Supplementary figures



Figure S1. Confirmation of endotoxin decontamination of the purified MAP1889c. BMDCs were incubated with LPS (100 ng/mL) or MAP1889c (10 μ g/mL) with or without pretreatment with polymyxin B (PmB) for 1 h. After 24 h, IL-10, TNF- α , and IL-12p70 production in the culture supernatants was analyzed by ELISA. All data are expressed as the mean ± SD (n = 3). * p < 0.05, and *** p < 0.001 for treatment compared with the difference between treatment data. *n.s.*, no significant difference.



Figure S2. Gating strategy for IL-10 cytokine from T cells for Figure 6E. (A) FMO control performed by excluding only the flour of interest. (**B**) After that, gating was selected based on the FMO bound line. Next, each of the CD4⁺ and CD8⁺ cells was selected. For cells in each of the CD4⁺ and CD8⁺ gates for IL-10 were then established.





Figure S3. Gating for transcription factor T-bet and GATA-3 from T cells for Figure 7B. (**A**) FMO control by excluding only the flour of interest. (**B**) After that, gating was selected based on the FMO bound line. Next, each of the CD4⁺ and CD8⁺ cells was selected. For cells in each of the CD4⁺ and CD8⁺ gates, individual gates for T-bet and GATA-3 were then established.



Figure S4. MAP1889c induces high levels of IL-10 production in BMDMs. The BMDMs were stimulated with MAP1889c (10 μ g/mL), LPS (100 ng/mL) for 24 h. IL-10, TNF- α , and IL-12p70 cytokines from the culture supernatants were measured by ELISA.