




# DENTO-VIRACTIS INSTRUGERM 52

Dilution:  
0,5%

<b>Description</b>	<p>Dento-Viractis Instrugerm 52 is recommended for the <b>pre-disinfection of medical devices.</b></p> 
<b>Scope of application</b>	<p>Use on medical devices</p>
<b>Commercial presentation</b>	<p>The 1 liter bottle with measuring device. The 5 liter can.</p>



DENTO-VIRACTIS

		Antimicrobial activity	Concentration	Contact time
<b>Bactericidal</b>	EN 13727	Pseudomonas aeruginosa, Staphylococcus aureus, Enterococcus hirae	0,5%	15 min
	EN 13697			
	EN 1276			
	EN 14561	Pseudomonas aeruginosa, Staphylococcus aureus, Enterococcus hirae	0,5%	15 min
<b>Yeasticidal</b>	EN 13624	Candida albicans	0,5%	15 min
	EN 13697			
	EN 1650			
	EN 14562	Candida albicans	0,5%	15 min
<b>Virucidal</b>	EN 14476	Herpes virus	0,5%	15 min
		Hepatite B		
		Hepatite C	0,5%	15 min
	EN 17111	Vaccina virus	0,5%	15 min



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<b>Instructions for use</b>	Concentration: 0,5 % (5 ml/L) =(25ml/5L). Contact time: 15 minutes. Temperature: 20°C.
<b>Precautions for use</b>	See Material Safety Data Sheet. <b>CAUTION</b> Causes severe eye irritation. Toxic to aquatic organisms, causes long-term adverse effects. <b>IF IN EYES:</b> Carefully flush with water for several minutes. Remove contact lenses if worn and if they can be. Remove contact lenses if present and easily removable. Continue rinsing.
<b>Notice Instruction Manual (2020/08/03)</b>	1st marketing: 2020-07. Intended for exclusive use by professionals for the pre-disinfection of invasive medical devices such as surgical instruments, operating room equipment, endoscopes endoscopes, ophthalmology and contactology equipment. Do not use the product after its expiration date. Close the canister tightly after use. Do not reuse the pre-disinfection bath (or follow local recommendations). Do not mix with other products. Packaging must be disposed of as hazardous waste under the sole responsibility under the sole responsibility of the waste holder. Do not dispose of Do not dispose of residues in sewers or waterways. Any serious incident with the device should be reported to the manufacturer and the and to the appropriate authority. Check the compatibility of the treated material with materials tested in corrosion studies COR010449 : compatibility with raw steel, stainless steel, aluminum, copper, endoscope sheaths, HDPE, polysulfone, PVC, silicone, tungsten, viton and brass; and sensitivity of bichromated steel steel, zinc-plated steel and zinc.
<b>Labelling</b>	 

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