

Fim Switch 96 well plate reader assay.

The Fim switch and GFP reporter strains were picked from agar plates and cultured in LB chloramphenicol broth for 4 hours and transferred to the wells of a fluorescence compatible 96 well plate. The total volume in each well was 150 μ l. For the co-culture experiments 140 μ l of the reporter strain was transferred to the selected wells followed by 10 μ l of the Fim switch strains. GFP fluorescence readings (excitation 395 nm, emission 509 nm) were taken every 10 minutes along with OD600 readings using a FLUOstar Omega microplate reader. The plate was incubated at 37°C with shaking in between readings. Each well was repeated in triplicate with LB chloramphenicol being used as a blank.