



The World's Largest Range of Diagnostic Kits for Analytical Chromatographic Systems (HPLC, GC, GC-MS, LC-MS/MS)



CONTROL IN PLASMA FOR TAMOXIFEN/ENDOXIFENE - LEVELS 1 and 2

Code LC92019

OBJECTIVE

These controls on blood 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 2-8 ° C. After reconstitution are stable for 1 week at 2-8°C and 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.

002	April 2019	April 2022

PACKAGING AVAILABLE:

- LC92017 CONTROL IN PLASMA FOR TAMOXIFEN/ENDOXIFENE - LEVEL 1 *5 x 1 ml*
- LC92018 CONTROL IN PLASMA FOR TAMOXIFEN/ENDOXIFENE - LEVEL 2 *5 x 1 ml*
- LC92019 CONTROL IN PLASMA FOR TAMOXIFEN/ENDOXIFENE - LEVELS 1 and 2 *2 x 5 x 1 ml*

CONCENTRATIONS
CODE LC92019

LOT		
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ANALYTE	MEASUREMENT UNIT	MEDIUM VALUE L1	RANGE L1	MEDIUM VALUE L2	RANGE L2
TAMOXIFEN	ng/ml	26,0	18,2 – 33,8	146,8	102,8 – 190,8
ENDOXIFENE	ng/ml	3,7	2,6 – 4,8	7,9	5,5 – 10,3

FOR IN VITRO DIAGNOSTIC USE ONLY

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The ONLY Italian Company that Researches, Develops and Produces
diagnostic KITS for HPLC, GC, GC-MS, LC-MS/MS

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

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Release N° 002

September 2019