AN ITALIAN COMPANY THAT RESEARCHES, DEVELOPS AND PRODUCES READY TO USE KITS
USING CHROMATOGRAPHIC TECHNIQUES FOR CLINICAL LABORATORIES

CONTROL LYOPHIL. IN SERUM FOR VITAMINS A/E - LEVELS 1 and 2

Code Z18019

OBJECTIVE
These controls in serum are used for the internal quality control and serve to monitor the accuracy and precision of analytical procedures devoted to the quantitative determination of analytes contained therein. These controls are lyophilized human matrix and are available in two different concentration ranges. Should be handled as if they were a real patient sample.

RECOVERY
Remove the metal seal and rubber stopper from the vial. Add exactly 1 ml of HPLC-grade water through the tube. Replace the rubber stopper, allow to stand for at least 5 to 10 minutes. Before use, mix by inverting the vial to dissolve the material until a homogeneous solution.

STORAGE AND STABILITY
The controls are stable for 36 months from the date of preparation if stored at -20 ° C. After reconstitution are stable for 1 day at 2 – 8 °C and 1 month at -20 ° C. **Store at the dark**. Do not use after the expiry date.

PRECAUTIONS
These controls in human matrix should be treated with care and treated as potentially infectious.

LOT		
027	July 2023	July 2026

PACKAGINGS AVAILABLE:

- Z18019 CONTROL IN SERUM FOR VITAMINS A/E - LEVELS 1 and 2 ***2 x 5 x 1 ml***

Release N° 002	**July 2019**

CONCENTRATIONS
CODE Z18019

LOT		
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ANALYTE	*MEASUREMENT UNIT*	*MEDIUM VALUE L1*	*RANGE L1*	*MEDIUM VALUE L2*	*RANGE L2*
VITAMIN A	**µg/ml**	**0,6**	**0,4 – 0,7**	**1,8**	**1,3 – 2,3**
VITAMIN E	**µg/ml**	**10,7**	**7,5 – 13,9**	**52,8**	**37,0 – 68,7**

This product fulfills all the requirements of Directive 98/79/EC of 27/10/1998 on in vitro diagnostic medical devices (IVD). The declaration of conformity is available upon request.

CE

FOR *IN VITRO* DIAGNOSTIC USE ONLY

produced by

COMPANY WITH MANAGEMENT SYSTEM CERTIFIED BY DNV
= ISO 9001 =
= ISO 13485 =

eureka Lab Division

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EUREKA srl – LAB DIVISION ,
Via Enrico Fermi 25, 60033 Chiaravalle (AN) ITALY
Tel. + 39 071 74 50 790
Fax. + 39 071 74 96 579
info@eurekaone.com
www.eurekakit.com

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