



EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. CE 00999
Issued To: **Varex Imaging Corporation**
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA

In respect of:

The design and manufacture of diagnostic X-Ray tubes, CT tubes and mammography tubes, and the design and manufacture of flat panel imaging components and systems.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):

Gary E Slack, Senior Vice President - Medical Devices

First Issued: **1995-11-15**

Date: **2020-01-22**

Expiry Date: **2024-05-26**

...making excellence a habit.™

Page 1 of 2

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI. This certificate was issued electronically and is bound by the conditions of the contract.

EC Certificate - Full Quality Assurance System

Supplementary Information to CE 00999

Issued To:

**Varex Imaging Corporation
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA**

NBOG Code	Device description	Intended purpose per IFU
Class IIb		
MD 1201	Diagnostic X-Ray Tubes	Intended to be used in Radiography.
MD 1201	CT Tubes	Intended to be used in Computed Tomography.
MD 1201	Mammography Tubes	Intended to be used in Mammography.
Class IIa		
MD 1201 MDS 7010	Flat Panel Imaging components – Radiography panels	Intended to be integrated into products by X-ray manufacturers for radiographic applications.
MD 1201 MDS 7010	Flat Panel Imaging components – Fluoroscopic panels	Intended to be integrated into products by X-ray manufacturers for fluoroscopic applications.

First Issued: **1995-11-15**

Date: **2020-01-22**

Expiry Date: **2024-05-26**

...making excellence a habit.™

Page 2 of 2

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI. This certificate was issued electronically and is bound by the conditions of the contract.

EC Certificate - Full Quality Assurance System

Certificate History

Certificate No: **CE 00999**
 Date: **2020-01-22**
 Issued To: **Varex Imaging Corporation**
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA

Date	Reference Number	Action
15 November 1995	-	First Issue.
17 September 1996	-	Addition of 'refurbishment' to the scope.
05 August 1998	-	Addition of sub-contractor 'VARIAN INTERAY' for 'The full refurbishment of X-ray tubes and the design and manufacture of amorphous-silicon flat panel imaging systems'.
26 January 2000	-	Addition of 'the design, development and manufacture of amorphous silicon flat panel imaging systems' to the scope.
23 February 2005	4660194	Certificate renewal. Scope update to improve regulatory compliance.
30 November 2005	4778102	Addition of Varian Medical Systems Deutschland GmbH (Germany) as a subcontractor.
11 December 2007	7145590	Clarification of scope, the activity refurbishment has been removed as it was misleading. The company will continue to fit replacement x-ray tubes in a previously manufactured tube housing assemblies Clarification of scope to specifically detail the components of amorphous silicon flat panel imaging systems.

...making excellence a habit.™

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.
 This certificate was issued electronically and is bound by the conditions of the contract.

EC Certificate - Full Quality Assurance System Certificate History

Certificate No: **CE 00999**
 Date: **2020-01-22**
 Issued To: **Varex Imaging Corporation**
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA

Date	Reference Number	Action
20 January 2010	7474386	Addition of VMS UK Ltd to significant subcontractor list as authorised representative. Certificate renewal.
13 April 2010	7506821	Addition of Varian Medical Systems X-Ray Products, California, as a design subcontractor.
09 May 2012	7819479	Company name changed from Varian Medical Systems to Varian Medical Systems, Inc. Design services moved from Varian Medical Systems, X-Ray Products, 2599 Garcia Avenue, Mountain View, CA 94043, USA to Varian Medical Systems Inc., X-Ray Products, 3120 Hansen Way- Bldg 4 M/S G-103, Palo Alto, CA 94043, USA.
23 January 2015	8253504	Certificate Renewal.

...making excellence a habit.™

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.
 This certificate was issued electronically and is bound by the conditions of the contract.

EC Certificate - Full Quality Assurance System

Certificate History

Certificate No: **CE 00999**
 Date: **2020-01-22**
 Issued To: **Varex Imaging Corporation**
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA

Date	Reference Number	Action
02 June 2015	8347872	Removal of the last four digits of the postal zip code from the legal manufacturer's address to match other certificates.
04 January 2017	8652652	Change of legal manufacturer name from Varian Medical Systems Inc to Varex Imaging Corporation. Removal of VMS UK Ltd and Varian Medical Systems, Inc – CA 94043 from the sub-contractors list and addition of Claymout. Change of name From Varian Medical Systems to Varex Imaging on the remaining subcontractors.
01 February 2018	8867468	Addition of Varex Imaging Equipment (China) as a Manufacture Sub-contractor.
24 March 2019	8940261	Change of EU Authorised Representative from Claymout to Arazy Group GmbH, Am Kalkhofen 8, Wöllstadt, 61206, Germany. Extension to scope, by removing 'amorphous-silicon'. Change of address of subcontractor Varex Imaging Deutschland AG, to Otto-Brenner-Str. 10, D 47877 Willich, Germany, following verification visit.
25 March 2019	8939261	Traceable to NB 0086.

EC Certificate - Full Quality Assurance System Certificate History

Certificate No: **CE 00999**
 Date: **2020-01-22**
 Issued To: **Varex Imaging Corporation**
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA

Date	Reference Number	Action
22 January 2020	9774581	Certificate renewal. EU Rep address changed from 'Am Kalkhofen 8, Wöllstadt, 61206, Germany' to 'Am Flughafen, The Squire 12, 60549 Frankfurt am Main, Germany'. Product table added to certificate. Correction of typographical error in the address of subcontractor Varex Imaging (China).
Non-significant changes approved after the 26th May 2021 as per the Transitional Provisions of MDR Article 120.3		
25 June 2021	3478796	Removed subcontractor "Varex Imaging Corporation, 3235 Fortune Drive, North Charleston, South Carolina, 29418 USA"
28 August 2024	30207540	Addition of subcontractor. Removal of subcontractor page.
14 April 2025	30377744	Addition of critical subcontractor.
24 April 2026	30680018	Change of critical subcontractor. Addition of manufacturer location.

24 April 2026

Varex Imaging Corporation
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA

To whom it may concern,

The transitional provisions specified in MDR Article 120(3) (as amended by (EU) 2023/607) prohibit Notified Bodies from issuing new certificates or amending, modifying, supplementing any existing MDD/AIMDD certificates from 26th May 2021.

The transitional provisions specified in IVDR Article 110(3) (as amended by (EU) 2024/1860) prohibit Notified Bodies from issuing new certificates or amending, modifying, supplementing any existing IVDD certificates from 26th May 2022.

This letter is to confirm that BSI has reviewed and approved the change(s) detailed in the table below. These changes do not represent a significant change in design or intended purpose under MDR Article 120(3) or under IVDR Article 110(3), as applicable, and as per the guidance provided in MDCG 2020-3/MDCG 2022-6.

The related certificate specified below continues to remain valid and devices can be placed on the market based on this certificate as long as the manufacturer complies with the conditions specified in Section 3c of Article 120 of MDR or in Section 3c of Article 110 of IVDR, as applicable.

Certificate	Directive and Annex	Reference Number	Changes approved
CE 00999	93/42/EEC Annex II excluding Section 4	30680018	Change of critical subcontractor. Addition of manufacturer location.

Should you have any queries concerning your certification, or if we can be of further assistance to you, please contact your BSI Scheme Manager.

Yours sincerely,

Graeme Tunbridge
Senior Vice President, Medical Devices

Varex Imaging Corporation
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA

21 Dec 2023

Notified Body Confirmation Letter
Reference: EU2023-607/ 751911

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:

Varex Imaging Corporation
1678 South Pioneer Road
Salt Lake City
Utah
84104
USA

SRN Number (if available): US-MF-000028124

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.,



Graeme Tunbridge
Senior Vice President, Medical Devices

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
XRyTubeRADS 0854344007	Class IIb excluding Class IIb implantable non-WET	N/A	CE 00999; NB Number: 2797
XRyTubeCTGL 0854344007	Class IIb excluding Class IIb implantable non-WET	N/A	CE 00999; NB Number: 2797
XRyTubeMAMMOBH 0854344007	Class IIb excluding Class IIb implantable non-WET	N/A	CE 00999; NB Number: 2797
Img_ReceptorSJM 0854344007	Class IIa	N/A	CE 00999; NB Number: 2797
Img_ReceptorPJF 0854344007	Class IIa	N/A	CE 00999; NB Number: 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
2023/12/21	Initial issue