

**Full Quality Assurance System**  
**Directive 93/42/EEC on Medical devices, Annex II excluding (4)**

CE Certiso Kft. (NB 2409) certifies that the following manufacturer's quality management system concerning to the listed devices and device categories meets the requirements of the related requirements of the directive.

Name of the manufacturer:

**Uscom Research, Development and Manufacturing Ltd.**

Headquarters: **1119 Budapest, Boglárka utca 17., Hungary**

Scope:

**Respiratory measurement and diagnostic devices,  
including spirometers and software**

The certificate covers the following devices:

| Description of the device  | Type              | Intended use                                  | Risk class |
|----------------------------|-------------------|---|------------|
| PC Spirometer              | SpiroSonic Flo    | pulmonary function diagnostics and monitoring | IIa        |
| Mobile Handheld Spirometer | SpiroSonic Smart  |   |            |
| Bluetooth Spirometer       | SpiroSonic Mobile |   |            |
| Mobile Spirometer          | SpiroSonic Air*   |   |            |

This certificate is valid only in case of successfully conducted annual surveillance audits.

ID number of the related audit report: **117-CE-171122**

Issue: 4

Issued: 21 May 2021

First issued: 18 June 2018

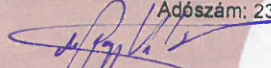
Start date of certified status: 17 October 2016, \*21 May 2021

Expires:

**17 June 2023**



**CE Certiso**  
Orvos- és Kórháztechnikai  
Ellenőrző és Tanúsító Kft.  
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Adószám: 23147049-2-13

  
**Valter PAPP, Dr.**  
General Manager