

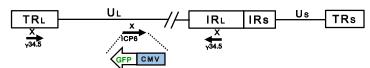
Product Description

Product Name:	HSV-GFP (Standard Titer, 0.25 mL)
Catalog Number:	OV3001
Lot Number:	OV-IM10
Shipping conditions: Storage conditions:	Dry ice Store at ≤ -70°C upon receipt, avoid subjecting to multiple freeze-thaw cycles
Species:	Herpes simplex virus-1 (HSV-1)
Virus strain:	Strain 17+
Transgene:	Green fluorescent protein (GFP)
Production Cell Line: Storage Media: Propagation:	Vero Serum-free media Inoculate producer cells (e.g. Vero) at an MOI of 0.01 for 2-3 hours. Harvest culture supernatant and cells when 80-90% of the cell monolayer is showing cytopathic effects (CPE); freeze-thaw 3 cycles and clarify by low-speed centrifugation. Centrifuge the supernatant at 14,000x g for 1h at 4°C.

HSV-GFP is a recombinant of the strain 17+ of Herpes simplex virus-1 (HSV-1). HSV-GFP contains a deletion of both copies of ICP34.5. The human cytomegalovirus (CMV) promoter followed by the GFP cDNA is inserted in the ICP6 reading frame (see diagram below). The purchased virus is ready for use in cell killing assays or may be amplified to generate additional virus stocks.

Resuspend the viral pellet in DPBS.

Genome:



References:

Campadelli-Fiume, et al. Rev. Med. Virol. (2011), 21(4):213-26. Li, et al. Cancer Gene Ther. (2013), 20(8): 478-485. Maneta, et al., Nat. Med. (1995), 1(9):938-43. Chou, et al., Science (1990), 250, 1262-1266. Kramm, et al., Human Gen. Therapy (1997), 8:2057-2068 Bolovan, et al., J Virol. (1994), 68(1): 48-55.

Safety Precaution:

All culture work with HSV should be performed by trained personnel and performed under BSL2 containment following NIH guidelines.

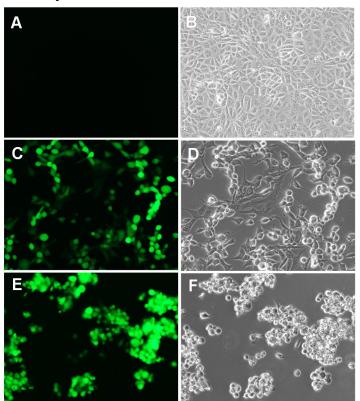


Certificate of Analysis

Testing performed by Imanis Life Sciences

Test description	Result
Titer (on Vero)	2.89x107 TCID ₅₀ units/mL
Sterility	Pass QC
Cytopathic Effect	Pass QC
Endotoxin Test	Pass QC
Fluorescence expression	Pass QC

Infectivity:



A&B: Mock-infected Vero cells C&D: HSV-GFP-infected Vero cells (MOI 0.01) at 24 h.p.i. E&F: HSV-GFP-infected Vero cells (MOI 0.01) at 48 h.p.i

Quality Control by: MT Quality Assurance by: RLV Effective Date: 06-Feb-2017



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