1	Supplementary Data
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3	Limited replication of human cytomegalovirus in a trophoblast cell line
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12	Supplementary Figure 1 Analysis of fluorescent protein expression in HFF and
13	SGHPL-4 cells. Low and high passage HFF and SGHPL cells (HFF passage 6, SGHPL-
14	4 passage 13) (A and C, B and D, respectively) were incubated in 0.5% (v/v) media for
15	24 hours before infection with an MOI of 0.5 with green fluorescent protein (GFP)
16	expressing virus Merlin(R1111)UL36GFP (green) or mock infected (grey). After 24 hours
17	uninfected and infected cells were analyzed for GFP expression using FACS. The
18	percentage of uninfected and infected cells detected in the FACS channel detecting GFP
19	in each condition is noted in each panel. The data presented in this figure is representative
20	of two independent experiments.
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22	Supplementary Figure 2 Replication of different HCMV strains in HFF and SGHPL-

4 cells. (A) Low passage HFF and SGHPL cells (HFF passage 6-10, SGHPL-4 passage

24 13-17) were incubated in 0.5% (v/v) media for 24 hours before infection with an MOI of 1 25 with the HCMV strains shown in the figure. After 96 hours post infection in 0.5% (v/v) media viral titre (p.f.u./ml) was determined by titration of viral supernatant on HFF cells. 26 27 Each data point represents the data from three independent experiments. The bar chart 28 and error bars represent the mean and standard deviation of that data, respectively. The 29 statistical difference between the indicated conditions was measured using an unpaired t 30 test (two-tailed) and is indicated above each figure. A statistically relevant difference was 31 where p = < 0.05. Not significant (ns). (B) Low and high passage HFF and SGHPL cells 32 (HFF passage 6, SGHPL-4 passage 14) were prepared for western blotting or incubated in 0.5% (v/v) media for 24 hours before preparation for western blotting. Proteins 33 34 recognized by the antibodies used in the experiment are indicated to the right of each 35 western blot panel. The presence of β -actin was assayed to assess the amount of cell 36 lysate assayed in each lane. The positions of molecular weight markers (kDa) are 37 indicated to the left of the figure. (C) Cells were infected with Merlin(R1111) as in (A) and 38 virus was harvested at the indicated time points. The data from three independent 39 experiments was presented. The bar chart and error bars represent the mean and 40 standard deviation of that data, respectively.

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Supplementary Figure 3 Replication of ZIKV in different cell lines. The cell lines indicated in the figure were incubated in 10% (v/v) media and infected with the ZIKV strain PE243 (MOI 0.1). In all experiments viruses were harvested at 48 hours post infection and viral titre (p.f.u./ml) was determined by titration of viral supernatant on Vero cells. The data from three independent experiments was presented. The bar chart and error bars

47 represent the mean and standard deviation of that data, respectively. The statistical 48 difference between the indicated conditions was measured using an unpaired t test (two-49 tailed) and is indicated above each figure. A statistically relevant difference was where 50 p=<0.05.



Supplementary Figure S1







Supplementary Figure S2





