

Anti-HHV6 gQ1 antibody, mouse monoclonal (119)

Product code	65-200
Size	50 μg
Storage	-20°C
Concentration	1.0 mg/ml
Buffer	PBS- with 50% glycerol
Purity	Purified IgG fraction with protein A from hybridoma cell culture medium.
Immunogen	His6-tagged recombinant gQ1 of HHV-6A encoding 3-422 amino acids expressed in <i>E. coli</i> .
Isotype	Mouse IgG1κ
Reactivity	gQ1 of HHV6A and HHV6B. However, this antibody is not recommended for IP of
	HHV-6B due to conformation specificity.
Special notes	N/A
Application	1. Western blotting (1/500~1/1,000 dilution)
	2. Immunoprecipitation (assay dependent)
	3. Imunofluorescence staining and Immunocytochemistry (1/100~1/3,200 dilution)
	4. Flow Cytometry (1/100)
	5. ELISA (assay dependent)
Background	Human herpesvirus 6 (HHV-6) is the common collective name for Human herpesvirus 6A (HHV-6A) and
	$Human\ herpesvirus\ 6B\ (HHV-6B).\ These\ closely\ related\ viruses\ are\ two\ of\ the\ nine\ herpesviruses\ known$
	to have humans as their primary host. HHV-6A and HHV-6B are double stranded DNA viruses within
	the betaherpesvirinae subfamily and of the genus Roseolovirus. HHV-6A and HHV-6B infects almost all
	of the human populations tested. The overall nucleotide sequence identity between HHV-6A and HHV-
	6B is 90% and they are now classified as distinct species.
	HHV-6A has been described as more neurovirulent, and as such is more frequently found in patients
	with neuroinflammatory diseases such as multiple sclerosis.
	HHV-6B primary infection is the cause of the common childhood illness exanthem subitum (also
	$known\ as\ roseola\ infantum\ or\ sixth\ disease).\ Additionally,\ HHV-6B\ reactivation\ is\ common\ in\ transplant$
	recipients, which can cause several clinical manifestations such as encephalitis, bone marrow
	suppression and pneumonitis.
	$\mathrm{gQ1}$ encoded by the U100 gene of HHV6 is glycoprotein, complexes with gH, gL and $\mathrm{gQ2}$ to form
	m HHV6Aligand to CD46 receptor and HHV6B ligand to CD134 receptor. It is expressed in two different
	forms: an 80-kDa form (gQ1-80K) and a 74-kDa form (gQ1-74K) - only gQ1-80K, but not gQ1-74K,
	forms the ligand complex with gQ2, gH, and gL. It associates with lipid rafts.Molecular Mass calculated
	from the gene is $59.9~\mathrm{kDa}$, $524~\mathrm{amino}$ acids.
Data Link	UniProtKB <u>Q69572</u> (GQ1_HHV6U)
Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC	
PROCEDURES. NOT FOR MILITARY USE.	



Data Images: 65-200 Anti-HHV6 gQ1 antibody, mouse monoclonal (119)

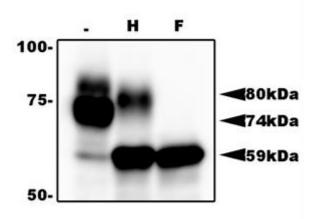
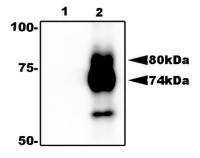


Fig.1. Identification of gQ1 in HHV-6A infected cells by western blotting using anti-gQ antibody (119).

T-cell line HSB-2 cells were infected with HHV-6A at

m.o.i of 0.1 and the cells were harvested at 72 h postinfection for lysate preparation for WB. gQ1 is detected as two glycoproteins with 80 kDa and 74 kDa molecular masses..

Sample lysates: (-) Non-treated. (H) Treated with end-glycosilase H. (F) Treated with peptide N-glycosidase F.



1. HSB-2 2. GS-infected HSB-2

Fig.2. Immunoprecipitation of gQ1 from HHV-6A infected cell lysate with anti-gQ1 antibody (119). gQ1 protein was precipitated from the lysate of infected HSB-2 cells by using agarose beads-conjugated anti-gQ1 antibody (119), and processed for western blotting by using anti-gQ1 antibody (119). Lane 1; HSB-2 cells. Lane 2; HSB-2 cells infected with HHV-6A GS strain



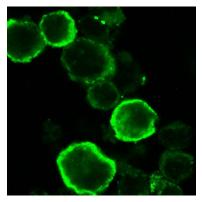


Fig.3. Immuno-staining of gQ1 in HHV6A infected HSB-2 cells by using anti-HHV6 gQ1 antibody.

The infected cells were harvested 3 days postinfection, fixed in cold acetone and immunostained with FITC-conjugated anti HHV6 gQ1 antibody (119). Specific immunofluorescence was observed with a confocal laser scanning.

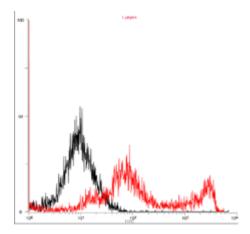


Fig.4. Flow cytometry analysis of gQ1 expression on the cell surface in a transient expression system.

293T cells were transfected with gQ1 expression plasmid. The cells were harvested on second day postinfection, fixed with 4% PFA, permeabilized with 0.1% Triton-X100, incubated with anti-gQ1 antibody (119) and then with FITC-conjugated anti-mouse IgG antibody. Histograms show fluorescence intensities measured in arbitrary units on a log scale (x axis) and relative cell numbers on a linear scale (y axis). Black line is control—and red line is gQ1 introduced cells.

References: This antibody has been described in Ref. 1 and used in Ref.1-7.

- 1. Mori Y. et al. (2003) The Human Herpesvirus 6 U100 Gene Product Is the Third Component of the gH-gL Glycoprotein Complex on the Viral Envelope. J. Virol. 77: 2452-2458. PubMed 12551983 WB, IP
- 2. Mori Y. et al (2003) Human Herpesvirus 6 Variant A Glycoprotein H-Glycoprotein L-Glycoprotein Q Complex Associates with Human CD46. J. Virol. 77: 4992-4999. <u>PubMed 12663806</u> WB, IP
- 3. Mori Y. et al. (2004) Discovery of a second form of tripartite complex containing gH-gL of human herpesvirus 6 and observations on CD46. J Virol. 78:4609-16. PubMed 15078943. WB, IP
- 4. Akkapaiboon P. et al. (2004) Intracellular processing of human herpesvirus 6 glycoproteins Q1 and



- Q2 into tetrameric complexes expressed on the viral envelope. J Virol. 78:7969-83. PubMed 15254169 WB, IP, IF
- 5. Huang H. et al (2006) Human herpesvirus 6 envelope cholesterol is required for virus entry. J Gen Virol. 87: 277-285. PubMed 16432012 WB, Flow Cyt.
- 6. Tang H. et al. (2010) Human herpesvirus 6 encoded glycoprotein Q1 gene is essential for virus growth. Virology. 407:360-7. PubMed 20863544 WB
- 7. Tang H. et al (2011) Human herpesvirus 6 glycoprotein complex formation is required for folding and trafficking of the gH/gL/gQ1/gQ2 complex and its cellular receptor binding. J Virol. 85:11121-30. PubMed 21849437 WB, IP, Flow Cyt.