CHROMSYSTEMS

EU-Declaration of Conformity

Manufacturer	Chromsystems Instruments & Chemicals GmbH
Address	Am Haag 12 82166 Gräfelfing, Germany
SRN (single registration number)	DE-MF-000010089

Order No.	Device Description	EMDN Code
Basic l	JDI-DI: 92922/XT: 425031792922XTR6	
92922/XT	MassTox® TDM Series A PARAMETER Set Antimycotic Drugs/EXTENDED in serum/plasma	W01010499
92051/XT	3PLUS1® Multilevel Plasma Calibrator Set MassTox® Antimycotic Drugs/EXTENDED	W0101050301
0252/XT	MassCheck [®] Antimycotic Drugs/EXTENDED Plasma Control Bi-Level (I + II)	W0101050299
0253/XT	MassCheck [®] Antimycotic Drugs/EXTENDED Plasma Control Level I	W0101050299
0254/XT	MassCheck [®] Antimycotic Drugs/EXTENDED Plasma Control Level II	W0101050299
92644/XT	Internal Standard Mix MassTox [®] Antimycotic Drugs/EXTENDED	W0101050399
92039/XT	Tuning Mix MassTox [®] Antimycotic Drugs/EXTENDED	W0101050399

Basic UDI-DI: 92111: 4250317921116F				
92111/200	MassTox [®] TDM BASIC Kit A, for 200 analyses	W01010499		
92111/1000	MassTox® TDM BASIC Kit A, for 1000 analyses	W01010499		
92001	Mobile Phase 1	W01019099		
92002	Mobile Phase 2	W01019099		
92003	Precipitation Reagent	W01019099		
92005	Extraction Buffer	W01019099		
92007	Dilution Buffer 1	W01019099		
92008	Dilution Buffer 2	W01019099		
92009	Rinsing Solution	W01019099		

92110	MassTox [®] TDM MasterColumn® Series A	W01019099
	Analytical column (equilibrated, with test chromatogram)	

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Device Intended Purpose	voriconazole in human serum or plasma samples via liquid chromatography mass spectrometry (LC-MS/MS).					
Risk Class	B, as per EU Regulation 2017/746, Annex VIII, Rule 6					
GMDN Code	64329, term "Multiple antifungal therapeutic drug monitoring IVD, kit, liquid chromatography/mass spectrometry (LC/MS)"					
Notified Body	TÜV Süd Product Service GmbH Ridlerstraße 65, 80339 Munich, Germany	Identification	No. 0123			
Conformity Assessment	Conformity assessment based on a quality management system and on assessment of technical documentation - Annex IX					
Declarations	Tox [®] Amministic DirightENTEMDED	aeoM xiM grám	AT . MARCOSP			
This EU declaration of conformity is issued under the sole responsibility of the manufacturer. The devices that are covered by the present declaration are in conformity with the In-Vitro Diagnostic Medical Devices Regulation (2017/746/EU) (IVDR). Following Common Specifications were considered as part of determining device conformity with						
the IVDR:		I MUL XONGED	MIGOUISIERSY			
	non Specifications exist for the concern	ed device.				
Additional information	n/a		N			
This EU declaration of conformity is issued by Gräfelfing, August 1 st , 2022 Michael Meier, Managing Director Gräfelfing, August 1 st , 2022 Dr. Ralf Fischer, PRRC						
COSCO D LANG						
EU declaration of conformity valid untilJuly 20th, 2027Version: 1.0						