



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

**CONTROLS LYOPHIL. IN PLASMA FOR COENZYME Q10 - LEVELS 1 and 2**  
**Code Z63019**  
**TO USE WITH THE KIT CODE Z63110**

**OBJECTIVE**

These controls in plasma 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 calibrators are stable for 36 months from the date of preparation if stored at -20 ° C. After reconstitution are stable for 1 month at -20 ° C. Do not use after the expiry date.

**PRECAUTIONS**

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

LOT		

**PACKAGING AVAILABLE:**



- Z63019 CONTROL IN PLASMA FOR COENZYME Q10 - LEVELS 1 and 2

2 x 5 x 1 ml

N° 003

06/2024

CONCENTRATIONS  
CODE Z63019

LOT		

<i>ANALYTE</i>	<i>MEASUREMENT UNIT</i>	<i>MEDIUM VALUE L1</i>	<i>RANGE L1</i>	<i>MEDIUM VALUE L2</i>	<i>RANGE L2</i>
Coenzyme Q10	µg/l				

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.



**FOR IN VITRO DIAGNOSTIC USE ONLY**



EUREKA S.R.L. LAB DIVISION  
Via M. D'Antona 28  
60033 Chiaravalle (AN) - Italy  
Tel +39 071 7450790  
eureka-support@sentinel.it  
www.eurekakit.com