



Declaration of Conformity

Manufacturer	Becton Dickinson GmbH Tullastrasse 8-12 69126 Heidelberg/Germany Tel.: +49-62 21-30 50 Fax: +49-62 21-30 52 16																																					
Conformity Assessment Procedure	IVD Directive 98/79/EC, Annex III																																					
Product(s):	<table><thead><tr><th>Product Name</th><th>Cat. No.</th></tr></thead><tbody><tr><td>Baird Parker Agar</td><td>255084</td></tr><tr><td>Columbia CNA Agar with 5% Sheep Blood</td><td>254007</td></tr><tr><td>Columbia CNA Agar with 5% Sheep Blood</td><td>254072</td></tr><tr><td>DNase Test Agar</td><td>255506</td></tr><tr><td>Enterococcosel Agar</td><td>254019</td></tr><tr><td>Gardnerella Selective Agar with 5% Human Blood</td><td>254094</td></tr><tr><td>Gardnerella Selective Agar with 5% Human Blood</td><td>257664</td></tr><tr><td>Group A Selective Streptococcus Agar (sSA)</td><td>254050</td></tr><tr><td>LBS Agar</td><td>255011</td></tr><tr><td>Mannitol Salt Agar</td><td>254027</td></tr><tr><td>Mannitol Salt Agar</td><td>254079</td></tr><tr><td>Middlebrook 7H10 Agar</td><td>254520</td></tr><tr><td>Middlebrook 7H10 Agar</td><td>254521</td></tr><tr><td>Modified CNA Agar + Crystal Violet + 5% Sheep Blood</td><td>255086</td></tr><tr><td>Neomycin Agar with 5% Sheep Blood</td><td>254444</td></tr><tr><td>Tellurite Agar (Hoyle)</td><td>256044</td></tr><tr><td colspan="2"></td></tr></tbody></table>	Product Name	Cat. No.	Baird Parker Agar	255084	Columbia CNA Agar with 5% Sheep Blood	254007	Columbia CNA Agar with 5% Sheep Blood	254072	DNase Test Agar	255506	Enterococcosel Agar	254019	Gardnerella Selective Agar with 5% Human Blood	254094	Gardnerella Selective Agar with 5% Human Blood	257664	Group A Selective Streptococcus Agar (sSA)	254050	LBS Agar	255011	Mannitol Salt Agar	254027	Mannitol Salt Agar	254079	Middlebrook 7H10 Agar	254520	Middlebrook 7H10 Agar	254521	Modified CNA Agar + Crystal Violet + 5% Sheep Blood	255086	Neomycin Agar with 5% Sheep Blood	254444	Tellurite Agar (Hoyle)	256044			
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We hereby declare that the above mentioned product(s) comply with the European In Vitro Diagnostic Directive and its relevant transposition into national laws of the member states into which we place the devices.																																						
Heidelberg/Germany,	March 14, 2016																																					
Name and Authority	Dr. Bernd Peschke Regulatory Affairs Manager G/S/A, Sicherheitsbeauftragter (Safety Officer)																																					
Signature																																						

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