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Proof of Authority

To use the CE mark on pressure equipment on the basis of internal production control plus supervised pressure equipment checks at random intervals according to module A2 of Decree 44/2016. (XI.28.) NGM and Directive 2014/68/EU of the European Parliament and Council

Certificate no.:	**H/Ü 23 0274**
Name and address of the manufacturer:	**Celitron Medical Technologies Kft.** **H-2600 Vác** **Nádas utca 2.** **Hungary**

The manufacturer is authorized on the basis of the results of the verification of the preconditions to use the CE mark along with the identification number of the notified body within the scope of the validity on the pressure equipment manufactured by the manufacturer as follows:

CE 1008

Scope of validity	**ISS AC575 Chamber**
Report No.:	E 122/2067/2024
Valid until:	12.11.2025
	(First issue of the Certificate: 02.02.2023)

Budapest, 03.12.2024

TÜV Rheinland InterCert Kft. TÜVRheinland® 10. Notified Body 1008

Ferenc Székely
Certifier

TÜV Rheinland InterCert Kft.
Industrial Services Business Unit
NoBo Number: 1008

TÜV Rheinland InterCert Kft. Ipari szolgáltatások üzletág – H-1143 Budapest, Gizella út 51-57.
tel.: +36-1/4611-100; fax: +36-1/4611-199; e-mail: tuv@hu.tuv.com, honlap: www.tuv.hu

Supervision report

Supervised pressure equipment checks at random intervals

TÜVRheinland®
Precisely Right.

Identification number of the notified body: 1008

		Report number	E122/2067/2024
MODUL:	**A2**	Internal production control plus supervised pressure equipment checks at random intervals	☒
	C2	Conformity to type based on internal production control plus supervised pressure equipment checks at random intervals	☐
	H1	Conformity based on full quality assurance plus design examination	☐
	E	Acceptance of pressure equipment of the III and IV categories according to the diagrams 1, 2, 3, 4, 5, acc.to 38. § and article 4. item (1) and (2)	☐
	D	Acceptance of pressure equipment of the III and IV categories according to the diagrams 1, 2, 3, 4, 5, acc.to 38. § and article 4. item (1) and (2).	☐

Manufacturer:/ Supplier:	Manufacturer's factory:
Celitron Medical Technologies Kft.	**Auth Impex Kft.**
Nádas utca 2.	**Budapesti országút, hrsz.: 4032/32**
H-2600 Vác	**H-7700 Mohács**

BASIS OF THE TEST:	**Decree 44/2016 (XI.28.) NGM, Directive 2014/68/EU**	
DOCUMENTS PRESENTED:	Technical documentation (Modul A2, C2, H1)	☒
	EU-type examination certificate (Module C2)	☐
	EC design examination certificate (Module H1)	☐
	Certificate of the QA certificate (Module E, D, H1)	☐

TESTS CARRIED OUT:		Fulfilled	Comments
1.	The aforementioned documents, reports and certificates are available	X	ISS AC575 Chamber Drawing Nr.: MWT150-100-00-21-AD-13
2.	The technical documentation fulfills the requirements of the Directive	X	-
3.	The tests have been carried out in the necessary scope according to art. 3.2 of Annex 3. / Annex I.	X	-
4.	Test of one or more pressure equipment Charge-No.: 00087.13.23044	X	Scope

RESULTS:

The aforementioned tests have been carried out according to the specified testing requirements.
The test documents are complete. In case of the modules A2, C2 the use of the CE 1008 mark on the aforementioned pressure equipment is approved until the next inspection.

Place: **Budapest** **Date:** **03.12.2024.**

TÜV Rheinland InterCert Kft. TÜVRheinland 18 Notified Body 1008

Attachment: -

Peter Bicsak - Expert

The most important data of the pressure equipment and the company data are stored for the smooth performance of the order. Your data are safe.

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