

Poster Communication 41

## Microbial Contamination in Saturated Saline Solution Used for Cadaver Preservation in Veterinary Anatomy Teaching

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### Abstract

**Background:** The use of animal cadavers is essential for veterinary anatomy education, requiring preservation methods that maintain tissue characteristics similar to those of living organisms. At CESPU, cadavers are preserved using a saturated saline (SS) solution, considered effective in maintaining tissue texture, color, and joint mobility [1,2]. However, its ability to control microbial growth, particularly for halotolerant strains, remains poorly studied [3]. Common bacterial species associated with animal cadavers include *Staphylococcus aureus*, *Bacillus* spp., *Enterococcus* spp., and *Escherichia coli* [4]. **Objectives:** This study aimed to detect fecal contamination indicators, namely *Enterococcus* spp. and *E. coli*, in the SS solution used for preserving animal cadavers in veterinary anatomy classes at CESPU. **Methods:** A total of 42 SS samples were collected from 7 cadavers (3 cats and 4 dogs), including samples obtained before immersion and after 7, 14, and 21 days of preservation. The SS solution was renewed weekly, with each cadaver immersed for 7 days before replacement. Samples were inoculated onto selective media: Slanetz-Bartley agar and Kanamycin Esculin Azide agar for *Enterococcus* spp., and MacConkey agar and Chromogenic Coliform agar for *E. coli*. Up to two typical colonies per sample were subcultured on brain heart infusion agar and identified using MALDI-TOF mass spectrometry. **Results:** Among the 21 SS samples collected before cadaver immersion, 1 showed bacterial growth, yielding two isolates: *Enterococcus hirae* and *E. coli*. Of the 21 samples collected after immersion, 15 showed microbial growth. Identified isolates included *Enterococcus faecalis* (n=2), *Enterococcus faecium* (n=3), *Enterococcus hirae* (n=1), *Enterococcus raffinosus* (n=1), and *Enterococcus* spp. (n=7). Enterococci were detected in samples collected at 7 (n=5), 14 (n=4), and 21 days (n=5). Additionally, two *E. coli* isolates were recovered from a single sample collected at 14 days. **Conclusion:** These findings indicate that the SS solution does not fully inhibit microbial growth during cadaver preservation. The detection of *Enterococcus* spp. and *E. coli*, particularly after immersion, suggests possible fecal contamination and highlights the need for regular

microbiological monitoring and enhanced biosafety measures to reduce occupational exposure and ensure a safer learning environment.

**Keywords:** veterinary microbiology; biosafety; cadaver preservation

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