

Information

This product contains a water-based disinfectant intended for disinfecting and cleaning general hard surfaces – 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⁽ⁱ⁾	Pseudomonas aeruginosa Staphylococcus aureus Enterococcus hirae, Candida albicans	Dirty
13727	30 Seconds	Enterococcus Hira Pseudomonas Aeruginosa Staphylococcus Aureus Enterococcus faecium Vancomycin-resistant (VRE) Staphylococcus aureus Methicillin-resistant (MRSA) Burkholderia cepacia	Dirty
13624	30 Seconds	Candida albicans	Dirty
1276	60 Seconds⁽ⁱ⁾	Pseudomonas aeruginosa Staphylococcus aureus Enterococcus hirae Escherichia coli	Dirty

(i) Minimum contact time designated by test standard.

Efficacy Test Results – Viruses

EN Test Method	Contact Time	Target Virus	Conditions
14476	30 Seconds	Modified Vaccinia virus, strain Ankara (MVA) ⁽ⁱⁱ⁾	Dirty
14476	60 Seconds	Murine Norovirus Adenovirus	Dirty

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

Coronavirus	Hepatitis C Virus (HCV)	Human Immunodeficiency Virus (HIV)
Poxviridae	Influenza Virus	Human T Cell Leukemia Virus (HTLV)
Herpesviridae	Measles Virus	Rabies Virus
Flavivirus	Hepatitis Delta Virus (HDV)	Rubella Virus
Paramyxoviridae	Hepatitis B virus (HBV)	
Filoviridae		

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 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%)
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%)
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%)

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