

**Center for Veterinary Biologics
and
National Veterinary Services Laboratories
Testing Protocol**

**Supplemental Assay Method for Potency Testing
Enterotoxigenic (K88 Pilus) *Escherichia coli* Bacterins**

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1. Introduction

This Supplemental Assay Method (SAM) for potency testing inactivated *Escherichia coli* bacterins employs a capture enzyme-linked immunosorbent assay (ELISA) for K88 pilus antigen. Relative potency is determined by comparing the K88 antigen content of the test bacterin to the K88 antigen content present in a nonexpired, suitably qualified reference bacterin.

2. Materials

2.1 Equipment/instrumentation

- 2.1.1 Micropipettors, to cover the range of 5.0 μ l to 1000 μ l
- 2.1.2 8- or 12-channel micropipettor, to cover the range of 50 μ l to 200 μ l
- 2.1.3 Orbital shaker
- 2.1.4 Automatic microplate washer (optional)
- 2.1.5 Microplate reader with dual wavelengths (490 nm and 650 nm)
- 2.1.6 Balance, validated from 150 mg to 15 g
- 2.1.7 Relative Potency Calculation Software (United States Department of Agriculture [USDA], Veterinary Services, Center for Veterinary Biologics-Laboratory [CVB-L]), current version

2.2 Reagents/supplies

- 2.2.1 96-well flat-bottom microtitration plates (Immulon 2, Dynatech Laboratories, Inc., or equivalent)
- 2.2.2 96-well microtitration plates suitable for making serial dilutions (transfer plate)

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- 2.2.3 Plate sealers
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3. Preparation for the test

3.1 Personnel qualifications/training

Technical personnel need a working knowledge of the use of general laboratory chemicals, equipment, and glassware; automated microplate washer and microplate reader; and data analysis software. They need specific training in the performance of this assay.

3.2 Preparation of equipment/instrumentation

3.2.1 Operate and maintain all equipment according to manufacturers' recommendations.

3.2.2 Validate the ELISA microplate reader according to the current version of GDOCSOP0005.

3.3 Preparation of reagent/control procedures

3.3.1 Carbonate coating buffer--NVSL media 20034

Na ₂ CO ₃	0.159 g
NaHCO ₃	0.293 g
Deionized water	q.s. to 100 ml

Adjust pH to 9.6 ± 0.1. Store at 2°-7°C up to 1 wk.

3.3.2 Phosphate buffered saline--NVSL media 10559

NaCl	8.00 g
KCl	0.20 g
Na ₂ HPO ₄	1.15 g
KH ₂ PO ₄	0.20 g
Deionized water	q.s. to 1 L

Adjust pH to 7.2 ± 0.1. Store at 20°-25°C no longer than 6 mo.

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3.3.3 Phosphate buffered saline with 0.05% Tween 20--
NVSL media 30179

PBS (see Section 3.3.2)	1000 ml
Tween 20	0.50 ml

Store at 20°-25°C no longer than 6 mo.

3.3.4 Phosphate buffered saline with 0.05% Tween 20
and 1.0% ovalbumin (conjugate diluent)

PBS-Tween 20 (see Section 3.3.3)	100 ml
Ovalbumin (Calbiochem 32467 or equivalent)	1.0 g

Add ovalbumin to the PBS-Tween 20 within 10 min of use.
Swirl gently to dissolve the crystals.

3.3.5 Phosphate elution buffer

KH ₂ PO ₄ (Mallinckrodt, Inc. 7100 or equivalent)	8.2 g
Deionized water	94 ml

Adjust pH to 9.3 ± 0.1, or other appropriate pH as
optimized for use with a specific bacterin. Store at
20°-25°C no longer than 1 mo.

3.3.6 Sodium desoxycholate elution buffer

Sodium desoxycholate (Difco Laboratories 0248-13-7 or equivalent)	0.5 g
PBS (see Section 3.3.2)	100 ml

Store at 2°-7°C for up to 30 days. Warm to room
temperature prior to use (the buffer gels at 2°-7°C).

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3.3.7 Citrate buffer (pH 5.0)--NVSL media 20033

Citric acid (Fisher Scientific A104-500 or equivalent)	5.26 g
Na ₂ HPO ₄ •7H ₂ O	6.74 g
Deionized water	q.s. to 1 L

Adjust pH to 5.0 ± 0.1. Store at 2°-7°C up to 60 days.
Use to prepare substrate solution (see **Section 3.3.8**).

3.3.8 Substrate solution (quantities for 1 plate)

Citrate buffer	12 ml
o-phenylenediamine (OPD) (Sigma P6787 or equivalent)	4 mg
30% H ₂ O ₂ (stabilized)	5 µl

Prepare within 15 min of use.

Caution: o-phenylenediamine is a carcinogen. See appropriate MSDS for precautions when handling this product.

3.3.9 Stop solution (2.5 M H₂SO₄)--NVSL media #30171

Concentrated (98%) H ₂ SO ₄	10 ml
Deionized water	90 ml

Add acid to water.

3.3.10 Monoclonal antibodies (MAb)

3.3.10.1 K88 antigen-capture monoclonal antibody (21BA1-1H1). Obtain MAb from the Center for Veterinary Biologics-Laboratory (CVB-L). Store the undiluted MAb at -70°C or less for long-term storage. It may be held at 2°-7°C for several weeks.

3.3.10.2 Horseradish peroxidase-labeled K88 antigen-indicator MAb (21BA1-1H1). Obtain conjugated MAb from the CVB-L. Store the conjugated MAb at 2°-7°C.

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3.3.11 Bacterins containing K88 antigen

3.3.11.1 Reference bacterin

3.3.11.2 Test bacterin(s)

CRITICAL CONTROL POINT: Ideally, the reference and test bacterins should be produced by the same Outline of Production. If reference formulation differs from that of the test bacterin, the assay must be validated to show that this does not adversely affect assay performance or accuracy of results.

3.4 Preparation of the sample

3.4.1 Antigen-elution treatments

Many bacterins do not require antigen-elution treatment prior to being serially diluted in twofold increments with PBS-Tween 20. Test representative batches of each adjuvanted product with and without each antigen-elution treatment to determine if the treatment specifically enhances the K88 antigen capture. If no enhancement of the K88 antigen capture can be demonstrated, test the bacterins without antigen-elution treatment. Treat the reference bacterin and the test bacterins by the same elution procedure. Alternate elution procedures, other than those described here, may be more appropriate for some bacterins.

3.4.1.1 Aluminum hydroxide adjuvanted bacterins

Bacterins adjuvanted with aluminum hydroxide may be treated with either sodium citrate or phosphate buffer prior to making serial twofold dilutions in PBS-Tween 20.

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1. Sodium citrate elution

Mix 1 g sodium citrate with 10 ml of bacterin (10% w/v). Place on an orbital shaker (100-120 rpm) overnight at 36°-38°C. Consider treated bacterin to be undiluted.

2. Phosphate buffer elution

Mix 1 ml of phosphate elution buffer with 1 ml bacterin. Place on an orbital shaker (100-120 rpm) overnight at 36°-38°C. Consider treated bacterin to be diluted 1:2.

3.4.1.2 Oil-adjuvanted bacterins

Mix 1 ml of sodium desoxycholate elution buffer with 1 ml of bacterin. Place on an orbital shaker (100-120 rpm) overnight at 20°-25°C. Consider treated bacterin to be diluted 1:2.

4. Performance of the test

4.1 Dilute the K88 antigen-capture MAb to the current use dilution in cold carbonate coating buffer, and place 100 μ l in each well of 96-well flat-bottom microtitration plates. Seal coated plates with plate sealers. Incubate coated plates overnight at 2°-7°C. Coated plates stored at 2°-7°C may be used for up to 5 days.

4.2 Make twofold dilutions of reference and test bacterins, using PBS-Tween 20 as a diluent. Add 125 μ l PBS-Tween 20 to each well of a clean microtitration plate (transfer plate). Place 125 μ l of bacterin in the first well of each row. Test each bacterin in at least 2 replicate rows. Test the reference bacterin and the test bacterin on the same plate.

Use a multichannel pipetting device to make serial twofold dilutions of each bacterin across the plate (125 μ l transfer volume). Reserve at least 2 unused wells on each plate to serve as blanks.

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The use of at least 7 serial twofold dilutions per bacterin is recommended. Ideally, the selected bacterin dilutions should delineate the sigmoid curve from antigen saturation to antigen extinction for each bacterin. The dilutions used for the reference and the test bacterin may differ.

4.3 Wash the coated ELISA plates 3 times with PBS-Tween 20. An automatic plate washer (250 μ l/well, 40-second soak cycle, 3 wash cycles) may be used, or the plates may be washed by hand. Tap the plates upside down on absorbent material to remove residual fluid.

4.4 Use a multichannel pipetting device to transfer the bacterin dilutions from the transfer plates to the coated ELISA plates (100 μ l/well). Seal the ELISA plates and incubate them on an orbital shaker (100-120 rpm) for 30 min (\pm 5 min) at 20°-25°C.

4.5 Wash the ELISA plates 3 times with PBS-Tween 20 as in **Section 4.3**.

4.6 Dilute the horseradish peroxidase-labeled K88 antigen-indicator MAb in conjugate diluent to the current use dilution, and add to each well (100 μ l/well). Seal the ELISA plates, and incubate on an orbital shaker (100-120 rpm) for 30 min (\pm 5 min) at 20°-25°C.

NOTE: Ovalbumin will stick to plastic. Prepare the working dilution of the conjugate in a glass vial. Transfer the diluted conjugate into pipetting trays immediately prior to the addition of the conjugate to the ELISA plate.

4.7 Wash the ELISA plates 3 times with PBS-Tween 20 as in **Section 4.3**.

4.8 Add freshly prepared substrate solution to each well (100 μ l/well). Incubate the ELISA plates on an orbital shaker (100-120 rpm) for 10 min (\pm 5 min) at 20°-25°C.

4.9 Stop the substrate color development by adding 100 μ l stop solution to each well.

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NOTE: The OPD substrate undergoes a color shift from yellow to orange when stop solution is added.

4.10 Read the ELISA plates using an ELISA reader with dual wavelengths (490 nm test, 650 nm reference). Calculate the mean absorbance for the blank wells. Subtract the mean absorbance of the blank wells from each bacterin test well absorbance value prior to data analysis.

5. Interpretation of the test results

5.1 Relative potency calculation method

5.1.1 Use the current version of the *Relative Potency Calculation Software* (RelPot) to calculate the relative potency of the test bacterin as compared to that of the reference bacterin.

5.1.2 Do not use bacterin dilutions with mean optical density (O.D.) values <0.050 (after subtraction of the mean O.D. of the blank) in the relative potency calculations.

5.1.3 Do not use regression lines with slopes > -0.150 in the relative potency calculations. Enter a minimum slope (Min Slope) assay parameter of 0.150 in the RelPot spreadsheet in place of the 0.000 default.

5.1.4 Enter the reference and test bacterin data, and execute the relative potency program as outlined in the current version of MVSAM0318.

5.1.5 Report the highest relative potency (RP) value included in the top scores from each test as the RP for the test bacterin.

5.2 Requirements for a valid assay

5.2.1 An assay must meet the validity requirements in the current version of MVSAM0318 to be considered valid.

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5.2.2 Lines determined by first-order linear regression of at least 3 contiguous points must have a correlation coefficient (r) of ≥ 0.95 .

5.2.3 The reference regression line and the test bacterin regression line must show parallelism (slope ratio ≥ 0.80).

5.2.4 Assays that are not valid may be repeated up to a maximum of 3 times. If a valid assay cannot be achieved with 3 independent assays, the test bacterin is unsatisfactory.

5.3 Requirements for a satisfactory test bacterin

5.3.1 To be considered satisfactory, a test bacterin must have an RP value of ≥ 1.0 . Test bacterins with RP values < 1.0 on a valid assay may be retested by conducting two independent replicate tests in a manner identical to the initial test. If both retests are valid and the reported RP values of both of the retests are ≥ 1.0 , the test bacterin is satisfactory.

6. Reporting of test results

Report the results of the test(s) as described in the current version of BBSOP0020.

7. Summary of revisions

This is a new SAM. The information contained in this document was previously available as a DRAFT SAM dated June 22, 1995. This document was rewritten to meet the current NVSL/CVB QA requirements, to clarify practices currently in use in the CVB-L, and to provide additional detail. No significant changes were made from the previous protocol.