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AIR, Test Code 1074-Total Coliforms/E. Coli

Air cultures are taken with an impactor style sampler (Andersen or SAS). This is a two plate protocol to select out gram-negative rods, such as *E. coli*, coliforms or non-fermenters (*Pseudomonas species*) in addition to a Total Bacterial Count with Identifications. This profile is particularly meaningful in assessing an indoor environment after a black water event in that it selects out for fecal gram-negative rods.

- 1. Calibrate each sampling pump or piece of equipment by following manufacturer's recommendations.
- 2. Before each run, thoroughly wipe each sampler stage with rubbing alcohol. Allow to dry. Make sure air passages are not blocked.
- 3. Load and immediately unload one set of sampling media in each sampler to serve as field blanks.
- 4. Label agar side of plate with identifier. Remove cover from media, load sampling media into sampler, and attach sampler to pump with flexible tubing or if using a SAS sampler screw the top back onto the sampler..NOTE: Take special care to prevent contamination of media during loading and unloading. Do not touch agar surface.
- 5. Sample at known preset flow for an accurately known time, e.g., 5 min. Rotary vane pump should run at 28. 3 lpm. (In heavily contaminated areas, sampling time may be adjusted) Both plates should be sampled at same total air volume.
- 6. Replace covers on sampling media. Tape plate or place each plate in separate bag, and pack securely for shipment (plates should be media side up).
- 7. If plates are going to be shipped back to the laboratory send them for overnight delivery in a cooler with an ice pack. If plates are not shipped that day keep the plates in the refrigerator until they are shipped the next day.
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REFERENCES:

Dillon, H. Kenneth, L. Hung, J. Miller, Field Guide for the Determination of Biological Contaminants in Environmental Samples.,

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