The Ubiquitous Colonizer Staphylococcus hominis Protects Host Skin from Opportunistic Staphylococcal Pathogens by Blocking Quorum Sensing. M Brown (PhD, GS) ${ }^{1}$, A Shahbandi ${ }^{2}$, D Todd ${ }^{2}$, N Cech $^{2}$, and A Horswill ${ }^{1,}$, ${ }^{1}$ Dept. of Immunology \& Microbiology, University of Colorado Anschutz Medical Campus, Aurora, CO, ${ }^{2}$ Dept. of Chemistry and Biochemistry, University of North Carolina at Greensboro, NC.

Commensal coagulase-negative staphylococci (CoNS) actively shape the skin barrier to resist colonization or infection by opportunistic pathogens, including Staphylococcus aureus, in a variety of mechanisms known as colonization resistance. The best characterized CoNS is Staphylococcus epidermidis, yet S. epidermidis is a frequent opportunistic pathogen that can actively degrade the skin barrier. We hypothesize that other commensal CoNS may have a greater protective role on the skin than previously appreciated, including the second most frequently isolated CoNS, Staphylococcus hominis. A potential S. hominis colonization resistance mechanism is the Accessory Gene Regulator (agr) quorum sensing system, which is ubiquitous among staphylococci. This two component system senses and responds to its autoinducing peptide (AIP) signal. In S. aureus, agr regulates virulence factor expression and inhibiting $S$. aureus agr has been proposed as an antibiotic alternative. We found that spent media from any $S$. hominis skin isolate was sufficient to inhibit $S$. aureus agr. We sequenced a hypervariable region of the agr locus and found that $S$. hominis makes at least six AIP variants. Using mass spectrometry, we identified and validated the structures of 5 of these AIPs. We found that synthetic $S$. hominis AIPs inhibit $S$. aureus and S. epidermidis agr signaling with varying degrees of potency, but the majority with nanomolar $\mathrm{IC}_{50}$. Together, these data suggest that $S$. hominis agr cross-talk with opportunistic staphylococcal pathogens may be one mechanism to protect the cutaneous barrier from damage.

