

VIROLOGY 5-7 September 2011 Baltimore, USA

International Conference and Exhibition on

Suppression of HIV-1 by DING

Shohreh Amini^{1,2}, Kamel Khalili¹ and Nune Darbinian-Sarkissian¹

¹Department of Neurocience and Center for Neurovirology, Temple University School of Medicine, USA

²Department of Biology, Temple University College of Science and Technology, Temple University, USA $P_{John's}$ Wort) that belongs to an emerging family of DING proteins. We have demonstrated that the interaction of p27SJ with HIV-1 Tat and C/EBP β greatly impacts the transcriptional activity of these two proteins that play a key role in replication of HIV-1 in microglia and macrophages. Further, we demonstrate that while the presence of p27SJ in cells interferes with the nuclear localization of C/EBP β , due to its phosphatase activity, expression of p27SJ interferes with phosphorylation of the carboxyl terminus of RNA polymerase II by cyclin T/ cdk9/Tat complex, an event that is critical for activation of the LTR promoter by Tat. Our results provide a new avenue for the development of therapeutic intervention against HIV-1 infection and AIDS.

This work was supported by grants from NIH awarded to S.A.