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For all users of mint Lesion™ versions from 3.8.6 up to and including 3.9.2

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Urgent Field Safety Notice

For the Attention of: All users of mint Lesion™ versions from 3.8.6 up to and including 3.9.2

Dear mint Lesion™ user,

We would like to inform you about a malfunction that may occur when using mint Lesion™ in one of the versions listed below with one of the reading templates listed below.

Information on affected devices

Affected medical device	mint Lesion™
Basic UDI-DI	426049588MINTLESIONSM

Affected mint Lesion™ device versions

Device Version	UDI-DI	UDI-PI
mint Lesion™ 3.8.6	04260495880389	(01)04260495880389(10)3.8.6(11)220414
mint Lesion™ 3.9.0	04260495880396	(01)04260495880396(10)3.9.0(11)230216
mint Lesion™ 3.9.1	04260495880396	(01)04260495880396(10)3.9.1(11)230502
mint Lesion™ 3.9.2	04260495880396	(01)04260495880396(10)3.9.2(11)231102

Affected components

The malfunction may occur when one of the following reading templates is used:

- Head and Neck cancer (TNM 8.0)
- Esophagus / Stomach (TNM 8)
- Colorectal cancer (TNM 8 / ESGAR Recommendations)
- PERCIST

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- PERCIST all targets
- Rapno LGG

> **Notice**
>
> Note: **mint Lesion™** allows to customize reading templates or to install custom-developed reading templates. Such templates may also be affected, if they contain eCRF sections with measurement questions.

Problem description

The malfunction is caused by a software error that is present in product versions 3.8.6, 3.9.0, 3.9.1, 3.9.2. The malfunction can occur in the following use scenario (**all** steps must apply):

1. The user activates one of the affected reading templates
2. The user assesses a follow-up time-point (e.g., re-staging or response assessment)
3. The user creates a measurement for an eCRF question listed below
4. The user switches to a different time-point or to a different patient record by leaving the "Read screen" without closing and restarting the application in-between.

Affected eCRF questions

Reading template	eCRF section	Question
Head and Neck cancer (TNM 8.0)	Further Size Information	• Third dimension diameter
	Tumor extension	• Depth of invasion
	Characterization	• Third dimension diameter
Esophagus / Stomach (TNM 8)	Further information	• Distance to esophagogastric junction • Craniocaudal Diameter

Reading template	eCRF section	Question
Colorectal cancer (TNM 8 / ESGAR Recommendations)	Local extent of disease	• Distance from cranial edge of tumor to anterior peritoneal reflection • Craniocaudal length • Distance to anocutaneous line • Distance to anorectal junction • Length of anal canal • Direct tumor infiltration: Minimum distance to the mesorectal fascia • EMVI: Minimum distance to the mesorectal fascia • Tumor deposits: Minimum distance to the mesorectal fascia • Extramural depth of invasion: measurement
	Relationship to mesorectal fascia	• Minimum distance to the mesorectal fascia
PERCIST	Lesion Properties	• CT/MRI Lesion Size
PERCIST all targets	Lesion Properties	• CT/MRI Lesion Size
Rapno LGG	Additional measurements	• Third perpendicular diameter • Cystic PPD

Please be aware you may use customized reading templates in your mint Lesion installation. Customized reading templates that contain eCRF questions of type "Measurement" are also affected.

Effects of the problem

After switching to a different time-point, different case or different patient, the sidebar area in the read screen will erroneously still display the eCRF section from the original patient and time-point that contains the measurement question, including the measurement value and other answers that apply to the originally selected patient and time-point. The section that applies to the current patient and time-point is displayed, too. The user interface does not allow to identify which section belongs to the current patient/time-point and which section contains outdated information. Figure 1 shows this situation.

The measurement questions could be displayed either as missing or with values from other time-points or patients. Figure 1 shows an example: The measurement question "Minimum distance to the mesorectal fascia" is erroneously shown twice, once for the current assesment context (marked with **2**), once for a prior assessment context (marked with **1**) .

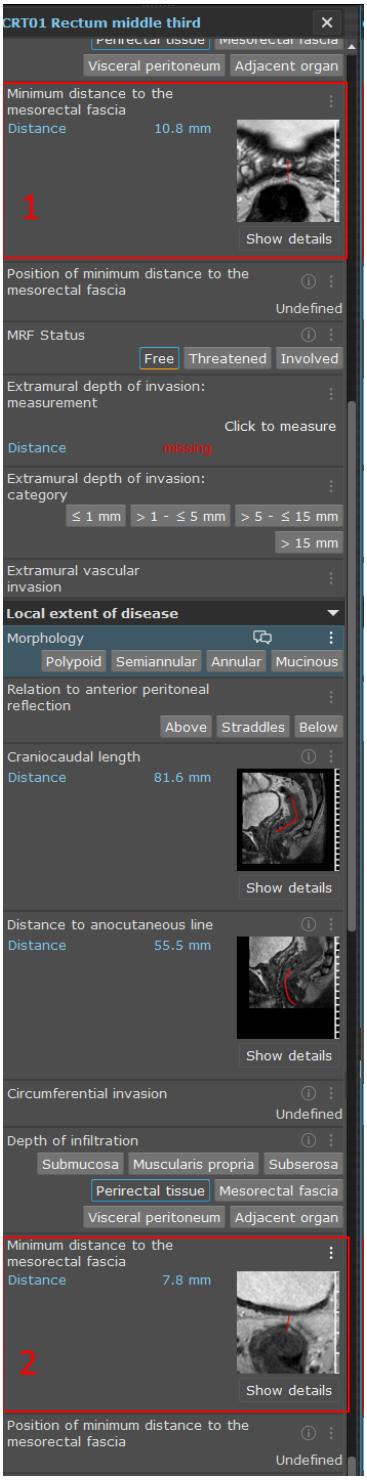

Figure 1: The measurement "Minimum distance to the mesorectal fascia" is displayed both for the current assessment context and erroneously additionally for a wrong assessment context

Notice

The wrong display of a section that belongs to a different time-point or case or patient is limited to the Read Screen. The Report Screen and all exported reports (e.g., PDF, CSV, XML, HTML, HL7, Word Add-In) are not affected by this malfunction. They all show correct information. If you use these reporting methods, your reports will contain correct information.

Actions to be taken by the user

Please read this information carefully and assess whether you are using an affected product version and affected reading templates. If that is the case, the malfunction may occur in your system. Please be aware that the malfunction may occur. When assessing follow-up time-points (e.g., re-staging or response assessment follow-up), scroll through the sidebar to check if duplicate sections are shown. If a section is shown twice, please close and restart the application. This will remove the out-of-context section from the sidebar.

If you are manually writing/dictating a radiological report based on the information shown in the read screen sidebar, take extra caution to identify duplicate sections. Do not use the information from the out-of-context section for reporting. Prefer to use the mint Lesion™ report screen or the integrated reporting capabilities to create a radiological report.

You can prevent this malfunction from occurring by restarting the application in between each assessment that is using one of the affected reading templates.

If you believe that this failure could have occurred in past use of mint Lesion™, please review the potentially affected radiological reports in your reporting application and take the necessary steps to correct them.

Actions being taken by the manufacturer

The error will be corrected with a software update. Mint Medical Support will contact you when the update is available to schedule the installation of the update on your system.

General Information

FSN Type	New Field Safety Notice	
Further advice or information already expected in follow-up FSN	Not planned	
Manufacturer information	Legal manufacturer name	Mint Medical GmbH
	Address	Mint Medical GmbH Burgstr. 61 69121 Heidelberg Germany
	Manufacturer Email	info@mint-medical.de

	Manufacturer Phone	+49 6221 64 79 76 0
	EUDAMED Single Registration Number (SRN)	DE-MF-000020202
	Person responsible for regulatory compliance (PRRC)	Dr. Jochen Neuhaus
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	PRRC Phone	(+49) 6221 32 18 018

The Competent (Regulatory) Authority of your country has been informed about this communication to customers.

Transmission of this Field Safety Notice

This notice needs to be passed on to all users of **mint Lesion™** within your organization. Please maintain awareness of this notice and resulting action for an appropriate period to ensure the effectiveness of the corrective action.

Please report all device-related incidents to the manufacturer, distributor or local representative, and the national Competent Authority if appropriate, as this provides important feedback.

Heidelberg, 2024-01-10

Dr. Jochen Neuhaus