Biosketch

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Dr. Levis received her PhD at Washington University in St Louis in 1988. She has worked at the US Food and Drug Administration since 1995. She is currently the Deputy Director of the Division of Viral Products in the Office of Vaccines Research and Review at CBER/FDA; a position she has held since 2006. Prior to this position, she served as the Regulatory Coordinator for the Division of Viral Products (2002-2006) and served as a Senior Staff Fellow in the Laboratory of Vector Borne Viral Diseases (1995-2002). Her initial research work at the FDA related to dengue virus replication. She then transitioned to be the lead subject matter expert and CMC reviewer for licensed rabies virus vaccine products and rabies vaccine and related products under development. Her laboratory work on rabies virus vaccines continued the collaborative efforts underway to develop an alternative, in vitro based potency test for rabies virus vaccines. This work is ongoing in the Laboratory of Methods Development and in collaboration with a global working group.