

Figure S1. Y57A Cj1386 protein is expressed in the $\Delta cj1386+cj1386^{Y57A}$ *C. jejuni* mutant construct at levels comparable to Cj1386^{WT} expression. Bacterial cultures were grown in MEMα media at 37°C under microaerophilic conditions. Bacterial cultures were pelleted, resuspended in PBS + protease inhibitor and soluble proteins were extracted following sonication. Two hundred and forty micrograms of lysates or 100 ng of purified Cj1386 were separated by SDS PAGE on a 14% polyacrylamide gel followed by immunoblotting. (Upper) Wild-type (WT), $\Delta cj1386$, $\Delta cj1386+cj1386^{WT}$ and $\Delta cj1386+cj1386^{Y57A}$ lysates and Cj1386 protein was detected using an anti-Cj1386 antiserum. (Bottom) Loading control of total protein contents as detected by an anti-Fur antiserum.

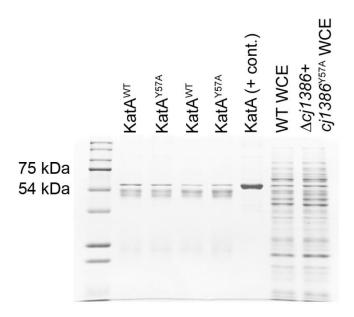


Figure S2. Immunoprecipitation of KatA from wild type and $\Delta cj1386+cj1386^{Y57A}$ *C. jejuni* strains. KatA was immunoprecipitated from prepared wild type and $\Delta cj1386+cj1386^{Y57A}$ whole cell extracts and eluted in 50 mM glycine, pH 2.8. Four microlitres of each immunoprecipitated sample, 1µg of purified KatA, and 5 µg of whole cell extract were separated on a 10% SDS-PAGE gel and visualized by coomassie staining.

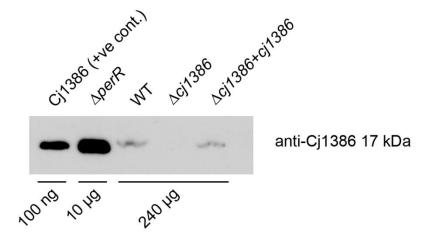


Figure S3. Cj1386 is expressed at low levels in wild-type *C.jejuni*. Bacterial cultures were grown in MEMα media at 37°C under microaerophilic conditions. Bacterial cultures were pelleted, resuspended in PBS + protease inhibitor and soluble proteins were extracted following sonication. Two hundred and forty micrograms of wild-type, $\Delta cj1386$, and $\Delta cj1386+cj1386$ lysate, 10 μg of $\Delta perR$ lysate, and 100 ng of purified Cj1386 protein were separated by SDS PAGE on a 14% polyacrylamide gel. Proteins were visualized by Western blotting using an anti-Cj1386 antibody.