

ADVANCED ACTIVE SPONGE

FOR URINE COLLECTION, TRANSPORT AND PRESERVATION



COPAN UriSponge™ is intended for collecting, transporting, and preserving urine specimens from the sampling site to the testing laboratory. UriSponge™- collected samples are processed using the clinical laboratory's standard operating procedures (SOPs) for the cultivation of uropathogenic bacteria and yeasts.

- **> NEEDLE-FREE:** Compared to competing products, UriSponge[™] simply collects urine using the high absorption capacity of an advanced sponge (applicator).
- **> PRACTICAL:** The sponge absorbs forthwith the correct amount of first-void urine sample. No binding sample volume and no need to dispose of biohazard material.

PLUS FEATURES

- HIGH-YIELD AGENTS: Ready-to-use preservatives are chemically incorporated inside the sponges. Zero-risk of overgrowth or overkill during transport.
- VERSATILE SYSTEM: UriSponge™ is fully compatible with Copan WASP®, a modular, open platform that fully automates specimen processing for Microbiology.
- > RELIABLE DESIGN: The internal conical shape enables centrifugation of urines, while the external cylindrical shape stands upright on the laboratory bench. Urisponge™ includes a leak-proof system.
- > OPTIMAL VIABILITY: UriSponge™ provides microorganisms viability at refrigerated and room temperature conditions up to 48 hours.





SAMPLE WORKFLOW



Please refer to complete instructions. Above pictograms are for marketing purpose only.



| Recovery of Urinary Tract Pathogenic Organisms in UriSponge™ per CLSI M40-A2 | |
|--|--------------------------------|
| C. albicans (ATCC® 24433) | <i>E. coli</i> (ATCC® 25922) |
| E. faecalis (ATCC® 29212) | P. aeruginosa (ATCC® 27853) |
| P. mirabilis (ATCC® 7002) | S. saprophyticus (ATCC® 15305) |

| Recovery of Additional Relevant Urinary Tract Pathogenic Organisms in UriSponge™ | |
|--|----------------------------------|
| C. freundii (ATCC® 8090) | C. glabrata (ATCC® MYA-2950) |
| E. cloacae (ATCC® 13047) | <i>M. morganii</i> (ATCC® 25829) |

^{*}The use of this device in association with other assays and/or instrumentation should be validated prior to use.

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