**DQS Medizinprodukte GmbH** | August-Schanz-Str. 21 | 60433 Frankfurt am Main

# Kaz Europe Sàrl

Q-Center

Route de la Chaux 4
1030 Bussigny
Switzerland

Date: 2024-05-24

## Notified Body Confirmation Letter
## Reference: 1000179480

To whom it may concern,

**Confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation EU 2023/607 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices**

This letter confirms that, DQS Medizinprodukte GmbH, a Notified Body designated against Regulation (EU) 2017/745 (MDR) and identified by the number 0297 on NANDO, has received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR with the following manufacturer:

# Kaz Europe Sàrl

Route de la Chaux 4
1030 Bussigny
Switzerland

SRN: CH-MF-000029980

The devices covered by the formal application and the written agreement mentioned above are identified in the Tables listed below: Table 1 identifies the devices for which an MDR application has been received, written agreement concluded and for which DQS Medizinprodukte GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive. Table 2 identifies the devices for which an MDR application has been received and a written agreement concluded, but DQS Medizinprodukte GmbH has not yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive.
In the case of devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry, or provided evidence that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 59(1) of MDR or Article 97(1) of the MDR respectively, by the 20 Mar 2023 for the relevant devices. 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 120.3c of MDR (as amended by (EU) 2023/607), are shown below:

- 26 May 2026 for Class III custom-made implantable devices

- 31 December 2027 for Class III devices and Class IIb implantable devices excluding Well-established technologies (WET - sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a measuring function
- 31 December 2028 for devices not requiring the involvement of a notified body under MDD but requiring it under MDR (e.g., class I devices that qualify as re-usable surgical instruments)

On behalf of the Notified Body,



**Viviana Indraccolo**
Regulatory Affairs Manager

**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:**

| Device name and Basic UDI-DI (as proposed by the manufacturer within the application) | MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage) | If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device | MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification |
|---|---|---|---|
| **Braun TempleSwipe thermometer (ref: BST200)** 76307593BST200LN | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun Sensian 5 Noncontact thermometer (ref: BNT300)** 76307593BN400GQ | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun Sensian 7 Noncontact thermometer (ref: BNT400)** 76307593BN400GQ | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ThermoScan 3 (ref: IRT3030)** 76307593IR3030G5 | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ThermoScan 5 (ref: IRT6030)** 76307593IR6000GH | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ThermoScan 6 (ref: IRT6515)** 76307593IR6000GH | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ThermoScan 7 (ref: IRT6520)** 76307593IR6000GH | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ThermoScan 7+ (ref: IRT6525)** 76307593IR6000GH | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun High Speed thermometer (ref: PRT1000)** 76307593PR1000J9 | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun Age Precision digital thermometer (ref: PRT2000)** 76307593PR2000JG | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |

| Device name and Basic UDI-DI (as proposed by the manufacturer within the application) | MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage) | If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device | MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification |
|---|---|---|---|
| **Braun ThermoScan Hygiene cap (ref: LF40/LF20)** 76307593LF4020DC | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ExactFit 1 (ref: BUA5000)** 76307593BU5000EG | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ExactFit 3 (ref: BUA6150)** 76307593BP6000CY | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ExactFit 5 connect (ref: BUA6350)** 76307593BP6000CY | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun Nasal aspirator 1 (ref: BNA100)** 76307593BNT100JS | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ThermoScan 5 IRT6020** 76307593IR6000GH | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun ExactFit 5 BP6200PHEMEAV1** 76307593BP6000CY | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |
| **Braun iCheck 7 BPW4500** 76307593BP4500DB | Class IIa | N/A | 381008 MR5 170773847 (NB 0297) |

**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:**

| Device name and Basic UDI-DI (as proposed by the manufacturer within the application) | MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage) | If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device | MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification |
|---|---|---|---|
| **N/A** | N/A | N/A | N/A |

**17Confirmation Letter Revision History**

| Date | NB internal reference traceable to each version of the letter | Action |
|---|---|---|
| 2024-05-17 | 1000178686 | Initial issue |
| 22024-05-23 | 1000179480 | Registration issues in system – New Internal Reference issued |