



3EC International a.s., Hraničná 18, 821 05 Bratislava, Slovakia  
Notified Body No. 2265

## EC CERTIFICATE

No. 2021-IVD/QS-007/A

issued in compliance with Directive 98/79/EC of the European Parliament and of the Council on in vitro diagnostic medical devices as amended, certifies that in vitro diagnostic medical devices according to Annex II, List B

**EIA Chlamydia pneumoniae REC IgA**  
**EIA Chlamydia pneumoniae REC IgG**  
(for detailed list refer to Annex if it is necessary)

manufactured by company

**TestLine Clinical Diagnostics s.r.o.**  
Křižikova 68, 612 00 Brno, Czech Republic


are manufactured under conditions fulfilling the quality system requirements of Annex IV, excluding (4 and 6), of the Directive 98/79/EC.

The Notified Body No. 2265 has performed an audit of the above products quality system and found that quality system meets the requirements. The quality system has been assessed, approved and is subject to continuous surveillance according to Annex IV, Section 5, of the Directive 98/79/EC. The detailed description of the system, requirements and measures applied by the manufacturer are presented in the Audit Report No. ICT\_131 and the Final protocol No. 320065-7/2021.

*This Certificate is issued under the following conditions:*

It applies only to the quality system maintained in the manufacture of the above referenced models of in vitro diagnostic medical devices and it does not substitute the design or type-examination procedures, if requested. The certificate remains valid until the manufacturing conditions or the quality system are changed but until May 26th, 2025 at the latest. The certificate validity is conditional upon positive results of regular surveillance audits and fulfillment of relevant legal and other requirements by manufacturer.



  
Dr. Katarina Tomin Srdošová  
Responsible to act on behalf of NB 2265

At Bratislava, on May 23rd, 2022  
This EC Certificate supersedes the EC Certificate No. 2021-IVD/QS-007

Certificate history:

<b>Revision</b>	<b>Date of issue</b>	<b>Application for Conformity Assessment of MD number</b>	<b>Description</b>
0	01.02.2021	320065	First issue of the Certificate
A	23.05.2022	320065	Issue of Certificate No. 2021-IVD/QS-007/A with extended validity until May 26th, 2025 based on Regulation (EU) 2022/112



## ANNEX TO EC CERTIFICATE No. 2021-IVD/QS-007/A

issued for the company

**TestLine Clinical Diagnostics s.r.o.**  
Křižíkova 68, 612 00 Brno, Czech Republic

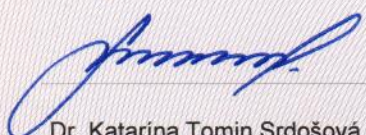
### List of *in vitro* diagnostic medical devices covered by the EC Certificate:

IVD MD Name	Variant
EIA Chlamydia pneumoniae REC IgA	EIA Chlamydia pneumoniae REC IgA
	SmartEIA Chlamydia pneumoniae REC IgA
EIA Chlamydia pneumoniae REC IgG	EIA Chlamydia pneumoniae REC IgG
	SmartEIA Chlamydia pneumoniae REC IgG

Page 1 of 1



At Bratislava, on May 23rd, 2022  
Valid until May 26th, 2025

  
Dr. Katarina Tomin Srdošová  
Responsible to act on behalf of NB 2265