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 UDI-DI: 42503176000B5Q		
6000-B	HPLC Reagent Kit Catecholamines in urine	W01010499
6000-BK	HPLC Basic Kit Catecholamines in urine	W01010499
5001	Mobile Phase	W01019099
5002	Mobile Phase	W01019099
6033	Analytes Mix	W0101050399
6009	Urine Calibration Standard	W0101050302
6004	Internal Standard	W0101050399
6055	Neutralisation Buffer	W01019099
6006	Elution Buffer	W01019099
6007	Sample Clean Up Columns	W01019099
6100	HPLC column	W01019099
Basic UDI-DI: 4250317102040502B		
0040	Endocrine Urine Control, Normal Range	W0101050207
0050	Endocrine Urine Control, Pathological Range	W0101050207

Device Intended Purpose	The Chromsystems assay "Catecholamines in urine" is an in vitro diagnostic medical device designed for professional users in clinical laboratories for the quantitative determination of noradrenaline (syn. norepinephrine), adrenaline (syn. epinephrine), and dopamine in human urine samples. Sample preparation is carried out manually, and analytic separation is done via high performance liquid chromatography with electrochemical detection (HPLC-ECD). The assay is intended to be used for patients in whom the urinary levels of noradrenaline, adrenaline and dopamine are of clinical importance, primarily as an aid to diagnosis and monitoring of suspected pheochromocytoma and paraganglioma.		
Risk Class	C, as per EU Regulation 2017/746, Annex VIII, Rule 3h		
GMDN Code	63823 "Multiple biogenic amine neurotransmitter IVD, kit, liquid chromatography"		
Notified Body	BSI Group The Netherlands B.V. Say Building, John M. Keynesplein 9 1066 EP Amsterdam The Netherlands	Identification No.	2797

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Conformity Assessment	Conformity assessment based on a quality management system and on assessment of technical documentation - Annex IX Chapters I and III
Certificates issued	EU Quality management System Certificate (IVDR) IVDR 838647
Declarations	
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:	
Not applicable as no Common Specifications exist for the concerned device.	
Additional information	n/a
This EU declaration of conformity is issued by	
Gräfelfing, March 23rd, 2026 Michael Meier, Managing Director	Gräfelfing, March 23rd, 2026 Dr. Ralf Fischer, PRRC
EU declaration of conformity valid until:	March 22nd, 2031 / Version: 2.0

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