

CERTIFICATE OF ANALYSIS

SERUM DILUTING BUFFER (FASDB)

IFA Serum Diluting Buffer

Catalog No.:	FASDB-100ML
Volume:	100 ml
Lot:	P120517-003
Expiration:	10 September 2016

Description:

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

Quality Control Method:

Indirect FA using VMRD 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 ERV (catalog no. PC-IFA-ERV-EQ, CJ-F-EQUG-1ML or 10ML, and SLD-IFA-ERV), Llama ERV (catalog no. PC-IFA-ERV-LL, CJ-F-CAMG-AP-1ML or 10ML, and SLD-IFA-ERV), Feline FVR (catalog no. PC-IFA-FVR, CJ-F-FELG-1ML or 10ML, and SLD-IFA-FVR), Porcine TGE (catalog no. PC-IFA-TGE, CJ-F-PORG-AP-1ML or 10ML, and SLD-IFA-TGE), Goat CDV Antiserum (catalog no. PAB-CDV, CJ-F-CAPG-1ML or 10ML and SLD-IFA-CDV), and Mouse MoAb 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 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.

Note: Microscopic precipitates may appear in this product and it is recommended that a short high speed centrifugation (approximately 10,000xg for 3 min) be performed to clarify it.

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):*

- Na₂HPO₄.....1.19 gm
- NaH₂PO₄.....0.22 gm
- NaCl.....8.55 gm
- BSA.....10.0 gm
- DI/dH₂O.....Q.S. to 1 liter

*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):

- Na₂CO₃.....11.4 gm
- NaHCO₃.....33.6 gm
- NaCl.....8.5 gm
- DI/dH₂O.....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.