Tequir, S.L.

Polígono Industrial El Oliveral, C/C,
s/n. 46190 - Ribarroja del Turia,
Valencia (Spain)

05.08.2024

Kiwa Belgelendirme Hizmetleri A.Ş.
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Notified Body Confirmation Letter Reference: MY-24-003147

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, Kiwa Belgelendirme Hizmetleri A.Ş Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number 1984 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:

Company Name*: Tequir, S.L.*

Address*: Polígono Industrial El Oliveral, C/C,*
s/n. 46190 - Ribarroja del Turia,
Valencia (Spain)

SRN Number*: ES-MF-000000783*

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 MDR application has been received, written agreement concluded and for which the NB 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 the NB 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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or provided evidence that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,

Mustafa Serkan Sevimli
Medical Devices Program Manager

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Mersis No: 0620007475700019

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 or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the preapplication 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
Femoral Implant "KEEP WALKING"	IIB	N/A	1984-MDD-19-615 Kiwa Belgelendirme Hizmetleri A.Ş #1984
Surgical Instruments for the Femoral Implant "KEEP WALKING"	IIA	N/A	1984-MDD-19-615 Kiwa Belgelendirme Hizmetleri A.Ş #1984

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 or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the preapplication 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
TRIAL IMPLANT	IIA	N/A	Self-declaration
Surgical devices for the Femoral Implant "KEEP WALKING" (DEPTH GUIDE; DIAMETER GUIDE; HEXAGONAL SCREWDRIVER; IMPACTOR; WASHER EXTRACTOR; JACOBS-HUDSON ADAPTOR; SILICONE HUDSON HANDLE; DYNAMOMETRIC HUDSON T-HANDLE; ANTI-ROTATION GUIDE; STEM EXTACTOR; BLOCKAGE SUPPORT)	IR	N/A	Self-declaration

Confirmation Letter Revision History

Date	Revision No	Action
05/08/2024	Rev00	Initial issue