Unit on MPA

TRANSASIA®

INSTALLATION & TRAINING REPORT

Customer Name : SRV Path Labs pvt Ltd

Department : LABORATORY

Address : B-2/28A, YAMUNA VIHAR, NEW DELHI

City : NEWDELHI Pin : 110053 State : NEW DELHI

Territory : DELHI-3 Zone : NZ-1

Telephone & Fax : NA email : NA

Contact Person : RAJIV MobileNo. : 9354381345

Instrument Model: ELITE H-580 Sr. No. : K1105213202 9

Installation Date : 14/04/2022

The above mentioned instrument has been satisfactorily installed by Service Engineer / Product Specialist of Transasia Bio-Medicals Ltd. Operational Training & User Maintenance of the instrument was provided to the following staff members on date (s) : 14/04/2022

Name (s)	Designation/ Dept.
1. Jyoti Gupta	Lab Technician
2. Sanjeev Jha	Sr. Lab Technician
3. Dharmen	Lab Technician

Test/ Parameters Demonstrated: CBC

Customer Comments (If any):

Engineer / Product Specialist : Application

Signature : Bhojraj / Rohit

Name : Bhojraj / Rohit Gupta

Date : 14/04/2022

Customer Signature :

Customer Name : Rajeev Jha

Designation : For SRV PATH LABS PVT. LTD.

Customer Seal :

DIRECTOR

Arbitration Clause

All disputes, Differences, controversies and questions directly or indirectly arising at any time under, out of, in connection with or in relation to the payment of this instrument / reagents / service shall be referred to the sole arbitrator to be appointed by Transasia Bio-Medicals Limited under the Arbitration and Conciliation Act 1996. The venue and seat of arbitration shall be at Mumbai. The award rendered by the arbitrator(s) shall be final and binding upon both Parties.

Erba Lachema s.r.o

DECLARATION OF CONFIRMITY and CALIBRATION

CE

The undersigned Erba Diagnostics Ltd., Unit 4, Block K, Vantage Suites, Central Park, Leopardstown, Dublin 18, manufacturer of diagnostic devices, with Company Quality Management System in compliance with standard ISO 13485,

declares, that the device:

Name:ELite580	K11052132029.

complies with all the essential requirements listed in Annex I of Directive 98/79/EC and obligations specified in Annex III, art. 2 – 5 of Directive 98/79/EC for In vitro Medical-Diagnostic Devices.

Therefore Erba Lachema s.r.o. declares and assures the following:

1. The a.m. device complies with the applicable provisions of Directive 98/79/EC.
2. The a.m. device is not included in the list A and B of the Annex II of Directive 98/79/EC.
3. The manufacturer declares to have established a procedure and to maintain it in order to assure the post-marketing surveillance, according to the Directive of 98/79/EC.
4. All mentioned products are Factory calibrated and Final QC passed at the time of release.

Date of issue: June-2021

Manish Airan
Representative of Quality Management

Erba Diagnostics Ltd.
Unit 4, Block K, Vantage Suites, Central Park, Leopardstown, Dublin 18
www.erbamannheim.com VAT Reg No : IE3517014VH

This is an electronically generated document requires no signatures.

Certificate of Conformity and Calibration