

### **CERTIFICATE OF ANALYSIS**

# Canine Herpesvirus Type 1 (CHV-1)

# FITC Conjugate

Catalog No.:	CJ-F-CHV-10ML
Volume:	10 ml
Lot:	P140407-003
Expiration:	03 January 2022
Agent:	Canine Herpesvirus Type 1 (CHV-1)
Strain:	NA

# CHV-1 FITC CJ P140407-003 400X

### Description:

CHV-1 polyclonal antiserum conjugated to fluorescein isothiocyanate (FITC). Canine origin. Ready to use. Liquid.

# Quality Control Method:

Direct FA using CHV-1 12 well slide (catalog no. SLD-IFA-CHV).

**Specific Reaction**: 2-3+ signal with trace background.

Other Comments: The raw material has also been screened by Indirect FA and does not react with

canine adenovirus type 1 and 2 (CAV-1 and 2), canine coronavirus (CCV), canine

distemper virus (CDV), canine parainfluenza virus type 2 (CPI-2), canine parvovirus (CPV), reovirus (REO), Rabies Recombinant Nucleoprotein (rNP), *Toxoplasma gondii*, vesicular stomatitis virus Indiana and New Jersey strains

(VSV).

### Pattern Of Fluorescence:

Cytoplasmic and nuclear fluorescence in rounded cells with occasional small syncytia.

### Intended Use:

This reagent is suitable for CHV-1 virus identification in cell cultures and in animal tissues.

### Storage:

This conjugate is provided in liquid form and should be stored at 2-7°C. DO NOT FREEZE! It should also be stored in the original container and/or in the dark. If conjugate becomes cloudy it should be discarded. This conjugate contains 0.09% sodium azide as a preservative.

### References:

Ledbetter EC, et al., Experimental reactivation of latent canine herpesvirus-1 and induction of recurrent ocular disease in adult dogs. Vet Microbiol.2009 Jul 2;138(1-2):98-105. Epub 2009 Mar 13.

P: 509.334.5815

F: 509.332.5356

# Recommended Staining Procedure for Indirect FA:

- 1. Warm slide to room temperature before removing from foil pouch.
- 2. Place diluted serum on the designated wells. Dilute serum in serum diluting buffer, pH 7.2 (catalog no. FASDB-100ML) however if high background due to anti-bovine IgG activity is present it may be advisable to use SSDB-100ML.
- 3. Incubate slide in humid chamber at 37°C for 30 minutes.
- 4. Using a wash bottle, gently rinse slide briefly in FA rinse buffer, pH 9.0 (catalog no. FARB-4X) and then soak for 10 minutes in FA rinse buffer, pH 9.0.
- 5. Drain slide and dry around wells by pressing blotter (included in pouch) to front surface. Place labeled anti-IgG or IgM on the wells.
- 6. Incubate as in step 3.
- 7. Rinse as in step 4.
- 8. Drain slide and dry back and edges with a paper towel. Do not allow stained surface to dry. Do not rinse with water.
- 9. Mount with mounting fluid [glycerol/FA rinse buffer, pH 9.0, (50/50)] (catalog no. FAMF-10ML) and view with good quality fluorescence microscope at 100X-250X. Confirmation may be made at 400X.

### Recommended Staining Procedure for Direct FA:

- 1. Warm slide to room temperature before removing from foil pouch.
- 2. Place direct FA conjugate on the designated wells.
- 3. Incubate slide in humid chamber at 37°C for 30 minutes.
- 4. Using a wash bottle, gently rinse slide briefly in FA rinse buffer, pH 9.0 (catalog no. FARB-4X) and then soak for 10 minutes in FA rinse buffer, pH 9.0.
- 5. Drain slide and dry back and edges with a paper towel. Do not allow stained surface to dry. Do not rinse with water.
- 6. Mount with mounting fluid [glycerol/FA rinse buffer, pH 9.0, (50/50)] (catalog no. FAMF-10ML) and view with good quality fluorescence microscope at 100X-250X. Confirmation may be at 400X.

# Serum Diluting Buffer (pH 7.2):\*

-	Na <sub>2</sub> HPO <sub>4</sub>	1.19 gm
-	NaH <sub>2</sub> PO <sub>4</sub>	0.22 gm
-	NaCl	8.55 gm
-	BSA	10.0 gm
-	DI/dH <sub>2</sub> O	Q.S. to 1 liter

<sup>\*</sup>This recipe makes 1 liter. If you need less, adjust recipe accordingly. Store at 2-7°C. Add 0.09% NaN₃ if diluted serum is not going to be used within one week.

# 4X FA Rinse Buffer (pH 9.0):

-	Na <sub>2</sub> CO <sub>3</sub>	11.4 gm
-	NaHCO <sub>3</sub>	33.6 gm
-	NaCl	8.5 gm
-	DI/dH <sub>2</sub> O	Q.S. to 1 liter

Final pH should be 9.0-9.5. This is a 4X concentrate and should be diluted 1/4 with DI/distilled water for use as a working buffer. Keep in a tightly stoppered container at room temperature. MOUNTING FLUID is made by mixing glycerol and FA rinse buffer, pH 9.0, in equal proportions.