DAC-SpectroMed S.R.L. Chisinau	Declaration of EC-Conformity	File No.	F-PS-10-05
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DAC-SpectroMed SRL

Nicolae Testemitanu str. 37, MD-2025, Chisinau, Republic of Moldova

hereby declares under its own responsibility that the *in-vitro diagnostics* medical devices (agent de curățare analizoarelor)

Clean Cell-DAC < 5%	100ml				
Clean Cell-DAC < 5%	500 ml	Clean Cell-DAC < 1%	100 ml		
Clean Cell-DAC < 5%	1000 ml	Clean Cell-DAC < 1%	500 ml	Clean Cell-DAC < 0,5%	100 ml
		Clean Cell-DAC < 1%	1000 ml	Clean Cell-DAC < 0,5%	500 ml
				Clean Cell-DAC < 0,5%	1000 ml

- are classified as neither listed in Annex II nor for self-testing according to EC Council Directive 98/79/EC from 27th October 1998 on in-vitro diagnostic medical devices;
- are in accordance with the Annex III of the EC Council Directive 98/79/EC from 27th October 1998 on in-vitro diagnostic medical devices.

DAC-SpectroMed SRL has a Quality System in place based on EN ISO 9001:2015 and EN ISO 13485:2016, issued by the IQNet&SRAC.

Our Authorized Representative in EU is Qarad EC-REP BV, with address at Pas 257, 2440 Geel, Belgium.

FIRMA STIINTIFICĂ DE PRODUCȚIE S.R.L. «DAC-SPECTROMED» IDNO 1002600042432 -2- REPUBLICA MOLDOVA, mun. CHISINĂU

General Manager
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01.11.2022

Specialist of Normative-Technical Documentation
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01.11.2022

Digitally signed by Efremova Iulia
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