

Efficacy Information v4

UWCL360C-M – Uniwipe, Midi Clinical Disinfection & Detergent



Wipes (360 Count)

Information

This product contains a water-based, alcohol-free disinfectant intended for cleaning and disinfecting non-invasive medical devices within healthcare environments – ensure surface compatibility prior to use.

The efficacy has been independently tested and this product meets the requirements of the BSI EN standards listed at the specified contact time.

See end of document for full names of the standards and minimum efficacy levels required.

Efficacy Test Results – Bacteria / Yeast

EN Test Method	Contact Time	Target Organisms	Conditions
16615	60 Seconds	Pseudomonas aeruginosa, Staphylococcus aureus Enterococcus hirae, Candida albicans	Dirty
1276	60 Seconds	Pseudomonas aeruginosa, Staphylococcus aureus Enterococcus hirae, Escherichia coli	Dirty
13727	30 Seconds	Pseudomonas aeruginosa, Staphylococcus aureus Enterococcus hirae, Burkholderia cepacia	Dirty
13697	60 Seconds	Pseudomonas aeruginosa, Staphylococcus aureus Enterococcus hirae, Escherichia coli	Dirty
13624	30 Seconds	Candida albicans*	Dirty

*

This test was performed as a step 1 test to prove basic efficacy in suspension before proving efficacy using the relevant surface test, the mixture used had not been extracted from a wipe.

Efficacy Test Results – Bacteria/ Yeast

14561	Pseudomonas aeruginosa, Staphylococcus aureus Enterococcus hirae	Dirty
14562	Candida albicans	Clean

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Efficacy Test Results – Viruses

EN Test Method	Contact Time	Target Virus	Conditions
14476	60 Seconds	Vaccinia virus**	Dirty
16777	60 Seconds	Vaccinia virus**	Clean
14476	30 Seconds	Vaccinia virus**	Clean
14476	60 seconds	Norovirus	Dirty

**This product therefore is effective against all enveloped viruses as defined in EN 14476:2013 + A2:2019 Annex A including:

Coronavirus
Poxviridae
Herpesviridae
Flavivirus
Paramyxoviridae
Filoviridae

Hepatitis C Virus (HCV)
Influenza Virus
Measles Virus
Hepatitis Delta Virus (HDV)
Hepatitis B virus (HBV)

Human Immunodeficiency Virus (HIV)
Human T Cell Leukemia Virus (HTLV)
Rabies Virus
Rubella Virus

EN Standards – Full Name & Detailed Information	
Full Name	Details
BS EN 14476:2013 + A2:2019	<p>Quantitative suspension test for the evaluation of virucidal activity in the medical area.</p> <p>Test method and requirements (Phase 2/Step 1)</p> <p>Result Required – 4 Log Reduction (99.99%)</p>
BS EN 13727:2012+A2:2015	<p>Quantitative suspension test for the evaluation of bactericidal activity in the medical area.</p> <p>Test method and requirements (phase 2, step 1)</p> <p>Result Required – 5 Log Reduction (99.999%)</p>
BS EN 1276: 2019	<p>Quantitative suspension test for the evaluation of bactericidal activity of chemical disinfectants and antiseptics used in food, industrial, domestic and institutional areas, Test method and requirements, (phase 2, step 1)</p> <p>Result Required – 5 Log Reduction (99.999%)</p>
BS EN 16615:2015	<p>Quantitative test method for the evaluation of bactericidal and yeasticidal activity on non-porous surfaces with mechanical action employing wipes in the medical area (4- field test). Test method and requirements (phase 2, step 2)</p> <p>Result Required for Bacteria – 5 Log Reduction (99.999%)</p> <p>Result Required for Yeast & Fungi – 4 Log Reduction (99.99%)</p>
BS EN 13697:2015+A1:2019	<p>Quantitative non-porous surface test for the evaluation of bactericidal and/or fungicidal activity of chemical disinfectants used in food, industrial, domestic and institutional areas. Test method and requirements without mechanical action (phase 2, step 2)</p> <p>Result Required for Bacteria – 4 Log Reduction (99.99%)</p> <p>Result Required for Yeast & Fungi – 3 Log Reduction (99.9%)</p>
EN 13624:2013	<p>Quantitative suspension test for the evaluation of fungicidal and yeasticidal activities of disinfectants intended for use in the medical area. (phase 2 step 1)</p> <p>Result Required for Yeast & Fungi – 4 Log Reduction (99.99%)</p>

EN Standards – Full Name & Detailed Information

Full Name	Details
BS EN 16777:2018	Quantitative non-porous surface test without mechanical action for the evaluation of virucidal activity of chemical disinfectants used in the medical area. Test method and requirements (phase 2/step 2). Result Required – 4 Log Reduction (99.99%).
BS EN 14561:2006	Quantitative carrier test for the evaluation of bactericidal activity for instruments used in the medical area. Test method and requirements (phase 2, step 2). Result Required – 5 Log Reduction (99.999%).
BS EN 14562:2006	Quantitative carrier test for the evaluation of fungicidal or yeasticidal activity for instruments used in the medical area. Test method and requirements (phase 2, step 2). Result Required - 4 Log Reduction (99.99%)

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