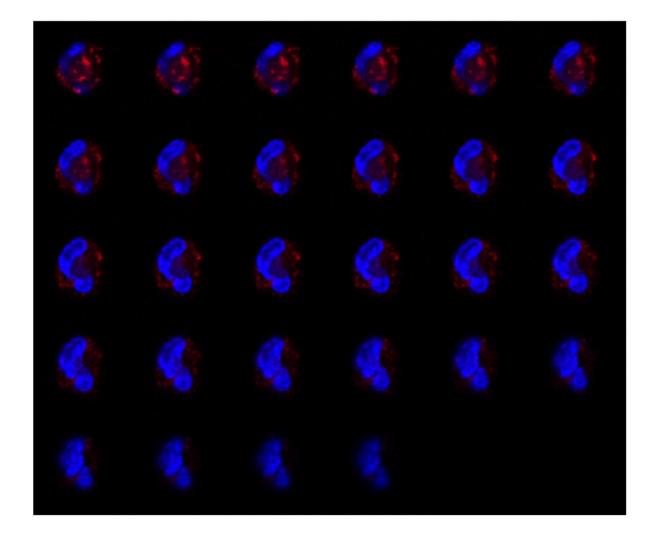


Supplementary Figure S1. Extracellular Tat protein uptake by activated endothelial cells after 30 minutes

exposure. HUVEC (blue plots) or IC-HUVEC (red plots) were incubated for 30 min in medium containing serial concentrations (1-1000 ng/mL) of biologically active Tat or its suspension buffer (PBS-0.1% BSA). The intracellular Tat content was evaluated by flow cytometry after staining with affinity-purified rabbit anti-Tat polyclonal Ab (or isotype control), as described in the Materials and Methods section. Non-permeabilized cells were also analyzed by intracellular staining and flow cytometry, with negative results. Results are expressed as the percentage of positive cells, as compared to isotype-stained samples. Box–plot of data obtained from three independent experiments and analyzed by the Mann-Whitney nonparametric test are shown.



Supplementary Figure S2. Extracellular Tat protein is efficiently taken up by activated endothelial cells. Images of serial sections of the cell analyzed in Fig. 1 B and C showing the presence of Tat (red) in the optical sections of the central region of the cell.