

# The body of evidence puts trophon® devices on top



BUESCHER, D. L., ET AL. (2016)

NGU, A., ET AL. (2015)

MINISTÈRE DES SOLIDARITÉS ET DE LA SANTÉ (2019)

SCOTT, D., ET AL. (2018)

JOHNSON S ET AL. (2013)

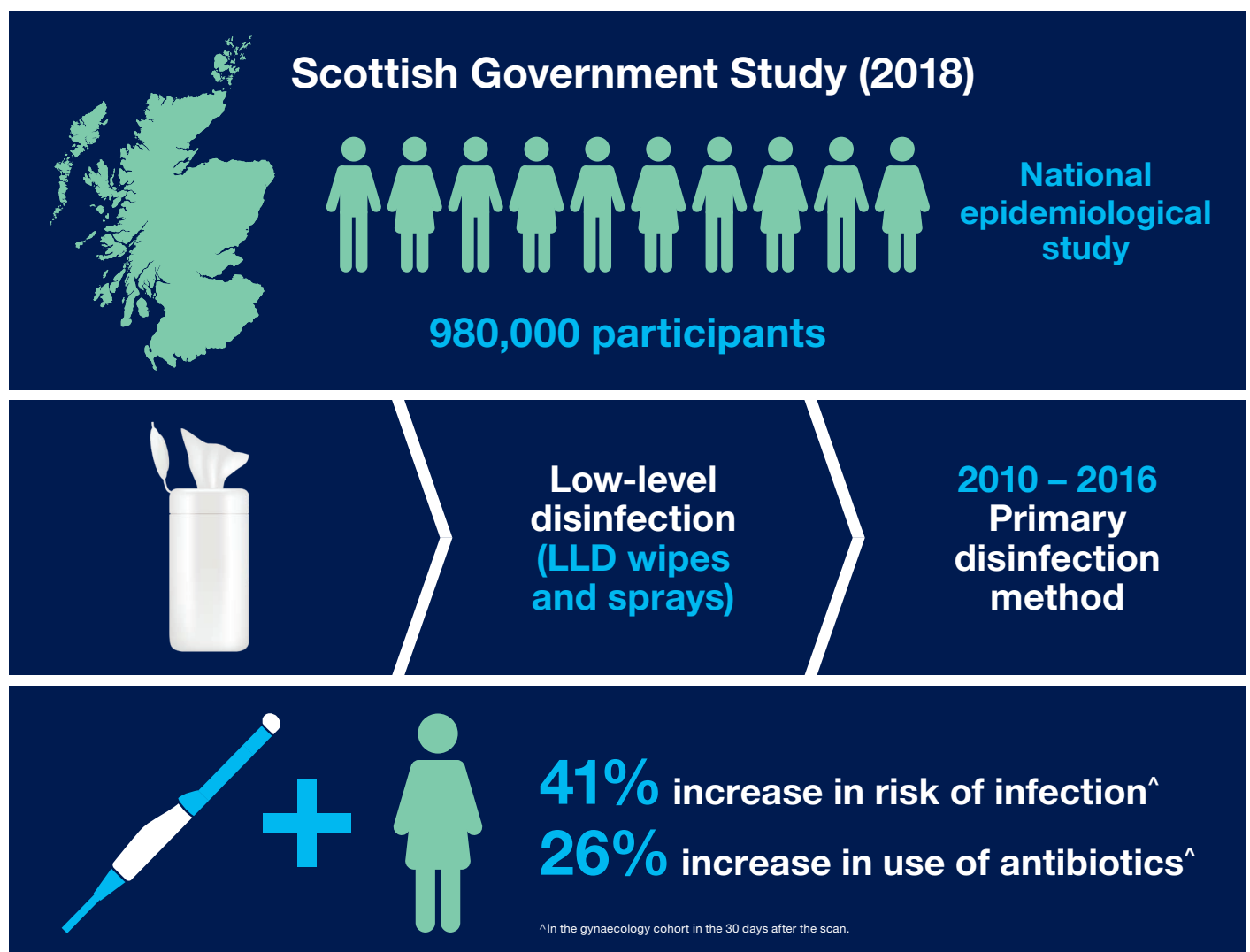
ALFA MJ. (2015)

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Infection Prevention. For Life.

# Inadequate transvaginal ultrasound probe disinfection leads to increased infection risk

- Studies have found >90% of ultrasound probes are contaminated after patient use, and can harbour serious pathogens like MRSA<sup>1-4</sup>
- Other studies have found low-level disinfection (LLD) leaves probes contaminated with bacteria and viruses, including chlamydia and HPV<sup>1-3,5</sup>
- A population level study by the Scottish government found gynaecological patients undergoing transvaginal scans were at a 41% increased risk of infection, where LLD was the standard of care<sup>6</sup>



**CONCLUSION:** “Hence, failure to comply with existing guidance on [high-level disinfection] of semi-invasive ultrasound probes will continue to result in an unacceptable risk of harm to patients.”<sup>6</sup>

# trophon technology outperforms other disinfection solutions

University Hospital Muenster, Germany. 2016.

## Disinfection of transvaginal probes in a clinical setting: comparative performance of automated and manual reprocessing methods<sup>2</sup>

**Comparison of disinfection methods of transvaginal ultrasound probes**

**Automated HLD\*  
(n=120)**

7 minute complete cycle  
Probe head and handle

**VS**

**LLD Wipes  
(n=120)**

1 minute contact time  
Probe head only (reflecting standard practice)

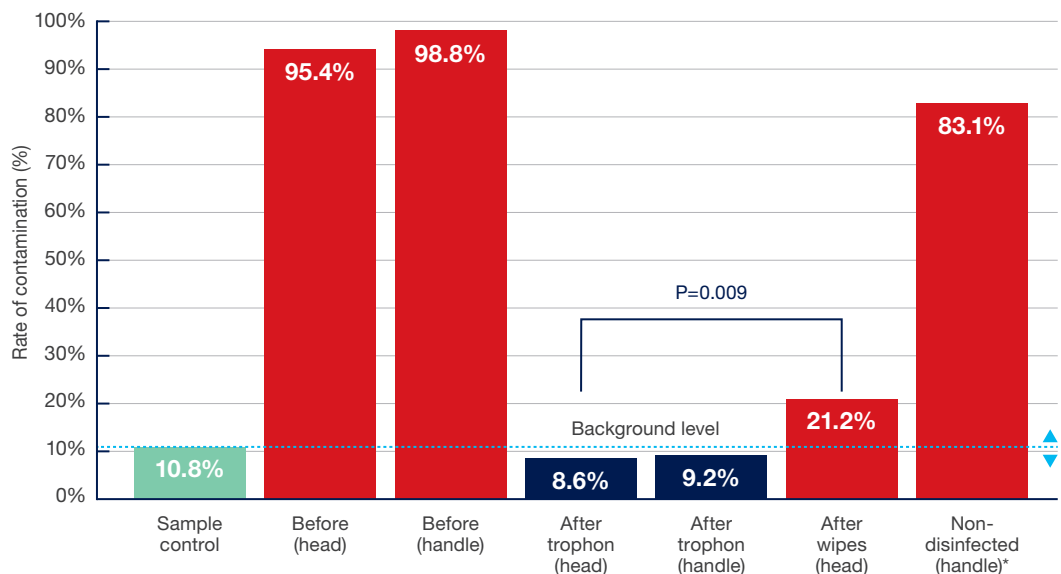
\*High level disinfection

## trophon device achieved disinfection below background levels on both heads and handles<sup>2</sup>

- Manual disinfection with LLD wipes failed to systematically eliminate bacteria to below background level<sup>2</sup>
- **Only trophon devices successfully eliminated bacteria; a group of organisms that should be inactivated by HLD and LLD.**

**Risk of contamination was almost 3X higher when compared with wipes vs trophon (95% CI 1.3-6.3)<sup>2</sup>**

Adapted from Buescher DL et al. 2016.  
\*Handles not disinfected according to standard facility practice.  
CI: Confidence Interval.

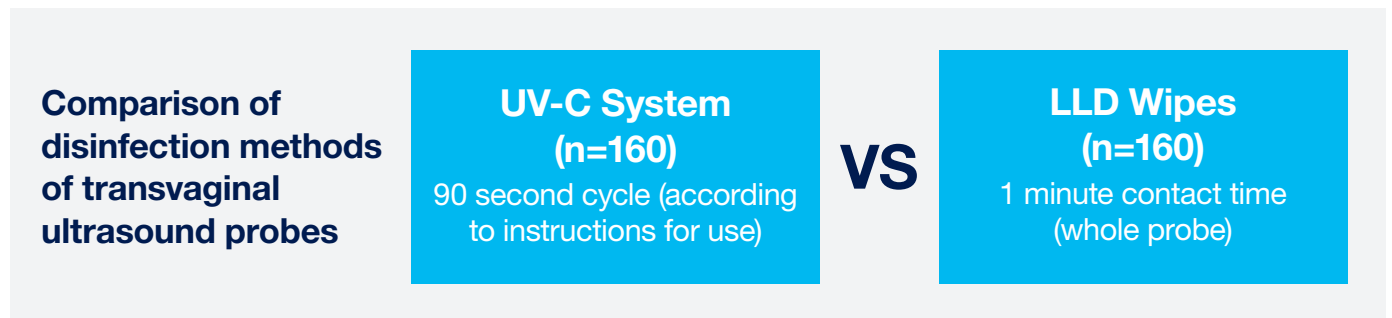


Considering the data from both studies, only trophon devices

# Automated UV-C light system demonstrates no significant difference in efficacy vs LLD wipes

University Hospital Muenster, Germany. 2019.

## Disinfection of transvaginal ultrasound probes by UV-C: Clinical evaluation of automated and manual reprocessing methods<sup>3\*</sup>

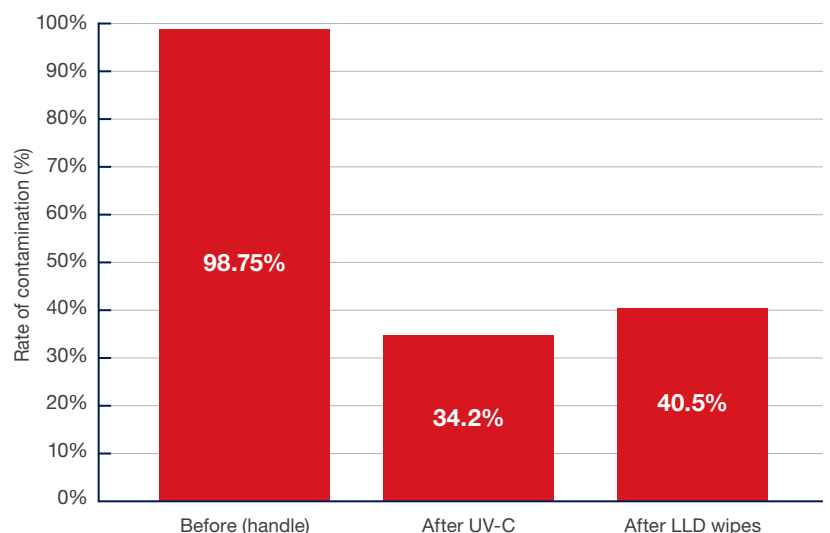


## 34.2% of probes were still contaminated following disinfection with UV-C and 40.5% with LLD wipes<sup>3</sup>

- Neither were able to systematically remove bacteria from probes and there was no significant difference in terms of effectiveness between UV-C and LLD wipes<sup>3</sup>
- **The UV-C device did not perform HLD effectively and there was no significant difference in efficacy vs LLD wipes.**

**This was the first published clinical study to use the UV-C device (AntiGermix AS1) strictly according to its instructions for use and without other disinfectants<sup>3</sup>**

Adapted from Schmitz J et al. (2019).  
US: Ultrasound.  
UV-C: Ultraviolet-C.



**systematically demonstrated superior efficacy vs LLD wipes<sup>2,3</sup>**

# Superior microbial efficacy with trophon technology

## trophon technology delivers a unique and fully automated device for high-level disinfection of both surface and endocavitary probes

- FDA-cleared and CE-marked as a high-level disinfectant device
- Demonstrated bactericidal, mycobacterial, fungicidal, virucidal and sporicidal disinfection efficacy in accordance with EN Standards, AOAC International Official methods and ASTM International Standards
- Nebulized H<sub>2</sub>O<sub>2</sub> mist particles reach and disinfect all challenging probe surfaces, including uneven grooves and crevices that can harbor pathogens
- Demonstrated to eliminate an extended range of clinically-relevant pathogens, including those that cause STIs, such as chlamydia, gonorrhoea, herpes, HIV, hepatitis A, B and C as well as HPV, *Clostridium difficile* spores and drug-resistant bacteria (MRSA and VRE), in addition to mandatory testing.



## trophon device - the global standard of care in ultrasound probe reprocessing

### Enhanced clinical workflows

- Optimized and configurable settings to deliver HLD at any point-of-care
- Minimal reprocessing hands-on time required

### Integrated digital traceability solutions

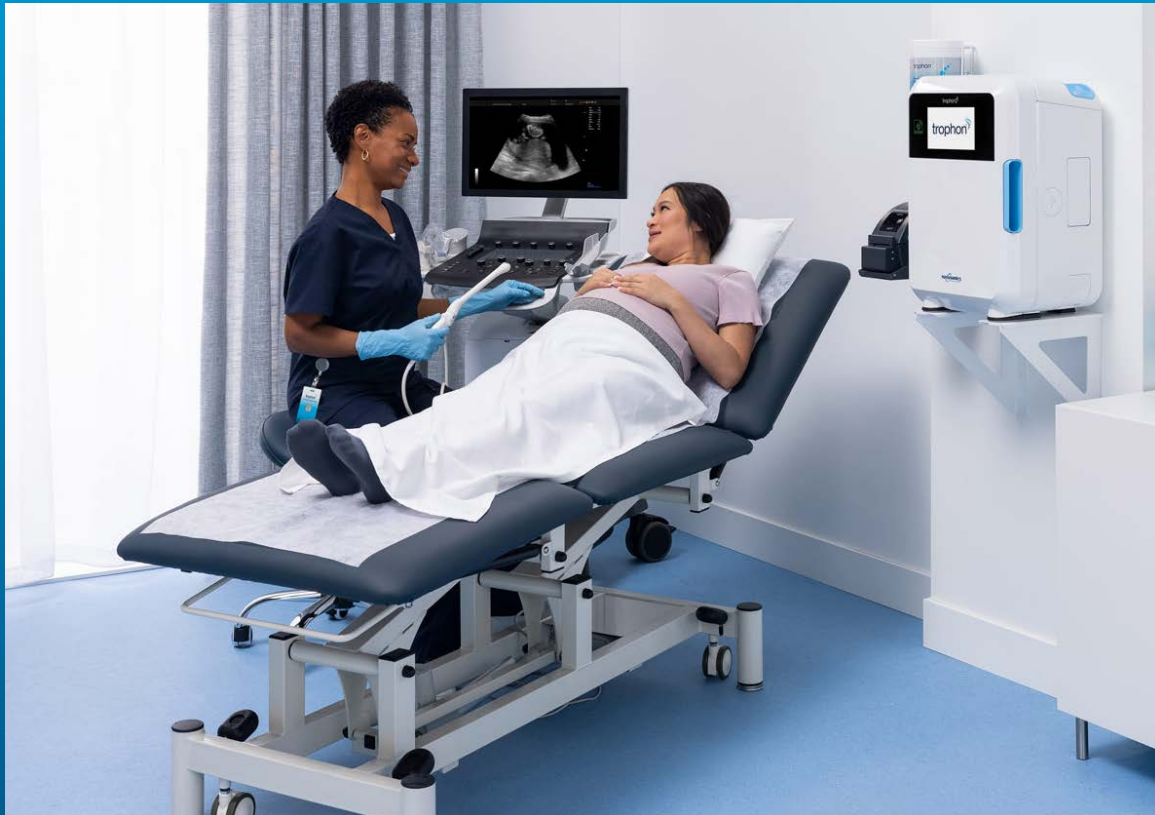
- Simplified workflows support audit-ready compliance
- Standardization to support facility-wide risk management

### Ultrasound manufacturers' reprocessing solution of choice

- Industry-leading compatibility program delivers rigorous testing
- Over 1,300 probes from 28 original equipment manufacturers approved and endorsed for trophon devices



# Nanosonics are the experts in ultrasound probe reprocessing



Over 34,000 trophon devices operating across  
thousands of hospitals in 30+ countries protect  
27 million patients each year.<sup>7</sup>

Contact a Nanosonics representative to discuss how trophon devices may be  
applicable to the different scenarios and workflows at your facility

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**References:** 1. M'Zali F, et al. PLoS One. 2014;9(4):e93368. 2. Buescher DL, et al. Ultrasound Obstet Gynecol. 2016 47(5): 646-651. 3. Schmitz J, et al. Ultraschall in Med. 2020;41(6):681-687. 4. Oide S, et al. J Med Ultrason (2001) 2019;46(4):475-479. 5. Leroy S, J Hosp Infect. 2013;83(2):99-106. 6. Scott D, et al. Ultrasound 2018;26(3):168-177. 7. Nanosonics Annual Report. December 2024

Always read the User manual before use and follow the instructions carefully to ensure proper usage of the medical device. The trophon® family includes a range of trophon devices which share the same core technology of 'sonically-activated hydrogen peroxide.' Nanosonics, trophon and NanoNebulant are trademarks of Nanosonics Limited. © 2025 Nanosonics Limited. All rights reserved. EM\_250203\_04\_COL0006. April 2025