

Supporting Information

Liu et al. 10.1073/pnas.1109796108

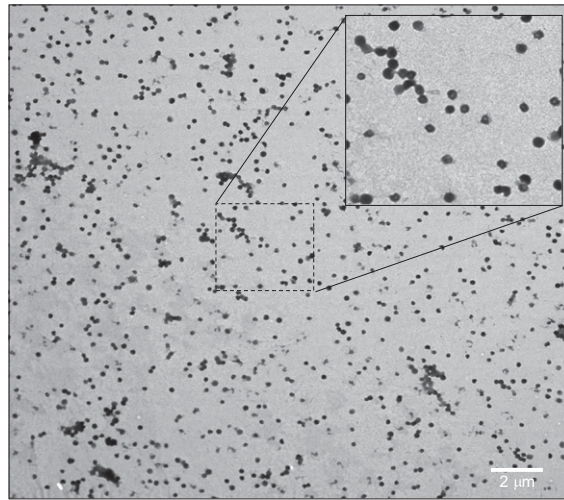


Fig. S1. Purified virus is substantially free of cellular debris. Purified virus was negatively stained with 1% uranyl sulfate and observed by electron microscopy. The expanded box is 3 μm in diameter.

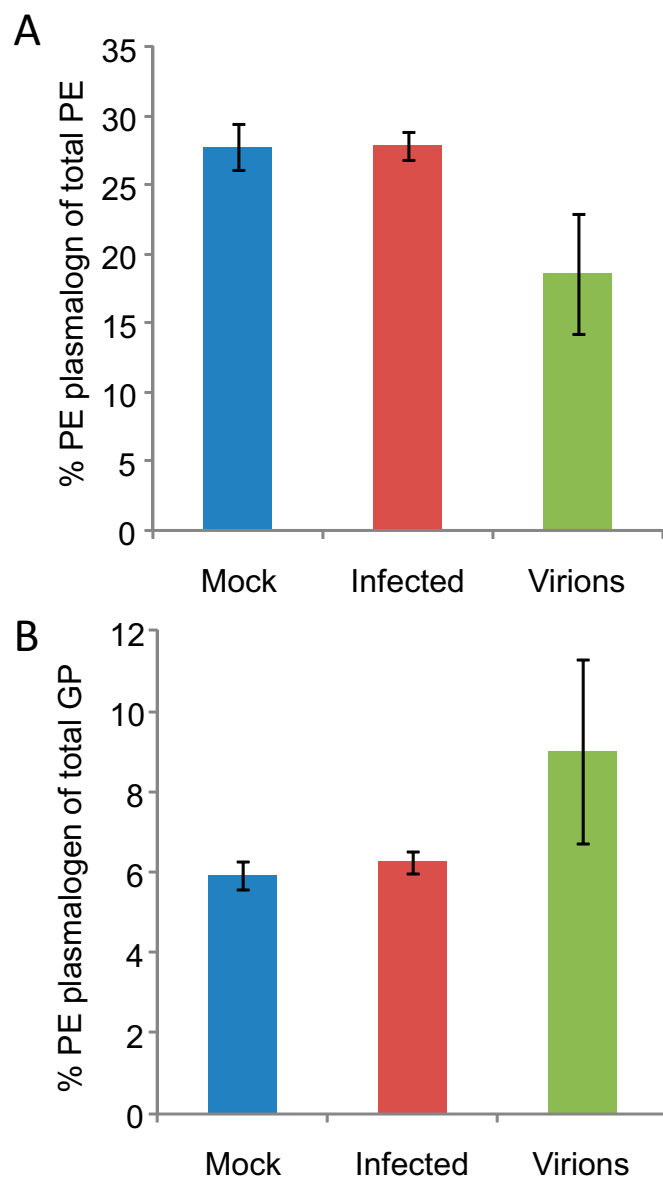


Fig. S2. Plasmalogen species in virions. (A) The pool of PE in virions has less representation by plasmalogen species than either mock- or HCMV-infected primary fibroblasts at 96 hpi ($P = 0.016$ by ANOVA; $P = 0.033$ for virion vs. uninfected post hoc, and $P = 0.010$ between virion vs. uninfected post hoc). (B) Virions are enriched for plasmalogen phosphatidylethanolamines (PEs) as a fraction of total glycerophospholipids (GP) compared with mock-infected or human cytomegalovirus-infected primary fibroblasts ($P = 0.026$ by ANOVA; $P = 0.043$ for virions vs. uninfected post hoc, and $P = 0.055$ for virion vs. infected post hoc). Error bars indicate SEM.

Dataset S1. Lipidomics data from time courses of mock- and human cytomegalovirus-infected primary fibroblasts

[Dataset S1](#)

Means and SEs from three independent experiments are shown.

Dataset S2. Glycerophospholipid profile of human cytomegalovirus virion

[Dataset S2](#)

Means and SEs from three independent experiments are shown.