

Hypersensitization of *Helicobacter pylori* to antibiotics through perturbation of bacterial glycan armor
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Bacterial glycans are a target of interest as a result of the role they play in pathogenesis.

They were grown in the presence or absence of three established glycan altering agents: polymyxin B nonapeptide (PMBN), vancomycin, and Bac-diNAc-OBn (BnBac). These glycan altering agents perturbed different parts of the *H. pylori* glycocalyx. By growing the *H. pylori* in the presence or absence of these inhibitors, we had an untreated (control) condition and three experimental conditions. After the bacteria was grown and treated with glycan-altering agents, an Epsilonometer test was performed to determine the minimum inhibitory concentration (MIC) of the FDA approved antibiotic, levofloxacin. In each experimental condition, the MIC value was found to be lower than that of the control condition, signifying that a lower concentration of levofloxacin was needed to kill the *H. pylori* when treated with glycan altering agents. These data suggested that altering the glycan armor of *pylori* successfully hypersensitized it to levofloxacin and that this could be a viable strategy for fighting bacterial infections in the future. Of the three glycan altering agents, BnBac inhibited glycoprotein biosynthesis the most effectively as it resulted in a lower MIC value and substantiated by W* n BT /F3 12 Tf 7(e) most effective as,