

# **Updated Field Safety Notice**

#### MR systems with 60cm wide bore

Quadrature Body Coil (QBC) seal adhesive failure may result in exposure of sharp edges

#### <Date>

#### Dear Customer,

Attached you will find an update to Philips 28-Dec-2023 Field Safety Notice relating to the potential for Quadrature Body Coil (QBC) seal adhesive to fail, creating sharp edges that may come in contact with patients.

#### **Summary of Updates:**

Philips has identified additional MR systems where the issue may occur. Section 3 of the Field
Safety Notice has been updated to include the additional model names and numbers (REF) listed
below. According to our records, affected product identified below has been distributed to your
facility.

Model	(REF) Numbers	
Enterprise 1.5T	781145	
Intera 0.5T Standard	781101	
Intera 1.0T Omni/Stellar	781102	
Intera 1.0T Power/Pulsar	781103	
Intera 1.5T	781195, 781295	
Intera 1.5T Achieva Nova-Dual	781108	
Intera 1.5T Master/Nova	781106	
Intera 1.5T Omni/Stellar	781104	
Intera 1.5T Power/Pulsar	781105	
Intera 1.5T R11	781170	
Intera 3.0T Quasar Dual	781150	
Intera CV	781107	
SmartPath to dStream for 1.5T*	782146	
SmartPath to dStream for 3.0T*	782145	

- Section 1 of the Field Safety Notice has been updated to include the number of reported adverse events Philips has received for this issue as of February 2025.
- Section 5 of the Field Safety Notice has been updated to include an additional field correction reference number (FCO78100615) for newly identified MR systems.
  - \* Note: SmartPath to dStream for 1.5T and SmartPath to dStream for 3.0T will be addressed via FCO78100573. The other MR systems listed above will be addressed via FCO78100615.

Please be assured that maintaining a high level of safety and quality is our highest priority. If you need additional information or support concerning this issue, please contact your local Philips representative.

Sincerely,

Akivia Rivera Garcia Head of MR Quality



# **Updated Field Safety Notice**

MR systems with 60cm wide bore

Quadrature Body Coil (QBC) seal adhesive failure may result in exposure of sharp edges

# This document contains important information for the continued safe and proper use of your equipment

Please review the following information with all members of your staff who need to be aware of the contents of this communication. It is important to understand the implications of this communication.

Please retain this letter for your records.

#### <Date>

Dear Customer,

Philips has identified an issue with the MR systems identified in Section 3 of this letter, that could pose a risk for patients and users. This Field Safety Notice is to inform you about:

#### 1. What the problem is and under what circumstances it can occur

The Quadrature Body Coil (QBC) seal adhesive may fail creating sharp edges that may come in contact with patients. The QBC seal may become loose as the patient table travels in a horizontal motion in and out of the system bore. The QBC seal (Figure 1) is a rubber seal that is glued between the cone cover and QBC cover and functions to prevent sharp edges of the QBC cover from contacting patients during an examination.

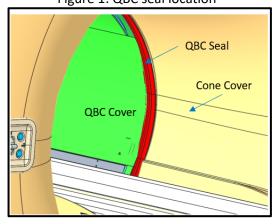


Figure 1. QBC seal location

Philips has received five (5) reports of adverse events associated with this issue: one patient received a cut on the hand, one patient's hair became entangled resulting in a scalp injury, and three patients received lacerations to their arm.

#### 2. Hazard/harm associated with the issue

If the QBC seal becomes loose during the scanning process, the risk to the patient may include one or more of the following: skin abrasions, bruises, lacerations, hair loss/entanglement, and tissue injury.

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#### 3. Affected products and how to identify them

#### **Identification of Impacted Systems:**

MR systems with 60cm wide bore are affected. Refer to Tables 1 and 2 for the system model names and model numbers (REF). The model name and model number (REF) can be found on the system label.

Table 1. Impacted MR Systems

Sample System	Label Example	Model	(REF) Numbers
MODEL: ACHIEVA 1.5T	Achieva 1.5T	781196, 781343, 781296	
REF   781343	REF   781343	Achieva 1.5T Conversion	781346, 781283
(01) 00884838055292		Achieva 1.5T Initial system	781178
Philips Medical Systems Nederland BV		Achieva 3.0T	781277, 781177, 781278,
Veenpluis 4-6, 5084 PC Best, The Netherlands			781344, 781345
RATED VOLTAGE RANGES		Achieva XR	781153, 781253
		Ingenia 1.5T CX	781262, 781261
		Ingenia 3.0T CX	781271, 782105
C € 854 <b>3</b> 10 🗵		Intera 1.5T Achieva Nova	781172
4598 011 13471	Intera 1.5T Achieva Nova-Dual	781173	
		Intera Achieva 1.5T Pulsar	781171
		SmartPath to dStream for 1.5T	781260, 782112
		SmartPath to dStream for XR and 3.0T	781270, 782113, 782129

Table 2. Additional impacted MR Systems

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Sample System	Label Example	Model	(REF) Numbers	
MODEL: ACHIEVA 1.ST  REF 781343  SN Interest.	Enterprise 1.5T	781145		
		Intera 0.5T Standard	781101	
(01) 00884838055292	(01) 0084433055392 Medical Magnetic Resonance System Communic Vision (Carlos Magnetic Resonance System Communic Vision (Carlos Magnetic Resonance System (Carlos Magnetic Resonance Carlos Magnetic Reso	Intera 1.0T Omni/Stellar	781102	
Philips Medical Systems Nederland BV Veenplais 4.6, 5684 PC Best, The Netherlands		Intera 1.0T Power/Pulsar	781103	
		Intera 1.5T	781195, 781295	
RATED VOLTAGE RANGES		Intera 1.5T Achieva Nova-Dual	781108	
		Intera 1.5T Master/Nova	781106	
		Intera 1.5T Omni/Stellar	781104	
C € 1344 <b>⑤</b> ∑	4598 011 13471	Intera 1.5T Power/Pulsar	781105	
		Intera 1.5T R11	781170	
		Intera 3.0T Quasar Dual	781150	
		Intera CV	781107	
		SmartPath to dStream for 1.5T	782146	
		SmartPath to dStream for 3.0T	782145	

#### **Intended Use:**

Philips Magnetic Resonance (MR) systems are Medical Electrical Systems indicated for use as a diagnostic device. This enables trained physicians to obtain cross-sectional images, spectroscopic images and/or spectra of the internal structure of the head, body or extremities, in any orientation, representing the spatial distribution of protons or other nuclei with spin.

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#### 4. Actions that should be taken by the customer / user in order to prevent risks for patients or users

- As part of the preparation before a patient scan:
  - 1. Inspect the QBC seal for separation between the cone cover and QBC cover.
  - If QBC seal is found detached or loose, Stop-use immediately.
  - 3. Contact your local Philips service representative.
- If QBC seal becomes loose during a patient scan:
  - 1. Immediately stop scanning and carefully remove patient from the system.
  - 2. Contact your local Philips service representative.
- Circulate this Field Safety Notice to all users of this device so that they are aware of the issue.
- Please retain this letter with your system(s) until a solution is installed on your system; ensure the letter is in a place likely to be seen/viewed.
- Please complete and return the attached response form to Philips promptly and no later than 30 days from receipt via email to: <Philips representative contact details to be completed by the Market/Business>. Completing this form confirms receipt of the Field Safety Notice, understanding of the issue, and required actions to be taken.

#### 5. The actions planned by Philips to correct the problem

Philips will contact you to schedule a time for a Field Service Engineer (FSE) to visit your site and replace your system's QBC Seal (reference FCO78100573, FCO78100615).

Please be assured that maintaining a high level of safety and quality is our highest priority. If you need any further information or support concerning this issue, please contact your local Philips representative.

Sincerely,

Akivia Rivera Garcia Head of MR Quality

Philips Reference: 2023-PD-MR-013 Rev C



# **Updated Field Safety Notice Response Form**

Reference: MR Systems Quadrature Body Coil (QBC) seal failure (reference FCO78100573, FCO78100615)

**Instructions:** Please complete and return this form to Philips promptly and no later than 30 days from receipt. Completing this form confirms receipt of the Field Safety Notice, understanding of the issue, and required actions to be taken.

required actions to be taken.
Customer/Consignee/Facility Name:
Street Address:
City/State/ZIP/Country:
Customer Actions:  • Follow the instructions provided in Section 4 of the Field Safety Notice.
We acknowledge receipt and understanding of the accompanying Field Safety Notice and confirm that the information from this notification has been properly distributed to all users of the affected systems.
Name of person completing this form:
Signature:
Printed Name:
Title:
Telephone Number:
Email Address:
Date (DD/MM/YYYY):

Please complete and return the response form to Philips promptly and no later than 30 days from receipt via email to: <Philips representative contact details to be completed by the Market/Business>.

Philips Reference: 2023-PD-MR-013 Rev C Page 5 of 5