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Standard Practice for Evaluation of Antimicrobial Agents as Preservatives for Invert Emulsion and Other Water Containing Hydraulic Fluids¹

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INTRODUCTION

Invert emulsion hydraulic fluids typically contain 60 % mineral oil and 40 % water (by volume). These fluids routinely are prepared using proprietary, oil-soluble, emulsifying agents, as well as other emulsifiable constituents. They are recommended for use where conditions indicate a low-cost, fire retardant product, compatible with water-based metal working fluids.

The high water content of these hydraulic fluids makes them susceptible to microbial attack. Uncontrolled microbial growth in these fluids can cause cartridge filter unit plugging, malodorous conditions, or general biodeterioration. Problem microorganisms associated with these fluids include bacteria and fungi.

The hydraulic system is essentially a closed one in which water of evaporation is added to maintain a fixed volume. The inclusion of an efficacious preservative in the water containing hydraulic fluids can prevent microbial growth and the resulting problems that follow.

1. Scope*

1.1 This laboratory practice is designed to evaluate the utility and effectiveness of antimicrobial agents intended to control microbial growth in invert emulsions and other water containing hydraulic fluids.

NOTE 1—Procedures for preparation of water soluble hydraulic fluids and recovery of organisms appear in Practice E2169.

1.2 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

1.3 *This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.*

¹ This practice is under the jurisdiction of ASTM Committee E35 on Pesticides, Antimicrobials, and Alternative Control Agents and is the direct responsibility of Subcommittee E35.15 on Antimicrobial Agents.

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2. Referenced Documents

2.1 ASTM Standards:²

- D1129 Terminology Relating to Water
- D4454 Test Method for Simultaneous Enumeration of Total and Respiring Bacteria in Aquatic Systems by Microscopy
- D5465 Practice for Determining Microbial Colony Counts from Waters Analyzed by Plating Methods
- E1326 Guide for Evaluating Non-culture Microbiological Tests Used for Enumerating Bacteria
- E2169 Practice for Selecting Antimicrobial Pesticides for Use in Water-Miscible Metalworking Fluids
- E2523 Terminology for Metalworking Fluids and Operations

3. Terminology

3.1 Terms used in this practice are defined in Terminologies D1129 and E2523.

² For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

*A Summary of Changes section appears at the end of this standard

4. Summary of Test Method

4.1 The antimicrobial agent to be evaluated is incorporated into an emulsion system by (a) addition to the aqueous phase employed in the preparation of the emulsion, (b) in doses to the formulated system, or (c) by other methods suitable for the test compound.

4.2 A heavy bacterial or fungal inoculum, or both, is then added.

4.3 The resulting mixture is aerated and passed over the surface of a simulated filter system for a minimum period of eight weeks either continuously or with shutdowns to simulate actual operations conditions.

4.4 The degree of microbial control is determined by periodically quantifying the bioburden in the emulsion by direct microscopic count (Test Method **D4454**), plate count (Practice **D5465**), or other appropriate method (Guide **E1326**) and visual observations for microbial fouling of the simulated filter surface.

NOTE 2—A knowledge of standard microbiological techniques is required for this procedure. It is also required that good laboratory practices be followed throughout these tests. This means appropriate containment for the microbiological systems being evaluated. The systems should be maintained in an enclosure so that during the aeration process the mists and aerosols generated do not contaminate the laboratory environment.

5. Significance and Use

5.1 This procedure is designed to determine the effectiveness of antimicrobial agents intended for microbial control in invert emulsions and other water containing hydraulic fluids.

6. Apparatus

6.1 *Air Supply*—Any air source which is free from organic vapors, organic matter, or other objectionable material may be used.

NOTE 3—If desired, air may be sterilized as follows:

Pack two 150-mm long drying tubes (bulb type) loosely with glass wool in a series with neoprene stoppers, glass tubing, and neoprene tubing. Wrap loosely in aluminum foil and steam sterilize at 15 to 20 psi for 30 minutes. Cool to room temperature while still wrapped. In-line pre-sterilization air filters are available from most local laboratory supply houses.

Insert into air line with bulbs on upstream side. Average lifetime in continuous use is two weeks. Discard sooner if upstream filter becomes wet or contaminated with oil.

6.2 *Colony Counter*—Any one of several types may be used.

6.3 *Incubator*—Any cabinet capable of maintaining a temperature of $35 \pm 1^\circ\text{C}$ may be used.

6.4 *Test Cabinet*—A large cabinet capable of maintaining a temperature of $35 \pm 1^\circ\text{C}$, able to house several two litre beakers, and into which an air line can be introduced.

6.5 *Sterilizer*—Any suitable steam sterilizer capable of producing the conditions of sterilization is acceptable.

6.6 *Simulated Filters:*

6.6.1 *Strainer*, 3-in. epoxy coated, ¼-in. mesh gutter strainer.³

6.6.2 *Screen*, 16 by 18 in. fiberglass screening material.

NOTE 4—Fiberglass mesh screening material (16 by 18 in.) is available from any local hardware dealer.

6.6.3 *Wire*, 20-gage, galvanized or stainless steel.

6.7 *Tubing*, ¼-in. ID Tygon.

NOTE 5—Tygon is available from most local laboratory supply houses.

6.8 *T-Connectors*, ¼-in. polypropylene.

6.9 *Laboratory Blender*—Any standard adjustable speed laboratory blender having a 2-L capacity glass or metal container is satisfactory.

6.10 *Hypodermic Needle*, 16-gage needle.

6.11 *Microscope*, Brightfield microscope equipped with 40× and 100× objectives.

6.12 *Labware:*

6.12.1 *Culture Dishes*—100 mm by 15 mm sterile culture dishes made of glass or plastic are required for making standard plate counts.

NOTE 6—Presterilized and disposable plastic petri dishes are available from most local laboratory supply houses.

6.12.2 *Bacteriological Pipettes of 1.1 or 2.2-mL capacity.*

NOTE 7—Presterilized and disposable 1.1-mL bacteriological pipettes are available from most local laboratory supply houses.

6.12.3 *Water Dilution Bottles*—Any sterilizable glass containers having a 150 to 200-mL capacity and tight closures may be used.

NOTE 8—Milk dilution bottles of 160-mL capacity having screw-cap closures are available from most local laboratory supply houses.

6.12.4 *Two-Liter Borosilicate Glass Beakers.*

6.12.5 *Bent Glass Rod.*

6.12.6 *Screw Cap Culture Tubes*, autoclavable, 15 by 150 mm.

6.13 *Water Bath*—Maintain at $46^\circ\text{C} \pm 2^\circ\text{C}$ to anneal agar based microbiological media.

6.14 *Aluminum Foil.*

7. Reagents and Materials

7.1 *Invert Emulsion Emulsifier.*⁴

7.2 *Paraffinic Mineral Oil.*

7.3 *Deionized or Distilled Water (>2 MOHM quality)*

³ The sole source of supply of the apparatus known to the committee at this time is Billy Penn Corp., Philadelphia, PA 19122. If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

⁴ The sole source of supply of a satisfactory emulsifier for the preparation of invert emulsion hydraulic fluids (Compound #5162) known to the committee at this time is the Lubrizol Co., Wickliffe, OH. If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

7.4 *Gentamicin Sulfate*.⁵

7.5 *Arlacel 80*.⁶

7.6 *Tween 60*.⁶

7.7 *Phosphate Buffer—For serial dilutions*.

7.8 *Mineral oil, sterile*.

7.9 *Microbiological Media*—General retrieval media consistent with good microbiological practices are acceptable. Examples are as follows:

7.9.1 *Soybean-Casein Digest Agar*, U.S.P. XIX, Medium II.

NOTE 9—Soybean-casein digest agar is available in dehydrated form from most laboratory media supply houses.

7.9.2 *Fluid Soybean-Casein Digest Medium*, U.S.P. XIX, Medium III.⁷

7.9.3 *Sabouraud Dextrose Agar*, U.S.P. XIX, Medium 20.⁷

7.9.4 *Sabouraud Dextrose Broth*, U.S.P. XIX, Medium 21.⁷

7.9.5 *Sulfate American Petroleum Institute (API) Agar*,⁶ for enumeration of sulfate reducing bacteria.

7.10 *Inoculum*:

7.10.1 The inoculum may vary according to the users' requirements. It may be either undefined or defined.

7.10.1.1 An undefined inoculum may consist of microorganisms isolated from a "spoiled" invert emulsion hydraulic fluid which exhibits microbiologically induced phase generation, or which is known to have caused plugging of a hydraulic system filter due to microbial slime, and grown in a nutrient medium.

7.10.1.2 An undefined inoculum may consist of the following: (1) equal volumes of fluid soybean-casein digest and "spoiled" (see 7.10.1.1) hydraulic fluid aerated at 35°C for 24 h (typically) until the bacterial count reaches 10⁹ CFU/mL, (2) equal volumes of sabouraud dextrose broth and "spoiled" (see 7.10.1.1) hydraulic fluid aerated at 35°C for 24 h (typically) or until fungal count reaches 10⁶ to 10⁷ CFU/mL, or (3) equal volumes of (1) and (2) if both bacteria and fungi are the desired test organisms.

7.10.2 A defined inoculum consisting of a mixed culture of specific microorganisms may also be used.

7.10.2.1 The defined inoculum may be prepared by isolating and identifying specific microorganisms from a "spoiled" (see 7.10.1.1) hydraulic fluid emulsion and culturing the bacterial

isolates in soybean-casein digest medium and the fungal isolates in sabouraud dextrose broth until there are 10⁹ CFU bacteria or 10⁶ to 10⁷ CFU fungi, or both, per mL, respectively.

7.10.2.2 Other microorganisms of particular interest (Rossmore and Szlatky)⁸ may be used such as: *Pseudomonas fluorescens*, *Pseudomonas cepacia*, *Klebsiella pneumoniae*, *Proteus mirabilis*, *Desulfovibrio desulfuricans*, *Aspergillus niger*, *Cephalosporium* sp., *Fusarium* sp., *Candida* sp.

7.10.2.3 Equal mixtures of any two of the above bacterial species or two of the above mold species, or both, plus the *Candida* species to provide a final titer of 10⁹ CFU bacteria, or 10⁶ to 10⁷ CFU fungi, or both, per mL, should be used as an inoculum for the emulsion system.

7.11 *Antimicrobial Agents*—The chemical agents to be evaluated as preservatives.

8. Preparation of Simulated Filters

8.1 Cut the epoxy-coated, ¼-in. mesh gutter strainers 16 by 18 in. mesh fiberglass screening material into 3 by 5 in. sections. Secure the screening to the strainers with 20-gage wire or with staples.

8.2 *Preparation of Aerators*—Cut tubing (see 6.7) into 13-in. sections. Bend tubing in a circle and connect both ends using a T connector (see 6.8). Connect third arm of T connector to a 20-in. length of tygon tubing. This tubing will be connected to the main air supply line. Using a hot 16-gage needle, carefully punch a series of holes, ½ in. apart, along the outer circumference of the tubing which forms the ring. Also punch similar holes ½ in. apart on the upper and lower surface of the tubing, at right angles to the holes previously punched. These holes allow the air from the air source to bubble up through the hydraulic fluid producing a cascading effect over the surface of the simulated filter.

9. Preparation of Microbiological Medium

9.1 Microbiological media should be prepared in accordance with manufacturer's instructions. Media to be augmented with antibiotics should be annealed in a 46 ± 2°C water bath before antibiotics are added. Antibiotics should be added just before pouring. Use 100 g gentamicin sulfate per mL to suppress bacterial growth on fungal recovery media.

10. Microbiological Methods

10.1 Solubilize the invert emulsion aliquot (see 7.1) according to the procedure of McConville, et al.,^{9,10} as follows:

10.1.1 Disperse 1 mL of the invert emulsion in 1 mL of Arlacel 80 and bring the volume up to 10 mL with 10 % Tween 60 solution.

⁵ The sole source of supply of gentamicin sulfate known to the committee at this time is as Garamycin Reagent Solution, available in two concentrations of 10 and 50 mg/mL, from the Schering Corp., Kenilworth, NJ 07033. If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

⁶ The sole source of supply of the reagents (Arlacel 80, Tween 60, and Sulfate API Agar) known to the committee at this time is Sigma Aldrich Co., St. Louis, MO 63178, <http://www.sigmaldrich.com>. If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

⁷ The sole source of supply of the media, available in dehydrated form, known to the committee at this time is Baltimore Biological Laboratories, Cockeysville, MD or Difco Laboratories, Detroit, MI. If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

⁸ Rossmore, H. W., and Szlatky, K., "Characterization of the Microbial Flora of Invert Emulsion Hydraulic Fluids," *Int. Biodetm. Bulletin*, Vol 13, No. 4), 1977, pp. 96–100.

⁹ McConville, J. F., et al., "Method for Performing Aerobic Plate Counts of Anhydrous Cosmetics Utilizing Tween 60 and Arlacel 80 as Dispersing Agents," *Applied Microbiology*, Vol 27, 1974, pp. 5–7.

¹⁰ Hoffman, N. M., "Hydraulic Fluid of 95-Percent Water," *Lubrication Engineering*, Vol 35, No. 2, 1979, pp. 65–71.