

TÜV Rheinland LGA Products GmbH • 51105 Köln

77 Elektronika Műszeripari Kft
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Hungary

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Date April 23, 2025

Notified Body Confirmation Letter

Reference. : 77ELE_PDQ0_HX_2025-04-11

To whom it may concern,

Confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation (EU) 2024/1860 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain in vitro diagnostic medical devices.

This letter confirms that **TÜV Rheinland LGA Products GmbH**, a Notified Body (NB) designated against Regulation (EU) 2017/746 (IVDR) and identified by the number **0197** on NANDO, has received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of IVDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of IVDR with the following manufacturer:

77 Elektronika Műszeripari Kft
Fehérvári út 98,
1116 Budapest
Hungary
SRN Number: HU-MF-000004266

The devices covered by the formal application and the written agreement mentioned above are identified in the tables below. Table 1 identifies the devices for which an IVDR application has been received, written agreement concluded and for which the NB is also responsible for appropriate surveillance under the applicable Directive 98/79/EC. Table 2 identifies the devices for which an IVDR application has been received and a written agreement concluded, but the NB has not yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive 98/79/EC.

In the case of devices covered by certificates issued under Directive 98/79/EC (IVDD) that expired after May 26, 2022 and before July 9, 2024, without having been withdrawn, this letter also confirms that the manufacturer either signed the written agreement under IVDR by the date of IVDD certificate expiry; or provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 54 of IVDR or Article 92 of the IVDR respectively, by July 9, 2024 for the relevant devices.

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Chairman of the
Supervisory Board

Dr.-Ing. Michael Fübi

The transition timelines that apply to the devices covered by this letter, subject to the manufacturer's continued compliance to the other conditions specified in Article 110.3c of IVDR (as amended by (EU) 2024/1860), are shown below:

- 31 December 2027 for devices covered by an IVDD certificate regardless of their risk class under the IVDR
- For devices not requiring the involvement of a notified body under the IVDD, but requiring it under the IVDR and for which a declaration of conformity was drawn up prior to 26 May 2022 in accordance with Directive 98/79/EC, the following dates apply:
 - 31 December 2027, for class D devices;
 - 31 December 2028, for class C devices;
 - 31 December 2029, for class B devices and for class A devices placed on the market in sterile condition

On behalf of the Notified Body

Rafał Byczkowski
Certification body

Table 1: Devices covered by this letter and for which the NB is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive 98/79/EC:

| Device name or Basic UDI-DI (under IVDR application) | IVDR Device classification (as proposed by the manufacturer and verified at the pre-application stage) | If the IVDR device is a substitute device, identification of the corresponding IVDD device | IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification |
|--|--|--|--|
| Dcont ETALON blood glucose meter Basic UDI-DI: 59973457DEK7V | Class C, self-testing | Dcont ETALON, AutoSense, AutoSense Plus, blood glucose meters | 7-029-400-2002 NB 1011 |
| Dcont NOVUM blood glucose meter Basic UDI-DI: 59973457DEK7V | Class C, self-testing | SensoCard, blood glucose meters | HL 1006099-1 NB 0197 |
| Dcont NOVUM blood glucose meter Basic UDI-DI: 59973457DEK7V | Class C, self-testing | N/A | 7-029-400-2002 NB 1011 |
| Dcont ETALON B blood glucose meter Basic UDI-DI: 59973457DEK7V | Class C, self-testing | SensoLite Nova, blood glucose meters | HL 1006099-1 NB 0197 |
| Dcont ETALON B blood glucose meter Basic UDI-DI: 59973457DEK7V | Class C, self-testing | N/A | 7-029-400-2002 NB 1011 |

| Device name or Basic UDI-DI (under IVDR application) | IVDR Device classification (as proposed by the manufacturer and verified at the pre-application stage) | If the IVDR device is a substitute device, identification of the corresponding IVDD device | IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification |
|---|---|---|---|
| Dcont ROLL blood glucose meter Basic UDI-DI: 59973457DEK7V | Class C, self-testing | SensoCard Plus, SensoLite Nova Plus blood glucose meters | HL 1006099-1 NB 0197 |
| Dcont ROLL blood glucose meter Basic UDI-DI: 59973457DEK7V | Class C, self-testing | Dcont MONDA, AutoSense Voice, blood glucose meters | 7-029-400-2002 NB 1011 |
| ETALON Teszt test strip Basic UDI-DI: 59973457ETC8X | Class C, self-testing | SensoCard, SensoLite Nova test strips | HL 1006099-1 NB 0197 |
| ETALON Teszt test strip Basic UDI-DI: 59973457ETC8X | Class C, self-testing | ETALON Teszt, Ideál Teszt, AutoSense, test strips | 7-029-400-2002 NB 1011 |
| KETO Cont blood ketone meter Basic UDI-DI: 59973457KKD94 | Class C, self-testing | N/A | 7-031-306-2009 NB 1011 |
| KETO Fit blood ketone meter Basic UDI-DI: 59973457KKD94 | Class C, self-testing | N/A | 7-031-306-2009 NB 1011 |
| KETO Cont Teszt test strip Basic UDI-DI: 59973457KTC9V | Class C, self-testing | N/A | 7-031-306-2009 NB 1011 |
| KETO Cont Teszt test strip Basic UDI-DI: 59973457KTC9V | Class C, self-testing | N/A | 7-031-306-2009 NB 1011 |
| KETO Cont Teszt test strip Basic UDI-DI: 59973457KTC9V | Class C, self-testing | N/A | 7-031-306-2009 NB 1011 |

Table 2: Devices covered by this letter and for which the NB is NOT responsible for appropriate surveillance of the corresponding devices under the applicable Directive 98/79/EC:

| Device name or Basic UDI-DI (under IVDR application) | IVDR Device classification (as proposed by the manufacturer and verified at the pre-application stage) | If the IVDR device is a substitute device, identification of the corresponding IVDD device | IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification |
|--|--|--|--|
| none | | | |

Confirmation Letter Revision History

| Date | NB internal reference traceable to each version of the letter | Action |
|------------|---|---------------|
| 2025-04-23 | 77ELE_CL1860_2025-04-23 | Initial issue |