

#### **CERTIFICATE OF ANALYSIS**

# FA Serum Diluting Buffer

Catalog No.:	FASDB-100ML
Volume:	100 ml
Lot:	P140123-003
Expiration:	17 March 2018

# Description:

Serum Diluting Buffer is PBS with 1% BSA, preserved with 0.09% sodium azide.

# Quality Control Method:

Indirect FA using positive controls, corresponding secondary conjugates and slides on one each of the following systems: bovine PI-3 (catalog no. PC-IFA-PI3, CJ-F-BOVG-AP-1ML or 10ML, and SLD-IFA-PI3), canine CDV (catalog no. PC-IFA-CDV-G, CJ-F-CANG-1ML or 10ML, and SLD-IFA-CDV), equine EHV-1/ERV (catalog no. PC-IFA-ERV-EQ, CJ-F-EQUG-1ML or 10ML, and SLD-IFA-ERV), llama EHV-1/ERV (catalog no. PC-IFA-ERV-LL, CJ-F-CAMG-AP-1ML or 10ML, and SLD-IFA-ERV), feline FHV-1/FVR (catalog no. PC-IFA-FVR, CJ-F-FELG-1ML or 10ML, and SLD-IFA-PCV2), caprine PCV Type 2 (catalog no. PC-IFA-PCV2, CJ-F-PORG-AP-1ML or 10ML, and SLD-IFA-PCV2), caprine CDV antiserum (catalog no. PAB-CDV, CJ-F-CAPG-1ML or 10ML and SLD-IFA-CDV), and murine MAb CDV-NP (catalog no. CDV-NP, CJ-F-MURG-AP-1ML or 10ML and SLD-IFA-CDV).

Specific Reaction: There is little to no difference between this lot of FASDB-100ML and the previous

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lot.

Other Comments: NA

#### Intended Use:

Serum Diluting Buffer should be used for diluting serum in IFA testing.

#### Storage:

Store at 2-7°C.

# 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 lite

<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 lite

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.