

# Cross-Infection risks in ultrasound examinations



Germitec

**REDUCING ULTRASOUND PROBE CROSS-INFECTION RISK CAN AID IN REDUCING HEALTHCARE-ASSOCIATED INFECTIONS**



**Among Healthcare-Associated Infections (HAIs), cross-infections due to the use of ultrasound probes in various medical procedures have become a major concern with converging evidence in research literature.**

## High Risk of infection following Low Level Disinfection

Various global guidelines recommend High Level disinfection for endocavity ultrasound probes that are classified as semi-critical/semi-invasive. However, some institutions and medical teams proceed to Low Level Disinfection (LLD) and may not be fully aware of the cross-infection risks after use of an ultrasound probe. However, LLD has been proven to be insufficient, as demonstrated by the results of a retrospective cohort study using linked national datasets from Scotland:<sup>6</sup>

41%

**Patients are 41% more likely to have positive bacterial cultures 30 days after a transvaginal scan where probes were low level disinfected and 26% (HR=1.26) more likely to be prescribed antibiotics.**

26%

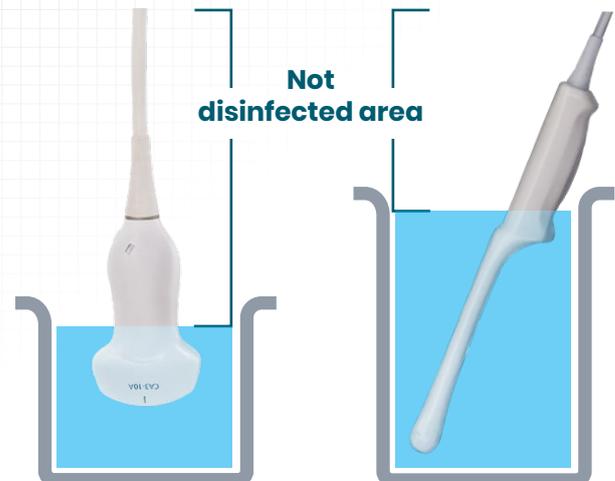
3.4x

For transrectal scans, patients were **3.4 (HR=3.4) times more likely to have positive bacterial cultures** and **75% (HR=1.75) more likely to be prescribed antibiotics** ( $p < 0.001$ ).

75%

### Probe Handles Infection Risk

Ultrasound probe handle is often forgotten in the disinfection process. In a prospective randomized controlled clinical study of two groups, one with an automated process and the other one with a manual process, it has been shown that more than 80% of probe handles that were not disinfected, had residual pathogens, including MRSA.<sup>5</sup> This stresses the fact that the probe handle should also be included in the disinfection.<sup>5</sup>



### Probe covers NOT 100% protection proof

Guidelines recommend the use of dedicated probe covers during endocavity examinations to reduce cross contamination. Research studies show that up to 13% of condoms and 5% of commercial covers have tears pre and post-transvaginal ultrasound examinations.<sup>1-4</sup> This highlights the reason why you need to HLD ultrasound probes to reduce cross contamination risk due to the high probability probe covers can have micro-tears.



**References:** **1.** Basseal,j. et al. (2019).Analysis of the integrity of ultrasound probe covers used for transvaginal examinations. Australasian College for Infection Prevention and Control. Dec. **2.** Vickery K, et al. Evaluation of an automated high-level disinfection technology for ultrasound transducers, Journal of Infection and Public Health, 2013. **3.** Masood J et al. Condom perforation during transrectal ultrasound guided (TRUS) prostate biopsies: a potential infection risk. Int Urol Nephrol 2007;39(4):1121-4. **4.** Leroy S. Infectious risk of endovaginal and transrectal ultrasonography: systematic review and meta-analysis. The Journal of hospital infection. 2013;83(2):99-106. NAN0046. **5.** Ngu,A, et.al Reducing Transmission Risk Through High-Level Disinfection of Transvaginal Ultrasound Transducer Handles. Infection Control & Hospital Epidemiology, 36, pp 581-584 doi:10.1017/ice.2015.12 **6.** Health Protection Scotland (HPS), NHS National Services Scotland (2017). NHS Scotland Risk Based Recommendations for the Decontamination of Semilnvasive Ultrasound Probes: Risk of infection following semi-invasive ultrasound procedures in Scotland, 2010 to 2016. Version 1.0.