

Supplementary Materials:

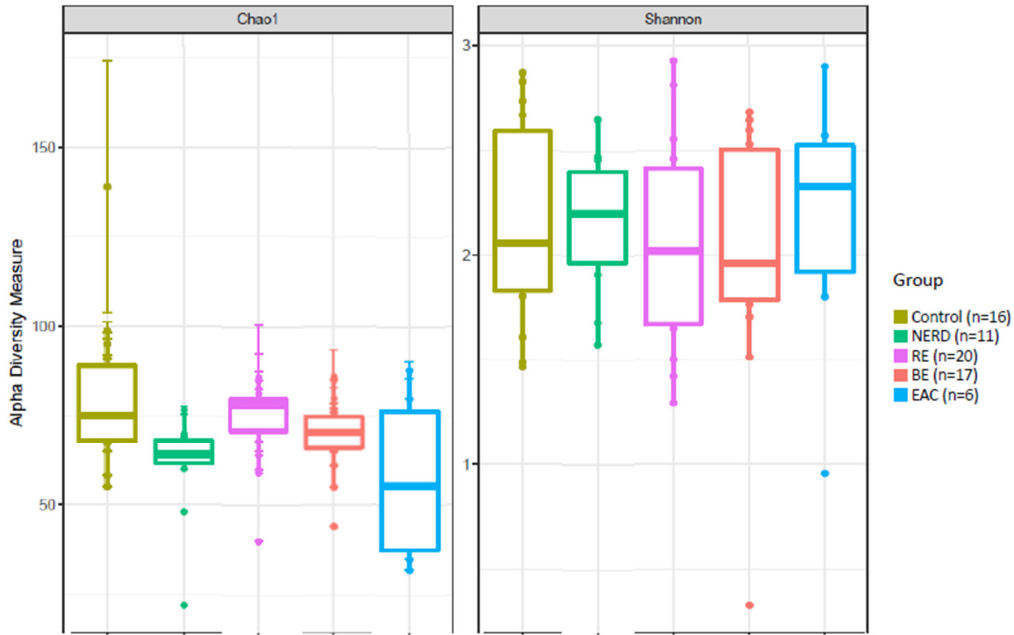


Figure S1. Alpha diversity analysis of control, non-erosive reflux disease (NERD), reflux esophagitis (RE), Barrett's esophagus (BE), and esophageal adenocarcinoma (EAC) using Chao1 richness estimator and Shannon diversity index.

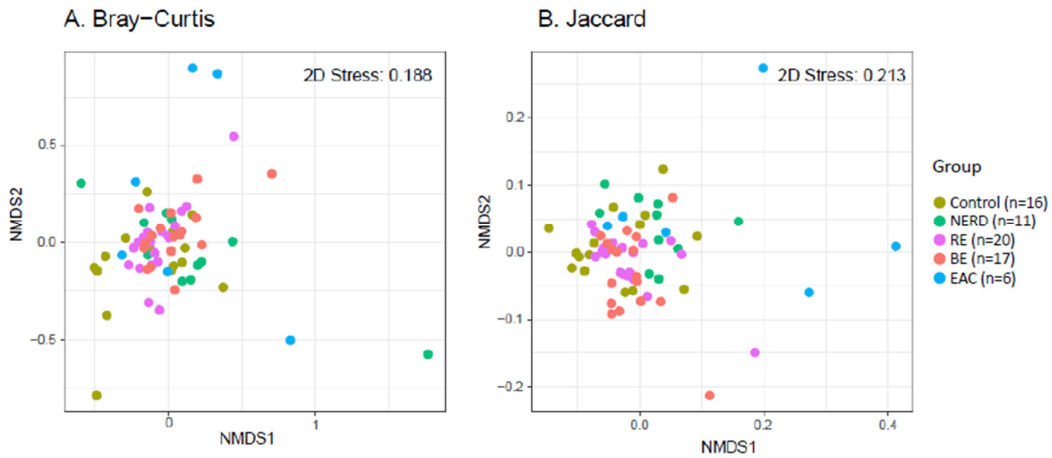


Figure S2. Non-metric multi-dimensional scaling (nMDS) plot for Bray-Curtis dissimilarity (abundance and composition) analysis and Jaccard (presence/absence) analysis of control, non-erosive reflux disease (NERD), reflux esophagitis (RE), Barrett's esophagus (BE), and esophageal adenocarcinoma (EAC).

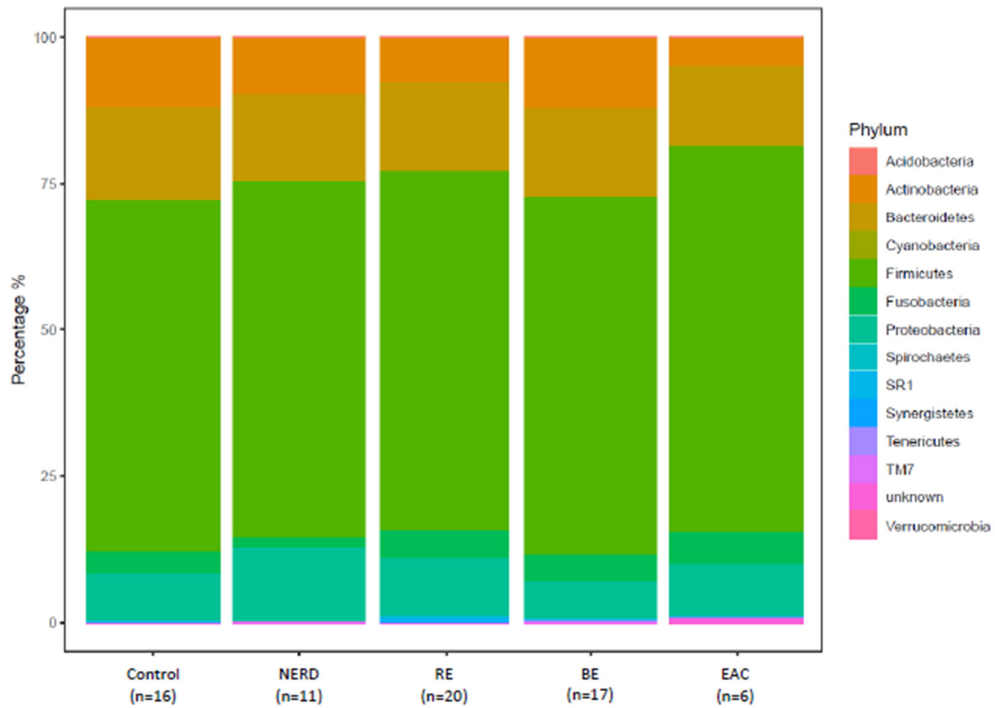


Figure S3. Microbiota phylum composition within each group: control, non-erosive reflux disease (NERD), reflux esophagitis (RE), Barrett’s esophagus (BE), and esophageal adenocarcinoma (EAC).

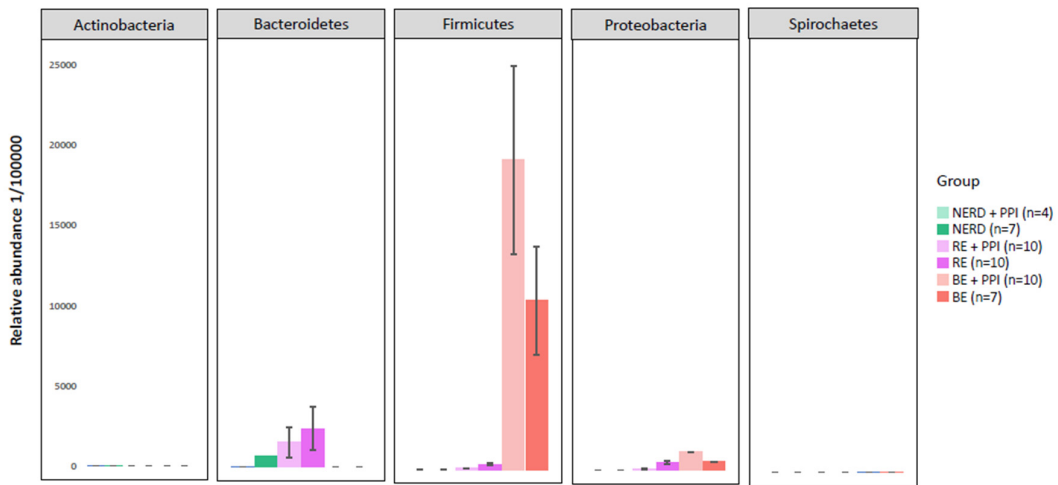


Figure S4. Differences in bacterial phylum composition between proton pump inhibitor treatment in non-erosive reflux disease (NERD), reflux esophagitis (RE), and Barrett’s esophagus (BE) disease phenotypes.

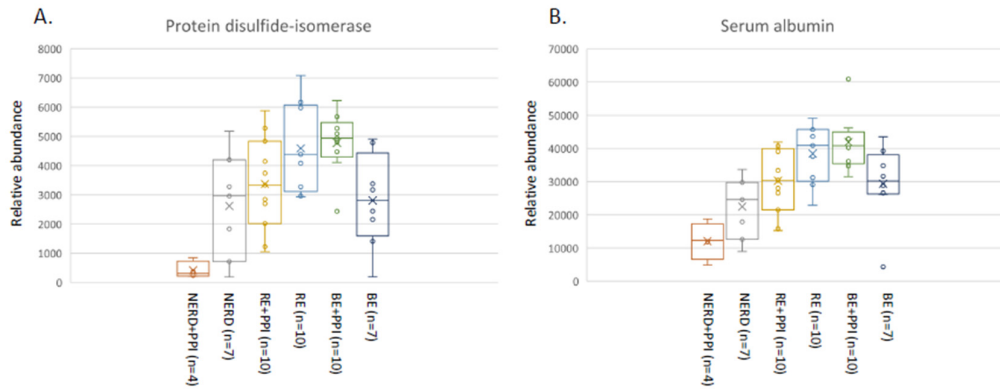


Figure S5. (A) protein disulfide isomerase and (B) serum albumin were found differentially expressed between proton pump inhibitor (PPI) treated and untreated mucosal proteomic samples in non-erosive reflux diseases (NERD), reflux esophagitis (RE) and Barrett’s esophagus (BE).