

Figure S1. A schematic illustration of the experimental design used in this study.

The cynomolgus macaques described in the initial study were used (7). After two years of rest (time= 0), animals were repeatedly boosted with a simultaneous intramuscular/ocular dose of LATV (filled triangles). Eight weeks after the second booster immunization, macaques were challenged ocularly with a ten fold higher dose of the virulent plasmid-bearing *C. trachomatis* strain A2497P+ (filled arrow). To evaluate the CD8+ T cell function in the LATV conferred protection, SP macaques received a subcutaneous injection of an anti-CD8 rhesus recombinant antibody (open squares). Three weeks following the initial injection, animals received a second dose of anti-CD8 antibody (open square) and simultaneously were ocularly challenged with 2 x10<sup>4</sup> IFU/eye (open arrow).

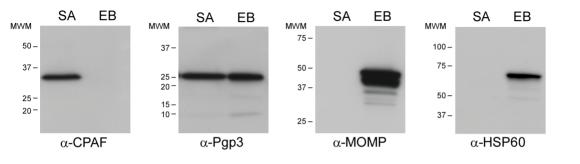


Figure S2. Western blots of *C. trachomatis* A2497 SA and EBs probed with monoclonal antibodies specific for chlamydial secreted and structural antigens. Western blotting of SA and EBs probed with Mabs specific to chlamydial protease activity factor (anti-CPAF), plasmid-encoded Pgp3 protein (anti-Pgp3), serovar A major outer membrane protein (anti-MOMP), and chlamydial heat shock protein 60 (anti-HSP60). Note that the secreted chlamydial protease CPAF is found primarily in the SA antigen preparation.

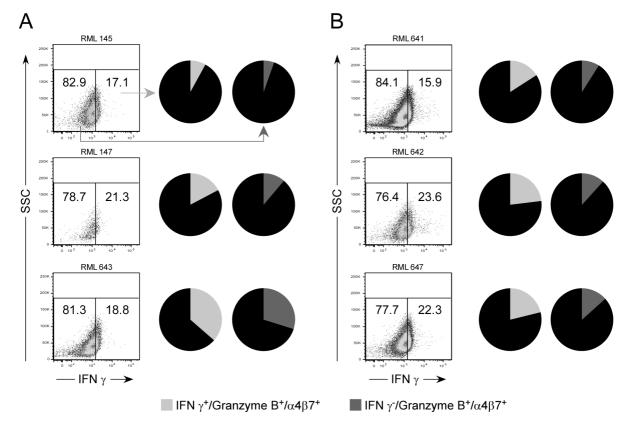


Figure S3. Chlamydial CD8<sup>+</sup> T cells responding to SA express IFN- $\gamma$ , granzyme B and α4β7 integrin. CFSE-PBMC antigen stimulated for five days and re-stimulated for additional six hours were subjected to a multi-color flow cytometry assay. Frequencies for CD8<sup>+</sup> antigen-specific IFN- $\gamma$ <sup>+</sup>, α4β7<sup>+</sup>, GraB<sup>+</sup> and IFN- $\gamma$ <sup>-</sup>, α4b7<sup>+</sup> GraB<sup>+</sup> responses were determined. The percentages of these T cell phenotypes found in PP (Fig. S3A) and SP (Fig. S3B) animals are shown in the pie charts. No differences in the frequency of individual surface markers or a combination of markers in SA stimulated CD8<sup>+</sup> T cells were detected between the groups.