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## Production and regulation of functional amyloid curli fimbriae by Shiga toxin-producing *Escherichia coli*

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Functional amyloid, in the form of adhesive fimbrial proteins termed curli, was first described in *Salmonella* and *Escherichia coli*. Curli fibers adhere to various host cells and structural proteins, interact with components of the host immune system, and participate in biofilm formation. Shiga toxin-producing *E. coli* (STEC) cause severe hemorrhagic diarrheal disease which can progress to hemolytic uremic syndrome in certain cases. STEC typically carry *stx1*, *stx2*, or both on lambdoid prophage. Although most STEC possess all of the genes required for functional curli production, curli expression is tightly regulated resulting in great variability in curli production among different STEC strains and serotypes. Curli and the exopolysaccharide cellulose are the major components of STEC biofilms. Both are controlled by the transcriptional regulator *CsgD*, whose expression is dependent on the RpoS sigma factor and enhanced by the *MlrA* transcription factor. We have shown that prophage insertions in the coding region of *mlrA*, often carrying *anstx* gene, are major barriers that limit *csgD* expression and curli production. This talk will discuss differences in *csgD*-dependent curli expression in various STEC serotypes and present new findings regarding prophage effects on *mlrA* expression and curli production.

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