

SUPPLEMENTARY MATERIAL

Figure S1

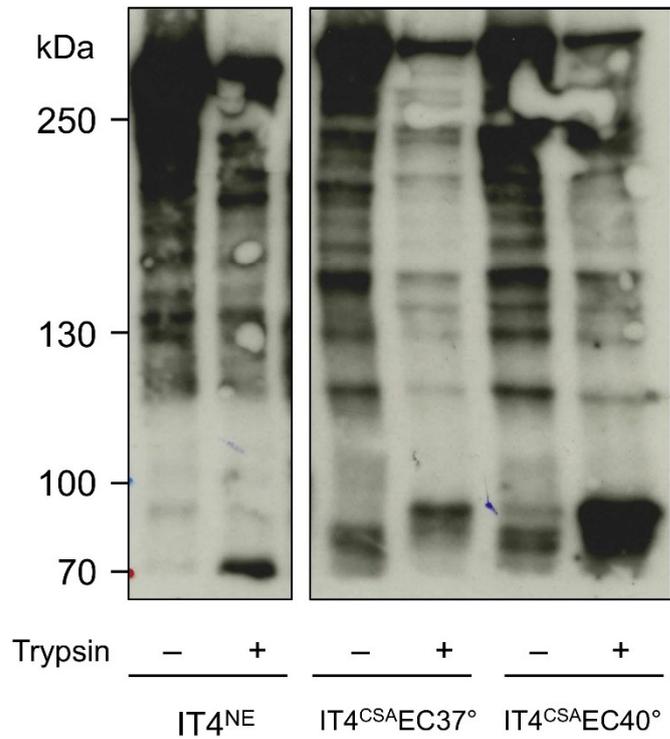


Figure S1: Western blot analysis for ATS was performed with trypsin-treated or untreated IEs and uninfected erythrocytes. All samples were solubilized in Laemmli sample buffer, separated by 6% SDS-PAGE, and analyzed by immunoblotting. Equivalents of 3×10^7 cells were loaded in each lane. The ATS antibody detects surface PfEMP1 as trypsin-cleaved intracellular ATS-residues.

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Figure S2

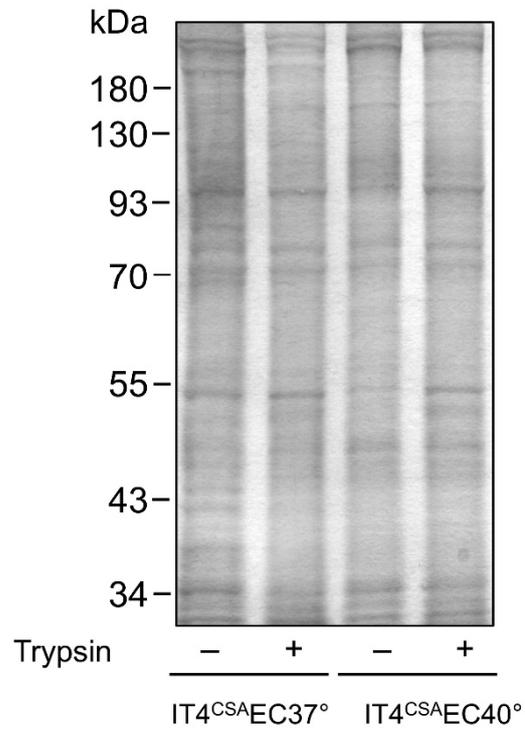


Figure S2: Coomassie-stained gel (10%) of membrane fractions (equivalents of 1×10^7 cells were loaded in each lane) of IT4^{CSA}EC37° and IT4^{CSA}EC40° parasite populations (control for Figure. 2A)