

Philips Image Guided Therapy Corporation  
9965 Federal Drive  
Colorado Springs, Colorado  
80921  
USA

December 14, 2023

**Notified Body Confirmation Letter**  
**Reference: EU2023-607/672713**

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, **BSI Group The Netherlands B.V.**, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number **2797** 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:

Philips Image Guided Therapy Corporation  
9965 Federal Drive  
Colorado Springs, Colorado  
80921  
USA  
SRN Number (if available): US-MF-000018632

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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or provided evidence that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 BSI Group The Netherlands B.V.,



BSI Scheme 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 or Basic UDI-DI (under MDR 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
<b>Refinity ST Rotational IVUS Catheter</b>	Class III	N/A	CE 742145; NB 2797 CE 742150; NB 2797
<b>Eagle Eye Platinum RX Digital IVUS Catheter</b>	Class III	N/A	CE 742145; NB 2797 CE 742147; NB 2797
<b>Eagle Eye Platinum ST RX Digital IVUS Catheter</b>	Class III	N/A	CE 742145; NB 2797 CE 742147; NB 2797
<b>Lead Locking Device (LLD, LLD 1, LLD 2, LLD 3, LLD E, LLD EZ, LLD Accessory Kit, Lead Cutter)</b>	Class IIa	N/A	CE 550209; NB 2797
<b>Visions PV .018 Digital IVUS Catheter</b>	Class IIa	N/A	CE 742145; NB 2797
<b>Reconnaissance PV .018 OTW Digital IVUS Catheter</b>	Class IIa	N/A	CE 742145; NB 2797
<b>Visions PV .014P RX Digital IVUS Catheter</b>	Class IIa	N/A	CE 742145; NB 2797 CE 742147; NB 2797
<b>Omniwire Pressure Guide Wire</b>	Class III	N/A	CE 742145; NB 2797 CE 735820; NB 2797
<b>GlideLight Laser Sheath</b>	Class III	N/A	CE 550209; NB 2797 CE 554849; NB 2797
<b>AngioSculpt PTA Scoring Balloon Catheter</b>	Class IIa	N/A	CE 632473; NB 2797
<b>TightRail, TightRail Sub-C Rotating Dilator Sheaths</b>	Class III	N/A	CE 550209; NB 2797 CE 611679; NB 2797
<b>Turbo-Elite Laser Atherectomy Catheter</b>	Class IIa	N/A	CE 550209; NB 2797
<b>Bridge Occlusion Balloon Catheter</b>	Class III	N/A	CE 550209; NB 2797 CE 650647; NB 2797
<b>ELCA Coronary Laser Atherectomy Catheter</b>	Class III	N/A	CE 550209; NB 2797 CE 554813; NB 2797

Device name or Basic UDI-DI (under MDR 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
<b>Turbo-Power Laser Atherectomy Catheter</b>	Class IIb excluding Class IIb implantable non-WET	N/A	CE 550209; NB 2797
<b>IntraSight System</b>	Class IIa	N/A	CE 742145; NB 2797
<b>IntraSight Mobile</b>	Class IIa	N/A	CE 742145; NB 2797
<b>Philips Laser System</b>	Class IIb excluding Class IIb implantable non-WET	N/A	CE 550209; NB 2797
<b>SyncVision System</b>	Class IIa	N/A	CE 742145; NB 2797
<b>Sterile Equipment Covers</b>	Class I device placed on the market in sterile condition	N/A	CE 742148; NB 2797
<b>Tack Endovascular System</b>	Class IIb implantable non-WET	N/A	CE 621167; NB 2797
<b>Visions PV .035 Digital IVUS Catheter</b>	Class III	N/A	CE 742145; NB 2797 CE 742557; NB 2797

**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 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

### Confirmation Letter Revision History

Date	Action
2023/12/14	Initial issue