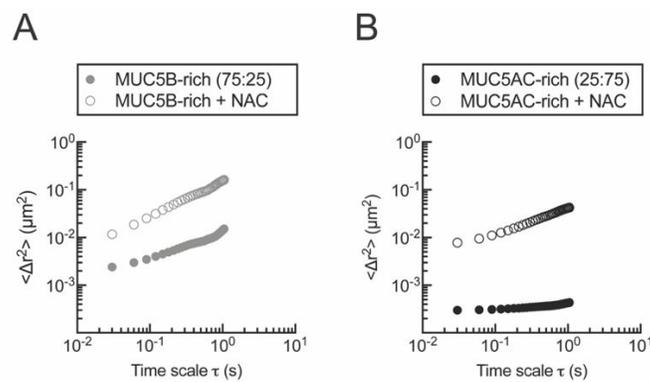


Fig. S3. (A) Ensemble average MSD ($\langle \Delta r^2 \rangle$) of 100 nm PEG-NP as a function of time scale (τ) in MUC5B-rich (75:25) gel before and after a 30-minute treatment with *N*-acetyl cysteine (NAC). (B) Ensemble average MSD ($\langle \Delta r^2 \rangle$) of 100 nm PEG-NP as a function of time scale (τ) in MUC5AC-rich (75:25) gel before and after a 30-minute treatment with NAC.