Efficacy Information v12

SKU.5833 – Uniwipe Clinical Clothwipes (100 Count)

Information

This product contains a water-based alcohol-free disinfectant intended for disinfecting nonporous hard surfaces within healthcare and other 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 / Fungi			
EN Test Method	**Contact Time**	**Target Organisms**	**Conditions**
1276	60 Seconds	Pseudomonas aeruginosa, Staphylococcus aureus Enterococcus hirae, Escherichia coli	Clean
13727	30 Seconds	Pseudomonas aeruginosa, Staphylococcus aureus Enterococcus hirae	Dirty

Efficacy Test Results – Bacteria			
EN Test Method	**Contact Time**	**Target Organisms**	**Conditions**
14561	60 Minutes	Pseudomonas aeruginosa, Staphylococcus aureus Enterococcus hirae	Dirty

Efficacy Test Results – Viruses			
EN Test Method	**Contact Time**	**Target Virus**	**Conditions**
14476	30 Seconds	Vaccinia virus*	Dirty
14476	5 Minutes	Norovirus**	Clean
14476	5 Minutes	Influenza 1 (H1N1)**	Clean

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

(Coronavirus / SARS-CoV-2 is an enveloped virus)

EN 14476 2013 + A2 2019 Annex A (informative – Enveloped viruses)
Reference: Van Regenmortel MHV et al.,Eds.: Virus Taxonomy, Classification and Nomenclature of Viruses, seventh report of the international committee on taxonomy of viruses. Academic Press, San Diego, 2000

Poxviridae	Hepatitis C Virus (HCV)	Human Immunodeficiency Virus (HIV)
Herpesviridae	Influenza Virus	Human T Cell Leukemia Virus (HTLV)
Flavivirus	Measles Virus	Hepatitis B virus (HBV)
Paramyxoviridae	Hepatitis Delta Virus (HDV)	Filoviridae (e.g. Ebola, Marburg)

**	These tests have been carried out using the mixture prior to impregnation on the wipe substrate

EN Standards – Full Name & Detailed Information	
Full Name	**Details**
BS EN 14476:2013 + A2:2019	Quantitative suspension test for the evaluation of virucidal activity in the medical area. Test method and requirements (Phase 2/Step 1) Result Required – 4 Log Reduction (99.99%)
BS EN 13727:2012+A2:2015	Quantitative suspension test for the evaluation of bactericidal activity in the medical area. Test method and requirements (phase 2, step 1) Result Required – 5 Log Reduction (99.999%)
BS EN 1276: 2019	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) Result Required – 5 Log Reduction (99.999%)
BS EN 16615:2015	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) Result Required for Bacteria – 5 Log Reduction (99.999%) Result Required for Yeast & Fungi – 4 Log Reduction (99.99%)
BS EN 13697:2015+A1:2019	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) Result Required for Bacteria – 4 Log Reduction (99.99%) Result Required for Yeast & Fungi – 3 Log Reduction (99.9%)
EN 13624:2013	Quantitative suspension test for the evaluation of fungicidal and yeasticidal activities of disinfectants intended for use in the medical area. (phase 2 step 1) Result Required for Yeast & Fungi – 4 Log Reduction (99.99%)

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%)

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