

EU DECLARATION OF CONFORMITY

Product

Product name/ Trade name	REF	Basic UDI-DI		
Substrat/ Substrate LINE <table border="1" style="display: inline-table; vertical-align: middle;"> <tr> <td>SUBS</td> <td>RTU</td> </tr> </table>	SUBS	RTU	WE200.02	++E654WE20002V02JE
SUBS	RTU			
Intended purpose				
Detection/Measurement	For use with LINE products from Virotech Diagnostic			
Function	Reagent for colour development of the LINE test sample by enzyme activity			
Specific information about	n/a			
Automated	n/a			
Type	n/a			
Specimen	Human Serum or Plasma/CSF			
Testing population	n/a			
Intended user	Specialised personnel in laboratories			

Manufacturer



Virotech Diagnostics GmbH
 Waldstrasse 23 A2
 63128 Dietzenbach
 Germany

SRN: DE-MF-000025733

We, as the manufacturer of the device(s) take sole responsibility for and hereby declare that the above-mentioned product(s) meet(s) the provisions of the following Regulation:

Regulation EU 2017/746 on *In vitro* Diagnostic Medical Devices

CLASSIFICATION	A
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Conformity Assessment Procedure	
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ANNEX II+III	apply
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Common Specifications	n/a
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Valid from lot: 20061

Valid until: 2023-08-31

<small>DocuSigned by:</small> <i>Marcello Salio</i> <small>B462E9960D47494...</small>	Dietzenbach	2022-12-15
Signature	Issued in	Date of issue
Marcello Salio	Managing Director	
Name	Function	