

Supplementary Figures

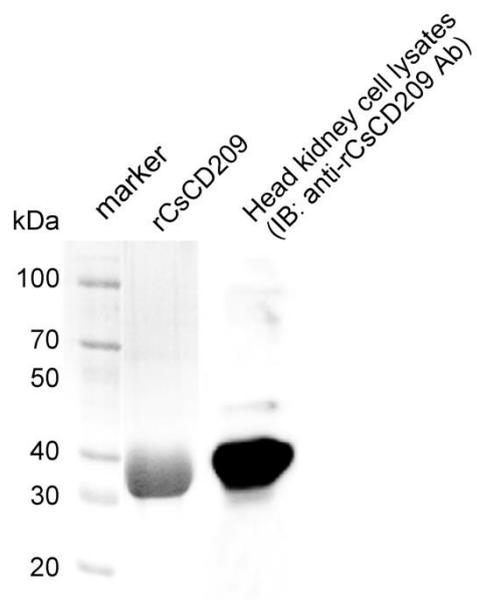


Figure S1. SDS-PAGE analysis of purified rCsCD209 and Western blot analysis of anti-rCsCD209 antibodies. rCsCD209 was expressed and purified by Ni-NTA chromatography, and further separated by SDS-PAGE. Western blot was performed to detect the CsCD209 in head kidney leukocytes by polyclonal antibodies against rCsCD209.

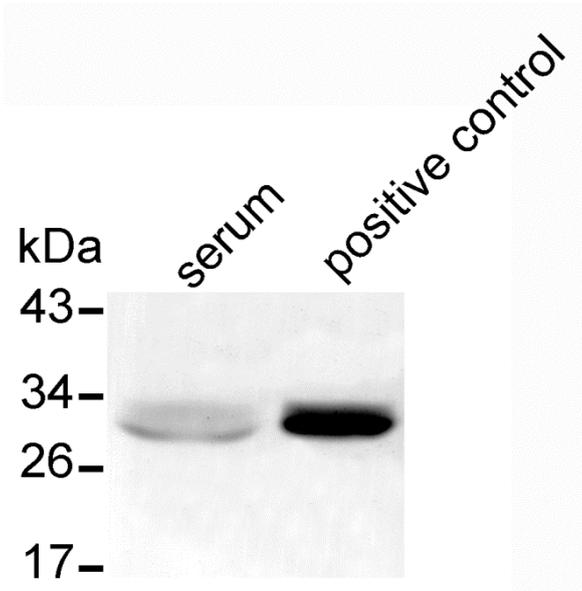


Figure S2. The soluble CsCD209 examined by Western blot. Serum protein was separated by SDS-PAGE, transferred to Polyvinylidene fluoride (PVDF) membrane and immunoblotted by anti-rCsCD209 antibody. Head kidney leukocyte proteins were used as a positive control.

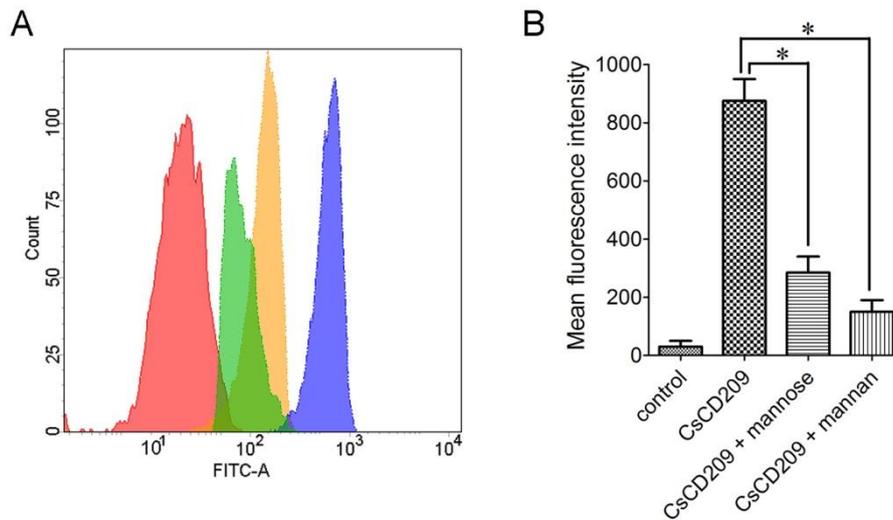


Figure S3. Effects of mannose and mannan on rCsCD29 binding to bacteria. **(A)** rCsCD29 was preincubated with or without mannose or mannan before incubating with *Edwardsiella tarda*. The cells were treated with FITC-labeled anti-His antibody, and rCsCD29-bacteria binding was determined by flow cytometry. Red histogram: control; blue histogram: rCsCD29; yellow histogram: rCsCD29⁺ mannose; green histogram: rCsCD29⁺ mannan. **(B)** The mean fluorescence intensity in **(A)** was statistically calculated. Results are means \pm SEM (n = 3), * $p < 0.05$.